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1987

# Cold-climate nitrification on plastic media trickling filters

Harvey Allen Gullicks *Iowa State University*

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Gullicks, Harvey Allen

COLD-CLIMATE NITRIFICATION ON PLASTIC MEDIA TRICKLING FILTERS

Iowa State University **PH.D.** 1987

University Microfilms International 300 N. Zeeb Road, Ann Arbor, MI 48106

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#### Cold-climate nitrification on plastic media

trickling filters

by

Harvey Allen Gullicks

A Dissertation Submitted to the

Graduate Faculty in Partial Fulfillment of the

Requirements for the Degree of

DOCTOR OF PHILOSOPHY

Department: Civil Engineering Major: Sanitary Engineering

Approved:

Signature was redacted for privacy.

In Charge of Major Work

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For the Major Department

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For the Graduate College

Iowa State University Ames, Iowa

1987

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Figure 48. Predicted NH<sub>4</sub><sup>+</sup>-N removal, kg/d  $\cdot$  m<sup>2</sup> of media surface, 176 versus applied hyraulic load and applied NH^^-N for nitrification of municipal secondary clarifier effluent **(BOD5** < 30 mg/1 and SS < 30 mg/1), wastewater temperature 10°C and media depths of 6-6.55 m of vertical or cross-flow plastic media (specific surface =  $89-98 \text{ m}^2/\text{m}^3$ ). No safety factor is implied. Allow additional surface area (25-50%) for bulk liquid percent of D.O. saturation less than 65% and/or higher specific surface area media to account for plugging

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#### DEDICATION

To my wife, Jean, and our sons, Donnie and Grant, whose love, patience and understanding have made both family life and professional growth possible and enjoyable, and to Dr. Donald Naismith, Shirley Naismith, Dr. Jonathan Berg, Laurie Berg, and Warren and Connie Dunham for their support throughout the difficult times.

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## NOMENCLATURE AND ABBREVIATIONS

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### Nomenclature



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## Abbreviations

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#### **INTRODUCTION**

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The trickling filter process was first used in the United States for wastewater treatment in 1908 (Tsugita et al., 1977). The trickling filter consists of a tower, containing a medium with a high specific surface area and a high void volume per unit medium volume. As wastewater is distributed to the top of the medium, it trickles down through the medium voids and over the medium surface area on which microorganisms are attached. The microorganisms attached to the medium are referred to as the biofilm or biomass. The succession of communities of microorganisms produced by microbial competition under various loading conditions and at various depths in the trickling filter will be addressed in a subsequent section. The microorganisms in the biofilm utilize the organic material and nutrients in the wastewater for growth, thereby removing or altering these contaminants.

The implementation of more stringent water quality legislation in 1972 has left many trickling filter plants designed before that time in violation of the discharge standards for biochemical oxygen demand (BOD), suspended solids (SS), and ammonium nitrogen  $(NH_A^+ - N)$ . Consequently, many designers have come to view the trickling filter process only as a pretreatment process. The activated sludge process has generally been accepted as the process-of-choice for nitrification, the biological conversion of  $NH_4^+$ -N to nitrite (NO<sub>2</sub><sup>--N</sup>) and nitrate  $(NO<sub>3</sub> - N)$ . Young (1974) reported that inflexibility in

trickling filter operation was responsible for the trickling filter's reputation for uncontrollable nitrification performance.

The advent of new, lightweight plastic media for trickling filters has improved the performance of trickling filters. Several studies have shown that trickling filters are capable of producing effluents with consistently low ammonia nitrogen concentrations even in cold climates when carbonaceous oxygen demand loadings are low (Sampayo, 1973, 1974, 1976, and 1981, Sampayo and Metcalf, 1984, Buddies and Richardson, 1973, Duddles et al., 1974, Baxter and Woodman, 1973, Bruce et al., 1975, and Gujer and Boiler, 1984). Gujer and Boller (1984) reported that tertiary nitrifying trickling filters do not require additional sedimentation tanks. The effluent can be directly filtered just as secondary effluent is directly filterable. The trickling filter should be considered to be a viable nitrification process.

Trickling Filter Application for Nitrification

Nitrification can be achieved in trickling filters in two processes. Either combined carbon oxidation-nitrification or separate stage processes may be used. These processes are distinguished on the basis of the BOD<sub>5</sub>/Total Kjeldahl Nitrogen (TKN) ratio for the wastewater being treated (Parker et al., 1975). If the ratio is less than 3.0 and the nitrogenous oxygen demand (NOD) is at least 60 percent of the total oxygen demand (TOD), the system is classified as a separate-stage process. If the ratio is greater than 5.0 and the

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NOD is less than 50 percent of the TOD, the system is classified as a combined carbon oxidation-nitrification process. Trickling filters treating wastewaters with ratios between 3.0 and 5.0 are considered to provide an intermediate degree of separation of the carbon oxidation and nitrification.

In general, the combined carbon oxidation-nitrification process is applied more frequently as a pretreatment to subsequent processes that provide an effluent meeting the necessary discharge requirements. Brown and Caldwell (1980) reported that the Stockton, California, forced-draft, combined carbon oxidation-nitrification trickling filter plant achieved 80-90 percent nitrification and removed approximately 50 percent of the organic nitrogen with 6.7 m. of vertical plastic media with a specific surface of 89 sq. m./cu. m. . Suwan arpa (1974) reported that heterotrophic organism growth accounted for substantial nitrogen uptake, but that  $NH_A^+$ -N was not the only nitrogen source for heterotroph growth.

When separate-stage nitrification is applied, staging, which is the operation of trickling filters in series, is often the selected design. Figure 1 demonstrates the schematic flow pattern used in staging of trickling filters. Staging allows a high quality nitrified effluent to be produced.

Plastic media, with large specific surface areas relative to rock and other media types, are generally selected for nitrifying trickling filters because their light weight and corrugated construction allows tall towers to be used and because their high void volumes help

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Figure 1. Staging of filters. After Tsugita et al. (1977)

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prevent clogging and improve air circulation through the tower. Plastic media may be classified as random, vertical or cross-flow media depending on the placement in the tower and on the hydraulic flow pattern they provide. Figure 2 demonstrates this classification system.

Average hydraulic loading rates of 0.34 to 1.7  $1/m^2$ .s are typically used for trickling filters filled with plastic media. Total BODg loading rates and soluble BODg concentrations must be low in order to achieve high degrees of nitrification. It is generally recommended that the total  $BOD<sub>5</sub>$  loading rate not exceed about 4 kg/1000  $\text{m}^2$  of specific surface area/d and that the soluble BOD<sub>5</sub> not exceed 20 to 30 mg/1 (Parker and Richards, 1985). Figures 3a and 3b demonstrate the onset of nitrification at soluble BOD<sub>5</sub> values less than 20 mg/1.

A model developed by Wanner and Gujer (1984) predicted that the activity of nitrifiers is dependent on the activity of heterotrophs. Their model predicted that nitrifying microorganisms did not exist in biofilms at soluble chemical oxygen demand (COD) concentrations greater than 27 mg/1.

Current Status of Design Procedures for Nitrification

The degree of nitrification achieved in a trickling filter depends on many operating parameters. The effects of variations in these operating parameters are discussed in detail in subsequent sections. The most important operating parameters are the wastewater

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Figure 3a. Nitrification versus soluble BOD<sub>E</sub> concentration. Parker and Richards (1985) After

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Nitrification versus soluble  $BOD_5$ . After Parker and Rich-<br>ards (1985) Figure 3b.

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temperature, pH, and alkalinity, the hydraulic loading rate, the  $NH_A^+$ -N concentration, the recycle ratio, and the organic loading.

There are several mathematical models (Williamson and McCarty, 1976a and 1976b, Suidan, 1986, Rittmann and McCarty, 1978, 1980a, 1980b, and 1981, Harremoes, 1982, and Strand, 1986) for trickling filters that attempt to include nitrification kinetics, the effects of some of the critical operating parameters, and recent fixed-film theoretical concepts. The models are complex and cumbersome, and a good deal of uncertainty is associated with the constants in the models. The current methods for the design of nitrifying trickling filters have been largely or entirely based on empirical data.

Figures 4, 5, 6, and 7 are examples of the types of design curves currently used for nitrification in plastic media trickling filters. The curves present an empirical relationship between the desired effluent  $NH_A^+$ -N concentration and the required surface area of media. Strictly speaking, these curves are applicable only to municipal wastewaters and the conditions under which the data were generated. The data were collected for plastic media trickling filters that followed primary treatment and secondary treatment for carbonaceous BOD removal. Secondary sedimentation was included before the nitrification towers. The curves were developed for specific plastic media types, characteristics, and depths, and the curves should not be extrapolated directly to other media types, characteristics, and depths, although the design engineer has few other options in the initial stages of design.

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EPA Process Design Manual design curve: Midland, Michigan Figure 4. data

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'Figure 5. EPA Process Design Manual design curve: Lima, Ohio data

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Figure 7. Filterpak ammonia loading curves. After Lewis (1984)

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Figure 4 contains some computational errors and Figures 4, 5, 6, and 7 contain conceptual shortcomings which leave their validity in question. Figure 4 contains points that were inaccurately calculated or plotted, and the effects of the independent  $NH_4^+$ -N loading rate parameters on the effluent  $NH_4^+$ -N concentration are obscured in all four figures. The excessive scatter of the data points clearly illustrates that the effects of the hydraulic loading rate and the influent  $NH_A^+$ -N concentration to the tower are not accounted for adequately. Figure 8 shows the effect of  $NH_4^+$ -N loading on  $NH_4^+$ -N concentration in the tower effluent.

#### Need for Research

The design of trickling filters for nitrification will likely continue to be based on empirical data in the foreseeable future. The currently available nitrification data, unlike the massive carbonaceous BOD<sub>5</sub> data base, do not provide a sufficient background for evaluation of all the constants needed for the mathematical models. There are many types of plastic media on the market; however, most of the existing data are for media developed more than a decade ago. The data base for the more recently developed cross-flow media is small and largely inaccessible for proprietary reasons. The basis for design must include the effects of  $NH_4^+$ -N loading parameters and the other operating variables if it is to be of use in optimizing and designing nitrification trickling filters.

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Figure 8. Ammonia loading vs. NH  $_{2}$ -N in tower effluent. After Adams et al.  $(1981)$   $3$ 

Gullicks and Cleasby (1986) proposed new empirical curves (Figures 9 and 10) that would account for the effects of  $NH_4^+$ -N loading parameters. These curves were based on the pilot studies conducted at Midland, Michigan, (Buddies and Richardson, 1973 and Buddies et al., 1974) and Bloom Township, Illinois (Baxter and Woodman, Inc. 1973). These curves still retained some of the objectionable limitations discussed above since they were based on pilot data developed with only one type of plastic medium and only one medium depth.

Parker (1984a and 1984b) has called for trickling filter research on trickling filter design models and on media research and design. He acknowledged the recent development of fairly complex biofilm models by the academic community, but observed that they have not been applied by the design community because of their complexity and unproven nature. Consultants have continued to use simplistic design equations and empirical design curves.

Parker reported that the available trickling filter BOD data base consists almost entirely of influent and effluent data only and that very few data from intermediate points exist. The lack of trickling filter nitrification profile data is even greater. Parker called for full-scale trickling filter towers to be built with intermediate sampling points to allow collection of profile data on full-scale units in the future. He stated that profile data would be obtained from a couple of towers in cooperation with the Environmental Protection Agency (EPA) in 1986.

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Figure 9. Predicted NH<sub>4</sub>  $^+$ –N removal, kg/d  $\cdot$  m $^2$  of media surface, versus applied hydraulic load and applied NH<sub>4</sub><sup>+</sup>-N for nitrification of municipal secondary clarifier effluent (BODs < 30 mg/L and SS < 30 mg/L), wastewater temperatures >  $14^{\circ}$ C, and 6.55 m of vertical plastic media (specific surface =  $88.6 \text{ m}^2/\text{m}^3$ ). After Gullicks and Cleasby (1986)

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Figure 10. Predicted  $NH_4^+$ –N removal, kg/d  $\cdot$  m $^2$  of media surface, versus applied hydraulic load and applied NHi\* -N for nitrification of municipal secondary clarifier effluent **(BOD5** < 30 mg/L and SS < 30 mg/L)-, wastewater temperatures 10-14°C, and 6.55 m of vertical plastic media (specific surface = 88.6 m'/m'). After Gullicks and Cleasby (1986)

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Most, if not all, of the available design models and curves include the total dry specific surface area term rather than an effective or wetted surface area term. In most cases, an increase in the specific surface area is assumed to give a proportional increase in the mass removal capacity of a given volume of media (refer to Figure 7). This is seldom a correct assumption (Brown and Caldwell, 1980, and Sarner, 1984), however, the data necessary to establish the correct relationship between media specific surface area and mass removal capacity for a given volume of media are lacking.

The effects of flow patterns within different media configurations and the effects of different wastewater application methods have recently received greater attention. Unfortunately, much of this information has been evaluated only by simplistic design equation methods rather than in conjunction with proposed fixed-film models. The media manufacturers have claimed that cross-flow media possess superior hydraulic residence times, surface wetting characteristics, and oxygen transfer characteristics. The superior hydraulic residence times of cross-flow media have been demonstrated by tracer test methods (Albertson and Davies, 1984, Albertson and Eckenfelder, 1984, Drury et al., 1986, Harrison and Daigger, 1985, Parker and Merrill, 1984, and Richards and Reinhart, 1986).

The surface wetting and oxygen transfer superiority claims are largely speculative and disputed at this time. Total wetted surface area and total oxygen transfer area undoubtedly increase with increases in specific surface area, but no data exist to show that

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cross-flow media wetted surface areas are greater than the wetted surface areas of vertical-flow media when both have the same total specific surface area. The work of Logan et al. (1986) suggests that cross-flow media provide better liquid phase mass transfer than vertical-flow media (and, therefore, better oxygen mass transfer) because mixing of the liquid film occurs at more configuration nodes.

The majority of pilot-scale trickling filter data have been collected using continuous dosing of wastewater to the filter. Few studies have included intermittent dosing of the wastewater even though full-scale plants typically are dosed intermittently by rotating distributors.

The majority of nitrification data available for trickling filters are for wastewater temperatures greater than 15°C. Cold climate data are needed to allow designers to extend trickling filter technology to cold-climate nitrification process designs.

## PURPOSE AND OBJECTIVES

The city of Ames, Iowa, plans to construct a new dual-stage trickling filter water pollution control plant (WPCP) to replace the existing, overloaded, conventional trickling filter plant. The new plant is expected to provide carbonaceous BOD removal in the first stage filter and nitrification in the second stage filter. Iowa State University obtained a grant from the city of Ames to study cold-climate, separatestage nitrification in a pilot-scale trickling filter filled with high specific surface, cross-flow plastic media. The experimental investigation reported herein was conducted at the existing Ames WPCP and was limited to separate-stage nitrification.

There has been a lack of nitrification data for trickling filters, particularly in cold climates. Ammonium profile data (concentration versus tower depth) in trickling filters are lacking. Few nitrification data are available for cross-flow media. The effective media surface area is unknown at this time, and researchers and designers use the dry specific surface area of the media in their equations and models. The recently developed, theoretically based fixed-film models are largely unproven due to the lack of available profile data. The majority of equations currently used in design assume first-order kinetics despite the fact that the upper portions of trickling filters often follow zeroorder kinetics (with respect to the contaminant substrate). Data from intermittently-dosed, pilot-scale plants are lacking.

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The investigation was undertaken to examine the effects of a range of hydraulic loading rates and influent  $NH_4^{+-}N$  concentrations on nitrification in cross-flow plastic media at wastewater temperatures ranging from 6° to 14°C. Limited data were collected at wastewater temperatures greater than 15°C to allow comparison of nitrification performance reported in the literature with that achieved in this experimental work.

The specific objectives of this study were to:

- 1. Determine the individual effects of hydraulic loading rate and influent  $NH_4^+$ -N concentration on areal nitrification rates.
- 2. Determine the effect of wastewater temperature on areal nitrification rates.
- 3. Compare continuously-dosed pilot plant and intermittently-dosed pilot plant performance.
- 4. Provide  $NH_4^+$ -N, DO,  $NO_2^--N + NO_3^--N$ , TKN, COD, pH, and wastewater temperature profile data in cross-flow plastic media.
- 5 Verify or disprove zero-order nitrification kinetics with respect to  $NH_{\Delta}^{+}$ -N concentrations, which is predicted at concentrations greater  $than 4$  mg/l.
- 6. Examine the effective or wetted specific surface area of the crossflow media, and if possible, establish a relationship between the hydraulic loading rate, the dry media specific surface area, and the wetted specific surface area.
- 7. Examine some of the theoretically-based, fixed-film models to determine if they adequately predict the nitrification performance observed in the pilot-scale plant.

Improve the design curves previously proposed by Gullicks and Cleasby (1986).

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### REVIEW OF LITERATURE

Nitrogen, in one or more of its numerous oxidation states, is a key component of life processes for all plants and animals. Nitrogen forms most important to environmental engineers are organic, gaseous, nitrite, nitrate, and ammonia nitrogen which possess oxidation states of  $-3$ , 0, +3, +5, and -3, respectively.

The conversion of ammonia to nitrite and nitrite to nitrate, a process "called nitrification, is accomplished biologically by two principal autotrophic bacteria genera, Nitrosomonas and Nitrobacter. Autotrophic nitrifying organisms derive their energy from the oxidation of inorganic nitrogen substances, rather than organic matter; and they utilize carbon dioxide as a carbon source for synthesis, rather than organic carbon.

# Nitrifier Growth

Nitrosomonas and Nitrobacter must both be available to achieve complete nitrification. Nitrosomonas can only convert ammonia to nitrite, while Nitrobacter can only convert nitrite to nitrate. The growth rate of <u>Nitrosomonas</u> is limited by the concentration of  $NH_A^+$ -N, while the growth of Nitrobacter is limited by the concentration of  $NO<sub>2</sub>$  -N. The kinetics of biological growth are usually described by the Monod kinetic equation:

$$
\mu = \hat{\mu} \frac{S}{K_s + S} \tag{1}
$$

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where:  $\mu$  = specific growth rate of microorganisms,  $\frac{mass~of~microorganism}{mass~of~biofilm~•~t}$ 

- $\hat{\mu}$  = maximum specific growth rate of maximum specific growth rate or mass of microorganism<br>microorganisms, the mass of biofilm . t
- $K_c$  = half velocity constant = growth limiting substrate concentration at half the maximum growth rate, mg/cm $^3$ . S = growth limiting substrate concentration,  $mg/cm^3$ .

The Haldane kinetic equation, which includes an inhibitory term in the denominator is gaining in popularity. It is expressed as follows (Rozich and Castens, 1986):

$$
\mu = \frac{\hat{\mu}S}{K_s + S + S^2/K_i}
$$
 (2)

where:  $K_i$  = Haldane inhibition constant, mg/cm<sup>3</sup>.

Figure 11 gives a graphical comparison of the Monod and Haldane kinetic growth rate equations. Rozich and Castens (1986) conducted a completely-mixed, two-stage, continuous culture study and reported that nitrification kinetics are represented more accurately by the Haldane equation. Table 1 gives values for  $\hat{\mu}$ ,  $K_c$ , and  $K_i$  calculated using a biomass mass balance equation and using a substrate mass balance equation. The Haldane equation gave comparable values for both mass balances, while the Monod equation did not. Allowable parameter ranges for which Haldane kinetics are valid are also given in Table 1.

Inhibition kinetics is of greater significance in activated sludge processes than in fixed-film processes where the rate-limiting reactant

 $\omega \propto \omega_{\rm{min}}$ 



Figure 11. Comparison of Monod (noninhibitory) and Haldane (inhibitory) growth rate models. After Rozich and Castens (1986)

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Table 1. Results of the growth model fits to nitrifier growth rate date. After Rozich and Castens (1986)

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may be oxygen rather than  $NH_{4}^{+}-N$ . The discussion in the subsequent section dealing with the determination of the flux-limiting reactant will make this point clear.

The maximum growth rate of Nitrobacter is much greater than that of Nitrosomonas.  $K_{\rm c}$  values, mg N/1, for both species are small. Thus, nitrite cannot accumulate to high concentrations in biological treatment systems under steady state conditions, and the conversion of  $NH_A^+$ -N to nitrite is the rate-limiting step for nitrifier growth.

Calculations based on thermodynamic theory predict that 0.29 mg Nitrosomonas will be synthesized per mg of ammonia nitrogen oxidized. Similarly, 0.084 mg Nitrobacter will be synthesized per mg of nitrite nitrogen oxidized. Experimentally measured yield values are considerably lower than these values probably because of microorganism maintenance functions (Haug and McCarty, 1971).

Nitrification usually occurs in aqueous carbonic acid systems at pH values less than 8.3. The consumption of carbon dioxide by the organisms results in some depletion of the dissolved carbon dioxide form (carbonic acid,  $H_2CO_3$ ). Simultaneously, the production of free acid  $(H^+)$  by the organisms proceeds immediately through reaction with bicarbonate (HCO<sub>3</sub><sup>-</sup>) to carbonic acid. Synthesis-oxidation equations have been developed for Nitrosomonas and Nitrobacter using representative yield values of 0.15 and 0.02 mg VSS per mg of nitrogen oxidized, respectively, and representative oxygen consumption measurements (Gujer and Jenkins, 1974 and Water Pollution Research Laboratory, England, 1971). These equations are:

 $\sqrt{2}$ 

55 NH<sub>4</sub><sup>+</sup> + 76 O<sub>2</sub> + 109 HCO<sub>3</sub><sup>+</sup> 
$$
C_5H_7NO_2
$$
 + 54 NO<sub>2</sub><sup>-</sup> + 57 H<sub>2</sub>O +  
\n104 H<sub>2</sub> CO<sub>3</sub>  
\n400 NO<sub>2</sub><sup>-</sup> + NH<sub>4</sub><sup>+</sup> + 4 H<sub>2</sub>CO<sub>3</sub> + HCO<sub>3</sub><sup>-</sup> + 195 O<sub>2</sub><sup>+</sup>  $C_5H_7NO_2$  +  
\n3 H<sub>2</sub>O + 400 NO<sub>3</sub><sup>-</sup>  
\noverall synthesis and oxidation reaction:

$$
NH_4^+ + 1.86 O_2 + 1.98 HCO_3^- \rightarrow 0.0206 C_5H_7NO_2 + 1.041 H_2O + 0.980 NO_3^- + 1.88 H_2CO_3
$$
 (5)

The synthesis-oxidation equation indicates that approximately 7 mg of alkalinity are consumed per mg of  $NH_4^+$ -N oxidized and that the  $H_2CO_3$  concentration increases. Thus, the pH of the system decreases. In an open system, such as the trickling filter, carbon dioxide is continually stripped from the wastewater retarding the pH depression as long as sufficient alkalinity was present initially in the wastewater. Gujer and Boiler (1984) reported that alkalinity was rate limiting at bulk liquid concentrations less than 1 meq/1 (50 mg/1 as  $CaCO<sub>3</sub>$ . Figure 12 shows that nitrification rates are significantly lower at pH values below 7.

### Trickling Filter Fixed-Film Theory

Sarner (1981), Richards (1984), Richards and Reinhart (1986), Muslu (1983), Harrison and Daigger (1985), Germain (1966), and Parker and Merrill (1984) have shown that the trickling filter behaves like a plug flow reactor with varying degrees of dispersion. That is.



E ATTACHED GROWTH REACTOR AT 22 C HUANG AND HOPSON (1974 a,b)

Figure 12. Effect of pH on nitrification rate. After Sawyer et al. (1973)



wastewater distributed to the top of the filter at any given time tends to flow as a unit from the top to the bottom of the tower.

Maier et al. (1967), Williamson and McCarty (1976a and 1976b), and Logan et al. (1986) have shown that the flow of wastewater over the surface of media in trickling filters at the hydraulic loading rates in common use is laminar. This implies that a nearly-stagnant layer exists next to the biofilm and that the bulk flow is smooth and streamlined as represented in Figure 13.

Substrate must diffuse through the bulk liquid layer, across the "stagnant" layer, and into the biofilm layer where microorganisms use it for growth. Oxygen must diffuse across the air-liquid interface, across the entire bulk liquid layer, across the "stagnant" layer, and into the biofilm layer where microorganisms utilize it. The bulk liquid layer is considered to be void of microorganisms in order to simplify the modeling of mass transport in trickling filters. This is a valid approximation in view of the relative populations of the microorganisms in the biofilm and in the liquid.

Diffusion through the bulk liquid and the "stagnant" layer at the biofilm interface is largely dependent on the liquid nearest the biofilm. Thus, diffusion through the bulk liquid and the "stagnant" layer is arbitrarily represented by diffusion through an effective or equivalent liquid diffusion layer.

Figure 14 is a schematic representation of the potentially ratelimiting phenomena in biofilm reactions and substrate concentration

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Figure 13. Schematic model of the trickling filtration process. Vaughan and Holder (1984). After

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Figure 14. Schematic showing the potentially rate-limiting phenomena in biofilm reactions and substrate concentration profiles. After Gullicks and Cleasby (1986)

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 $\label{eq:3} \sqrt{p_{\rm e}^2} \left| \frac{1}{2 \pi \epsilon_{\rm eff}^2} \right|$ 

profiles. The biofilm thickness is greater than the active biomass layer in nearly all situations, except when excess sloughing or hydraulic shearing of the biofilm occur. The rate of substrate utilization is usually flux-limited (i.e., mass transport limited). The limitation of the rate of substrate utilization by the ability of the microorganisms to process the reactants, a process called metabolism limitation, typically occurs only when excess sloughing or hydraulic shearing of the biofilm occur so that all reactants fully penetrate the biofilm at concentrations high enough to sustain growth.

#### Flux-Limitation of Substrate Uptake

Harris and Hansford (1976) demonstrated that the rate of substrate uptake in trickling filters could be flux-limited by the substrate, or by oxygen, or by both the substrate and oxygen simultaneously depending on which reactant is depleted first within the biofilm depth as shown in Figure 15. Simultaneous substrate and oxygen flux limitation occurs over a narrow region of the normal range of bulk liquid substrate concentrations applied in fixed-film nitrification processes (Rittmann and Dovantzis, 1983). Therefore, consideration of a single reactant as the flux-limiting species simplifies the mathematics necessary for modeling, and the error, in any case, probably is not greater than the uncertainties related to measurement of the many parameters associated with the Monod or Haldane kinetic equations, or for that matter, the measurement of the constants associated with fixed-film modeling and design equations.

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Mass transfer in the liquid phase can be represented by Equation 6 when a single reactant is rate-limiting (Williamson and McCarty, 1976a).

$$
J = -AD_w \frac{dS}{dZ} = -AD_w (S_s - S_o)/L_1 + L_2)
$$
 (6)

where

 $J =$  mass transfer rate, mg/d

- $A = cross sectional area perpendicular to the direction of mass$ transfer, cm'
- $D_{\mu}$  = diffusivity of the limiting reactant, either oxygen or substrate, in the wastewater, cm'/d
- $S_{0}$  = concentration of the limiting reactant in the bulk liquid. mg/cm'
- $S_{\rm g}$  = concentration of the limiting reactant at the biofilm-liquid interface, mg/cm'
- $L_1+L_2$  = the effective liquid diffusion layer thickness, cm.  $L_2$  is relatively constant.  $L_1$  varies with hydraulic
	- loading.

 $Z =$  distance in the direction of mass transfer, cm.

The utilization or rate of uptake of the single limiting reactant in the biofilm is proportional to the biomass density and follows a saturation functional pattern commonly represented by Monod kinetics. The ammonium oxidation rate can be related to the Nitrosomonas growth rate for any single limiting reactant as follows;

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$$
\frac{dS_f}{dt} = -k S_f X_f / (K_s + S_f) = -\frac{\hat{\mu}_N}{Y_N} \frac{v_M}{v_M M W_N} \frac{X_f S_f}{K_s + S_f}
$$
 (7)

where:

- $S_f$  = concentration of the limiting reactant, either oxygen or substrate, in the biofilm, mg/cm'
	- $t =$  time, d
- $k =$  maximum specific rate of utilization of the limiting reactant, mass of reactant/mass of biofilm • d  $K_{c}$  = limiting reactant half-velocity coefficient, mg/cm<sup>3</sup>  $X_{\epsilon}$  = biofilm density, mg/cm<sup>3</sup>  $Y_{\text{M}}$  = specific nitrifier organism yield coefficient, mass of nitrifiers mass N oxidized  $\hat{\mu}_M$  = maximum specific growth rate of nitrifiers, mass of nitrifiers mass of biofilm • d  $v =$  stoichiometric coefficient of limiting reactant in Eq. 5  $v_N$  = stoichiometric coefficient of nitrogen in Eq. 5  $MW = molecular weight of limiting reactant$  $MW_{M}$  = molecular weight of nitrogen Molecular diffusion in the biofilm is related to the limiting

reactant gradient by Fick's second law.

$$
\frac{dS_f}{dt} = D_f \frac{d^2S_f}{dZ^2} \tag{8}
$$

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Where:

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 $D_f$  = the diffusivity of the limiting reactant in the biofilm  $(cm<sup>2</sup>/d).$ 

Consequently,

$$
\frac{d^{2}S_{f}}{dZ^{2}} = \frac{dS_{f}}{dt} \cdot \frac{1}{D_{f}} = -kX_{f}S_{f}/(D_{f} \cdot (K_{s} + S_{f}))
$$
\n(9)

Analytical solutions to this differential equation are not available except for limiting cases of the Monod equation where  $S_s \gg K_s$ or  $S_s \ll K_s$ . Where flux limitation exists, the solution must satisfy the following boundary equations for deep biofilms:  $S_f = S_g$  at Z = 0 (relative to the biofilm surface;  $Z = L_1 + L_2$  relative to the axes of Figure 14);  $dS_f/dZ = 0$  at  $Z = L_e$  (relative to the biofilm surface); and  $S_f = S_l$  at Z = L<sub>e</sub> (relative to the biofilm surface) where L<sub>e</sub> is the active biofilm thickness (cm) and  $S_t^{\dagger}$  is some limiting reactant concentration equal to a small fraction of the corresponding halfvelocity coefficient.

Williamson and McCarty (1976a and 1976b) demonstrated that the electron acceptor, oxygen, will be both the flux-limiting and metabolism-limiting reactant species if the following conditions are satisfied:

$$
S_{oa} < \frac{D_{fd} \cdot v_a \cdot MU_d}{D_{fa} \cdot v_d \cdot MU_d} S_{od}
$$
 (10)

 $S_{fa}$  < K<sub>sa</sub>  $S_{fd}/K_{sd}$ (11) where:

 $S_{0a}$  = bulk liquid oxygen concentration, mg/1  $S_{\text{odd}}$  = bulk liquid NH<sub>4</sub><sup>+</sup>-N, (substrate) concentration, mg/1  $D_{f_a}$  = diffusivity of oxygen in the biofilm, cm<sup>2</sup>/d  $D_{\text{f-d}} =$  diffusivity of NH<sub>A</sub><sup>+</sup>-N in the biofilm, cm<sup>2</sup>/d  $v_a$  = stoichiometric coefficient of oxygen in Eq. 5  $v_{\mathcal{A}}$  = stoichiometric coefficient of NH<sub>A</sub><sup>+</sup>-N in Eq. 5  $S_{f_a}$  = oxygen concentration at any depth in the biofilm, mg/1  $S_{fd} = NH_4^+ - N$  (substrate) concentration at any depth in the biofilm, mg/1  $K_{ca}$  = oxygen half-velocity coefficient, mg/1  $K_{sd}$  = NH<sub>4</sub><sup>+</sup>-N half-velocity coefficient, mg/1

 $MW_a$  = molecular weight of oxygen

 $MW_{d}$  = molecular weight of nitrogen

Parker and Merrill (1984) showed that BOD removal in a trickling filter plant at Malmo followed zero-order kinetics in the upper zone and first-order kinetics in the lower zone, as predicted by the equations of Williamson and McCarty. Richards and Reinhart (1986) obtained similar results for pilot-scale trickling filters at Atlanta, Georgia.

Using the Williamson and McCarty equations 10 and 11, Gullicks and Cleasby (1986) showed that the electron acceptor, oxygen, was both

the metabolism-limiting and the flux-limiting reactant for bulk liquid  $NH_A^+$ -N concentrations greater than 4 mg/L even when the bulk liquid was nearly saturated with dissolved oxygen. Therefore, in most domestic wastewater treatment trickling filter applications, oxygen will be the limiting reactant through a substantial portion of the media depth or even the entire media depth in some cases.

In those depths of the nitrification tower where the  $NH_4^+$ -N concentration is greater than 4 mg/L, the oxidation of  $NH_4^+$ -N will be a linear function of filter depth (zero-order with respect to  $NH_4^+$ -N concentration) if the bulk liquid oxygen concentration is assumed to be constant with filter depth. Nearly constant oxygen concentration with depth is probable if adequate air ventilation is provided so that the partial pressure of oxygen is essentially the same at all depths in the tower.

Atasi and Borchardt (1984) have summarized the bulk liquid substrate kinetics that would be observed for various biofilm kinetics and for partial or full penetration of the substrate in the biofilm. They concluded that zero-order substrate kinetics are observed in the bulk liquid when the intrinsic biofilm reaction rate is pseudo-zeroorder  $(K_s < S)$  and the substrate fully penetrates the biofilm.

Nitrification has been reported at D. 0. concentrations as low as 0.5 mg/1, but at much lower rates than at higher D. 0. concentrations (Gulp and Heim, 1978 and O'Shaughnessy et al., 1984). Wuhrmann (1960) demonstrated that activated sludge plants achieved higher nitrification efficiency at D. 0. concentrations of 4-7 mg/1 than at

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1 mg/1 when other operating conditions were similar. The Ministry of Technology (1965) demonstrated that the rate of nitrification in activated sludge plants was about 10 percent higher with D. 0. concentrations of 4-8 mg/1 than with a D. 0. concentration of 2 mg/1.

Schoberl and Engel (1964) used a polarographic laboratory technique and reported that growth of Nitrosomonas was oxygen-limited at D. O. concentrations of 0.9 mg/1 and that growth of Nitrobacter was oxygen-limited at D. 0. concentrations of 2 mg/1. They also found that the limiting D. 0. concentration increased as the concentration of nitrifier cells increased. This implies that limiting D. 0. concentrations in fixed-films may be higher than those in suspended growth reactors. Meyerhof (1917) and Ulken (1963) also found Nitrobacter to be more sensitive to oxygen depletion than Nitrosomonas.

Bulk liquid D. 0. concentrations are considerably higher than 0.5 mg/1 in nitrifying trickling filters with low organic loadings, but D. 0. concentrations in the biofilm, which have not been measured, could drop below 0.5 mg/1 before fully penetrating the biofilm. Brown and Caldwell (1980) reported improvement in nitrification in a biofilter when forced draft was increased.

Near-linear removal of  $NH_4^+$ -N with depth is evident in the literature for  $NH_4^+$ -N concentrations greater than about 4 mg/L (Buddies and Richardson, 1973, O'Shaughnessy et al., 1984, Huang and Hopson, 1974b, Wilderer et al., 1982, and Parker and Richards, 1985). Bruce et al. (1975) reported that ammonia mass removal rates in their nitrifying trickling filters did not change when the influent ammonia

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concentrations were increased from 20-40 mg/1 over a four week period. Gujer and Boiler (1984) reported that nitrification in a 138 cm dual media (expanded shale and sand) final filter was oxygen-limited at  $NH_A^+$ -N influent concentrations greater than 1.7 mg/l when the superficial velocity was 10 m/hr. Examples of zero-order bulk liquid  $NH_A^+$ -N kinetics observations are shown in Figures 16-19.

O'Shaughnessy et al. (1984) reported that the specific rate of  $NH_A^+$ -N uptake, k (mg N/mg VSS  $\cdot$  d), was best described by the following least squares regression equation:

$$
k = 0.947 (1 - exp (-1.35C))
$$
 (12)

where

k = specific rate of  $NH_4^+$ -N uptake for <u>Nitrosomonas</u> at 28°C. mg N/mg VSS • d C = concentration of  $NH_4^+$ -N, mg/1

Equation 12 predicts zero-order (constant  $k = 0.95 d^{-1}$ ) nitrification at  $NH_4^+$ -N concentrations greater than 4 mg/1.

If nitrification rates in fixed-film reactors at known  $NH_A^+$ -N and D. 0. bulk liquid concentrations are compared to the nitrification rate at some arbitrary reference condition, some rate-limitation conclusions can be established. Monod kinetics imply that  $NH_4^+$ -N bulk liquid concentrations should have a direct effect on nitrification



Figure 16. Effect of artificial ventilation and occasional flushing respectively on the nitrification efficiency. After Wilderer et al. (1982)

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Figure 17. Nitrifying tower nitrogen species profiles. After Duddles and Richardson (1973)



Figure 18a. Nitrifying tower nitrogen species profiles. After Parker and Richards (1985)

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Figure 18b. Nitrifying tower nitrogen species 'profiles. After Parker and Richards (1985)

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Figure 19. Correlation between influent and effluent ammonia concentration in a nitrifying dual media filter after a trickling filter. Above a removal of 1.7 mg NH<sub>4</sub>-N/l nitrification becomes oxygen limited (superficial velocity 10 m/h, total depth of filterbed 135 cm; dual media sand and expanded slate). After Gujer and Boiler (1984)

 $\sqrt{g_{\rm g}}/g_{\rm c}$  ,

rates unless oxygen or some other reactant is the rate-limiting reactant. This is true whether the process is flux-limited or metabolism-1imited.

Bulk liquid  $NH_4^+$ -N and D. O. reference concentrations of 15 and 5 mg/1, respectively, were selected in Table 2. Table 2 shows that near linear removal of  $NH_A^+$ -N with tower depth could only be observed if oxygen were the rate-limiting reactant. It cannot be discerned from bulk liquid  $NH_A^+$ -N concentration versus tower depth plots whether oxygen flux-limitation or oxygen metabolism-limitation is predominant.

The use of inhibition kinetics (the Haldane equation) for oxygen flux-limited conditions appears to offer little, if any, improvement over Monod kinetics. The Haldane inhibition kinetics offer some improvement for  $NH_{4}^{+}-N$  limitation (really  $NH_{4}^{+}-N$  inhibition) conditions at high bulk liquid  $NH_A^+$ -N concentrations (Rozich and Castens, 1986). However,  $NH_4^+$ -N concentrations high enough to cause inhibition are not likely to exist in domestic wastewater trickling filter nitrification processes. At low  $NH_4^+$ -N concentrations, Monod and Haldane kinetics predict essentially the same nitrifier growth rates for  $NH_A^+$ -N limitation conditions.

Effects of Operating Parameters on Nitrification in Trickling Filters

In most cases, the rate at which trickling filters can process substrate is controlled by the factors that control the flux of one or more reactant species. Flux of reactants is controlled by the following factors:

$NH_{4}$ <sup>+</sup> -N Conc. Bulk Liquid mg/l	D.O. Conc. Liquid $mg/1$	Rate of Areal NH <sub>4</sub> <sup>+</sup> -N Uptake	Implication of Observed Rate of Areal $NH_A$ <sup>+</sup> -N
15	5	Reference Level	Reference Level
20	5	a) No Change	Oxygen Limited
		b) Increase	Ammonium-Limited
15		a) Increase	Oxygen-Limited
		b) No Change	Limitation by Species Other Than O <sub>2</sub> or NH <sub>4</sub> <sup>--N</sup>

Table 2. Expected areal rate response for oxygen or substrate limitation conditions

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- 1. The biofilm area, thickness and density available for mass transport and utilization.
- 2. The concentration gradient of the reactants from the bulk liquid to the biofilm.
- 3. The coefficients of molecular diffusion for the reactants in the bulk liquid and in the biofilm.
- 4. The mean detention time of wastewater flow through the trickling filter.

Each of the above factors is controlled by one or more operating parameters. A number of additional operating parameters are important because they affect competition between the various microorganisms that inhabit the trickling filter. Table 3 lists the operating parameters that may affect the performance of nitrification trickling filter towers.

#### Wastewater temperature effects

Maier et al. (1967), Harris and Hansford (1976), Benzie et al. (1963), Antonie (1976), and others demonstrated that increasing the temperature of the wastewater feed causes greater substrate removal for the fixed-film process. This would be expected whether the process is flux-limited or metabolism-limited. The Nernst-Einstein equation for coefficients of diffusion adequately describes the effect of temperature.

$$
C = D_{\omega} \ \mu/T \tag{13}
$$
Operating Parameter		Effect on Nitification Rate, mass/time	<b>Effect on Percent</b> <b>Removal</b>	Factors in Observed Effects
1.	Wastewater tem- perature	Rate increase with temperature increase	Increase with tem- perature increase	Diffusivity coeffi- cients, biofilm activity
2.	Hydraulic loading rate	Rate increase with hydraulic loading increase	Decrease with hydraulic loading increase	Liquid diffusional resistance, hydrau- lic retention time, biofilm surface activity, scouring, percent active bio- film area
3.	Ammonium concen- tration	Rate increase for greater ammonium concentration up to 4 mg/1	Decrease with greater ammonium concentration	Substrate diffusional driving force
4.	Suspended solids	Rate decrease for greater suspended solids	Decrease with greater suspended solids	Competition for sub- strate by adsorption. percent active biofilm area
5.	Alkalinity	Rate decrease for alkalinity less than 1 meg/1	Decrease with alkalin- ity less than $1 \text{ meq}/1$	pH buffering
	$6.$ pH	Rate decrease for decreasing pH. $0$ pti- mum $pH$ 8.3	Decrease with decreas- ing pH	Biofilm activity
7.	Toxicants	Rate decrease	Decrease	Biofilm activity

Table 3. Effects of operating parameters on nitrification in trickling filters

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 $\label{eq:2} \frac{1}{\sqrt{2}}\sum_{i=1}^n\frac{1}{\sqrt{2\pi\sigma_i^2}}\int_{\mathbb{R}^n} \frac{1}{\sqrt{2\pi\sigma_i^2}}\frac{1}{\sqrt{2\pi\sigma_i^2}}\frac{1}{\sqrt{2\pi\sigma_i^2}}\frac{1}{\sqrt{2\pi\sigma_i^2}}\frac{1}{\sqrt{2\pi\sigma_i^2}}\frac{1}{\sqrt{2\pi\sigma_i^2}}\frac{1}{\sqrt{2\pi\sigma_i^2}}\frac{1}{\sqrt{2\pi\sigma_i^2}}\frac{1}{\sqrt{2\pi\sigma_i^2}}\frac{1}{\sqrt{2\pi\$ 

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where:

where:

- $C = constant$
- $D_{\text{tot}}$  = molecular diffusion coefficient in wastewater, cm<sup>2</sup>/s
- $\mu$  = dynamic viscosity of wastewater, g/cm  $\cdot$  s
- $T =$  absolute wastewater temperature,  $\alpha$ K

Molecular diffusion coefficients are seen to be directly proportional to the absolute temperature and inversely proportional to the dynamic viscosity. The diffusion coefficients increase as the temperature increases leading to an increase in the removal of substrate.

The mean detention time is also a function of the viscosity. The Nusselt equation for laminar flow down an inclined plane demonstrates that mean detention time must decrease at higher wastewater temperatures since velocity of flow increases at lower viscosities.

$$
\delta = \left(\frac{3V\mu}{\rho g \sin\theta}\right)^{1/3} \alpha \frac{V}{W \cdot velocity}
$$
 (14)

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 $V = volumetric$  flow rate,  $cm^3/s$  $\mu$  = dynamic viscosity, g/cm • s  $p =$  density of water, g/cm<sup>3</sup>  $g =$  acceleration of gravity,  $cm/s<sup>2</sup>$  $\delta$  = bulk liquid film thickness, cm  $\theta$  = angle of inclination from horizontal

w = width of plane, cm

For a constant mass loading to the system, increasing the wastewater temperature results in a reduction in the time for mass transfer and disregarding the effect on diffusion coefficients, the substrate mass removal and percent removal should decrease. Of course, the increase in flow velocity also decreases the effective liquid diffusion layer thickness  $(L_1 + L_2)$  of Figure 14), thus increasing the concentration gradient from the bulk liquid to the biofilm. The detention time reduction and the concentration gradient increase have opposing effects on the removal of substrate. Cleasby (1985) has shown that consideration of temperature effects on diffusion and detention time, while neglecting concentration gradient effects, adequately predicted substrate removal for data from the inclined plane, fixed-film study of Maier et al. (1967).

The net result of temperature-related diffusion coefficient and flow velocity effects on substrate removal can be summarized as follows:

At greater wastewater temperatures, the flat plate substrate removal increases predicted from diffusion coefficient increases are reduced approximately in proportion to the decrease in mean detention time. The more irregular biofilm surfaces associated with corrugated plastic media may reduce the wastewater temperature effects on mean detention time.

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The effects of wastewater temperature, influent substrate concentration, and hydraulic loading rate on mass removal of substrate, percent removal, bulk liquid thickness, bulk liquid velocity, and detention time have been clearly summarized by DeBoer (1984) and Cleasby (1985) in Tables 4 and 5 using data from the glucose removal, inclined plane biofilm study of Maier et al. (1967). It is important to clarify that bulk liquid film thickness is not synonymous with boundary layer thickness and that the majority of the resistance to mass transfer in the bulk liquid is associated with the boundary layer located along the biofilm surface.

#### Substrate loading rate effects

The most important operating variable from the plant operator's perspective is the substrate loading rate. The substrate loading rate can be varied by changing either the hydraulic loading rate or the influent substrate concentration, or both.

Harris and Hansford (1976), Sarner (1984 and 1986), and Maier et al. (1967) demonstrated that increasing either the hydraulic loading rate or the influent substrate concentration (at concentrations equal to or below the value at which oxygen is limiting) increased the mass removal per unit area per unit time. However, the percent removal efficiency was reduced. At all substrate concentrations in the oxygen-limitation region the rate of substrate removal was nearly constant for any constant hydraulic loading rate (Harris and Hansford,

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Table 4. Analysis of data from Maier flat plate biofilm study DeBoer (1984) . After

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Influent Glucose Concentration mg/1	<b>Flow</b> Rate $ml/10$ min	Observed Ratio of removal at $37.3^{\circ}/10.6^{\circ}$	Predicted Ratio <sup>a</sup> of removal at $37.3^{\circ}/10.6^{\circ}$
27.5	300	1.40	1.71
	600	1.25	1.64
65.5	300	1.58	1.71
	600	1.76	1.64
127.5	300	1.61	1.71
	600	2.00 1.60 Avg.	1.64 1.67 Avg.

Table 5. Observed and predicted effects of temperature on removal using the data from Table 4. After Cleasby (1985)

 $a$ Predicted from Diffusion Coeff (37.3°/10.6°) x Mean Detention (37.3°/10.6°).

 $\mathcal{A}^{(k)}$  and  $\mathcal{A}^{(k)}$ 

1976). The oxygen-limited constant rate of substrate removal was zero-order with respect to the substrate concentration.

If the substrate or oxygen concentration in the bulk liquid is increased, the concentration gradient from the liquid to the biofilm increases; therefore, the mass transport of substrate or oxygen to the biofilm increases. If the hydraulic loading rate is increased, the boundary layer thickness associated with laminar flow at the 1iquid-biofiIm interface is reduced. The effect is equivalent to an increase in the concentration gradient from the liquid to the biofilm if the liquid concentration is held constant. The increased gradient increases the mass transport of substrate or oxygen to the biofilm.

In addition, increasing the hydraulic loading rate increases the wetted surface area of the media providing more biofilm area for mass transport and utilization (Onda et al., 1968, and Crine, 1986). Crine (1986) predicted that an upper limit, less than complete wetting, exists for wetted area.

Brown and Caldwell (1980) defined the minimum wetting rate as "the minimum total hydraulic loading to ensure complete wetting of the media surface, which allows the media to be fully effective in biological treatment." Manufacturers typically recommend minimum wetting rates of  $0.34-0.55$   $1/m^2$   $\cdot$  s for plastic media to promote good wastewater flow distribution to the media. Data correlating wetted surface area with hydraulic loading rate are lacking. An upper limit on wetted surface area at typical minimum wetting rates is unlikely

based on air stripping data (Onda et al., 1968). An upper limit on wetted surface area at higher hydraulic loading rates cannot be ruled out, however.

Recycle causes a complex interaction of the factors that affect the substrate loading rate. Parker and Richards (1985) and Brown and Caldwell (1980) reported that recycle was beneficial to wastewater treatment in trickling filters. Germain (1966), Richards (1984), Richards and Reinhart (1986), and Albertson and Davies (1984) reported that recycle has little effect on percent removal efficiency. Albertson and Davies (1984) presented data from pilot studies at Huntingdon, Pennsylvania and the Chino Basin, California and concluded that higher wetting rates rather than recirculation accounted for the better performance of the trickling filter processes.

It is logical that recycle should have little effect on treatment efficiency if the opposing effects of the reduced bulk liquid concentrations versus the decreased boundary layer thickness, increased wetted area, and improved biofilm shearing and flushing are considered. Nitrification may be improved by the dilution of BOD concentrations in the influent which would reduce competition for oxygen available in the biofilter for nitrification (Culp and Heim, 1978). In addition, the reduction in the retention time in each pass through the tower is offset by the additional passes through the tower so that the total time opportunity for mass transfer is similar to that without recycle.

Recycle has been credited with reduction of toxic materials or inhibitors and dampening of shock loadings (Oleszkiewicz, 1980). Recycle may be beneficial at low hydraulic loadings to maintain the minimum wetting rate and for severe oxygen limitation situations. Recycle reduces psychoda fly and clogging problems by improving the flow distribution at low influent hydraulic loads. Benzie et al. (1963) and Benzie (1969) showed that recycle may be detrimental in winter climates because of wastewater cooling. They reported that the efficiency of trickling filters was reduced when the air and sewage temperatures were equal and that recirculation helped eliminate this drop in efficiency.

A recirculation design offering flexible operation may be beneficial for achieving denitrification. Gujer and Boiler (1984) reported that summer nitrite concentrations in nitrifying wastewater treatment process effluents are higher than winter nitrite concentrations because of the relative growth rates of Nitrobacter and Nitrosomonas. The growth rate of Nitrobacter compared to that of Nitrosomonas is higher in winter than in summer. Nitrite concentration data of Ouddles and Richardson (1973) were in agreement.

Nitrite can be as toxic to aquatic life as ammonia. Therefore, summer designs of nitrifying fixed-film processes should consider effluent nitrite concentrations, and may be controlled by effluent nitrite concentrations. Wanner and Gujer (1984) predicted that denitrification in nitrifying fixed-film processes was possible only when recycle allowed conditions in the upper part of the media to be

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favorable. Young (1974) supported this position and cited the opposing environmental conditions necessary for nitrification and denitrification.

Conditions are favorable when oxygen concentrations and pH deep in the film are low, oxidized nitrogen is present, and organic substrate is present. This would imply that recycle in separate-stage nitrification should direct effluent from the nitrifying biofilter back to the top of the carbon oxidation biofilter during summer operation.

Brown and Caldwell (1980) reported that recirculation increased the bulk liquid D. 0. concentration of the trickling filter influent. However, they could not verify whether an increase in recirculation, an increase in forced draft, or a combination of these factors was responsible for the observed improvement in nitrification performance at Stockton, California. They reported short circuiting of final clarifiers to be a problem during morning hours of winter operation.

The short circuiting was determined to be caused by temperature gradients in the clarifiers. The forced draft convection operation was modified so that fewer fans were operated during the cooler night hours and additional fans were operated during the day hours to minimize temperature fluctuation in the trickling filter effluent.

Johnson and van Durme (1986) reported that recirculation of biological sludge through biofilters resulted in oxygen-limitation. The reported rate of soluble BOD removal increased with increased oxygen transfer rate, which increased with increased total hydraulic loading

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rate. They reported the oxygen transfer rate to be 0.0031 kg  $0<sub>2</sub>/d$  . m' of media surface area \* m of media depth/unit wetting rate above  $0.5 L/m^2$  • s.

Most of the pilot-scale trickling filter studies conducted in the past have used continuous or nearly continuous dosing of the media with wastewater rather than the periodic dosing typical of large rotary distributors used in most full-scale trickling filters. Lumb and Barnes (1948), Tomlinson and Hall (1955), Craft and Ingols (1973), Richards (1984), Richards and Reinhart (1986), Hutchinson (1975), Cook and Katzberger (1977), and Bruce and Merkens (1973) have reported that periodic dosing provided better trickling filter performance. Improved performance was most pronounced at low hydraulic loading rates.

Periodic dosing reduces the retention time in the media, and a decrease in the removal of substrate would logically be expected. However, it has been observed that periodic dosing and continuous dosing with identical media and average hydraulic, loading rates provided essentially.equal treatment (Richards and Reinhart, 1986). thus, decreased retention time must be compensated for by other factors.

These factors are likely to include the decrease in resistance to diffusion of reactants into the biofilm due to the reduction in the liquid-biofiIm boundary layer thickness and/or the increased bulk liquid turbulence, the additional shearing of the biofilm resulting in a more active exposed biomass, improved distribution of wastewater

flow, and better oxygen transfer from the bulk liquid to the biofilm. Richards and Reinhart (1986) and Sarner (1984 and 1986) pointed out that liquid retention time and biofiIm-reactant contact time are not synonymous since substrate-biofiIm sorption may be significant and that contact time may not be significantly changed when retention time decreases. They failed, however, to recognize and account for the potential increase in the effective biofilm surface area that periodic dosing would be expected to produce. The increase in the effective biofilm surface area is likely to be the single largest contributor to the observed performance improvement of periodic dosing versus continuous dosing.

## Competition between microorganism species

Wilderer et al. (1982) showed that nitrification in trickling filters can be upset by inadequate consideration of competition between microorganism populations within the media. They demonstrated that separate biological carbonaceous and nitrogenous BOD treatment processes operated in series favor establishment of a succession of biocommunities for effective organic carbon removal and nitrification. Wanner and Gujer (1984) reached the same conclusion and showed that carbon oxidizing heterotrophs quickly overgrow nitrifiers in trickling filters if carbonaceous BOD loadings are not low. Wild et al. (1971) reported that instantaneous, short-term increases or decreases in BOD loading did not affect the rate of nitrification in activated sludge

plants. Similar responses to instantaneous short-term BOD loading changes would be expected for trickling filter plants.

The purpose of the carbonaceous BOD removal stage is conversion of organic carbon into biomass and subsequent separation of the biomass before the nitrification stage. Wanner and Gujer (1984) found that when competition for a common reactant existed, the faster growing species may overgrow the slower growing species at high substrate concentrations and may be washed out of the biofilm at low substrate concentrations. Carbonaceous BOD and biomass carryover to the nitrification stage promotes establishment of heterotrophs and protozoa populations in the nitrification stage. Nitrification begins only when carbonaceous BOD and biomass in the liquid phase have been reduced to levels where the nitrifying bacteria population is favored by the filter environment. Figures 3a and 3b (Parker and Richards, 1985) demonstrated that nitrification did not begin until the soluble  $BOD<sub>5</sub>$ concentration fell below 20 mg/L.

Wanner and Gujer (1984) found that the faster growing species tend to exist near the biofilm surface and that the slower growing species tend to exist at greater depths in the biofilm. Thus, slow growing species can be maintained in the biofilm even when total biomass growth rates exceed the maximum growth rate of the slow growing species. This no doubt influences the relative stability observed for trickling filter processes with loading variations. However, at long-term, high organic substrate concentrations, nitrifier populations may be totally

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displaced from the biofilm. Wanner and Gujer (1984) predicted that nitrifiers could be totally displaced from the biofilm at soluble organic substrate concentrations greater than 27 mg COD/1.

### Particulate effects

Sarner (1981, 1984, and 1986) and Zimmerle et al. (1982) reported that soluble substrate removal in trickling filters is impeded by suspended or colloidal matter in the liquid feed. Sarner (1984 and 1986) suggested that particulate matter produced in the upper depths of trickling filters may be a factor in the lower removal rates observed in the lower depths of trickling filters. Suspended and colloidal matter may adhere to the biofilm surface reducing the active area for mass transport and exerting additional oxygen demand. Particulates may compete with the microorganisms for the reactants through adsorption, subsequently releasing the reactants to the effluent by desorption.

The importance of intermediate clarification between the carbonaceous oxidation and nitrification stages is evident. However, in winter climates the cooling effects of clarification should be considered in process design. Wall and Peterson (1986) have developed a model for winter heat loss from uncovered clarifiers that may be of use in this evaluation.

## Ventilation effects

Proper ventilation of the media is necessary to maintain high dissolved oxygen concentrations in the bulk liquid phase. This is especially important if oxygen is the limiting species. Brown and Caldwell (1980) reported improved nitrification when forced draft ventilation was increased at the Stockton, California plant.

Most trickling filters rely on natural convection and whatever small benefit the wind may provide, in the case of uncovered filters, to cause air movement through the biofilter. The use of covers for trickling filters in cold climates is increasing; therefore, wind effects in cold climates are often absent. Convection currents may be either upward or downward depending on the relative temperatures of the air and the wastewater. The Recommended Standards for Sewage Works (1978) recommended that underdrains should flow not more than 1/2 full at peak flow to provide free air circulation.

The manufacturers of modern plastic media have made many advancements in configuration and specific surface that have increased detention time, air flow, and available contact surfaces. Forced convection is usually not considered necessary except for very tall plastic media towers and towers containing relatively small random media. The benefits of artificial ventilation were evident in the work of Wilderer et al. (1982) with small rock media. Figure 16 clearly demonstrates the oxygen limitation of nitrification in trickling filters and the improvement of nitrification with artificial ventilation.

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#### Shock load attentuation

Cook and Herning (1978) reported that trickling filters have shock load attenuation capacities resulting from the harboring of "starved" bacteria at the lower depths. Discussion of their work by Greenfield and Moodie (1979) pointed out that the bacteria at lower depths were not necessarily starved, but that the bacterial response to shock loads could be explained by Monod kinetics. They showed that oversizing of trickling filters may not be a cost-effective means of providing shock load attenuation capacities. All trickling filters have shock load attenuation capacities except those obeying zero-order bulk liquid substrate kinetics (oxygen-limited) throughout the entire biofilm depth.

These observations have important implications. It can be concluded that shock attenuation capacities in nitrifying trickling filters may be provided by oversizing the biofilter so that nitrifier populations in the lower depths are not oxygen-limited and can respond according to Monod kinetics to shock loads. Alternatively, artificial ventilation can be provided so that oxygen-limited biofilter performance can be improved during shock loads. Recycle may be beneficial by increasing bulk liquid D.O. concentrations, by diluting any BOD shock load that may accompany the  $NH_4^+$ -N shock load, by increasing the wetted surface area, by shearing and flushing away less active biofilm, and by improving the mass transfer of reactants in the liquid phase.

Recycle may be detrimental by diluting the influent  $NH_4^+$ -N concentration so that oxygen-limited nitrification does not exist through the

entire biofilter depth. Total  $NH_A^+$ -N uptake rates are higher for oxygen-limited conditions than for  $NH_4^+$ -N limitation conditions. Recycle also can be detrimental by reducing wastewater temperatures during winter operations.

Randtke and McCarty (1977) conducted 24-hour studies of nitrogen and organic loading variations at three activated sludge plants. They found that the primary effluent peak hourly flow, ammonia concentration, ammonia mass loading, and soluble COD mass loading to be 123-139 percent, 136-177 percent, 151-240 percent, and 139-173 percent of the 24-hour average, respectively. They found the secondary effluent peak hourly flow, ammonia concentration and ammonia mass loading to be 123- 139 percent, 123-138 percent and 122-142 percent of the 24-hour average, respectively.

Randtke and McCarty (1977) concluded that peak nitrogen and organic substrate concentrations coincided with the peak hydraulic loads during the 24-hour diurnal variation observation period. Thus, maximum oxygen demand was being exerted at the time when hydraulic residence was at its minimum. Their comparison of secondary effluent versus primary effluent ammonia mass loading peaks provides evidence of the beneficial effects of flow equalization. Negligible change in ammonia concentration was observed in primary treatment.

Stenquist et al. (1978) presented data from several studies that showed that primary treatment had no effect on ammonia concentrations, but that it did reduce organic nitrogen concentrations somewhat. They

presented peak hourly ammonia loading versus peak hourly hydraulic load correlations similar to Randtke and McCarty.

Since nitrification is approximately zero-order with respect to bulk liquid  $NH_A^+$ -N through the majority of the tower depth, significant  $NH_A^+$ -N may reach the lower end of the tower only during peak load hours for properly sized biofilter towers. Biofilters, especially oxygenlimited ones, are not ideal for handling shock loadings. Therefore, shock load attenuation is better provided by forced draft variability (flexibility). Digester supernatant should not be recycled during peak load hours because of its significant  $NH_A^+$ -N content. Digester supernatant should be recycled during nonpeak hours.

The data of Duddles and Richardson (1973), Sampayo (1981), Sampayo and Metcalf (1984), and Baxter and Woodman, Inc. (1973) indicated relatively stable biofilter nitrification treatment performance during normal diurnal loading variation. These biofilters were not oxygen limited in the lower media depths during nonpeak hours. Intervals between peak hour loadings were evidently short enough to maintain nitrifier populations capable of treating the peak load. Seasonal shock loading variability may be of more concern. While response to diurnal loading fluctuations appears to be quite rapid, response to seasonal shock loads may require a substantial acclimation period.

## Inhibitory conditions

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The hydrogen ion concentration has a strong influence on the rate of nitrification. Nitrification rates decrease significantly as the

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pH drops into the acidic range. Figure 12 demonstrated the effect of pH on the nitrification rate. The pH range of 8 to 9 is considered ideal for nitrification. It is important that sufficient alkalinity (either natural or added) be available to prevent the pH from falling into the acidic range.

Painter (1970) published a literature review of inorganic nitrogen metabolism in microorganisms. It included a list of toxicants and the concentrations at which toxic effects were observed for nitrification processes. Vismara (1982) also has reviewed a number of nitrification toxicants. He found that the thresholds for toxic effects from chloride and magnesium were 10 and 50 mg/L, respectively. The threshold for toxic effects from ammonium nitrogen ranged from 436 to 1000 mg/L and was pH dependent.

#### Media characteristics

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Media usually fall into the following categories; 1) rock or slag, 2) wood lattice, 3) plastic random, 4) plastic vertical tubes, 5) corrugated plastic vertical-flow, and 6) corrugated plastic crossflow. Plastic media have been specified for most new trickling filter construction in the last ten to twenty years, and have been employed in rehabilitating older rock trickling filters to improve their performance. Media configuration and increased specific surface areas are the most significant developments to happen in trickling filter

technology in several decades. The new plastic media are light enough and strong enough to allow stacking to depths up to 10 m.

Porter and Smith (1979), Albertson and Davies (1984), Albertson and Eckenfelder (1984), Harrison and Daigger (1985), Richards (1984), Richards and Reinhart (1986), Parker and Merrill (1984), Parker and Richards (1985), Sarner (1984), and Drury et al. (1986) have studied the performance of the various types of media. Porter and Smith (1979) found that random media was more efficient than vertical-flow media and that vertical-flow media was more efficient than vertical tube media when performance was compared at similar hydraulic application rates per unit specific surface area. The majority of the investigators found that cross-flow media performed better than other types of media, including plastic vertical-flow media.

They have usually supported their conclusions based on a comparison of modified Velz equation, first-order  $k_{20}$  kinetic coefficient (treatability) values for carbonaceous BOD removal and on comparison of hydraulic residence times. The modified Velz equation is discussed in greater detail in the next section of this literature review. Hydraulic residence time has been described by the following empirical equation;

$$
t = C a_c D/q^n
$$
 (15)

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where

t = average hydraulic residence time, minutes.

 $q = \text{cross sectional hydraulic loading rate, gpm/ft}^2$ .

- $d =$  media depth. ft.
- n = dimensionless constant dependent on media properties and wastewater distribution.
- $a_{s}$  = specific surface area, ft<sup>2</sup>/ft<sup>2</sup>.
- $C =$  constant,  $(ga)/(ft^2)^n$  min<sup>1-n</sup>.

Adams et al. (1981) demonstrated the methodology appropriate for determination of  $k_{20}$  and n values. Liquid distribution at the top of the tower can affect the values observed. Values of  $n < 0.3$  imply that retention time is independent of flow rate, while n values > 0.7 imply dependence of retention time on flow rate (Albertson and Eckenfelder, 1984).

Cross-flow media were found to have longer hydraulic residence times and higher  $k_{20}$  values than vertical-flow media (Harrison and Daigger, 1985, Sarner, 1984, Albertson and Eckenfelder, 1984, Drury et al., 1986, Parker and Merrill, 1984, Richards, 1984, Richards and Reinhart, 1986). Vertical media were found to have small n values indicating that the hydraulic residence time in vertical media is somewhat insensitive to hydraulic load (Harrison and Daigger, 1985, Richards, 1984, and Sarner, 1984). Cross-flow media n values were higher indicating that hydraulic residence times would be significantly longer at lower hydraulic loading rates (Harrison and Daigger,

 $\mathcal{F}_\mathbf{c}$  and

1985, Richards, 1984, and Sarner, 1984). Rock and random media were found to have very high n values, explaining why they performed well at low hydraulic loading rates, but performed poorly at high hydraulic loading rates (Harrison and Daigger, 1985, Richards, 1984, and Sarner, 1984).

Over the normal hydraulic loading range cross-flow media have generally been shown to have longer hydraulic residence times than vertical media. Harrison and Daigger (1985) have shown that n values for clean plastic media are significantly higher than n values for media containing biofilm growth. Thus, hydraulic residence times in trickling filters with established biofilms are less sensitive to hydraulic loading. Balakrishnan and Eckenfelder (1969) found that biofilms could increase retention time by 200-500%, but that n values for clean media and for media containing biofilm were not significantly different.

Researchers have attributed the superior performance of cross-flow media compared to other media to: 1) improved distribution of the wastewater and better wetting properties at lower flow rates, 2) longer hydraulic retention times, and 3) better oxygen transfer characteristics. Richards (1984) and Richards and Reinhart (1986) showed that oxygen transfer efficiency alone could not explain the difference in performance between cross-flow and vertical media, but they acknowledged that it could play a significant role in treatment of high strength wastewaters.

Most researchers have recognized that the clean media specific surface area and the effective biofilm area are not the same entities.

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Furthermore, they generally recognize that the effective biofilm area must vary with hydraulic loading rate. Unfortunately, most equations, models and curves available for predicting trickling filter performance use the assumption of a constant biofilm area equal to the clean media specific surface area.

This assumption can lead to erroneous conclusions relating to hydraulic retention time and the effects of dosing as previously discussed. In addition, it can lead to erroneous values for other constants in the design equations. This assumption has been defended by citing the lack of data and understanding in predicting the effective biofilm area with varying hydraulic loading rate.

Onda et al. (1968) reported the following equation to predict the. effective surface area for air stripping towers:

$$
\frac{a_{w}}{a_{t}} = 1 - \exp \left[-1.45 \left(\frac{\sigma c}{\sigma}\right)^{0.75} \left(\frac{L}{a_{t} \mu_{L}}\right)^{0.1} \left(\frac{L^{2} a_{t}}{\rho_{L}^{2} g}\right)^{-0.05} \left(\frac{L^{2}}{\rho_{L} \sigma a_{t}}\right)^{0.2} \right] (16)
$$

 $\sqrt{S_{\rm{B}}}/T_{\rm{B}}$ 

where:  $a_{\mu}$  = wetted specific surface area, m<sup>2</sup>/m<sup>3</sup>  $a_+$  = total specific surface area, m<sup>2</sup>/m<sup>3</sup>  $\sigma_{c}$  = critical surface tension of the packing material, kg/hr<sup>2</sup>  $\sigma$  = surface tension of the liquid, kg/hr<sup>2</sup> L = superficial mass velocity of liquid,  $kg/hr \cdot m^2$  of tower cross section

 $\mu_1$  = dynamic viscosity of the liquid, kg/m • hr

 $q =$  gravitational acceleration m/hr<sup>2</sup>

 $p_1$  = liquid density, kg/m<sup>3</sup>

This equation has been shown to have an accuracy of ±20 percent for randomly packed towers. This investigator correlated the equation's performance for the air stripping of trichloroethylene on cross-flow plastic media with specific surface areas of 223-403 m'/m'. The observed  $k_1$  a data of Amy and Cooper (1986) were plotted versus the predicted  $k_1 a_\omega^*$ . An R-squared value of 0.84 was found for the relationship. Figure 20 presents the ratio of the wetted specific surface area and the total specific surface area versus hydraulic loading rate and wastewater temperature as predicted from the Onda equation for two media with different specific surface.areas.

The predicted wetted specific surface areas are in excellent agreement with the upper part of the range suggested by Hosono et al. (1980) and with the lower part of the range suggested by Gullicks and Cleasby (1986) based on the oxygen-limited work of Maier et al. (1967). Figure 20 clearly demonstrates that hydraulic loading rate has a great influence on the wetted specific surface area and that wastewater temperature has a lesser impact on the wetted specific surface area.

Mills and Dudukovic (1981) proposed the following equation to predict the external wetting efficiency for conditions typical of random media (< 0.004 m in diameter) trickle flow operations;

$$
\eta = \tanh \left[ 1.09 \left( \frac{L}{a_t \mu_L} \right)^{0.28} \left( \frac{L^2 a_t}{\rho_L^2 g} \right)^{0.21} \left( \frac{L^2}{\rho_L \sigma a_t} \right)^{-0.16} \left( \frac{6(1-\epsilon)}{\epsilon} \right)^{0.03} \right] (17)
$$

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Figure 20. Wetted specific surface area based on Onda's correlations

where:  $\varepsilon$  = media bed porosity

 $\eta$  = external wetting efficiency =  $a_w/a_t$ 

all other terms are the same as those in Eq. 16.

Crine (1986) reported that the Mills-Dudukovic equation is consistent with percolation theory analyses of trickle flow operations. Figure 21 shows that external wetting efficiencies predicted by equation 17 are approximately twice as great as Onda equation predictions. Media specific surface area had little effect on predicted wetting efficiencies. Maximum external wetting efficiencies of approximately 80% are predicted by equation 17 at normal trickling filter hydraulic loading rates.

The minimum liquid wetting velocity for which stable liquid film trickling over a solid surface does not exist can be predicted from percolation theory (Crine, 1986). At liquid velocities less than the minimum wetting velocity, isolated rivulet flow occurs. Crine (1986) reported the following relationship to predict the minimum liquid wetting rate:

$$
\frac{a_w}{a_t} = \left(\frac{Re_t}{Re_M}\right)^{1 - a_w/a_t}
$$
 (18)

where:  $Re_{\parallel}$  = Reynolds number at the liquid velocity applied.

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 $Re^M$  = Reynolds number at the minimum liquid wetting velocity. Albertson and Eckenfelder (1984) showed that biofilm thickness could affect the aerobic surface area. They demonstrated that a 4 mm thickness of biofilm would cause a 12 percent reduction of surface area



Figure 21. External wetting efficiency based on percolation theory



in cross-flow media with a total specific surface area of  $98 \text{ m}^2/\text{m}^3$ . Surface area loss due to biofilm growth is dependent on media configuration as well as biofilm thickness. Albertson and Eckenfelder (1984) predicted that media with the highest proportions of converging sheets (i.e., narrow opening, high sheet and mixing point densities) would lose more surface area than media with low proportions of converging sheets. Of course, excessive biofilm buildup at mixing points or sheet convergence points would result in even greater loss of effective surface area by partially or completely plugging access to a portion of the media sheets.

### Trickling Filter Models

Velz (1948) and Germain (1966) established the use of first-order kinetics in the design of trickling filters for the removal of carbonaceous BOD. The Velz equation was modified by Schulze (1960) to include specific surface area and temperature effects. Total substrate concentrations were used in the original first-order models; however, current practice is to use soluble substrate concentrations (Albertson and Eckenfelder, 1984 and Harrison and Daigger, 1985). The Schulze equation can then be defined as follows:

$$
S_e = S_i \exp\left[-k_{20} a_s \ D \theta \frac{(T-20)}{q}\right]
$$
 (19)

where:  $S_i$  = soluble substrate concentration in influent wastewater flow without recirculation, mg/1

> $S_e$  = soluble substrate concentration in settled effluent, mg/l  $Q = Influent$  wastewater flow without recirculation, gpm/ft<sup>2</sup>

> > $\mathcal{P}^{\mathcal{C}}_{\mathbf{r}}$  ,  $2^{n-1}$

 $D =$  media depth. ft.  $\theta = 1.035$  $T =$  wastewater temperature,  $^{\circ}C$  $a^{\prime}_{s}$  = media specific surface area, ft<sup>2</sup>/ft<sup>3</sup> n = dimensionless hydraulic rate constant  $k_{20}$  = soluble substrate treatability coefficient @ 20°C.  $(gpm/ft^2)^n$ 

The Schulze equation does not include recirculation. The modified Velz equation, which includes recirculation, has gained widespread acceptance in carbonaceous trickling filter design. The modified Velz equation is defined below:

$$
S_e = S_b \exp\left[-k_{20} a_s \, b \, (T-20)/\rho_T n\right] \tag{20}
$$

where:  $S_{\rho}$  = soluble substrate concentration in settled effluent, mg/l  $S_b$  = soluble substrate concentration applied =  $(S_i + S_gR)/(R+1)$ , mg/1

 $R =$  recycle ratio = recycle hydraulic load/influent (feed) hydraulic load

 $Q_T$  = total applied hydraulic load, gpm/ft<sup>2</sup>

Other symbols were defined for equation 19.

The validity of first-order modeling of substrate removal in trickling filters is questionable. The  $k_{20}$  values used in the above equations are not independent of media depth as often assumed. Albertson and Eckenfelder (1984) and Albertson and Davies (1984) showed that

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 $k_{20}$  values must be converted to a standard media depth. They recommended the following equation for this conversion:

$$
k_{20} = k_D (D/20)^{0.5}
$$
 (21)

where:

 $k_{20}$  = the treatability coefficient for 20 ft. media depth.

 $k_{\text{D}}$  = the treatability coefficient for media depth, D.

Rittmann (1984) reported that the apparent first-order reaction coefficient was a function of the influent substrate concentration when detention time was held constant. An increase in the substrate concentration yielded a decrease in the apparent first-order reaction coefficient. The apparent first-order reaction coefficient was also dependent on the hydraulic load or detention time when the influent con-, centration was held constant. An increase in the detention time caused a decrease in the apparent first-order reaction coefficient.

Richards and Reinhart (1986) reported that the apparent first-order reaction coefficient was dependent on the media characteristics, including media configuration. Furthermore, k<sub>20</sub> values for nitrification have not been studied extensively, and data scatter, even for carbonaceous BOD removal, has been quite significant due to the variability of wastewaters treated.

Detention time studies have been conducted on many of the new plastic media (Sarner, 1984, Harrison and Daigger, 1985, Albertson and Davies, 1984, Richards and Reinhart, 1986, and Parker and Merrill,

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1984). These studies have helped to explain the differences in performance of the media and have established that the n values in the equations above depend not only on media configuration, media depth, hydraulic load, and initial distribution, but also on the nature of the biofilm established on the media (Harrison and Daigger, 1985).

#### Williamson and McCarty model

Williamson and McCarty (1976a and 1976b) set up iterative computer solutions to equate the flux of the limiting reactant through the liquid phase with the flux of the limiting reactant into the biofilm. They published their solutions in graphical form allowing the use of the model without computer assistance. Their model is based on Monod kinetics and is applicable to either substrate or oxygen diffusion limited cases.

The primary drawbacks to the model are the uncertainties associated with the Monod constants and coefficients, the biofilm density, and the liquid phase effective diffusion layer thickness and the time-consuming nature of the graphical iterative solution which must be applied at numerous depths in the trickling filter. The computer solution eliminates the time-consumption disadvantage of this model. Suidan (1986) published a similar dimensionless graphical solution for plug-flow, deep biofilm reactors.

# Rittmann and McCarty models

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Rittmann and McCarty (1978) proposed a variable order model based on Monod kinetics and using dimensionless variables. At low bulk liquid

substrate concentrations (much less than the appropriate half velocity coefficient) the model is first-order with respect to the substrate concentration. At high bulk liquid substrate concentrations (much greater than the appropriate half velocity coefficient) the model is one-half order with respect to the substrate concentration. For other bulk liquid substrate concentrations, the kinetics range from one-half to first order.

Rittmann and McCarty (1980b) reported that organic materials may persist in low concentrations  $(S^{\dagger}_{min})$  because insufficient energy is available to sustain microorganisms. Steady-state biofilm thickness approached zero as the bulk liquid substrate concentration approached  $\mathsf{s}_{\textsf{min}}$ . For bulk liquid concentrations of the limiting substrate between S<sub>min</sub> and 3.2 mg/l, Rittmann and McCarty (1980a) recommended their steady-state biofilm kinetics model.

Rittmann and Dovantzis (1983) reported that dual substrate nitrification rate limitation would occur within the  $NH_{\text{A}}^{+}-N$  interfacial concentration range of 1 to 5 mg/1 for an interfacial D.O. concentration of 2 mg/1. Available bulk liquid D.O. concentration data (Brown and Caldwell, 1980) suggest that higher D.O. concentrations exist. Therefore,  $NH_A^+$ -N will be the single limiting reactant only when the bulk liquid NH<sub>A</sub><sup>+</sup>-N concentration approaches 1 mg/l; and oxygen will be the single limiting reactant for all  $NH_4^+$ -N bulk liquid concentrations greater than about 5 mg/1.

These models share many of the drawbacks of the Williamson and McCarty model relating to the uncertainty of constants and coefficients.

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They are, however, readily amenable to simple computer or hand calculator solution. The models are not applicable to oxygen-limited cases which comprise the majority of fixed-film nitrification applications. Harremoes zero-order model

Harremoes (1982) developed a simplified mixed microbial film nitrification model using zero-order kinetics in the biofilm for both the carbon oxidizing microbes and the nitrifiers. The population of nitrifiers was assumed to increase exponentially with increasing depth into the biofilm, giving rise to a nonuniform ammonia removal per unit volume of biofilm. Nitrifiers have a small growth rate when compared to the growth rate of heterotrophs which oxidize carbon. Therefore, nitrifiers are quickly submerged beyond the penetration depth of oxygen in biofilms growing fast on carbonaceous substrate.

Harremoes showed that nitrification in fixed-film systems is nearly always oxygen flux-limited. Nitrification in fixed-films was represented by two potential substrate profile cases: Case 3, in which only oxygen partially penetrated the biofilm, and Case 6, in which only ammonia fully penetrated the biofilm. For these cases, Harremoes showed that the removal rate of nitrogen per unit surface area can be predicted from the following set of equations. Appropriate parameter estimates have been included from a variety of sources.

Case 3 
$$
B_0 < B_s
$$
  
\n $(1 - B_0) e^{-B_0} = 1 - B_0^{*2} (1 - \frac{B_0^{*2}}{B_0^{*2}})$  (22)

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Case 6  $B_0 > B_s$ 

$$
(1 - Bo) eBo = 1 - Bo*2 (1 - Cs2)
$$
 (23)

 $r_{an}$  = ammonium removal rate per unit surface area

$$
= (e^{B_0} - 1) \left(\frac{\chi *_{V_f}}{\gamma_n}\right) \tag{24}
$$

where:

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$$
B_{s} = \frac{\mu L_{s}}{v_{f}} \qquad B_{o} = \frac{\mu L_{o}}{v_{f}} \qquad B_{o}^{*} = \left(\frac{v_{n} \mu D_{o} S_{o}^{*}}{f_{n} \chi^{*} v_{f}^{*}}\right)^{\frac{1}{2}}
$$

$$
B_0' = \left(\frac{2D_0 S_0 * \mu^2}{f_s K_s f^V f}\right)^{\frac{1}{2}} L_s = \left(\frac{2D_s S_s * \frac{1}{2}}{K_s f}\right)^{\frac{1}{2}}
$$

$$
C_s^2 = \frac{f_s D_s S_s^*}{D_s S_0^*} \qquad v_f^2 = \frac{Y_s K_s f L_s}{\gamma}
$$
  
\n
$$
S_s^* = \text{bulk liquid organic carbon concentration, g soluble } B00/m^3
$$
  
\n
$$
S_0^* = \text{bulk liquid dissolved oxygen concentration, g/m}^3
$$
  
\n
$$
D_o = \text{diffusivity of oxygen,} \qquad 15 \times 10^{-10} \text{ m}^2/\text{s, } 67\% \text{ (range 35–80%) D} \qquad \text{in water Chen and Bungay (1981)}
$$
  
\n
$$
D_s = \text{diffusivity of organic matter, } 5.8 \times 10^{-10} \text{ m}^2/\text{s, glucose } \text{Maier et al. (1967)}
$$
  
\n
$$
f_s = \text{stoichiometric ratio of oxygen to B0D, 0.5 g O}_2/\text{g B0D} \qquad \text{Harremoes (1982)}
$$

$$
f_n = \text{stoichiometric ratio of oxygen to NH_4^+ - N. 4.24 g O_2/gN}
$$
\n
$$
\text{Parker et al. (1975)}
$$

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 $K_{\rm sf}$  = zero-order specific rate of organic substrate oxidation in the biofilm, g BOD/m' • s

$$
\gamma
$$
 = biofilm density, 4 x 10<sup>4</sup> to 5 x 10<sup>4</sup> g VSS/m<sup>3</sup> William—  
\nson and McCarthy (1976a and 1976b)

 $K = \frac{sf}{st} = k_e$  = specific rate of organic substrate uptake in biofilm,  $1.5 \times 10^{-4}$  g BOD/g VSS • s, Williamson and McCarty (1976a and 1976b)

 $\mu$  = nitrifier growth rate, s<sup>-1</sup> = k<sub>n</sub>Y<sub>n</sub>

$$
k_n = \text{specific rate of NH}_4^+ - N \text{ uptake in biofilm, } 2.3 \times 10^{-5}
$$
  
g N/g VSS • s, Williamson and McCarthy (1976a and 1976b)

$$
Y_n
$$
 = yield coefficient for nitrifiers, 0.166 g VSS/gN,  
 Parker et al. (1975)

$$
Y_s
$$
 = yield coefficient for heterotrophs, 0.5 g VSS/g BOD  
 Parker et al. (1975)

$$
X^* =
$$
 specific nitrifier concentration at the biofilm surface, fraction of  $\gamma$ , say 5000 g VSS/m<sup>3</sup>

 $L_{0}$  = penetration of oxygen into biofilm

 $L_{g}$  = penetration of organic substrate into biofilm

For the most simplified condition when no organic carbon exists,  $S_{s}^{*} = 0$ , Harremoes showed that the rate of removal of nitrogen per unit area can be expressed in the form of a half-order solution (with respect to D.O. bulk liquid concentration) as follows:

$$
r_{\rm an} = 2^{0.5} B_0^* \left(\frac{v_f^{\chi*}}{\gamma_n}\right) = \left(\frac{2 D_0 S_0^* \chi^* (0.5)}{f_n \gamma_n}\right) \tag{25}
$$

Assuming a constant D.O. bulk liquid concentration and constant specific nitrifier concentration at the biofilm surface at all depths in the tower, the above equation implies zero-order  $NH_4^+$ -N response with tower depth. Harremoes found no data in the literature that could be used to verify his model, and he strongly urged research of the type reported in this dissertation.

#### Strand's combined carbon and ammonia oxidation model

Strand (1986) proposed a mathematical model for mixed microbial biofilms based on Monod kinetics and Fickian diffusion for organic carbon uptake and based on zero-order kinetics for oxygen and nitrogen uptake within the biofilm. The computer solution of the approximation of the nonlinear equation for organic carbon uptake provided definition of the organic carbon profile from the biofiIm-water interface into the biofilm depths. Nitrification was assumed to begin at the distance relative to the biofiIm-water interface where the heterotrophic and nitrifying organism growth rates were equal. Strand presented the model in graphical form making it simple to use.

The primary drawback to this model was its dependence on parameter values from the literature which exhibited wide variation in the reported ranges of the parameter values. The model adequately predicted the ammonium flux for rotating biological contactor data from four

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sources (Stratta and Long, 1981, Mueller et al., 1980, Watanabe et al., 1980, and Weng and Molof, 1974).

#### First-order media configuration model

Logan et al. (1986) incorporated media configuration differences directly into a model based on first-order microbial kinetics for substrate-limited trickling filters. Substrate removal differences for different media were due only to differences in media geometries rather than implying changes in microbial kinetics for different media. The finite difference model adequately predicted soluble BOD removal based on comparison with observed data from several studies.

The model can be adapted to use in oxygen-limited situations. In fact, even without incorporating the modifications necessary for oxygenlimited applications, the soluble BOD removal predicted by the model was in good agreement with oxygen-limited data reported by Hutchinson (1975).

The importance of liquid phase mass transfer in trickling filter performance was demonstrated by the successful performance of this model. Cross-flow media were shown to be superior to vertical-flow media because cross-flow media provided more frequent disruption (intermittent mixing) of the liquid film (and, therefore, the boundary layer of highest mass transfer resistance adjacent to the biofilm).

## EXPERIMENTAL

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#### Materials and Equipment

The trickling filter pilot plant used in this study was located at the Ames Water Pollution Control Plant (WPCP). A schematic diagram of the pilot-scale, separate-stage, trickling filter plant is shown in Figure 22. The pilot plant influent was the Ames WPCP secondary clarifier effluent (final effluent). Subsequent to June 16, 1986 this influent wastewater was pretreated to control carbonaceous oxygen demand and suspended solids loadings to the nitrification biofilter. The nitrification biofilter performance prior to June 16 may have been slightly impaired by carbonaceous oxygen demand and solids carryover from the Ames WPCP final clarifiers.

#### Pretreatment process

Ames WPCP secondary clarifier effluent was distributed continuously at 1.6 to 1.9  $\frac{\mu}{m^2}$  . s to the top of a 2.44 m high x 0.61 m x 0.61 m pretreatment biofilter containing 45-degree, cross-flow plastic media with a specific surface area of  $98 \text{ m}^2/\text{m}^3$ . Carbonaceous oxygen demand was reduced in this step. Wastewater samples were collected at the top, 1.22 m depth, and 2.44 m depth of the pretreatment biofilter. Clarification at approximately 40,740  $L/m^2$  · d followed the pretreatment tower to reduce suspended solids loadings on the subsequent separate-stage, pilot-scale, nitrifying biofilter.

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Figure 22. Ames WPCP pilot-scale separate-stage nitrifying biofilter schematic

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#### Separate-stage nitrifying biofilter

Subsequent to June 16, 1986, the pretreatment process effluent was pumped to a constant head tank above the nitrifying biofilter. Prior to that, secondary effluent had been used as the nitrifying biofilter influent. When recycle was in use, nitrifying biofilter effluent was pumped to a separate constant head tank above the nitrifying biofilter. Separate streams of pretreated wastewater and recycled nitrifying biofilter effluent flowed at constant, preset rates into a plexiglas dosing siphon which measured 0.30 m x 0.30 m in cross section.

The dosing siphon had a 0.10 m diameter x 0.10 m high bell and 0,05 m diameter discharge piping. The siphon breaker consisted of a weighted air intake mounted on a spring-stiffened, flexible tube. This allowed the siphon breaker to bounce when the siphon break occurred, which provided stable dosing cycles and reduced fouling of the air intake by biomass accumulation.

The dosing siphon allowed wastewater to be applied to the nitrifying biofilter on an intermittent basis rather than as continuous dosing. The operation of dosing siphons was discussed in detail by Babbitt and Baumann (1958). The cycle time and dosing volume of the dosing siphon were determined by the influent flow rate, the height of the siphon breaker, and the height of the discharge leg of the siphon trap.

During some data collection periods, the  $NH_4^+$ -N concentrations in the wastewater were supplemented to maintain bulk liquid concentrations above 4 mg/L throughout the nitrifying biofilter depth. A chemical feed pump, operated at constant flow rates, was used to add concentrated

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 $NH_{\Delta}$ Cl solution directly to the wastewater stream as it entered the dosing siphon. Turbulence from the stream(s) of wastewater entering the siphon tank and from the siphon tank discharge cycle provided good mixing of the  $NH_ACl$  and wastewater.

The discharge from the dosing siphon occurred in a short period of time (approximately 8 to 20 seconds depending on the influent flow rate and preset dosing volume). The discharge spilled onto a  $0.61$  m  $\times$   $0.61$ m plexiglas distribution plate with 0.07 m high sidewalls. The distribution plate had sixteen, equally-spaced 0.0095 m diameter orifices and thirty-three 0.0064 m diameter orifices located such that the distance between orifices was 0.076 m.

The total area of the 0.0064 m orifices equalled the total area of the 0.0095 m orifices. The 0.0095 m orifices were fitted with target nozzles to spread the wastewater discharged by the orifices over the media surface below. This arrangement gave very satisfactory distribution of wastewater over the media. Total discharge time (dosing siphon plus distribution plate) was 15 to 30 seconds, depending on the influent flow rate and preset dosing volume.

The dosing siphon and distribution plate were very reliable. Maintenance was required only on a weekly to bi-weekly basis to clean the biomass accumulation from the dosing siphon bell and the distribution plate orifices. Malfunctions and clogging were not observed with this weekly to bi-weekly cleaning schedule.

The nitrifying biofilter consisted of four welded steel sections, each measuring 0.61 m x 0.61 m x 1.22 m tall, stacked end to end for a

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total tower height of 4.88 m. The nitrifying biofilter was filled with B. F. Goodrich 60-degree, cross-flow media with a specific surface area of 138  $m^2/m^2$ . Short circuiting of wastewater flow along the tower walls was prevented and redistribution was promoted by securely wrapping each media section with plastic sheeting, by rotating the media orientation 90° at 0.61 m depth intervals, and by providing sidewall baffles or skirts at 1.22 m depth intervals to redistribute any sidewall flow back onto the media.

Wastewater samples were collected at the dosing siphon discharge, at 1.22 m, 2.44 m, 3.66 m, and 4.88 m depths, and, in some cases, following a 30 minute simulated clarification.

#### Pilot Plant Operational Procedures

#### Start-up operations

Funding for the project was obtained from the city of Ames in late October, 1985. Construction of the uninsulated building to house the pilot plant was completed in late November, and the media arrived in late December. The plant was assembled and hydraulically functional (except for the pretreatment process) on January 21, 1986.

The nitrifying microorganism population in the biofilter was established in cold temperatures. Antonie (1976) discussed the difficulties associated with cold climate RBC nitrification start-up. Heaters were used to maintain building interior ambient temperatures of 7° to 12° C.

Recycle of the biofilter effluent was employed, and the effluent collection tank at the base of the biofilter was heated with a livestock tank heater so that the temperature of the recycled wastewater was raised from approximately 6°C to final temperatures of 8° to 11°C. Recyle accounted for 50 to 80 percent of the total hydraulic load during the start-up operations. Dosing cycle length was 40 to 80 seconds during the start-up operations. The average total hydraulic loading rate was maintained between  $0.45$   $1/m^2$  · s and  $0.56$   $1/m^2$  · s.

The key to the successful cold weather start-up was the opportunity to seed the biofilter influent with effluent from a full-scale nitrifying biofilter located in Huxley, Iowa. A small sampling pump was used to introduce small amounts of nitrified effluent into the dosing siphon tank at constant time intervals for about six to eight hours per day from February 24 to March 1, 1986.

Prior to the seeding operations, nitrification in the biofilter was detectable, but the rate of nitrification was low and was not improving rapidly. Significant improvement in the rate of nitrification was measurable in the biofilter after only a few days following the initial seeding.

The hydraulic loading was then adjusted to the desired average hydraulic loading and the desired recycle ratio for the first data collection period. The system was allowed to remain at this hydraulic loading condition for two weeks with occasional sample collection and testing to establish the approach to "pseudo-steady state". The first "pseudo-steady state" data collection was conducted after March 13, 1986.

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#### Post-start-up operations

When hydraulic loading and/or influent  $NH_A^+$ -N concentration (supplementation instituted) conditions were changed for subsequent data collection periods, an acclimation period of 2-4 weeks was allowed with occasional sampling and testing to establish the approach to the new "pseudo-steady state" condition. The chronology of operating procedures, conditions, and changes and equipment malfunctions is summarized in Table 6.

Table 7 lists the data collection periods, the hydraulic loading conditions in effect for each, the wastewater temperature for each, and the range of influent  $NH_A^+$ -N and TKN concentrations for each.

## Hydraulic residence tests

Hydraulic residence tests were conducted on two occasions for continuously and intermittently-dosed wastewater applications at average hydraulic loading rates of  $1.2$   $1/m^2$  . Because of the cyclic nature of the hydraulic loading provided by the dosing siphon, special procedures were used to determine the average hydraulic residence times. The adequacy of flow distribution within the biofilter media was determined prior to hydraulic residence testing.

The water level in the dosing siphon was monitored on a continuous basis using an ISCO 1870 flow meter in the level mode. The chart time was synchronized with the operator's watch. The cyclic rise and fall of the water level in the dosing siphon was recorded relative to real time. The volume of influent wastewater per cycle, V<sub>APP</sub>, was equal to







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# Table 6. (Continued)



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the flow rate into the dosing siphon tank multiplied by the time interval between cyclic discharges.

A collector of known cross-section (9.0 cm diameter) was used to collect effluent from the media at the 4.88 m depth. The collected effluent drained directly into a 1000 mL graduated cylinder. The cumulative volume of effluent collected was recorded at time intervals of six to ten seconds. The effluent of three to four dosing siphon tank cycles was collected in a single 1000 mL cylinder.

Effluent collection with this equipment indicated uniform wastewater distribution in the media. The average rate of effluent collection was plotted versus real time. The volume of effluent collected,  $V_{\text{COL}}$ , for a single dosing siphon tank discharge was determined as the area under the effluent collection rate versus real time curve. Comparison of  $V_{\text{COLL}}$  to  $V_{\text{APP}}$  confirmed that the superficial velocities at various points beneath the tower were within ten to twenty percent of the theoretically applied superficial velocity.

When intermittent dosing was in use, a 500 ml volume of a known concentration of  $NH_4C1$  brine (tracer) was introduced onto the distribution plate immediately prior to discharge of the dosing siphon tank, and the time was recorded. This volume allowed fairly uniform spread of the tracer over the distribution plate and onto the media immediately before the dosing siphon tank discharge occurred. Introduction of the tracer in other locations or by other methods was found to be unsatisfactory for intermittent dosing conditions. When continuous dosing was in use.

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the tracer was introduced directly into the dosing siphon discharge pipe.

Biofilter effluent was collected at twenty to thirty second intervals for eight to ten minutes following tracer introduction. Each of the samples was analyzed for  $NH_4^+$  and Cl<sup>-</sup>. The NH<sub>4</sub><sup>+</sup> and CL<sup>-</sup> concentrations were plotted versus real time. The cumulative recovered mass was determined with time from these plots, and the average hydraulic residence time was determined to be the time required to recover 50% of the chloride tracer mass introduced.

Since the biofilter was operated in an oxygen-limited mode throughout its entire depth (i.e., constant total  $NH_4^+$  mass removal across the tower depth, independent of the bulk liquid  $NH_4^+$ -N concentration), the  $NH_4^+$  tracer was used to determine the average NH $_4^+$  contact time. For this condition,  $NH_4^+$ -N loss due to nitrification did not affect the  $NH_4^+$ -N contact time calculation. The average  $NH_4^+$  contact time was defined as the time required to recover 50% of the  $NH_4^+$  tracer mass introduced.

# Sample collection, preservation, and analysis

The sampling ports on the nitrification biofilter were constructed such that the main wastewater access slots at any two sampling depths were not vertically aligned. Thus, flow collected by an upper sampling port did not reduce the hydraulic loading rate for wastewater entering sampling ports at greater depths. Figure 23 demonstrates the staggered orientation of the sampling port access slots.

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Figure 23. Nitrifying biofilter sampling port construction

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Samples were collected simultaneously at all sampling points during any given profile sampling of the nitrifying biofilter. Samples were usually collected at various depths in the pretreatment biofilter simultaneously to allow some comparison of the two types of media in use.

Prior to sample collection, the sampling ports were flushed clean in the following manner. The sampling ports were opened and closed many times over a ten minute period to flush accumulated solids out of the sampling ports. The sampling ports were then left open for an additional ten minutes to assure that they were flushed clean of any stagnant water and solids. Clean, five-liter collection buckets were attached to the sampling ports to simultaneously collect discharge from the ports for a period of time ranging from 15 to 45 minutes. The sample collection times at all sampling ports were essentially equal  $(t 2 to 3 minutes)$ .

When complete biofilter profile data were not deemed necessary or practical, discrete samples were collected at constant time intervals at the nitrifying biofilter dosing siphon tank and the biofilter effluent using ISCO 2100 automatic samplers. A composited sample was collected simultaneously at the eight feet depth using an N-CON automatic sampler set at three equal volume samples per hour.

Samples were collected at one to eight hour intervals during data collection periods depending on sampling objectives and operator convenience. Upon collection, the samples were transferred to clean sample bottles when necessary and were preserved and analyzed in accordance with Table 8. When composited samples were required, the discrete

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Analytical Parameters	<b>Preservation Methods</b>	Analytical Method Used				
pH	Determined immediately in field	Method #423 in Standard Methods <sup>a</sup>				
Temperature	Determined immediately in field					
D.0.	Determined immediately in field	Hach Chemical Company version of the azide modification of the Winkler Method #421 B. in Standard Methods <sup>8</sup> , for 60 mL sample size				
$NH_{4}^+$ -L	4°C. $H_2SO_4$ to pH < 2 <sup>b</sup>	Ammonia selectivę electrode method #417 E. in Standard Methods'				
$NO_2$ + $NO_3$ - N	4°C, $H_2SO_a$ to pH < 2 <sup>b</sup>	Automated cadmium reduction method #418 F. in Standard Methods'				
<b>TKN</b>	4°C, $H_2SO_a$ to pH < 2 <sup>b</sup>	Colorimetric, semiautomated block digester AA II, EPA Method 351.2 (semiautomated phenate method)				
Filterable TKN	$0.45\mu$ filtration, $4^{\circ}$ C, $H_2SO_4$ to pH < 2					
<b>COD</b>	4°C, $H_2SO_a$ to pH < 2 <sup>b</sup>	Dichromate reflux method #508 in Standard Methods				

Table 8. Sample preservation and analytical methods

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Standard Methods for the Examination of Water and Wastewater. 16th ed. Washington, D.C. American Public Health Association, 1985.

<sup>D</sup>Samples were not acidified until after filterable TKN, filterable COD, SS, VSS and alkalinity liquid fractions were removed for analyses when applicable.

Table 8. (Continued)

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samples taken at any given sampling point were combined on a flowproportional basis to account for sampling interval variations, properly preserved and analyzed.

The significant analytical parameters in this study were the  $NH_4^+$ -N, TKN, filterable TKN,  $NO_2^-$  +  $NO_3^-$ -N, COD, filterable COD, pH, temperature, SS, VSS, alkalinity, and DO of the bulk liquid. The pH, temperature, DO, and NH $_{A}^{+}$ -N analyses were performed by the operator. The remaining parameter analyses were performed by the Engineering Research Institute laboratory personnel. The analytical methods used for the analytical parameters are summarized in Table 8.

The analytical schedule for samples was generally related to sampling point location as shown in Table 9. However, the operator exercised discretion in deviation from this schedule when he felt that the understanding of process variables and biofilter performance were likely to be enhanced or when analytical costs and operator convenience were considerations.

### Characteristics of Pilot Plant Influent

The values of significant parameters in composited samples of the Ames WPCP secondary clarifier (final) effluent were determined three times per week during the pilot study. These analyses were performed and reported by the Ames WPCP personnel.

The properties of the wastewater influent to the nitrifying biofilter portion of the pilot plant were usually different from those of the Ames WPCP secondary clarifier effluent because of the pretreatment portion of the pilot plant,  $NH_4^+$ -N supplementation, and recycle.

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Table 9. Sampling point analytical schedule

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<sup>a</sup>pH and D.O. measurements were made during some, but not all, sample collection periods.

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Detailed reporting of the secondary effluent analytical parameters is, therefore, not appropriate in this dissertation. Pertinent secondary effluent DO and BOD<sub>5</sub> concentrations reported by the Ames WPCP personnel during the study period are included in the pretreatment tower data section in the Appendix.

## Huxley, Iowa Full-Scale Plant Sampling

The city of Huxley operates a new 2.4x10 $^6$  1/d average design flow separate-stage trickling filter plant, which began operation in May, 1985. The minimum initial wastewater flow was expected to be about 5.4x10 $<sup>5</sup>$  1/d. The minimum design flow was 1.3x10 $<sup>6</sup>$  1/d. The covered</sup></sup> nitrifying biofilter is 7.62 m in diameter and 4.27 m deep. It is filled with 98  $m^2/m^3$  cross-flow media. The rotary distributors are rated for maximum and minimum flow rates of 79 L/s and 28 L/s, respectively. A 16 L/s recirculation capacity is available.

Approval was obtained to sample the nitrifying biofilter influent and effluent during the months of January, February, and March, 1986. The samples were analyzed for  $NH_A^+$ -N and temperature, and occasionally for TKN, COD, SS, and pH. The samples were collected at preset, constant time intervals using ISCO 2100 automatic samplers. All samples were preserved immediately by the cold ambient temperatures and by acidification for TKN,  $NH_4^+$ -N, and COD analyses.

The low flow conditions and low  $NH_4^+$ -N plant loadings yielded little useful oxygen-limited data. Most influent  $NH_4^+$ -N concentrations were very low. In addition, plant records were not always sufficient to evaluate the use of recycle. Ammonium-limited data were obtained.

Only the oxygen-limited nitrification data will be presented in subsequent presentations.

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#### RESULTS AND ANALYSIS

This nitrifying biofilter study was noteworthy for the following reasons:

- 1. It involved successful start-up in cold temperatures.
- 2. It involved a comparison of two types of wastewater application, i.e., intermittent dosing and continuous dosing, at identical average hydraulic loading rates.
- 3. Bulk liquid sampling was performed at many tower depths, and chemical parameter concentration profiles along the tower depth were constructed.
- 4. Bulk liquid chemical parameters monitored included D.O.,  $NH_A^+-N$ ,  $NO_2^- + NO_3^- -N$ , TKN, filterable TKN, COD, filterable COD, pH, SS, VSS, and alkalinity.
- 5. 60 degree cross-flow structured media with a specific surface area of 138 sq. m/cu. m and 45 degree cross-flow structured media with a specific surface area of 98 sq. m/cu. m were used in the nitrifying biofilter and the pretreatment biofilter, respectively.
- 6. Nitrification performance at bulk liquid  $NH_A^+$ -N concentrations greater than 3 to 4 mg/1 throughout the tower depth was emphasized.
- 7. The project was performed with equipment and operational budgets of \$2,000 and \$6,000, respectively, and with a one-person staff.

Typical results for the variety of hydraulic and mass loadings studied will be presented. Observations of conditions which may have influenced the nitrification performance will be discussed. The

experimental and calculated data for this investigation have been included in the appendix.

It was desirable to evaluate the data at a common wastewater temperature. The common temperature selected was 10°C. Nitrification rates observed at temperatures other than 10°C were converted to a 10°C basis by the use of the Nernst-Einstein equation presented in the Review of Literature section. The effects of temperature-related viscosity changes on the hydraulic residence time were not considered for two reasons. First, hydraulic flow patterns over the surface of structured, plastic media would not be the same as flow patterns over flat, inclined planes like those used in the studies evaluated in the Review of Literature section. Second, biofilm growth on the plastic media would alter the viscosity effects.

Oxygen flux-limited nitrification rates approaching  $8 \times 10^{-4}$  to 10  $\times$  10<sup>-4</sup> kg N/d/sq. m of specific surface area were expected at 10°C. This prediction was based on conversion of the 11.7 x 10<sup>-4</sup> kg N/d/sq. m oxygen flux-limited, average nitrification rate, reported by Gullicks and Cleasby (1986), to 10°C by the use of the Nernst-Einstein equation.

## Pretreatment Biofilter Performance

Analytical parameter concentrations versus depth in the pretreatment tower are presented in Table 10. The pretreatment tower was 2.44 m tall and was filled with 45 degree cross-flow media with a specific surface area of 98 sq. m/cu. m. Wastewater flow to the pretreatment. tower consisted entirely of secondary effluent from the final clarifiers

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Date	Avg. Hyd. Load $1/m^2$ • s	Water Temp. $^{\circ}$ C	Depth in Tower, m	$NH4 + -N$ mg/1	<b>TKN</b> mg/1	$NO2 +$ $NO_3 - N$ mg/1	<b>TKN</b> $< 0.45 \mu$ mg/1	<b>COD</b> mg/1	<b>COD</b> $< 0.45 \mu$ mg/1	Alk mg/1 as CaCO <sub>3</sub>	SS <sub>1</sub> mg/1
7/31/86	1.90	22.5	$\bf{0}$	8.6							
			1.22	7.4							
			2.44	6.3				$\sim$			
11/11/86	1.72	16.3	$\mathbf{0}$	19.7	22.3			92			
			1.22	18.9							
			2.44	17.1	23.4	6.4		133			
11/12/86	1.72	13.2	$\mathbf{0}$	17.2	20.4			181			
			1.22	<b>17.1</b>	19.9			116			
			2.44	15.9	19.4	5.5		130			
11/13/86	1.70	15.8	$\mathbf{0}$	14.2	20.6			121			
			1.22	14.9	21.9		16.8		47		
			2.44	16.2	22.8	5.5		109			
11/19/86	1.66	15.5	$\mathbf{0}$	19.0	27.1			84			
			1.22	19.0	26.2						
			2.44	17.0	23.9			72			

Table 10. Analytical parameter concentrations versus depth in the pretreatment tower. 45° cross-flow media with a specific surface area of 98 m'/m'. No recycle

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 $\label{eq:2} \frac{1}{\sqrt{2}}\sum_{\mathbf{k}\in\mathbb{Z}^d}\left(\frac{1}{\sqrt{2}}\right)^{\frac{1}{2}}\left(\frac{1}{\sqrt{2}}\right)^{\frac{1}{2}}$ 

 $\mathcal{L}^{\text{max}}_{\text{max}}$  and  $\mathcal{L}^{\text{max}}_{\text{max}}$  and  $\mathcal{L}^{\text{max}}_{\text{max}}$ 

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Table 10 (Continued)

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at the Ames WPCP. Recycle was not used. Wastewater was dosed continuously at hydraulic loading rates ranging from 1.4 to 1.9 1/sq. m • s.

The pretreatment tower was put into operation on June 16, 1986. The pretreatment tower was constructed to provide carbonaceous oxygen demand reduction ahead of the pilot-scale nitrifying biofilter. Nitrification was apparent in the pretreatment tower by July 14, 1986. Near steady-state nitrification was observed by July 30, only six to seven weeks after start-up. The pretreatment tower nitrification performance was monitored simultaneously with the performance of the pilot-scale nitrifying biofilter through January 27, 1987.

Figure 24 demonstrates that the rate of nitrification, converted to a common 10°C basis by the use of the Nernst-Einstein equation, in the pretreatment tower was independent of influent bulk liquid  $NH_A^+$ -N concentration at  $NH_4^+$ -N concentrations greater than 6 mg/1, and was, therefore, flux-limited by oxygen. At influent  $NH_4^+$ -N concentrations less than 6 mg/l, the concentration of  $NH_A^+$ -N in the tower effluent was less than 4 mg/1, and nitrification was flux-limited by substrate.

As expected with biological fixed-film processes, there was substantial data scatter. The data scatter was due primarily to relatively poorer performance of the top section of the pretreatment tower when compared to the performance of the lower section of the tower. It would be easy to assume that carbonaceous oxygen demand carryover from the secondary clarifiers was responsible for the poor performance of the top tower section; however, this was not the case. It will be shown that depleted D.O. concentrations in the secondary effluent had stronger



Pretreatment tower nitrification rate versus influent<br> $NH_4$ <sup>--N</sup> Figure 24.

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influence on the low or nonexistent nitrification rate in the top section.

Pilot-Scale Nitrifying Biofilter Performance

The nitrifying biofilter consisted of four sections, each 1.22 m tall, stacked end to end for a total height of 4.88 m. The tower was filled with 60 degree cross-flow media with a specific surface area of 138 sq. m/cu. m. Prior to June 16, 1986, secondary effluent from the Ames WPCP was distributed directly to the top of the nitrifying biofilter, with varying amounts of recycled nitrifying biofilter effluent. Subsequent to June 16, 1986, secondary effluent from the Ames WPCP was pretreated to assure good carbonaceous oxygen demand removal before it was distributed onto the nitrifying biofilter media, again with varying amounts of recycled nitrifying biofilter effluent.

Intermittent dosing was used during most of the study, however, some data were collected for continuous dosing conditions. When intermittent dosing was used, the periodic dosing cycle length ranged from 30 to 70 seconds, depending on the average total hydraulic loading rate and the dosing siphon control elected by the operator. The cyclic flow volume was distributed in 15 to 25 seconds of the cycle, depending on the same dosing siphon operational conditions.

#### 0.46-0.51 1/sq. m \* s average hydraulic load

The rate of nitrification, converted to a common 10°C basis by the use of the Nernst-Einstein equation, was plotted versus influent  $NH_A^+$ -N concentration for average hydraulic loading rates of 0.46-0.51

1/sq. m • s in Figure 25. The rates of nitrification at  $NH_4^+$ -N concentrations greater than 4 mg/1 were generally lower than expected. However, some data indicate that the anticipated rates of nitrification discussed previously in this section for  $NH_A^+$ -N concentrations greater than 4 mg/1 can be achieved when influent liquid conditions and the tower environment are suitable.

Recycle, expressed as percent of total flow, did not appear to affect nitrification performance; however, recycle modifies factors such as bulk liquid D.O., carbonaceous oxygen demand, toxicity, etc. Thus, for example, the net influent bulk liquid D.O. is not necessarily higher when a greater percentage of recycle is used; but, were recirculation not in effect the D.O. of the bulk liquid would definitely be lower. This would explain why tower performances with recirculation rates of 31 and 59 percent of the total flow were similar.

#### 0.55-0.59 1/sq. m \* s average hydraulic load

Analytical parameter bulk liquid concentrations versus depth in the nitrifying biofilter are shown in Figures 26-28 for average total hydraulic loading rates of 0.57 to 0.58 1/sq. m • s and various recycle and wastewater temperature conditions. Figures 26 and 27 represent cold climate data, while Figure 28 demonstrates the enhanced performance at warmer wastewater temperatures. Nitrification performance at the lower temperatures was somewhat lower than would be expected if all influent conditions, except temperature, were equal. However, wastewater

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Nitrification rate versus influent  $NH_A^+$ -N concentration for<br>average total hydraulic loading rates of 0.46-0.51  $1/m^2$  · s.<br>All data points were for recycle conditions of either 30-31%<br>or 58-59% of the total flow. For ni Figure 25.

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Analytical parameter concentrations versus depth in tower,<br>4/3/86 Figure 26.

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Figure 27. Analytical parameter concentrations versus depth in tower, 4/10/86



Analytical parameter concentrations versus depth in tower,<br>7/30/86 - 7/31/86 Figure 28.

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conditions at colder temperatures will be shown to be less satisfactory for nitrification.

All sections of the biofilter did not perform equally well at any given time. During this study, the 2.44 to 3.66 m tower section often performed poorly when compared to the sections immediately above and below it. Similarly, the top section often performed poorly relative to the section immediately below it. However, Figures 26 and 27 show that the poorer performance of these sections was transient and that other sections also exhibited this phenomenon on different occasions.

The performance of individual tower sections was probably affected by variation of influent bulk liquid characteristics and by partial plugging of the media. Partial plugging of the media would affect liquid distribution and the flow of air through a given section of media. Partial plugging in a tower section would probably not affect the performance of other sections of the tower because both air and water flow would be redistributed in adjacent tower media sections. Partial plugging which occurred at the middle of a tower section, where individual 0.61 m structured media units were stacked could affect both water and air flow in the 0.61 m high units above and below the plugging point.

Figures 29 and 30 show the rate of nitrification, converted to a common 10°C basis by the Nernst-Einstein equation, versus influent bulk liquid  $NH_A^+$ -N concentration for average total hydraulic loads ranging from 0.55 to 0.59 1/sq. m • s. The rate of nitrification was observed to be independent of influent  $NH_A^+$ -N bulk liquid concentration at

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Figure 29. Nitrification rate versus influent  $NH_4^-$ -N concentration for average total hydraulic loading rates of  $0.55-0.59$   $1/m^2$  · s. Some data, usually for the 0-1.2m and 2.4-3.7m depth sections, with nitrification rates less than 50% of the nitrification rates in adjacent tower sections were deleted

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Figure 30. 4.88 m biofilter average nitrification rates versus influent  $NH_A^+$ -N concentrations for average total hydraulic loading rates of 0.55-0.59 1/m' • s

 $NH_4^+$ -N concentrations greater than 5 to 8 mg/1. Thus, oxygen or some reactant other than  $NH_4^+$ -N is the rate-limiting reactant above those concentrations.

Influent bulk liquid characteristics were significantly different during the cold weather data collection period than they were during the warm weather data collection period. The average bulk liquid percent of D.O. saturation was lower in the cold weather period. The average bulk liquid COD concentration was higher in the cold weather period. Thus, cold weather performance was lower than warm weather performance, even when converted to the same 10°C temperature basis.

Data scatter was substantial, but understandable, in view of the D.O. and COD variability. However, it was apparent that nitrification rates could approach those expected  $(8 \times 10^{-4}$  to 10 x 10<sup>-4</sup> kg/d • sq. m of total surface area as developed previously in this section) if influent liquid conditions and the tower environment were suitable. In general, the data presented in Figure 30 reflect the lower performance of the top section of the tower, and, in many cases, the 2.44 to 3.66 m depth tower section, as well.

## 1.13-1.29 and 0.67-0.8 1/sq. m • s average hydraulic loads

Figures 31 and 32 show profiles of analytical parameter concentrations versus tower depth for average total hydraulic loading rates of approximately 1.2 1/sq.  $m \cdot s$  and wastewater temperatures of 11 to 13°C. These figures are significant in that they demonstrate denitrification in the upper section of the tower, as evidenced by the disappearance of

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Analytical parameter concentrations versus depth in tower. Figure 31.  $11/12/86$ 

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Figure 32. Analytical parameter concentrations versus depth in tower, 11/13/86

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 $NO<sub>2</sub> + NO<sub>3</sub> - N$ . Thus, D.O. in the bulk liquid appears to have been inadequate for nitrification in the top tower section and possibly in short supply in the other tower sections as well. Wanner and Gujer (1984) predicted correctly that denitrification could occur in the upper sections of combined carbon oxidation-nitrification trickling filters when recycle was instituted. Apparently, denitrification is possible in the nitrifying biofilter stage of separate-stage systems, as well, when  $NO_2$  +  $NO_3$  -N concentrations are high enough.

Wanner and Gujer (1984) predicted that soluble COD concentrations greater than 27 mg/1 would exclude nitrifiers from the biofilm because of competition with heterotrophs. While the filterable fraction of COD in the influent to the tower exceeded 27 mg/1, the apparent nitrification and negligible change in total COD through the tower depth suggests that the prediction of Wanner and Gujer is not correct.

Figure 33 contains analytical parameter concentration versus tower depth data for an average total hydraulic load of  $1.28$  1/sq. m  $\cdot$  s, but without intermittent dosing. At first glance, this figure could easily be misinterpreted. It would appear that intermittent dosing provided slightly better nitrification than continuous dosing. However, D.O. deficiency may be the real reason for the nearly identical performance of the intermittently-dosed (Figures 31 and 32) and continuously-dosed (Figure 33) towers. Recycle, which was in use in Figures 31 and 32, would have the effect of increasing D.O. in the bulk liquid, and may be responsible for the equivalent performance of the two dosing methods.

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Analytical parameter concentrations versus depth in tower.<br>1/25/87 Figure 33.

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Nitrification was evident in Figure 33 even at filterable COD concentrations of 66 mg/1. This indicated that the 27 mg/1 nitrifier extinction COD threshold predicted by Wanner and Gujer (1984) is incorrect. However, some competition between heterotrophs and nitrifiers was suggested by the total COD decrease with increased tower depth. Competition between heterotrophs and nitrifiers was not apparent in Figures 31 and 32.

Nitrification rate versus influent  $NH_4^+$ -N bulk liquid concentration data for average total hydraulic loading rates of 1.13 to 1.27 1/sq. m • s are shown in Figure 34. The data collected from 11/11/86 to 11/13/86 represented tower performance approximately one to two weeks after the onset of cold weather. The higher rates of nitrification (circled data points) approached, and in some cases, exceeded the expected values  $(8 \times 10^{-4}$  to 10 x  $10^{-4}$  kg/d  $\cdot$  m<sup>2</sup>) for cold weather performance.

A gradual decline in nitrification performance in the tower was noted with time, as evidenced by the data for 12/15/86 to 12/19/86, and corresponded to a gradual decline in secondary effluent D.O. concentrations. The decline of the D.O. concentrations in the Ames WPCP final clarifier effluent is documented in their records.

The pump supplying pretreated secondary effluent began to lose efficiency subsequent to 11/13/86. The total hydraulic loading had dropped from 1.2 down to 0.8 1/sq. m • s by 11/19/86. Analytical parameter bulk liquid concentrations versus depth in the tower on 11/19/86 are shown in Figure 35. Recirculation, expressed as a percent of the

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Figure 34. Nitrification rate versus influent  $NH_A^+$ -N concentration for average total hydraulic loading rates of 1.13-1.27  $1/m^2$ : s in nitrifying biofilter,  $60^\circ$  cross-flow media, 138 m<sup>2</sup>/m<sup>3</sup>

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Figure 35. Analytical parameter concentrations versus depth in tower, 11/19/86

total flow, was 44% on 11/19/86, while recirculation on 11/13/86 was only 32%. A dramatic increase in nitrification performance occurred, as shown by comparison of Figures 32 and 35.

Again, nitrification was noted at filterable COD concentrations greater than the nitrifier elimination threshold of Wanner and Gujer (1984). Another noteworthy observation from Figure 35 is that sloughing from the 1.22 to 2.44 m tower section was documented. Two related phenomena were noted. First, the increase in TKN, as expected with an increase in solids due to sloughing, was accompanied by an increase in  $NH_A^+$ -N. This may be an indication of adsorption of NH $_A^+$ -N onto the solids. The degree of adsorption did not appear to be high. Second, the solids associated with the sloughing from the 1.22 to 2.44 m tower section appeared to be retained in the 2.44 to 3.65 m tower section based on the TKN data.

Further evidence of adsorption of  $NH_4^+$ -N by solids in the biofilter process was observed in comparisons of the  $NH_A^+$ -N concentrations of unsettled biofilter effluent samples versus biofilter samples that had been settled for 30 to 60 minutes. The  $NH_A^+$ -N concentrations in the unsettled biofilter effluent samples were typically 0.3 to 0.6 mg/1 higher than those in the settled samples.

Subsequent to 11/19/86, recirculation was discontinued. A plot of nitrification rate versus  $NH_A^+$ -N bulk liquid concentration for average total hydraulic loads of 0.7-0.8 1/sq. m ' s is presented in Figure 36. Analytical parameter concentrations versus depth in the tower are shown in Figures 37 and 38 for two dates following the discontinuation of

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Nitrification rate versus influent  $NH_4^+$ -N concentration for<br>average total hydraulic loading rates of 0.7-0.8 l/m<sup>2</sup> · s Figure 36.

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Figure 37. Analytical parameter concentrations versus depth in tower, 11/21/86

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Figure 38. Analytical parameter concentrations versus depth in tower,  $12/3/86$ 

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recirculation. These figures demonstrate that a rapid decrease in nitrification performance, corresponding to the discontinuation of recirculation, occurred. Figure 36 also shows that the rate of nitrification was independent of bulk liquid  $NH_{4}^{+}-N$  concentrations greater than 5.6 mg/1.

Figures 37 and 38 show that the nitrifier-elimination soluble COD threshold is greater than the 27 mg/1 proposed by Wanner and Gujer (1984). The figures demonstrate that the total COD exhibited some fluctuation in concentration with increasing depth in the tower, but that the COD fraction smaller than 0.45 microns remained relatively constant at about 33 mg/1.

Some competition between heterotrophs and nitrifiers appeared to exist in some sections of the tower, based on the decline in total COD concentrations with increased tower depth in Figure 38. However, the decrease in nitrification performance subsequent to 11/19/86 appears to be unrelated to COD concentrations in the bulk liquid since both the total and filterable fractions of COD in the tower influent were essentially constant for Figures 35-38 related to the discussion above.

The average hydraulic loading rate was increased to 1.15 1/sq. m • s, with a recirculation rate of 38 percent of the total flow, on 12/4/86. Data were collected two to three weeks later. These data are plotted on Figure 34. They indicate that the rate of nitrification was independent of bulk liquid  $NH_4^+$ -N concentration at NH $_4^+$ -N concentrations greater than 6 mg/1.

The data in Figure 34 also show that the nitrification rate observed five weeks earlier at essentially the same hydraulic loading conditions could not be regained at this high hydraulic loading, at least when intermittent dosing was in use. Thus, part of the gradual loss of nitrification performance observed between 11/11/86 and 11/13/86 and between 11/19/86 and 12/4/86 could have been due to scouring or washout effects of intermittent dosing at relatively high, total average hydraulic loads of  $0.8$  to  $1.2$   $1/sq$ . m  $\cdot$  s.

Intermittent dosing was discontinued on 12/19/86. The total average hydraulic loading was not altered. Figures 33 and 39 indicate that nitrification performance was regained by the discontinuation of intermittent dosing. In fact, the rate of nitrification in the lowest section of the tower returned to the levels observed in early November. The return of nitrification performance occurred despite the fact that the fraction of influent COD smaller than 0.45 microns was higher in January, 1987 than it was in November and December, 1986.

The effects of hydraulic loading rate on cold-climate nitrification were further demonstrated by reducing the hydraulic loading rate from 1.25 down to 0.62 1/sq. m \* s on January 27, 1987, while maintaining continuous, rather than intermittent, dosing. Comparison of February 13, 1987 nitrification performance (Figure 40) with that of January 25, 1987 (Figure 33) clearly shows an improvement in nitrification performance at the lower hydraulic loading rate.

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Nitrification rate versus influent  $NH_A^+$ -N concentration for Figure 39. average total hydraulic loading rates $^{4}$ of 1.21-1.28 l/m $^{2}\cdot$  s and continuous dosing. Average nitrification performance of the bottom half of the biofilter and the top half of the biofilter have been shown by the upper and lower solid lines, respectively

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Analytical parameter concentrations versus depth in tower.<br>2/13/87 Figure 40.

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## Operating Parameter Effects

The performance of nitrifying biofiIters can be affected by many operating parameters. These include; 1) wastewater temperature, 2) wastewater pH, 3) wastewater alkalinity, 4) wastewater suspended solids, (5) carbonaceous oxygen demand of the wastewater, 6) wastewater dissolved oxygen concentrations, 7) hydraulic loading, 8) method of wastewater application, 9) media configuration and specific surface area, 10) toxicity of the wastewater, 11)  $NH_A^+$ -N concentrations < 4 mg/1, and 12) previous operation history.

The influences of most of these operating parameters were observed at some point during the nitrification study at the Ames WPCP. The project budget and staffing were insufficient to investigate possible toxicity effects. Distinguishing which of the operating parameters is most influential at any given time is a difficult task. Many of the observations of this study are contrary to those previously reported in the literature.

## Wastewater temperature effects

The effects of wastewater temperature on the rate of nitrification are well documented in the literature. The rate of nitrification would be expected to increase with increasing temperature whether the fixedfilm process is metabolism-limited or diffusion-limited. Previous discussions have established that the Nernst-Einstein equation adequately predicts the effects of temperature. Results of this study neither confirm or refute the validity of the Nernst-Einstein theory.

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When other factors such as carbonaceous oxygen demand and dissolved oxygen effects were not apparent, the transformation of nitrification rate data to a common 10°C basis by the Nernst-Einstein equation appeared to be acceptable (reference Figure 29).

## pH and alkalinity effects

The pH measurements in this study were observed to range from 7.1 to 8.1 during cold weather data collection periods. The pH increased slightly from the nitrifying biofilter influent to the effluent, an observation contrary to theoretical considerations. The nitrification process produces hydrogen ions which react instantaneously with bicarbonate in the wastewater to form carbonic acid. The consumption of alkalinity in the nitrifying biofilter was observed and was generally close to the theoretical value of 7 mg  $CaCO<sub>3</sub>$  per mg N oxidized. The decrease in alkalinity and the resulting presence of carbonic acid would be expected to produce a lower pH.

The increase of pH from the nitrifying tower influent to the effluent has been observed in many studies (Brown and Caldwell, 1980, Baxter and Woodman, Inc., 1973, Duddles and Richardson, 1973, and Sampayo and Metcalf, 1984). Air stripping of carbon dioxide has been credited with these observed pH changes. Consumption of carbon dioxide by autotrophic microorganisms may account for part of the observed pH changes.

The observed pH changes from the biofilter influent to the effluent during summer data collection were in agreement with theoretical expectations. The pH was observed to range from 7.0 to 7.7 for summer

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nitrification data and to decrease slightly from the influent to the effluent of the tower. Figure 28 indicates that nitrification did not appear to be hindered at alkalinity concentrations greater than 30 mg/1.

#### Suspended solids effects

Sarner (1981 and 1986) reported that solids adversely affect the removal of soluble substrate in trickling filters. The influent suspended solids concentrations in the Ames WPCP study ranged from 30 to 60 mg/1. The Ames study data within this narrow suspended solids range neither support nor refute Sarner's conclusion.

Evidence that  $NH_A^+$ -N may be adsorbed onto solids in the biofilter process has been presented in discussions related to Figure 35. Sloughing of solids onto which  $NH_A^+$ -N has been adsorbed would have an adverse impact on biofilter nitrification performance. However, the impact would likely be small unless massive sloughing occurred.

## Carbonaceous oxygen demand effects

The effects of carbonaceous oxygen demand on the rate of nitrification are usually evaluated in terms of the concentration of soluble carbon (expressed as  $BOD_E$ , COD, or TOC) and the ratio of carbon to TKN. These approaches were both examined in this study.

Plots of the rate of nitrification, converted to a common 10°C basis by the use of the Nernst-Einstein equation, versus influent total COD:influent total TKN and versus influent 0.45 micron filterable COD: influent 0.45 micron filterable TKN or total TKN (as indicated) were constructed for the hydraulic loading conditions used in this study.

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Figure 41. Nitrification versus COD:TKN ratios for average total hydraulic loading rates of 0.46-0.6 1/m' • s

Figure 41, constructed for average total hydraulic loading rates of 0.46-0.61 1/sq. m • s with intermittent dosing, indicates that the ratio of COO:TKN can have an effect on the rate of nitrification at these hydraulic loading conditions.

Similarly, Figure 42 demonstrates that the ratio of COD;TKN appears to influence the rate of nitrification at average total hydraulic loading rates of 0.67-0.8 1/sq. m • s with intermittent dosing. Figure 43 demonstrates that at higher hydraulic loading rates without intermittent dosing, the ratio of COD: TKN exhibits little, if any, effect on the rate of nitrification. In fact, low CODrTKN ratio data existed for which the rate of nitrification was zero, indicating that some factor, other than carbonaceous oxygen demand, was influencing the rate of nitrification.

Nitrification rates were plotted versus the ratios of total influent COD: total influent TKN and versus 0.45 micron filterable C0D;0.45 micron filterable TKN for average total hydraulic loading rates of 1.13-1.28 1/sq. m s with intermittent dosing. However, these plots revealed no relationship between CODiTKN ratios and the rate of nitrification, and, therefore, have not been included in this presentation.

The observation that C0D;TKN ratios have a greater effect at lower hydraulic loading rates is consistent with mass transport expectations. At low hydraulic loading rates, the mass transfer of organic and nonorganic molecules into the biofilm is dependent on both liquid and biofilm phase resistance or diffusion. At high hydraulic loading rates,the resistance may be essentially a biofilm diffusion resistance,

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Figure 42: Nitrification rate versus COD:TKN.ratios for average total hydraulic loading rates of 0.67-0.8 1/m' • s



Figure 43. Pretreatment tower nitrification rate versus COD;TKN ratios for average total hydraulic loading rates of 1.66-1.8  $1/m^2$  • s

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and mass transport of both organic and nonorganic molecules would be expected to increase.

The diffusivities of smaller (nonorganic) molecules such as  $NH_A^+$ -N and oxygen are greater than those of organic molecules. In denser media, such as biofilms, the relative diffusivities of oxygen and  $NH_A^+$ -N compared to organic molecules are greater than the relative diffusivities of oxygen and  $NH_4^+$ -N compared to organic molecules in water. Thus, at high hydraulic loading rates, the relative mass transport of oxygen and  $NH_A^+$ -N compared to the mass transport of organic molecules would be greater. Under these relative mass transport conditions, the rate of nitrification would be expected to show less response to bulk liquid variations in COD concentrations typical of separatestage nitrification (i.e., low COD concentrations).

Wanner and Gujer (1984) have reported that nitrifiers would be displaced completely from biofilms if the soluble fraction of COD in the wastewater was greater than 27 mg/1. To test the effect of the filterable fraction of COD in the wastewater on the rate of nitrification, the rate of nitrification, converted to a common 10°C basis by the use of the Nernst-Einstein equation, was plotted versus the 0.45 micron filterable fraction COD bulk liquid concentration. The plots were constructed for each of the following hydraulic loading conditions:  $1) < 0.6$  1/sq. m • s with intermittent dosing, 2) 1.13-T.27 1/sq. m • s with intermittent dosing, 3) 0.67-0.8  $1/sq$ . m  $\cdot$  s with intermittent dosing, 4) 1.2-1.28  $1/sq$ . m  $\cdot$  s without intermittent dosing, and 5) 1.7-1.9  $1/sq$ . m • s (pretreatment tower) without intermittent dosing.

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No relationship was found to exist between the filterable COD concentration and the rate of nitrification for any of the above hydraulic loading conditions. Nitrification rates were often high when the 0.45 micron filterable COD concentration was in excess of 60 mg/1. Conversely, the rate of nitrification was often low when the concentration of 0.45 micron filterable COD was 34 mg/1 or less. The plots have, therefore, not been included in this presentation.

## Dissolved oxygen effects

It has been hypothesized in earlier discussions that the D.O. concentration in the wastewater was an important factor in the observed nitrification performance in a number of circumstances. However, because of the limited financial and personnel resources available for this study, the D.O. data base was not large.

Chen and Bungay (1981) used microprobes in their study of external mass transfer in fixed film processes. Figure 44 clearly demonstrates their finding that the concentration of oxygen at the biofilm surface can be appreciably lower than the concentration in the bulk liquid. Bungay et al. (1969) used microprobe techniques to measure D.O. concentrations within biofilms. They determined that respiration ceased, i.e., oxygen was depleted, at depths into the biofilm of 50 to 150 microns depending upon the substrate concentration in the biofilm. Thus, the concept of a thin active layer was established. Chen and Bungay (1981) also found that the active layer can be defined by depletion of either the electron donor or the electron acceptor.

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Figure 44. Oxygen concentration profile up to and. through the biofilm on a rock from a trickling filter. Data from Chen and Bungay (1981)

In order to test the effect of D.O. on the rate of nitrification, the rate of nitrification, converted to a common 10°C basis by the use of the Nernst-Einstein equation, was plotted versus the percent of D.O. saturation in the bulk liquid at the prevailing water temperature when the sample was collected. The percent of D.O. saturation was selected rather than the D.O. concentration because the investigator felt that it would better reflect the oxygen mass transfer potential across both the air-liquid and 1iquid-biofiIm interfaces. The data appear in Figure 45.

The rates of nitrification observed in the pretreatment tower (45 degree cross-flow media) at hydraulic loading rates of 1.63-1.91 1/sq. m • s were plotted against the percent of D.O. saturation of the Ames WPCP final effluent, as reported by Ames WPCP personnel. The relationship between the nitrification rate and the D.O. saturation was observed to be a straight line having an R-squared value of 0.87. The relationship is significant at the 0.001 level.

The data collected from the nitrifying biofilter (60 degree crossflow media) were plotted for hydraulic loading rates ranging from 0.57- 0.78 1/sq. m • s with intermittent dosing (Figure 45). These data are less reliable than the Ames WPCP data because the samples for D.O. analysis had to be collected from the tower sampling ports. The nitrifying biofilter samples were, therefore, subject to higher aeration. Nonetheless, the data indicate that D.O. in the wastewater did affect the rate of nitrification.



Figure 45. Nitrification rate versus bulk liquid D.O.

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The two sets of data for the nitrifying biofilter should overlap if all conditions were equal since the rates of nitrification were converted to a common 10°C basis. However, the COD:TKN ratio effects, which have been shown to be active at low hydraulic loading rates, influenced the two data sets. Figure 45 suggests that nitrification rates can be affected significantly by bulk liquid D.O. levels less than 60 to 65% of saturation. Figure 45 also suggests that nitrification may be totally nonexistent at 45 to 50% D.O. bulk liquid saturation. Thus, for normal wastewater operating temperatures of 10 to 25°C, nitrification rates can be reduced by bulk liquid D.O. concentrations less than 7 to 5.5 mg/1, respectively.

Another observation of possible significance was that the D.O. concentrations in the Ames WPCP final effluent were quite low at times. Effluent from trickling filter towers would generally not be expected to have D.O. concentrations as low as those observed in the final clarifier effluent. Continued biomass metabolism during detention in the final clarifier reduces the D.O. concentration of the wastewater. Records at the Ames WPCP indicate that the oxygen depletion in the final clarifiers can exceed 1 mg/1 of D.O. This might have significance in recommendations regarding intermediate clarifiers between the first and second stage filters in separate-stage nitrification systems.

Brown and Caldwell (1980) reported poor nitrification in the trickling filter plant at Stockton, California, despite effluent D.O. concentrations which were reported to be generally in excess of 5 mg/1. The same study reported that the nitrogen balance indicated the loss of

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total nitrogen from the influent to the effluent. This nitrogen loss was composed of  $NO<sub>9</sub>$ -N and  $NO<sub>3</sub>$ -N which were consistently low, despite significant ammonia and organic nitrogen removal in the secondary treatment process. The disappearance of  $NO<sub>2</sub>$  -N and  $NO<sub>3</sub>$  -N was most likely indicative of oxygen deficiency in the trickling filter since the nitrogen balance could not be rectified by the calculated nitrogen assimilation in effluent solids from the tower.

Increasing the rate of recirculation and the number of forced draft fans increased the nitrification performance of the plant. It was not known whether it was the increased forced draft or the increased recirculation that had the greatest impact on the rate of nitrification. However, both the increased forced draft and the increased recirculation would increase the D.O. concentration in the tower influent.

# Hydraulic loading and application method effects

Figures 25, 29 and 34 show that the rate of nitrification observed in this study increased with increasing average hydraulic loading, except in the period of time subsequent to 11/13/86. The influence of depleted D.O. concentrations has been shown to have affected nitrification subsequent to 11/13/86. This was true for both the pretreatment tower and the nitrifying biofilter.

There is evidence in this study that high hydraulic loading rates, greater than 0.8-1.0 1/sq. m • s, have the potential of reducing or totally eliminating the nitrification activity in the upper section(s) of biofiIters when influent D.O. concentrations are low. Similar
observations were reported by Richards and Reinhart (1986). Intermittent dosing, which produces higher instantaneous hydraulic loadings and shear forces, is disadvantageous in this regard. The thicker biofilms and D.O. deficiencies associated with cold-weather operation may make biofilms more susceptible to shearing in winter months.

There is strong evidence that high hydraulic loadings, particularly when intermittent dosing is in use and cold temperatures prevail, prevent the return of nitrification activity once the nitrification activity in the upper sections has been depleted. This can be verified by inspection of the nitrifying biofilter data of 12/14/86 to 12/19/86 (Figures 31 and 34), 1/19/87 to 1/27/87 (Figures 33 and 39), and 2/13/87 (Figure 40).

Intermittent dosing was expected to reduce the hydraulic residence time in the nitrifying biofilter due to the increase in instantaneous superficial velocities. To test this theory, tracer analyses were performed at similar average hydraulic loadings of 1.12 to 1.23 1/sq.  $m \cdot s$  for both continous and intermittent dosing cases. NH<sub>A</sub>Cl was used as the tracer. The average hydraulic residence time was defined as the time required to collect 50% of the applied Cl<sup>-</sup> in the biofilter effluent. Biofilter effluent background concentrations of Cl<sup>-</sup> were accounted for.

The average contact time of  $NH_A^+$  was defined as the time required to collect 50% of the applied  $NH_4^+$ . Biofilter effluent background  $NH_4^+$ concentrations were accounted for. Since effluent  $NH_4^+$ -N concentrations

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were greater than 4 mg/l, a constant  $NH_4^+$ -N mass removal across the biofilter depth, independent of the bulk liquid  $NH_A^+$ -N concentration, was assumed. For this condition, the influent  $NH_4^{+-}N$  mass increase due to spiking would produce an equal effluent  $NH_A^+$ -N mass increase.

The results of the hydraulic residence tests are shown in Figures 46 and 47. The average hydraulic residence times for average hydraulic loadings of 1.2 1/sq. m • s were 92 seconds and 89 seconds, respectively, for intermittently-dosed and continuously-dosed conditions. The average  $NH_{4}^{+}-N$  contact time was, for all practical purposes, the same as the average hydraulic residence time for both types of wastewater application. These results are in agreement with literature reports that intermittent dosing and continuous dosing provide essentially equal trickling filter performance (Richards and Reinhart, 1986 and Albertson and Davies, 1984).

The average hydraulic residence times determined in this study are in good agreement with the biofilm-free hydraulic residence times reported for 3 m of 60° cross-flow media by Harrison and Daigger (1985). The residence times determined in the Ames WPCP study were only 1/5 of their reported values for media with biofilm present, despite the fact that active nitrifying biofilm was present in the Ames WPCP study when the hydraulic residence tests were run. The hydraulic residence times reported by Richards and Reinhart (1986) for 3 m of 60° cross-flow media with biofilm present agree fairly well with those reported by Harrison and Daigger (1985). Both of the literature studies involved heterotrophic biofilms, while the present study involved a nitrifier biofilm.



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Figure 46. Hydraulic residence test on December 19, 1986



Figure 47. Hydraulic residence test on January 12, 1987

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Thus, it appears that nitrifier biofilms have little effect on hydraulic residence times.

Richards (1984) and Richards and Reinhart (1986) reported that intermittent dosing produced shorter hydraulic residence times than continuous dosing for hydraulic loading rates greater than 0.5 1/sq. m • s. Inspection of their published data for 6.1 m towers containing 60 degree, cross-flow media with specific surface areas of 98 sq.m/cu. m (Richards, 1984), reveals that hydraulic residence times for average hydraulic loading rates of 0.5 1/sq. m • s, but with different application methods, agree within 7 percent or less.

The results of this study indicate that intermittent dosing had little effect at low hydraulic loading rates and at high wastewater temperatures and corresponding higher influent percentages of D.O. saturation (Figures 28, 29 and 45). However, intermittent dosing, which more closely resembles the rotary distributor application of full-scale trickling filters, appears to be detrimental for cold-weather operation (Figures 31-40). These detrimental effects of intermittent dosing stiould be minimized by operating at lower average hydraulic loading rates in winter months as evidenced by Figures 33, 34, 35, 36, 39 and 40.

#### Media effects

Partial plugging of the 60° cross-flow media in the nitrifying biofilter, particularly at the 2.44-3.66 m depth, was suspected at various times during the study. However, the tower has not been disassembled.

and these suspicions have not been verified. Partial plugging of similar 60° cross-flow media was reported by Richards and Reinhart (1986). That reported incidence indicated that plugging occurred primarily at the contact points within the media pack where two corrugated plastic sheets touched and at the interface between stacked pack units. Richards and Reinhart (1986) also reported that partial plugging of the media occurred to a greater extent in the winter months, a finding largely substantiated by the transient, reduced performance of various tower sections in this study, as well.

Partial plugging of cross-flow media channels would not prevent redistribution of water and air flow to other unrestricted channels. Nonetheless, overall efficiency in partially plugged media sections would be reduced due to lower effective media surface area, higher hydraulic loading per unit of effective media surface area, and lower oxygen transfer efficiency.

The performance of the bottom section of 45° cross-flow media in the pretreatment tower was very consistent, both in warm and cold weather. The top section performed much the same as that of the nitrifying biofilter, reflecting the effects of high hydraulic loading and/or D.O. deficient wastewater. Plugging potential of the 45° cross-flow media could not be evaluated because of the design of the tower, i.e., the individual media packs were not stacked in direct contact and had no common interfaces where plugging would occur.

The effective media specific surface area is a poorly understood parameter in trickling filter processes. Estimated effective surface areas may differ by 100%, depending on the method used to calculate the area. The Onda equation agrees fairly well with the work of Hosono et al. (1980) and the lower range of the calculations of Gullicks and Cleasby (1986). The percolation theory, discussed by Crine (1986), agrees closely with the reported effective areas calculated by Atkinson and Abdel Rahman Ali (1976).

Nitrification rates predicted by Wanner and Gujer (1984) and calculated using the models of Strand (1986) and Harremoes (1982) for wastewater conditions similar to those at the Ames WPCP are summarized in Table 11. These values are applicable to 100% effective surface areas. Some typical nitrification rates observed in the Ames WPCP study are shown in Table 12. The Ames WPCP nitrification rates in Table 12 have been compared to the theoretical maximum nitrification rates predicted by the Strand and Harremoes models in order to estimate the percent of effective media surface area for various hydraulic loading rates.

These effective surface area percentages have been compared to the percentages of effective surface area predicted by the Onda et al. (1968) and percolation theory (Crine, 1986) curves presented in Figures 20 and 21, respectively. The percolation theory relationship appears to give the best agreement with maximum observed effective media surface area percentages, and, therefore, probably provides a better estimate of the wetted specific surface area. However, the average effective media surface area for the entire tower depth was usually intermediate between the Onda and percolation theory predictions.

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Converted to 10°C basis by the use of the Nernst-Einstein equation.

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bMeasured at the biofilm surface.

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Table 12. Ames WPCP nitrification rates, corresponding effective area based on maximum predicted nitrification rates, and predicted effective areas by the Onda correlations and percolation theory

	Influent Conc.
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a<sub>Not converted to 10°C basis because of range of constants</sub> assumed.

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 $^{b}$ Based on 4/18/86 data. Total COD 100-125 mg/l.

 $c_{D.0.}$  Conc. on 12/4/86.



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# **Table 12. (Continued)**





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### Previous operating history effects

The effects of previous operating conditions on nitrification performance cannot be separated from the discussion of the effects of the factors already discussed. It has been documented in this study that the nitrifying biofilter performed satisfactorily at high hydraulic loading rates for about one week to ten days after the onset of cold weather. The performance of the upper section of the pretreatment tower dropped off sooner. Thereafter, the nitrifying biofilter performance dropped off, probably due to both the documented reduction in bulk liquid D.O. saturation and the high shear forces associated with high hydraulic loads.

The difficulty experienced in the Ames WPCP study with reestablishing nitrification performance lost during cold weather at high hydraulic loading rates is a significant finding. The operation of nitrification towers at low hydraulic loading rates would help prevent the initial loss of performance and has been shown to promote the return of satisfactory nitrification performance during cold weather operation.

## Modification of the Design Curves of Gullicks and Cleasby (1986)

Data from this study, the work of Duddles and Richardson (1973), Baxter and Woodman, Inc. (1973), Richards (1984), Richards and Reinhart (1986), Parker and Richards (1985), Sampayo and Metcalf (1984), and Sampayo (1973) have been assembled in the Appendix. The rates of

nitrification in the appendix have been adjusted to a common 10°C basis by the use of the Nernst-Einstein equation. The data have been used to construct a nitrifying biofilter design curve (Figure 48) for wastewater temperatures of 10°C similar to those presented by Gullicks and Cleasby (1986). Only Ames WPCP data are shown in Figure 48.

The 10°C nitrification rates were plotted versus both the total applied  $NH_{4}^{+}-N$  concentrations and the total applied hydraulic loading rates, including recirculated  $NH_4^+$ -N and flow. Winter discharge standards for  $NH_4^+$ -N are usually less stringent than 3 mg/1, and it has been shown that nitrification rates above 3-4 mg/1 are relatively independent of  $NH_A^+$ -N concentration. The design curve was constructed using data for which the effluent concentrations from the biofilters (or sections of biofilters) generally exceeded 3 mg  $NH_A^+$ -N/1. Some data for which the effluent concentrations from biofilters (or sections of biofilters) were less than 3 mg  $NH_A^+$ -N/1 were included in cases where oxygen still appeared to be the rate-limiting reactant.

The new curve can be used in exactly the same manner described by Gullicks and Cleasby (1986). The new design curve was constructed primarily with data from towers which contained 6-6.55 m of plastic media with specific surface areas of  $89-98$  m<sup>2</sup>/m<sup>3</sup>. No safety factor is implied in the design curve.

Additional surface area should be provided by increasing the tower media depth to account for the desired safety factor and to account for bulk liquid percent of D.O. saturation less than 65% and/or the partial plugging potential of media with specific surface areas greater than

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Figure 48. Predicted NH $_{\Lambda}^+$ –N removal, kg/d • m $^2$  of media surface, versus applied hydraulic load and applied  $NH_A^+$ -N for nitrification of municipal secondary clarifier effluent (BOD5 < 30 mg/1 and SS < 30 mg/1), wastewater temperature 10°C and media depths of 6-6.55 m of vertical or cross-flow plastic media (specific surface =  $89-98$  m<sup>2</sup>/m<sup>3</sup>). No safety factor is implied. Allow additional surface area (25-50%) for bulk liquid percent of D.O. saturation less than 65% and/or higher specific surface area media to account for plugging

98 m'/m'. If the percent of D.O. saturation is less than 65%, the secondary effluent could be aerated prior to application onto the plastic media so that additional media surface area would not have to be provided.

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#### **DISCUSSION**

The experimental equipment, procedures, assumptions, and results used in the investigation and analysis of nitrification in separatestage nitrifying biofilters at the Ames WPCP were included in the initial presentations of those topics and will not be repeated. There are aspects of this investigation that require further discussion and development in order to clarify some of the observations and results and to keep the conclusions in the proper, limiting perspective.

#### General Discussion

The primary data obtained in the Ames WPCP study were nitrogen parameter concentrations at various depths in a structured, plastic-media, separate-stage nitrifying biofilter. A variety of secondary effluent application rates and conditions were examined. Additional chemical parameters were analyzed, within the monetary and personnel constraints of the project budget, to assist in the evaluation of the nitrification performance of the nitrifying biofilter.

### Experimental considerations

The existing Ames, WPCP is an overloaded facility. Therefore, the secondary effluent from this conventional trickling filter plant exhibited greater variability than the secondary effluent applied in nitrifying biofilter studies examined in the literature. The overloaded condition of the Ames plant had an impact on the carbonaceous oxygen demand (measured by the COD test in this study), the suspended solids.

and the D.O. concentrations in the secondary effluent that served as the influent to the pilot-scale nitrification tower.

Wastewater, including secondary effluent, at any plant exhibits seasonal and diurnal variability in pollutant concentrations. The overloaded condition of the Ames plant increased the seasonal secondary effluent variability, however, diurnal variability was not observed to be excessive. The seasonal fluctuation of the population of Ames, revolving around the academic calendar at Iowa State University, also affected the seasonal variability of the load on the existing Ames WPCP, and, subsequently, the variability of the secondary effluent pollutant concentrations from the plant.

The seasonal variability of the influent to the pilot-scale nitrifying biofilter made it necessary to supplement the  $NH_4^+$ -N concentration in the biofilter influent with  $NH_4C1$  during data collection periods when Iowa State University was not in session. Data collected during March, 1986 did not involve  $NH_4^+$ -N supplementation. The low  $NH_4^+$ -N concentrations in the Ames WPCP secondary effluent during the second and third weeks of March, 1986 may have retarded the pilot nitrifying biofilter's initial approach to pseudo-steady state.

Alkalinity, COD, pH, suspended solids, and D.O. measurements were performed throughout the project duration, within the project budgetary and personnel restrictions, to assist in evaluation of the nitrification performance of the pilot-scale nitrifying biofilter. Manual sampling was performed simultaneously at the sampling points of both the pretreatment filter and the nitrifying biofilter whenever possible. When

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manual sampling was not possible for the investigator, automated sampling was instituted at selected nitrifying biofilter sampling points. Sampling probe location was critical during automatic sampling of the nitrifying biofilter effluent collection tank to prevent disturbance of sediments in the collection tank during the backflushing of the sampling tube and resulting high solids in the collected samples.

Analytical parameters associated with suspended solids, such as COD and TKN, were occasionally biased in automatic sampling at this sampling point when the backflushing of the sampling tube disturbed sediments in the effluent collection tank. Soluble parameters, such as  $NH_{4}^{+}-N$  and  $NO<sub>2</sub> + NO<sub>3</sub>$ -N would not be significantly affected by the increased suspended solids occasionally captured by the automated sampler. D.O. measurements for samples, taken manually from the nitrifying biofilter sampling ports, as well as those reported by the Ames WPCP personnel for the secondary effluent, proved to be critical to proper evaluation of the pilot-scale nitrifying biofilter performance.

Previous studies from the literature did not measure the D.O. concentration in the influent to the nitrifying biofilters and rarely measured the D.O. concentration in the effluent from nitrifying biofilters. The D.O. concentrations in the effluent from nitrifying biofilters, which were reported in some studies, were similar to those measured in the Ames study. However, previous investigators failed to associate the reported D.O. values with the observed nitrification performance. In retrospect, additional D.O. concentration measurements would have been desirable in the evaluation of nitrification at the Ames

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WPCP; however, that fact was not apparent until a detailed evaluation of the project data was completed.

Considerable effort was expended to prevent or minimize short circuiting through the Ames nitrifying biofilter, as described in the Experimental section. However, the possibility of short circuiting effects cannot be completely ruled out in any study.

The structured, plastic media used in the Ames study, nitrifying biofilter had a higher specific surface area and more complicated geometry than media used in the majority of nitrifying biofilter studies in the literature. Partial plugging of the 60 degree, cross-flow media, with a specific surface area of 138 sq. m/cu. m, was suspected at various times and at various depths in the biofilter. The construction of the biofilter tower available for the Ames study was not suitable for examination of the media at various depths to verify or refute the partial plugging suspicions without disassembly. The implications of partial plugging of the media in the biofilter have been presented in the Results and Analysis section and will not be repeated here.

There were brief periods of unexplained low nitrification performance during this study. No direct evidence of toxicity was observed. The effects of toxicity could not be evaluated extensively in the Ames study because of the limited project budget. Ames is not a heavily industrialized city, however, the research facilities associated with Iowa State University and government agencies and the industries located in Ames cannot be ruled out as potential sources of toxicity.

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#### CONCLUSIONS AND RECOMMENDATIONS

The Ames WPCP pilot-scale, separate-stage nitrifying biofilter study was similar to studies reported in the literature which applied settled secondary effluent to the media in the biofilters. However, the overloaded condition of the Ames WPCP allowed the examination of nitrification in separate-stage biofilters at slightly greater organic loadings than previous studies. The Ames study was an important contribution to the separate-stage, nitrifying biofilter data base for a number of reasons previously outlined in the Results and Analysis section. It emphasized cold-climate performance and oxygen flux-limited operating conditions.

Many important conclusions can be drawn from the results of the Ames study. Some of these observations have not been reported in previous literature or are contrary to findings reported in previous literature. The significant findings from the Ames study were:

- 1. D.O. is critical to nitrifying biofilter performance when oxygen flux limitation conditions exist. When the percent of D.O. saturation drops below 60 to 65 percent, nitrification performance will be diminished. When the percent of D.O. saturation drops below 45 to 50 percent, nitrification may be totally eliminated. This finding suggests that tower influent aeration and/or forced draft ventilation of nitrifying biofilters may be desirable.
- 2. Carbonaceous oxygen demand has.greater impact on nitrification performance at low hydraulic loading rates than at high hydraulic loading rates.

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- 3. The bulk liquid filterable (< 0.45 micron) COD concentration necessary to cause elimination of nitrifier populations from the biofilm is greater than 60 mg/1. However, competition between heterotrophs and nitrifiers can occur at somewhat lower filterable COD concentrations resulting in partial reduction in nitrification performance.
- 4. Nitrification is not limited by bulk liquid alkalinity concentrations greater than 30 mg/1 as  $CaCO<sub>3</sub>$ .
- 5. Above bulk liquid  $NH_4^+$ -N concentrations of about 4 mg/l, nitrification in biofilters is essentially zero-order relative to bulk liquid  $NH_A^+$ -N concentrations and linear with respect to tower depth and is, therefore, oxygen flux-limited.
- 6. Denitrification can occur in the upper levels of separate-stage, nitrifying biofilters when bulk liquid  $NO_2^- - N$  and  $NO_3^- - N$  concentrations are high, when filterable COD concentrations as low as 50 mg/1 exist in the bulk liquid, and when bulk liquid 0.0. concentrations are low.
- 7. Nitrification performance is improved by recirculation when bulk liquid D.O. concentrations in the influent wastewater stream are inadequate for maximum nitrification performance. Temperature changes in the wastewater resulting from recirculation should be considered in cold-climate operation when recycle decisions are being made. However, the effects of small temperature changes can be predicted adequately by the Nernst-Einstein equation when the percent of D.O. saturation in the bulk liquid is greater than 65

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percent and carbonaceous oxygen demand does not foster heterotroph competition within the biofilm.

- 8. When intermediate clarification is used between the carbon oxidation stage biofiIters and the nitrifying stage biofiIters, the detrimental effect of clarification due to dissolved oxygen reduction must be considered, as well as the beneficial effects of carbonaceous oxygen demand and suspended solids loading reductions. Relatively short detention in the intermediate clarifiers may be desirable to minimize dissolved oxygen reduction. This would also reduce heat loss effects in winter operation.
- 9. Hydraulic loading rates greater than  $0.8$  1/sq. m  $\cdot$  s are detrimental to stable operation of nitrifying biofiIters in cold-weather operation. This is particularly true when intermittent dosing is used and when bulk liquid D.O. concentrations are low. Low bulk liquid D.O. concentrations, though not necessarily below typical summer D.O. concentrations, are likely to be the case in the colder weather conditions. In cold weather, first-stage biological treatment is impaired and continued metabolism in intermediate clarifiers results in greater reductions in D.O. concentrations. This, coupled with reduced diffusivity of D.O. in the bulk liquid and biofilm in cold weather, results in reduced oxygen mass transfer into the biofilm.
- 10. If nitrification performance in sections of the nitrifying biofilter is lost during cold-weather operation, high hydraulic loading rates will prevent the return of nitrification performance. Low

hydraulic loading rates are desirable to prevent the initial loss of nitrification performance in cold weather, as well as to assist in restoring nitrification performance if it is temporarily lost.

- 11. Nitrifier populations can be established in fixed-film reactors even in cold-weather start-up operations provided a treated effluent from another plant containing active nitrifiers is used for seeding.
- 12. Near-steady state operation of nitrifying biofiIters can be established in six to seven weeks in warm weather operation. Two to three additional weeks may be required in cold weather.
- 13. Actively nitrifying biofilms respond quite rapidly to environmental changes. The majority of acclimation to new environmental conditions occurs within one to two days.
- 14. The 60-degree, cross-flow media with a specific surface area of 138 sq. m/cu. m used in this study appeared to have a tendency toward partial plugging. The partial plugging appeared to be transient, both in regard to tower depth location and longevity at any location. Partial plugging may persist at a given depth for fairly long periods of time. The biofiIter used in this study has not been disassembled, and confirmation or negation of the partial plugging hypothesis has not been made.
- 15. The cold weather empirical design curve (Figure 10) proposed by Gullicks and Cleasby (1986) was revised to reflect additional data from this pilot-scale study and the literature. The revised design curve (Figure 48) presents an empirical relationship between the

area! rate of nitrification and the total applied hydraulic load and the applied  $NH_A^+$ -N concentration, including recirculated flow and  $NH_A^+$ -N, for 10°C secondary effluent. The design curve was constructed primarily with data from towers which contained 6-6.55 m of plastic media with specific surface areas of 89-98 m'/m'. No safety factor is implied in the design curve. The new design curve can be used in exactly the same manner described by Gullicks and Cleasby (1986) for the use of Figure 10. Additional surface area should be provided by increasing the media depth to account for the desired safety factor and to account for bulk liquid percent of D.O. saturation less than 65% and/or the partial plugging potential of media with specific surface areas greater than 98  $m^2/m^3$ . Nitrification performance conformed well with the previous design curve (Figure 10) when continuous secondary effluent dosing was used. When intermittent dosing was used, the rate of nitrification was less than that predicted by the previous design curve (Figure 10).

16. Percolation theory (Figure 21) predicted the maximum effective media surface areas observed in this study quite well. However, the average effective media surface areas observed in this study were intermediate between the effective areas predicted by the Onda correlations (Figure 20) and by the percolation theory (Figure 21).

The Ames study has been beneficial to improving the design and operation of nitrifying biofilters. Additional studies are desirable to further expand the design and operation knowledge of nitrifying biofilters. Topics requiring further research include:

- 1. More detailed studies of the effects of bulk liquid D.O. concentrations on the nitrification performance of biofiIters, including forced draft ventilation effects and influent aeration effects,
- 2. Detailed studies of the effect of intermediate clarifier detention time on the nitrification performance of biofiIters,
- 3. Evaluation of partial plugging potential and its effects on nitrification performance,
- 4. Additional evaluation of the effect that intermittently-dosed versus continuously-dosed wastewater application has on nitrification, and comparison of full-scale nitrifying biofiIter performance to pilot-scale performance for intermittent versus continuous wastewater dosing, and
- 5. Additional cold-climate nitrification evaluation.

Careful attention should be given to the design of pilot-scale biofilters used in researching these topics. The design should allow direct visual observation of the media and removal of media segments at various depths in the tower. Bulk liquid sampling should be possible at various depths along the tower so that nitrification performance can be related to individual segments of the tower. The effects of D.O., temperature, carbonaceous oxygen demand, pH, and alkalinity should all be considered.

Some of the effects of D.O. concentrations on nitrification performance may be studied effectively on a laboratory scale. The study

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of other D.O. effects, particularly related to intermediate clarifica tion and forced draft ventilation, could best be evaluated with pilot scale or full-scale facilities.

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**APPENDIX: EXPERIMENTAL AND CALCULATED DATA** 

**Some of the terms used in the appendix merit explanation. The term total flow means the influent flow plus the recycle flow, when recycle was in use. The term DIO/DT is the Nernst-Einstein correction (multiplier) for temperature effects. Influent concentrations were the parameter concentrations influent to the specified section of the biofilter tower. The term oxidized N was used to represent the change in the concentrations of the various specified nitrogen forms for any given section of the biofilter tower. Thus, oxidized nitrogen is really a misnomer that was used in order to facilitate the spreadsheet organization.** 

**The data of Buddies and Richardson (1973) and Baxter and Woodman, Inc. (1973) were the data from the Midland, Michigan and Bloom Township, Illinois studies, respectively, that were used in construction of the original design curves of Gullicks and Cleasby (1986). However, in this appendix, the data have been converted to a common 10 degree C basis by the Nernst-Einstein equation.** 

**The influent-effluent data from the Ames WPCP study were often collected on the same day as the biofilter profile data. However, these data represent distinct sampling times when complete profile data were not collected simultaneously.** 

Ames WPCP

## Pilot-Scale Nitrifying Biofilter

## Tower Profile Data

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**Source: AMES Date: 031086 Avg. Total Flow, 1/ sq. m x s: .468 Recycle, percent total flow: 59 Periodic Dosing Cycle, sec: 73**  Distribution Time, sec/cycle: **Avg. Distributed Flow, 1/sq. m x s: 1.3 Media Spec. Surf. Area, sq.m/cu. m: 137.8 •nda % Wetted Area: 26 Water Temp., C: Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/11 Media Influent Oxidized 10 C**  Concentration **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 4 1.3 .0003126 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N 2.7 1.1 **1.2-2.4 NH4-N 2.7 1.1 .0002645 TKN** 0 **N02+N03 0 Filt. TKN O 2.4-3.6 NH4-N 1.6 .6 .0001443 TKN** 0 **N02+N03 0 Filt. TKN**<br>
NH4-N
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1
4 **3.7-4.9 NH4-N 1 .4 .0000962 TKN** 0 **N02+N03 0 Filt. TKN 0** 

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**Source: AMES Date: 031186**<br>**Avg. Total Flow, 1/ sq. m x s: 031186 Avg. Total Flow, 1/ sq. m x si .458 Recycle, percent total flow: 59 Periodic Dosing Cycle, secs 73**  Distribution Time, sec/cycle: 27<br>Avg. Distributed Flow, 1/sq. m x s: 1.3 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. ms 137.8 Onda % Wetted Areas 26 Water Temp., Cs 10 DIO/DT: 1 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. 0. O., mg/1: Media Influent Oxidized 10 C Concentration Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 1.4 .5 .0001177 TKN O N02+N03 0**  Filt. TKN<br>NH4-N .9 **1.2-2.4 NH4-N .9 .5 .0001177 TKN O N02+N03 O Filt. TKN**<br>NH4-N .4 .2 **2.4-3.6 NH4-N .4 .2 .0000471 TKN** 0 **N02+N03 O Filt. TKN**<br>NH4-N .2 .1 **3.7-4.9 NH4-N .2 .1 .0000235 TKN O N02+N03 O**  Filt. TKN

 $\sim 10^{11}$  MeV  $\sim 10^{11}$ 

**Sources AMES Date: 031386**  Avg. Total Flow, 1/ sq. m x s: **Recycle, percent total flow: 59**  Periodic Dosing Cycle, sec: 73<br>Distribution Time, sec/cycle: 77<br>27 Distribution Time, sec/cycle: **Avg. Distributed Flow, 1/sq. m x s: 1.3 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 26 Water Temp., C: 8.8 DIO/DT: 1.043 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C pth Concentration Nitr** $\begin{array}{ccc} \text{min} & \text{min} \ \text{min} & \text{min} \ \text{min} & \text{min} \end{array}$ **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 1.8 .8 .0001963 TKN O N02+N03 0 Filt. TKN**<br>NH4-N 1 .6 **1.2-2.4 NH4-N 1 .6 .0001473 TKN O N02+N03 O Filt. TKN O 2.4-3.6 NH4-N .4 .2 .0000491 TKN** 0 **N02+N03 0**  Filt. TKN<br>NH4-N .2 **3.7-4.9 NH4-N .2 -.1 -.0000245**  TKN 0 **N02+N03 0 Filt. TKN 0** 

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**Source: AMES Date: 031386 Avg. Total Flow, 1/ sq. m x s: .458 Recycle, percent total flow: 59 Periodic Dosing Cycle, sec: 73**  Distribution Time, sec/cycle: **Avg. Distributed Flow, 1/sq. m x a: 1.3**  Media Spec. Surf. Area, sq.m/cu. m: **137.8**<br>Onda % Wetted Area: 26 **Onda % Wetted Area: 26 Water Temp., C: 8.8 DIO/DT: 1.043 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C Concentration Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 3.2 1 .0002454 TKN O N02+N03 0 Filt. TKN** 0<br> **NH4-N** 2.2 1 .0002454 **1.2-2.4 NH4-N 2.2 1 .0002454 TKN O N02+N03 0 Filt. TKN O 2.4-3.6 NH4-N 1.2 .6 .0001473 TKN** 0 **N02+N03 O**  Filt. TKN<br>NH4-N **3.7-4.9 NH4-N .6 .2 .0000491 TKN 0 N02+N03 O**  Filt. TKN

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Source: AMES **Dates 031386 Avg. Total Flow, 1/ sq. m x st .458 Recycle, percent total flow: 59 Periodic Dosing Cycle, sec: 73 Distribution Time, sec/cycle: 27 Avg. Distributed Flow, 1/sq. m x s: 1.3 137.8 Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: 26 1.036**  9 **DIO/DTJ Water Temp., C: COD, mg/1 : Infl. Filt. COD, mg/1: Infl. Infl. BODS, mg/1 : Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: Infl. Alk, mg CaC03/l: pH: 7.9 Infl. D. 0. mg/1 : Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 3.5 1 .0002438 TKN** 0 **N02+N03 0 Filt. TKN** 2.5 1 **1.2-2.4 NH4-N 2.5 1 .0002438 TKN** 0 **N02+N03 0 Filt. TKN**<br>
NH4-N 1.5 .9 **2.4-3.6 NH4-N 1.5 .9 .0002194 TKN** 0 **N02+N03 0 Filt. TKN O 3.7-4.9 NH4-N .6 0 0 TKN** 0 **N02+N03 0 Filt. TKN 0** 

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 $\sqrt{\mathbf{g}}_k^{\prime}$   $\mathbf{g}$   $\mathbf{g}_k$ 

Source: AMES **Dates** 031386 **Avg. Total Flow, 1/ sq. m x s: .458 Recycle, percent total flow: 59**  Periodic Dosing Cycle, sec: 73<br>Distribution Time, sec/cycle: 73 **Distribution Time, sec/cycle: 27 Avg. Distributed Flow, 1/sq. m x s: 1.3 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 26 Water Temp., C: 9 DIO/DT: 1.036 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: 8 Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C Concentration Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 3.4 1 .0002438 TKN O N02+N03 0 Filt. TKN 0 1.2-2.4 NH4-N 2.4 1 .0002438 TKN O N02+N03 O Filt. TKN 0 2.4-3.6 NH4-N 1.4 .8 .000195 TKN O N02+N03 0**  Filt. TKN 0<br>
NH4-N 0.6 1 0000244 **3.7-4.9 NH4-N .6 .1 .0000244 TKN O N02+N03 O**  Filt. TKN

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**Source: AMES** Date: 031386<br>
Avg. Total Flow, 1/ sq. m x s: 031386 **Avg. Total Flow, 1/ sq. m x si .458 Recycle, percent total flow: 59**  Periodic Dosing Cycle, sec: 73<br>Distribution Time, sec/cycle: 773 Distribution Time, sec/cycle: **Avg. Distributed Flow, 1/sq. m x s: 1.3 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 26 Water Temp., C: 9 DIO/DT: 1.036 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: 8 Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized .10 C Concentration m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 3.2 .9 .0002194 TKN O N02+N03 0 Filt. TKN O 1.2-2.4 NH4-N 2.3 1 .0002438 TKN O N02+N03 0 Filt. TKN**<br>
NH4-N 1.3 .7 **2.4-3.6 NH4-N 1.3 .7 .0001707 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N .6 .1 **3.7-4.9 NH4-N .6 .1 .0000244 TKN O N02+N03 0**  Filt. TKN

 $\omega \propto \omega_{\rm{eff}}$  .

**Sources AMES Date: 031386 Avg. Total Flow, 1/ sq. m x s: .458 Recycle, percent total flows 59**  Periodic Dosing Cycle, sec: 73<br>Distribution Time, sec/cycle: 73 **Distribution Time, sec/cycle:** 27<br> **Avg. Distributed Flow, 1/sq. m x s: 1.3** Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. ms 137.8 Onda % Wetted Areas 26 Water Temp., C: Infl. COD, mg/ls Infl. Filt. COD, mg/ls Infl. B0D5, mg/ls Infl. Filt. BODS, mg/ls SS, mg/ls VSS, mg/ls pHs Infl. Alk, mg CaC03/l: Infl. D. O., mg/ls Media Influent Oxidized 10 C Concentration Nitrogen m mg/1 mg/1** kg/d **sq.m tot. surface area 0-1.2 NH4-N 3.2 1 .0002396 TKN** 0 **N02+N03 0 FiIt. TKN O 1.2-2.4 NH4-N 2.2 1.1 .0002635 TKN O N02+N03 0 Filt. TKN**<br>
NH4-N 1.1 .7 **2.4-3.6 NH4-N 1.1 .7 .0001677 TKN 0 N02+N03 0 Filt. TKN O 3.7-4.9 NH4-N .4 0 O TKN O**  TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 Filt. TKN

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**Source: AMES Avg. Total Flow, 1/ sq. m x s: Recycle, percent total flows Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s\* Date: 031486 .461 59 73 27 1.3 Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: 8.7 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. B0D5, mg/1: DlO/DTs SB, mg/1: pH: Infl. D. 0. mg/1 : 8 VSS, mg/1: Infl. Alk, mg CaC03/l: 137.8 26 1.047 Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 1.6 .6 .0001488 TKN** 0 **N02+N03 0 Filt. TKN**<br>
NH4-N 1 .6 **1.2-2.4 NH4-N 1 .6 .0001488 TKN** 0 **N02+N03 0 Filt. TKN O 2.4-3.6 NH4-N .4 .2 .0000496 TKN O N02+N03 0 Filt. TKN** 0<br>
NH4-N 12 0 0 **3.7-4.9 NH4-N .2 0 0 TKN O**  TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 **Filt. TKN 0** 

 $\omega(t) \lesssim \omega$  )



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**Sources AMES Dates 031986 Avg. Total Flow, 1/ sq. m x sa .485 Recycle, percent total flows 30 Periodic Dosing Cycle, sees 67 Distribution Time, sec/cycles 23**  Avg. Distributed Flow, 1/sq. m x st **Media Spec. Surf. Area, sq.m/cu. ms 137.8 Onda % Wetted Areas 27 Water Temp., Cs 7.4 DlO/DTs 1.094 Infl. COD, mg/ls Infl. Filt. COD, mg/ls Infl. B0D5, mg/ls Infl. Filt. B0D5, mg/ls SS, mg/ls VSS, mg/ls pHs Infl. Alk, mg CaC03/l: Infl. D. O., mg/ls 10 C Nit. Rate kg/d sq.m tot. surface area 0-1.2 NH4-N 6.3 1.2 .0003271 TKN O N02+N03 0 Filt. TKN**<br>NH4-N 5.1 1.6 **1.2-2.4 NH4-N 5.1 1.6 .0004362 TKN O N02+N03 O Filt. TKN** 3.5 1 **2-4-3.6 NH4-N 3.5 1 .0002726 TKN O N02+N03 0 Filt. TKN**<br> **RH4-N** 2.5 .7 **3.7-4.9 NH4-N 2.5 .7 .0001908 TKN** 0 **N02+N03 0 Filt. TKN O 0-4.9 NH4-N 6.3 4.5 .0003067 TKN** 0 **N02+N03 0 Filt. TKN Media Influent Oxidized Concentration Nitrogen m mg/1 mg/1** 

 $\omega(r) \lesssim \sqrt{r}$ 

**Source: AMES Dates** 031986 **Avg. Total Flow, 1/ sq. m x s: .501 Recycle, percent total flow: 31**  Periodic Dosing Cycle, sec: **67**<br>Distribution Time. sec/cycle: **67 1999 Distribution Time, sec/cycle: 23**  Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda 7, Wetted Area: 27 Water Temp., C: 8.7 DlO/DTr 1.047**  Infl. COD, mg/l: **Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: 7.9 Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C**  Concentration **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 8 1.2 .0003234 TKN O N02+N03 0 Filt. TKN**<br>NH4-N 6.8 1.8 **1.2-2.4 NH4-N 6.8 1.8 .0004851 TKN O N02+N03 O Filt. TKN** 0<br> **NH4-N** 5 1.4 .0003773 **2.4-3.6 NH4-N 5 1.4 .0003773 TKN O N02+N03 O Filt. TKN**<br>NH4-N 3.6 .5 **3.7-4.9 NH4-N 3.6 .5 .0001348 TKN O N02+N03 O**  Filt. TKN **0-4.9 NH4-N 8 4.9 .0003301 TKN O N02+N03 0 Filt. TKN** 

 $\omega(\lambda) \geq \omega(\lambda)$ 

**Source: AMES Date: 031986**<br>Avg. Total Flow, 1/ sq. m x s: **Date: 031986 Avg. Total Flow, 1/ sq. m x s: .507 Recycle, percent total flows 32 Periodic Dosing Cycle, sec: 67 Distribution Time, sec/cycle: 23**  Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area:** 27<br> **Water Temp., C:** 2.4 **D**10/DT: 1.058 Water Temp., C: **Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1 : VSS, mg/1: pH: 7.8 Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 7 1.2 .0003307 TKN** 0 **N02+N03 0**  Filt. TKN **1.2-2.4 NH4-N 5.8 1.7 .0004685 TKN** 0 **N02+N03 0 Filt. TKN** 0<br> **NH4-N** 4.1 1.1 .0003032 **2.4-3.6 NH4-N 4.1 1.1 .0003032 TKN O N02+N03 0 Filt. TKN 0 3.7-4.9 NH4-N 3 .4 .0001102 TKN O N02+N03 O**  Filt. TKN **0-4.9 NH4-N 7 4.4 .0003032 TKN** 0 **N02+N03 0**  Filt. TKN

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**Source: AMES Date: 032086 Avg. Total Flow, 1/ sq. m x s: .501 Recycle, percent total flow: 31 Periodic Dosing Cycle, sec: 65 Distribution Time, sec/cycle: 23**  Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 27 Water Temp., C: 7 DIO/DT: 1.109 Znfl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/li Infl. D. O., mg/1: Media Influent Oxidized 10 C Concentration Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 6.3 2.8 .0007993 TKN O N02+N03 0 Filt. TKN**<br>NH4-N 3.5 1.2 **1.2-2.4 NH4-N 3.5 1.2 .0003426 TKN O N02+N03 0 Filt. TKN O 2.4-3.6 NH4-N 2.3 1 .0002855 TKN O N024.N03 O**  Filt. TKN<br>NH4-N **3.7-4.9 NH4-N 1.3 .2 .0000571 TKN** 0 TKN<br>
NO2+NO3 0<br>
Filt. TKN 0 Filt. TKN **0-4.9 NH4-N 6.3 5.2 . 0003711 TKN** 0 **N02+N03 0 Filt. TKN 0** 

 $\omega\propto\omega_{\rm{min}}$ 

**Sources AMES Date: 032086**  Avg. Total Flow, 1/ sq. m x s: **Recycle, percent total flow: 32 Periodic Dosing Cycle, sec: 65 Distribution Time, sec/cycle: 23**  Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 27 Water Temp., C: 7.8 DIO/DT: 1.079 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: 8 Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C Concentration Nitrogen Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 2.6 .7 .0001983 TKN O N02+N03 0 Filt. TKN O 1.2-2.4 NH4-N 1.9 1 .0002833 TKN O N02+N03 0 Filt. TKN O 2.4-3.6 NH4-N .9 .5 .0001416 TKN** O **N02+N03 0 Filt. TKN O 3.7-4.9 NH4-N .4 .1 .0000283 TKN O N02+N03 O**  Filt. TKN **0-4.9 NH4-N 2.6 2.3 .0001629 TKN O N02+N03 0 Filt. TKN O** 

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**Sources AMES Dates 032086 Avg. Total Flow, 1/ sq. m x ss .503 Recycle, percent total flows 31 Periodic Dosing Cycle, secs 65**  Distribution Time, sec/cycle: **Avg. Distributed Flow, 1/sq. m x s: 1.4 Media Spec. Surf. Area, sq.m/cu. ms 137.8 Onda % Wetted Areas 27 Water Temp., Ci 8 DlO/DTs 1.072 Infl. COD, mg/ls** 105 **Infl. Filt. COD, mg/ls Infl. BODS, mg/ls Infl. Filt. BODS, mg/li SS, mg/ls VSS, mg/ls pHs Infl. Alk, mg CaC03/li Infl. D. O., mg/ls Media Influent Oxidized Concentration m mg/1 mg/1 10 C Nit. Rate kg/d sq.m tot. surface area**   $0 - 1.2$ **1.2-2.4 2.4-3.6 3.7-4.9 NH4-N TKN N02+N03 Filt. TKN 6.5 11 9.4 5 10.7 10.3 3.5 8**  12.2 **2.4 7.3 1.5 .3 .9 1.5 2.7 1.9 1.1 .7 .4 .6 .0004156 .0000831 .0002493**  O **.0004156 .000748 .0005264**  O **.0003047 .0001939 0**  O .0001108 .0001662 **0**  O **0-4.9 NH4-N TKN N02+N03 Filt. TKN 6.5 11 9.4 4.5 4.3 4. 1 .0003117 .0002978 .000284**  O

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**Source: AMES Date: 033186**<br>Avg. Total Flow, 1/ sq. m x s: **Date: 033186 Avg. Total Flow, 1/ sq. m x ss .583 Recycle, percent total flow: 27 Periodic Dosing Cycle, sec: 59**  Distribution Time, sec/cycle: 18<br>Avg. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 31 Water Temp., C: 16.9 DIO/DT: .812 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. BODS, mg/1s Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1:**  Ŀ, **Media Influent Oxidized 10 C Concentration Nitrogen Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 12.7 .9 .0002189 TKN O N02+N03 0 Filt. TKN**<br>NH4-N 11.8 1.8 **1.2-2.4 NH4-N 11.8 1.8 .0004378 TKN O N02+N03 0**  Filt. TKN 0<br> **NH4-N** 10 1.3 0003162 **2.4-3.6 NH4-N 10 1.3 .0003162 TKN O N02+N03 0 Filt. TKN**<br>NH4-N 0.7 1.4 **3.7-4.9 NH4-N 8.7 1.4 .0003405 TKN O N02+N03 0 Filt. TKN O 0-4.9 NH4-N 12.7 5.4 .0003284 TKN** 0 **N02+N03 0**  Filt. TKN

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**Source: AMES Dates** 040286<br>
Avg. Total Flow. 1/ sq. m x s: **Dates** 040286 **Avg. Total Flow, 1/ sq. m x si .591 Recycle, percent total flow\* 27 Periodic Dosing Cycle, sees 59 Distribution Time, sec/cycle:** 18<br> **Avg. Distributed Flow.** 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. mi 137.8 •nda % Wetted Areas 31**  Water Temp., C: **Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1s Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C**  Concentration **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 17.1 1.1 .0003003 TKN O N02+N03 O Filt. TKN O 1.2-2.4 NH4-N 16 2.3 .0006279 TKN O N02+N03 0 Filt. TKN O 2.4-3.6 NH4-N 13.7 .3 .0000819 TKN** 0 **N02+N03 0 Filt. TKN O 3.7-4.9 NH4-N 13.4 1.2 .0003276 TKN** 0 **N02+N03 0**  Filt. TKN **0-4.9 NH4-N 17.1 4.9 .0003344 TKN O N02+N03 0 Filt. TKN O** 

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**Source: AMES Date: 040286 Avg. Total Flow, 1/ sq. m x si .577 Recycle, percent total flows 25**  Periodic Dosing Cycle, sec:<br>Distribution Time, sec/cycle: **59 18 18 Distribution Time, sec/cycle:** 18<br> **Ava. Distributed Flow. 1/sa. m x s: 1.9** Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. mi 137.8 Onda % Wetted Areas 31 Water Temp., Cs 12.7 DlO/DTs .922 Infl. COD, mg/ls Infl. Filt. COD, mg/ls Infl. BODiS, mg/ls Infl. Filt. B0D5, mg/ls SS, mg/ls VSS, mg/ls pHs Infl. Alk, mg CaC03/ls Infl. D. O., mg/ls Media Influent Oxidized 10 C Concentration m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 15.4 1.3 .0003553 TKN O N02+N03 0 Filt. TKN O 1.2-2.4 NH4-N 14.1 2.4 .000656 TKN O N02+N03 O**  Filt. TKN 0<br>NH4-N 11.7 .8 .0002187 **2.4-3.6 NH4-N 11.7 .8 .0002187 TKN O N02+N03 O Filt. TKN**<br>NH4-N 10.9 2 **3.7-4.9 NH4-N 10.9 2 .0005467 TKN O N02+N03 O**  Filt. TKN **0-4.9 NH4-N 15.4 6.5 .0004442 TKN** 0 **N02+N03 O**  Filt. TKN

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Sources AMES **Dates** 040286 **Avg. Total Flow, 1/ sq. m x ss .577 Recycle, percent total flow: 25**  Periodic Dosing Cycle, sec:<br>Distribution Time, sec/cycle: **59 18 18** Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.9 **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Areas 31 Water Temp., C: Infl. COD, mg/1I Infl. Filt. COD, mg/ls Infl. B0D5, mg/ls Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/ls pHs 7.8 Infl. Alk, mg CaC03/ls Infl. D. O., mg/ls 5.2 Media Influent Oxidized 10 C . Concentration Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 14.2 1.1 .0003036 TKN O N02+N03 0 Filt. TKN O 1.2-2.4 NH4-N 13.1 2.3 .0006348 TKN O N02+N03 0 Filt. TKN O 2.4-3.6 NH4-N 10.8 .7 .0001932 TKN O N02+N03 O Filt. TKN** 10.1 2.3 **3.7-4.9 NH4-N 10.1 2.3 .0006348 TKN O N02+N03 O**  Filt. TKN **0-4.9 NH4-N 14.2 6.4 . 0004416 TKN O N02+N03 O**  Filt. TKN

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**Sources AMES Date: 040386**  Avg. Total Flow, 1/ sq. m x s: **.578 .578** .578 **Recycle, percent total flows 24 Periodic Dosing Cycle, sect 59**  Distribution Time, sec/cycle: 18<br>Avq. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. ms 137.8 Onda y. Wetted Areas 31**  Water Temp., C: **Infl. COD, mg/ls Infl. Filt. COD, mg/ls Infl. B0D5, mg/ls Infl. Filt. BODS, mg/lt SS, mg/ls VSS, mg/ls pHs 7.7 Infl. Alk, mg CaC03/ls Infl. D. O., mg/ls 6.1 Media Influent Oxidized 10 C Depth Concentration Nitrogen<br>mmg/l** mg/l **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 8.3 .7 .0001942 TKN O N02+N03 0 Filt. TKN**<br>NH4-N 7.6 1.8 **1.2-2.4 NH4-N 7.6 1.8 .0004993 TKN** 0 **N02+N03 0 Filt. TKN 0 2.4-3.6 NH4-N 5.8 .8 .0002219 TKN O N02+N03 0 Filt. TKN O 3.7-4.9 NH4-N 5 2 .0005547 TKN O N02+N03 0**  Filt. TKN **0-4.9 i\IH4-N 8.3 5.3 . 0003675 TKN O N02+N03 0 Filt. TKN O** 

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**Source: AMES Date: 041086**<br> **Avq. Total Flow. 1/ sq. m x s: 041086 Avg. Total Flow, 1/ sq. m x ss .572 Recycle, percent total flow: 25**  Periodic Dosing Cycle, sec: **59 and Cycle Control Control** 59<br>Distribution Time. sec/cycle: **59 and Control Control** 18 **Distribution Time, sec/cycles** 18 **18**<br> **Ava. Distributed Flow. 1/sa. m x s:** 1.9 Avg. Distributed Flow, 1/sq. m x si **Media Spec. Surf. Area, sq.m/cu. mi 137.8 Onda % Wetted Areas 31 Water Temp., Ci 11.5 DlO/DTi .956 Infl. COD, mg/ls Infl. Filt. COD, mg/ls Infl. B0D5, mg/ls Infl. Filt. BOD5, mg/li SS, mg/ls VSS, mg/li pHi 7.7 Infl. Alk, mg CaC03/li Infl. D. O., mg/ls Media Influent Oxidized 10 C Concentration m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 12.6 1.2 .0003371 TKN O N02+N03 0 Filt. TKN** 11.4 1.6 **1.2-2.4 NH4-N 11.4 1.6 .0004495 TKN O N02+N03. 0 Filt. TKN**<br> **NH4-N** 9.8 1.7 **2.4-3.6 NH4-N 9.8 1.7 .0004776 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N 8.1 1.1 **3.7-4.9 NH4-N 8.1 1.1 .0003091 TKN O N02+N03 O Filt. TKN 0 0-4.9 NH4-N 12.6 5.6 .0003933 TKN** 0 **N02+N03 O**  Filt. TKN

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**Source: AMES Date: 041086**<br> **Avg. Total Flow, 1/ sq. m x s: 041086 Avg. Total Flow, 1/ sq. m x ss .589 Recycle, percent total flows 28**  Periodic Dosing Cycle, sec:<br>Distribution Time. sec/cycle: **59 18 18** Distribution Time, sec/cycle:<br>Avg. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **137.8 Media Spec. Surf. Area, sq.m/cu. ms Onda % Wetted Areas 31 Water Temp., Cs 12.7 DlO/DTs .922 Infl. COD, mg/Is Infl. Filt. COD, mg/li Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1s VSS, mg/1s pHs 7.7 Infl. Alk, mg CaC03/ls**  7 **Infl. D. D. mg/1 s Media Influent Oxidized 10 C Depth Concentration Nitrogen Millips**<br>mg/1 mg/1 mg/1 **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 14 1.5 .0004185 TKN O N02+N03 0 Filt. TKN**<br>NH4-N 12.5 2.3 **1.2-2.4 NH4-N 12.5 2.3 .0006417 TKN O N02+N03 O Filt. TKN** 0<br> **NH4-N** 10.2 1.4 .0003906 **2.4-3.6 NH4-**<br>TKN **TKN** 0 TKN 0<br>
NO2+NO3 0<br>
Filt. TKN 0 **Filt. TKN O 3.7-4.9 NH4-N 8.8 1.4 .0003906 TKN** O **N02+N03 O**  Filt. TKN **0-4.9 NH4-N 14 6.7 .0004674 TKN O N02+N03 O**  Filt. TKN

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**Source: AMES Dates 071486 Avg. Total Flow, 1/ sq. m x s: .586 Recycle, percent total flows 0 Periodic Dosing Cycle, sec: 58 Distribution Time, sec/cycle: 18**  Avg. Distributed Flow, 1/sq. m x s: 1.9 **Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1:**  21 **DIO/DT: pH: Infl, D. 0., mg/1: Infl Alk, mg CaC03/li VSS, mg/1: 137.8 32 .723 Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 2.2 1.1 .0002394 TKN O N02+N03 0 Filt. TKN 0 1.2-2.4 NH4-N 1.1 .7 .0001524 TKN O N02+N03 0 Filt. TKN 0 2.4-3.6 NH4-N .4 .3 .0000653 TKN** 0 TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 **Filt. TKN 0**  1 NH<br> **12+NO3**<br> **5.7-4.9 NH4-N .1**<br> **3.7-4.9 NH4-N .1**<br> **3.7-4.9 NH4-N .1**<br> **1**<br> **5.7-4.9** NH4-N .1 **TKN O N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 2.2 2.1 .0001143 TKN O N02+N03 0 Filt. TKN 0** 

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**Source: AMES Date: 071486 Avg. Total Flow, 1/ sq. m x s: .586 Recycle, percent total flow: 0**  Periodic Dosing Cycle, sec: 58<br>Distribution Time, sec/cycle: 58<br>18 **Distribution Time, sec/cycle: 18 Avg. Distributed Flow, 1/sq. m x s: 1.9 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Areas 32 Water Temp., C: Infl. COD, mg/ls Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/ls VSS, mg/ls pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/ls Media Influent Oxidized 10 C Concentration Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 3.7 1.8 .0003918 TKN** 0 **N02+N03 0 Filt. TKN 0 1.2-2.4 NH4-N 1.9 1.2 .0002612 TKN 0 N02+N03 0 Filt. TKN 0 2.4-3.6 NH4-N .7 .4 .0000871 TKN** 0 **N02+N03 O**  Filt. TKN 0<br>NH4-N 03 1 0000218 **3.7-4.9 NH4-N .3 .1 .0000218 TKN** 0 **N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 3.7 3.5 .0001905 TKN** 0 **N02+N03 0**  Filt. TKN

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**Source: AMES Date: 072186 Avg. Total Flow, 1/ sq. m x s: .591 Recycle, percent total flow: 0**  Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: 18<br>Avg. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 32 Water Temp., C: 22.5 DIO/DT: .695 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1:**  Media Influent **Oxidized** 10 C<br>Depth Concentration Nitrogen Nit.Rate epth Concentration Nitrہe<br>m mg/l mg/l **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 11.3 1.8 .0003799 TKN O N02+N03 0 Filt. TKN**<br> **NH4-N** 9.5 2.3 **1.2-2.4 NH4-N 9.5 2.3 .0004854 TKN O**  TKN 0<br>
NO2+NO3 , 0<br>
Filt. TKN 0 **Filt. TKN O 2.4-3.6 NH4-N 7.2 .4 .0000844 TKN O N02+N03 0 Filt. TKN**<br>  $NA4-N$  6.8 1.2 **3.7-4.9 NH4-N 6.8 1.2 .0002532 TKN** 0 **N02+N03 0**  Filt. TKN **0-4.9 NH4-N 11.3 5.7 .0003007 TKN O N02+N03 0 Filt. TKN O** 

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**Sources AMES Date: 072286 Avg. Total Flow, 1/ sq. m x s: .573 Recycle, percent total flow: 0 Periodic Dosing Cycle, sec: 58**  Distribution Time, sec/cycle: 18<br>Avg. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 32**  Water Temp., C: **Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C Concentration Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 16.2 2.5 .0004983 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N 13.7 3.4 **1.2-2.4 NH4-N 13.7 3.4 .0006777 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N 10.3 .6 **2.4-3.6 NH4-N 10.3 .6 .0001196 TKN** 0 **N02+N03 0 Filt. TKN**<br> **NH4-N** 9.7 2.1 **3.7-4.9 NH4-N 9.7 2.1 .0004185 TKN** 0 **N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 16.2 8.6 .0004285 TKN** 0 **N02+N03 0 Filt. TKN 0** 

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 $\sim 200$  MeV  $\sim$ 

**Source: AMES Date: 072386**<br>**Avg.** Total Flow, 1/ sq. m x s: **Date:** 072386 **.** 556 **Avg. Total Flow, 1/ sq. m x s: .556 Recycle, percent total flows** (1999)<br>Periodic Dosing Cycle, secs (1999) 1999 (1999) 1999 **Periodic Dosing Cycle, sec: 58**  Distribution Time, sec/cycle: **Avg, Distributed Flow, 1/sq. m x si 1.9 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Areas 32 Water Temp., Ci 24.5 DlO/DTs .658 Infl. COD, mg/ls Infl. Filt. COD, mg/lt Infl. B0D5, mg/ls Infl. Filt. BODS, mg/li SS, mg/ls VSS, mg/ls pHs Infl. Alk, rag CaC03/l: Infl. D. O., mg/ls Media Influent Oxidized 10 C Concentration m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 15.5 3.3 .0006203 TKN O N02+N03 0 Filt. TKN**<br>
NH4-N 12.2 4.3 **1.2-2.4 NH4-N 12.2 4.3 .0008083 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N 7.9 .7 **2.4-3.6 NH4-N 7.9 .7 .0001316 TKN** 0 **N02+N03 0 Filt. TKN 0 3.7-4.9 NH4-N 7.2 3 .0005639 TKN O N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 15.5 11.3 .000531 TKN** 0 **N02+N03 0 Filt. TKN 0** 

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**Source: AMES Date: 073086 Avg. Total Flow, 1/ sq. m x s: .578 Recycle, percent total flows O Periodic Dosing Cycle, sec: 58**  Distribution Time, sec/cycle: 18<br>Avg. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, aq.m/cu. m: 137.8 Onda % Wetted Area: 32**  Water Temp., C: **Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: 7.4 Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C Concentration Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 15.3 3.9 .0007841 TKN** 0 **N02+N03 0 Filt. TKN 0 1.2-2.4 NH4-N 11.4 4.7 .0009449 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N 6.7 1.8 **2.4-3.6 NH4-N 6.7 1.8 .0003619 TKN** 0 **N02+N03 0**  Filt. TKN 0<br>
NH4-N 4.9 3.8 000764 **3.7-4.9 NH4-N 4.9 3.8 .000764 TKN O N02+N03 O Filt. TKN 0 0-4.9 NH4-N 15.3 14.2 .0007137 TKN O N02+N03 0 Filt. TKN 0** 

 $\omega(t) = \omega(t)$ 

**Source: AMES Dates Dates 073086 Avg. Total Flow, 1/ sq. m x s: .568 Recycle, percent total flows 0 Periodic Dosing Cycle, sect 58**  Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.9 **Media Spec. Surf. Area, gq.m/cu. m: 137.8 Onda % Wetted Area: 32 Water Temp., C: 24.6 DIO/DT: .656 Infl. COD, mg/li Infl. Filt. COD, mg/li Infl. B0D5, mg/ls Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/li 6.4 Media Influent Oxidized 10 C Concentration m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 20.8 5.2 .0009955 TKN O**  TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 **Filt. TKN**<br>NH4-N 15.6 3.7 **1.2-2.4 NH4-N 15.6 3.7 .0007083 TKN O N02+N03 0 Filt. TKN 0 2.4-3.6 NH4-N 11.9 2.3 .0004403 TKN** 0 **N02+N03 O Filt. TKN**<br>NH4-N 9.6 4.2 **3.7-4.9 NH4-N 9.6 4.2 .0008041 TKN O N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 20.8 15.4 .000737 TKN** 0 **N02+N03 O Filt. TKN 0** 

 $\omega\propto\omega_{\rm{min}}$ 

**237** 

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**Source: AMES Dates 073086 Avg. Total Flow, 1/ «q. m x s: .568**  Recycle, percent total flow: Periodic Dosing Cycle, sec:<br>Distribution Time, sec/cycle: **58 18 18** Distribution Time, sec/cycle:<br>Avg. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 32 Water Temp., C: 24.3 DIO/DT: .662 Infl. COD, mg/lt Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/lt pHi 7.2 Infl. Alk, mg CaC03/l: Infl. D. 0., mg/lt 7 Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 18.1 4 .0007728 TKN** 0 **N02+N03 0 Filt. TKN O 1.2-2.4 NH4-N 14.1 4.9 .0009466 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N 9.2 3 **2.4-3.6 NH4-N 9.2 3 .0005796 TKN** 0 **N02+N03 O Filt. TKN**<br>NH4-N 6.2 4 **3.7-4.9 NH4-N 6.2 4 .0007728 TKN** 0 **N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 18.1 15.9 .0007679 TKN O** 

**N02+N03 O** 

 $\omega \propto \omega_{\rm{max}}$ 

Filt. TKN

**Source: AMES Date: 073186 Avg. Total Flow, 1/ sq. m x** st **.558 Recycle, percent total flow: 0 Periodic Dosing Cycle, sec: 59**  Distribution Time, sec/cycle: 18<br>Avg. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, aq.m/cu. m: Onda % Wetted Area: Water Temp., C; 22 COD, mg/1: Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: Infl. DIO/DT: SS, mg/1: pH: Infl. D. 0., mg/1I VSS, mg/1: Infl. Alk, mg CaC03/l 137.8 32 .705 Media Influent Oxidized 10 C pth Concentration Nitr** $m = \frac{m g}{1 - m g}$ **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 14.7 3.3 .000667 TKN O N02+N03 O Filt. TKN**<br>NH4-N 11.4 5.1 **1.2-2.4 NH4-N 11.4 5.1 .0010308 TKN O N02+N03 0**  Filt. TKN 0<br>NH4-N 06.3 2.4 0004851 **2.4-3.6 NH4-N 6.3 2.4 .0004851 TKN O N02+N03 O Filt. TKN 0 3.7-4.9 NH4-N 3.9 3 .0006064 TKN** 0 **N02+N03 0**  Filt. TKN **0-4.9 NH4-N 14.7 13.8 .0006973 TKN O N02+N03 0 Filt. TKN 0** 

 $\sim 1000$  MeV  $\sim 1000$ 

**Source: AMES Dates** 073186<br> **Avg. Total Flow, 1/ sq. m x s: Dates** 073186 Avg. Total Flow, 1/ sq. m x s: **Recycle, percent total flow: 0**  Periodic Dosing Cycle, sec: **Distribution Time, sec/cycle: 18 Avg. Distributed Flow, 1/sq. m x s: 1.9 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 32**  Water Temp., C: **Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 20.6 3.7 .0007438 TKN** 0 TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 **Filt. TKN<br>NH4-N** 16.9 4.7 **1.2-2.4 NH4-N 16.9 4.7 .0009449 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N 12.2 2.5 **2.4-3.6 NH4-N 12.2 2.5 .0005026 TKN O**  TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 **Filt. TKN**<br>NH4-N 9.7 4.3 **3.7-4.9 NH4-N 9.7 4.3 .0008645 TKN O N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 20.6 15.2 .0007639 TKN** 0 TKN 0<br>
NO2+NO3 0<br>
Filt. TKN 0 **Filt. TKN 0** 

 $\omega\propto\omega_{\rm{eff}}$  .

**240** 

 $\Delta\sim 10^{-1}$  as

**Source: AMES Dates 0730-073186 Avg. Total Flow, 1/ sq. m x si .566 Recycle, percent total flow: 0 Periodic Dosing Cycle, sec: 59**  Distribution Time, sec/cycle: **Avg. Distributed Flow, 1/sq. m x s: 1.9 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 32 Water Temp., C: 23.5 DIO/DT: .677 Infl. COD, mg/1: 49 Infl. Filt. COD, mg/1: 33 Infl. B0D5, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: 34 VSS, mg/1: 26 pH: Infl. Alk, mg CaC03/l: 148 Infl. D. 0., mg/1: Media Influent Oxidized Concentration m mg/1 mg/1 10 C Nit. Rate kg/d sq.m tot. surface area**   $0 - 1.2$ **1.2-2.4 2.4-3.6 3.7-4.9 NH4-N TKN N02+N03 Filt. TKN 17.8 20.4 13.8**  16.6 **9.2 13.3 6.7 11.4 4 3.8 4.6 3.3 2.5 1.9 3.8 4.4 .0007875 .0007481**  O **0 .0009056 .0006497 0 0 .0004922 .0003741 0**  O **.0007481 .0008662 0 0 0—4.9 NH4-N TKN N02+N03 Filt. TKN 17.8 20.4 14.9 13.4 .0007334 .0006595 0 0** 

 $\sim 1000$  MeV  $\sim 100$ 



**Source: AMES Dates 111286**<br> **Avg. Total Flow, 1/ sq. m x s: Dates** 1.201 **Avg, Total Flow, 1/ sq. m X s: 1.201 Recycl e. percent total flows 28 Periodic Dosing Cycle, sec: 31**  Distribution Time, sec/cycle: **Avg. Distributed Flow, 1/sq. m X s: 2.1 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % L Wetted Area; 33 Mat er Temp., Cl 11.5 DIO/DT: .956 Infl. COD, mg/1: 91 Infl. Filt. COD, mg/1: 53 Infl. BODS, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: vas. mg/1 : pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C Concentration Nitrogen**<br>mg/1 mg/1 **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 12.4 0** 0 NH4−N 12.4 0<br>
TKN 16.6 .2 .000118<br>
NO2+NO3 9.9 −.9 −.0005309 **N02+N03 9.9 -.9 -.0005309 Filt. TKN 13.8 0 1.2-2. 4 NH4-N 12.4 2. 1 .0012388 TKN 16.4 1.9 .0011208 N02+N03 9 2 .0011798 Filt. TKN** 0<br> **NH4-N** 10.3 .4 .000236 **2.4-3. 6 NH4-N 10.3 .4 .000236 TKN 14.5 0 N02+N03 11 0 Filt. TKN** 0<br>
NH4-N 9.9 2.9 .0017107 **3.7-4. 9 NH4-N 9.9 2.9 .0017107 TKN 0 N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 12.4 5.4 .0007964**  TKN 16.6 5.7 .0008406<br>
NO2+NO3 9.9 4.3 .0006342 **N02+N03 9.9 4.3 .0006342 Filt. TKN 0** 

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**Source: AMES Date: 111386**<br>**Avg. Total Flow. 1/ sq. m x s: Date:** 111386 **Avg. Total Flow, 1/ sq. m x s: 1.128**<br>Recycle. percent total flow: 32 **Recycle, percent total flow: 32**  Periodic Dosing Cycle, sec: 34<br>Distribution Time. sec/cycle: 34<br>18 **Distribution Time, sec/cycle: 18 Avg. Distributed Flow, 1/sq. m** X SI **2.1 Media Spec. Surf. Area, sq.m/cu . m: 137.8 Onda '/ Wetted Area: 32 Water Temp., C: 12.8 DIO/DT: .919 Infl. COD, mg/1: 93 Infl. Filt. COD, mg/ls 47 Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: vss. mg/1 : pHs Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Dxidized 10 C**<br>Depth Concentration Nitrogen Nit.Rate **Concentration** Nitrogen **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 12.1 .5 .0002663**  TKN 15.8 .3 .0001598<br>NO2+NO3 10.4 -.9 -.0004793 **10.4** −.9 −.0004793<br>12.7 0 **Filt. TKN 12.7** 0<br>
NH4-N 11.6 1.6 0008522 **1.2-2. 4 NH4-N 11.6 1.6 .0008522 TKN IS.S .8 .0004261 N02+N03 9.5 1.9 .001012 Filt. TKN 0 2.4-3. 6 NH4-N 10 1.5 .0007989 TKN 14.7 0 N02+N03 11.4 0**  Filt. TKN 0<br>
NH4-N 8.5 .4 .000213 **3.7-4. 9 NH4-N 8.5 .4 .000213 TKN** 0 TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 **Filt. TKN 0 0-4.9 NH4-N 12. 1 4 .OOOS326**  TKN 15.8 -1.9 -.000253<br>NO2+NO3 10.4 4.2 .0005592 **N02+N03 10.4 4.2 .0005592 Filt. TKN 0** 

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**Source: AMES Avg. Total Flow, 1/ sq. m x s: Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: Date: 111986 .786 44 49 20 1.9 Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: 14 DIO/DT: pH: Infl D. O., mg/1I Infl Alk, mg CaC03/l: VSS, mg/1: 137.8 31 .885 103 36**  Media Influent **Oxidized 10 C**<br>Depth Concentration Nitrogen **Nit.** Rate **pth Concentration Nitr** $m = \frac{mq}{1}$ **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 15.7 5. 1 .0018227**  TKN 38.1 21.7 .0077555<br>NO2+NO3 11.9 2.8 .0010007 **N02+N03 11.9 2.8 .0010007 Filt. TKN 34** 0<br>NH4-N 10.6 -1.4 -.0005004 **1.2-2. 4 NH4-N 10.6 -1.4 -.0005004 TKN 16.4 -13.2 -.0047176 N02+N03 14.7 1.2 .0004289 Filt. TKN 0 2.4-3. 6 NH4-N 12 4.8 .0017155 TKN 29.6 17.4 .0062187**<br>**NO2+NO3 15.9 1.8 .0006433 N02+N03 15.9 1.8 .0006433 Filt. TKN 0 3.7-4. 9 NH4-N 7.2 2.8 .0010007 TKN 12.2 3.7 .0013224 N02+N03 17.7 2.9 .0010364 Filt. TKN 0 0-4.9 NH4-N 15.7 11.3 .0010096 TKN 38.1** 30 **.0026805**  0.0007773<br>^ **Filt. TKN 0** 

 $\omega\propto\omega_{\rm{min}}$ 

**Source: AMES Date: 112186**  Avg. Total Flow, 1/ sq. m x s: **.798 .798 .798 .798 Recycle, percent total flow: 0 Periodic Dosing Cycle, sec: 45 Distribution Time, sec/cycle: 20**  Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 30 Water Temp., C: 13.8 DIO/DT: .891 Infl. COD, mg/1: 70 Infl. Filt. COD, mg/1: 33 Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1\* SS, mg/13 VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C Concentration m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 9.6 1 .0003653 TKN 16.7 2.1 .0007672**<br>**NO2+NO3 7.3 1 .0003653 N02+N03 7.3 1 .0003653 Filt.** TKN 13.5 0<br>
NH4-N 8.6 1.1 .0004018 **1.2-2.4 NH4-N 8.6 1.1 .0004018 TKN 14.6 1 .0003653 N02+N03 8.3 1.4 .0005114 Filt. TKN 0 2.4-3.6 NH4-N 7.5 1.8 .0006576**  TKN 13.6 3.6 .0013151<br>
NO2+NO3 9.7 2.5 .0009133 **N02+N03 9.7 2.5 .0009133 Filt. TKN 0 3.7-4.9 NH4-N 5.7 2 .0007306**  TKN 10 1.5 .000548<br>NG2+NG3 12.2 2.5 .0009133 **N02+N03 12.2 2.5 .0009133 Filt. TKN 0 0-4.9 NH4-N 9.6 5.9 .0005388**  TKN 16.7 8.2 .0007489<br>NO2+NO3 7.3 7.4 .0006758 **N02+N03 7.3 7.4 .0006758 Filt. TKN 0** 

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**Source: AMES Date\* 112586 Avg. Total Flow, 1/ @q. m X s: .669 Recycle, percent total flow: 0 Periodic Dosing Cycle, sec: 57 Distribution Time, sec/cycle: 23**  Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda 7 1 Wetted Area: 30 Water Temp., C: 14.7 DIO/DT: .865 Infl. COD, mg/1: 65 Infl. Filt. COD, mg/l: 25 Infl. BODS, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1** : **VSS, mg/1 : pHs Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C Concentration Nitrogen Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 13 1.4 .0004163**  TKN 15.8 1.8 .0005352<br>NO2+NO3 6.6 1.4 .0004163 **1.4** .0004163 .0004163 .0004163 0 **Filt. TKN 13.6**<br> **NH4-N** 11.6 1.7 **1.2-2. .4 NH4-N 11.6 1.7 .0005054 TKN 14 1.8 .0005352 N02+N03 8 2 .0005946 Filt. TKN 0 2.4-3. .6 NH4-N 9.9 1.5 .000446 TKN 12.2 0 N02+N03 10 0 Filt. TKN**<br>NH4-N 8.4 2.3 **3.7-4. .9 NH4-N 8.4 2.3 .0006838 TKN** 0 **N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 13 6.9 .0005129 TKN 15.8 8.4 .0006244 N02+N03 6.6 7.8 .0005798 Filt. TKN 0** 

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**Source: AMES Date: 120386**  Avg. Total Flow, 1/ sq. m x s: **.781** .781 .791 **Recycle, percent total flow: 0**  Periodic Dosing Cycle, sect **50 a.m. 1986**<br>Distribution Time, sec/cycle: 1986 1988 1988 1988 **Distribution Time, sec/cycle:** 23<br> **Avg. Distributed Flow, 1/sq. m x s: 1.7** Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda 7 L Wetted Area: 30 Water Temp., Cl 12.4 DIO/DT: .931 Infl. COD, mg/1: 109 Infl. Filt. COD, mg/1: 34 Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SS, mg/la VSS, mg/11 pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1s Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 16.2 .5 .0001868**  TKN 19.6 .6 .0002241<br>NO2+NO3 8.3 .2 .0000747 **1.3** .2 .0000747<br>17 1.2 .0004483 **Filt. TKN 17 1.2**<br>NH4-N 15.7 .9 **1.2-2. 4 NH4-N 15.7 .9 .0003362 TKN 19 1.5 .0005604 N02+N03 8.5 1.2 .0004483 Filt. TKN 15.8** 0<br> **NH4-N 14.8** .3 .0001121 **2.4-3. 6 NH4-N 14.8 .3 .0001121**  TKN 17.5 -.4 -.0001494<br>NO2+NO3 9.7 .3 .0001121 **N02+N03 9.7 .3 .0001121 Filt. TKN** 0<br>
NH4-N 14.5 1.5 .0005604 **3.7-4. 9 NH4-N 14.5 1.5 .0005604 TKN 17.9 1.6 .0005977 N02+N03 10 1.8 .0006724 Filt. TKN 0 0-4.9 NH4-N 16.2 3.2 .0002989 TKN 19.6 3.3 .0003082 N02+N03 8.3 3.5 .0003269 Filt. TKN 0** 

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**Source: AMES Date: 012587**<br> **Avg. Total Flow, 1/ sq. m x s: 1.278 Avg. Total Flow, 1/ sq. m X s: 1.278 Recycle, percent total flow: 0 Periodic Dosing Cycle, sec: Distribution Time, sec/cycles**  Avg. Distributed Flow, 1/sq. m x s: 1.278 **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Omda % 1 Wetted Area: 27 Water Temp., C: 11.4 DIO/DT: .959 Infl. COD, mg/1: 140 Infl. Filt. COD, mg/1: 66 Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SS, mg/li S7 VSS, mg/11 pHi Infl. Alk, mg CaC03/l: 161 Infl. D. 0., mg/1: Media Influent Oxidized 10 C Concentration Nitrong/1 m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 13.7 -. 1 -.000063 TKN 19.7 .3 .0001889 N02+N03 7.5 0 Filt. TKN 16**<br>NH4-N 13.8 1.1 **1.2-2.4 NH4-N 13.8 1.1 .0006927**<br>TKN 19.4 1.8 .0011335 **TKN 19.4 1.8 .0011335 N02+N03 0 Filt. TKN** 12.7 .5 **2.4-3. ,6 NH4-N 12.7 .5 .0003148 TKN 17.6 .5 .0003148 N02+N03** 8.5 0<br>
Filt. TKN 14.7 0 **Filt. TKN 14.7**<br>NH4-N 12.2 2.3 **3.7-4. ,9 NH4-N 12.2 2.3 .0014483 TKN 17.1 2.6 .0016372 N02+N03 0 Filt. TKN 0 0-4.9 NH4-N 13.7 3.8 .0005982 TKN 19.7 5.2 .0008186 N02+N03 7.5 3.8 .0005982**  Filt. TKN

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**Source: AMES** Date: 012787<br>
Avg. Total Flow, 1/ sq. m x s: 1.207 **Avg. Total Flow, 1/ sq. m x s: 1.207 1.207 1.208 1.20** Recycle, percent total flow: **Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, l/sq. m x s: 1.207 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 26 Water Temp., C: 11.2 DIO/DT: .965 Infl. COD, mg/1: 106 Infl. Filt. COD, mg/l: Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: 46 VSS, mg/1: pH: Infl. Alk, mg CaC03/l: 147 Infl. D. O., mg/1: Media Influent: Oxidized<br>Depth Concentration Nitrogen Philary Concentration**<br>millong/1 **m mg/1 mg/1 10 C Nit. Rate kg/d sq.m tot. surface area 0-1.2 NH4-N 6. 8 .5**   $11.4$ <br> $8.2$ **N02+N03 8.2 Filt. TKN 8.8 1.2-2. 4 NH4-N 6.3 .7**  11.6 **N02+N03 Filt. TKN 2.4-3. 6 NH4-N 5.6 1.1 9.5 1.1 9.3 N02+N03 9.3 Filt. TKN 7.4 3.7-4. 9 NH4-N 4.5 2. 1 TKN 8.4 3.5 N02+N03 Filt. TKN 0-4.9 NH4-N 6.8 4.4**  11.4 6.5<br>8.2 3.4 **N02+N03 8.2 3.4**  Filt. TKN **.0002992 -.0001197 0 0 .0004189 .0012567 0**  O **.0006583 .0006583**  O **o .0012567 .0020945**  O **o .0006583 .0009725 .0005087 .0005386** 

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**Source: AMES Dates 021387 Avg. Total Flow, 1/ sq. m x s: .62 Recycle, percent total flow# 0 Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow» 1/sq. m x si .62 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 21**  Water Temp., C: **Infl. COD, mg/1 : Infl. Filt. COD, mg/1: 59 Infl. BODS, mg/1: Infl. Filt. BOD5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C pth Concentration Nitr** $m = 2/1$ **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 13.2 -.2 -.0000555 TKN** 0 **N02+N03 7.8 1.2 .0003329 Filt. TKN 17.3 1.5**<br>NH4-N 13.4 3.4 **1.2-2.4 NH4-N 13.4 3.4 .0009433 TKN** 0 **N02+N03 9 0 RO2+NO3** 9<br> **Filt. TKN 16.8** 0<br> **RH4-N** 10 .5 .0001387 **2.4-3.6 NH4-N 10 .5 .0001387 TKN** 0 **N02+N03 0 Filt. TKN** 0<br> **NH4-N** 9.5 4.6 .0012763 **3.7-4.9 NH4-N 9.5 4.6 .0012763 TKN O N02+N03 14.4 3.6 .0009988**  Filt. TKN **0-4.9 NH4-N 13.2 8.4 .0005827 TKN** 0 **N02+N03 7.8 10.2 .0007075**  Filt. TKN

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## Pilot-Scale Nitrifying Biofilter Influent/Effluent Data

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**Source, AMES NIT.TOWER INFL/EFFL Dates 031486 Avg. Total Flow, 1/ sq. m x et .461 Recycle, percent total flow: 59**  Periodic Dosing Cycle, secs<br>
Distribution Time, sec/cycles<br>
27 **Distribution Time, sec/cycles** 27<br>
Avg. Distributed Flow, 1/sq. m x s: 1.3 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, aq.m/cu. mi 137.8 Onda % Wetted Areas 26 Water Temp., C: Infl. COD, mg/ls Infl. Filt. COD, rag/li Infl. B0D5, mg/ls Infl. Filt. B0D5, mg/li**  SS, mg/l: VSS, mg/l:<br>pH: **Infl.** Alk, mg Ca **pHi Infl. Alk, mg CaC03/l: Infl. D. O., mg/ls Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 2.6 1.8 .0002233 TKN O N02+N03 0**  Filt. TKN 0<br>NH4-N 18.5 000062 **2.4-4.9 NH4-N .8 .5 .000062 TKN O TKN**<br> **NO2+NO3** 0<br> **Filt.** TKN 0 **Filt. TKN**<br>NH4-N 2.6 2.3 **0-4.88 NH4-N 2.6 2.3 .0001426 TKN O N02+N03 O**  Filt. TKN

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**Source; AMES NIT.TOMER ZNFL/EFFL Date: 031986 Avg. Total Flow, 1/ sq. m x s: .517 Recycle, percent total flows 32**  Periodic Dosing Cycle, sec: **Distribution Time, sec/cycles 23**  Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. ms 137.8 Onda % Wetted Areas 27 Water Temp., Ci 8.7 DIO/DT: 1.047 Infl. COD, mg/li Infl. Filt. COD, mg/li Infl. BODS, mg/li Infl. Filt. BODS, mg/li SS, mg/lt VSS, mg/li pHi Infl. Alk, rag CaC03/ls Infl. D. O., mg/li Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 rag/1 kg/d sq.ra tot. surface area 0-2.4 NH4-N 7.3 O TKN 0 N02+N03 0 Filt. TKN O 2.4-4.9 NH4-N 0 TKN O**  TKN 0<br>**NO2+NO3** 0<br>Filt. TKN 0 **Filt. TKN**<br>NH4-N 7.3 4.7 **0-4.88 NH4-N 7.3 4.7 .0003269 TKN** 0 **N02+N03 O**  Filt. TKN

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**Source: AMES NIT.TOWER INFL/EFFL Date: 032086 Avg. Total Flow, 1/ sq. m x s: .503 Recycle, percent total flow: 31**  Periodic Dosing Cycle, sec: **65**<br>Distribution Time. sec/cycle: 65 **Distribution Time, aec/cycle: 23 Avg. Distributed Flow, 1/sq. m x s: 1.4 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area:** 2.5 **Water Temp., C: 7.5 DIO/DT: 1.09 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1s VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 6.1 O TKN O**  TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 **Filt. TKN O 2.4-4.9 NH4-N 0 TKN O N02+N03 0 Filt. TKN**<br> **NH4-N** 6.1 4.3 **0-4.88 NH4-N 6.1 4.3 .0003029 TKN** 0 **N02+N03 O Filt. TKN O** 

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**Source: AMES NIT.TOMER INFL/EFFL Date: 032086**  Avg. Total Flow, 1/ sq. m x s**: .506 .506 .506 .506 Recycle, percent total flow: 31 Periodic Dosing Cycle, sect 65 Distribution Time, sec/cycle: 23**  Avg. Distributed Flow, 1/sq. m x s: Media Spec. Surf. Area, sq.m/cu. m: 137.8 **Onda % Netted Area: 27 Water Temp., C: 7 DIO/DT: 1.109 Infl. COD, mg/lt Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. BODS, mg/li SS, mg/1: VS8, mg/lt pH: Infl. Alk, mg CaC03/lt Infl. D. O., mg/1I Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 3.6 0 TKN O N02+N03 O Filt. TKN O 2.4-4.9 NH4-N 0 TKN O N02+N03 0. Filt. TKN O 0-4.88 NH4-N 3.6 3 .0002163 TKN O N02+N03 0** 

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Source: AMES NIT.TOWER INFL/EFFL. Date: 032086<br>Avg. Total Flow, 1/ sq. m x s: 032086 **Avg. Total Flow, 1/ sq. m x si .503 Recycle, percent total flow: 31**  Periodic Dosing Cycle, sec: **Distribution Time, sec/cycle: 23**  Avg. Distributed Flow, 1/sq. m x s: 1.4 **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 27 Water Temp., Ci 12 DIO/DT: .942 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/lr VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized Concentration m mg/1 mg/1 10 C Nit. Rate kg/d sq.m tot. surface area 0-2.4 2.4-4.9 0-4.88 NH4-N TKN N02+N03 Filt. TKN NH4-N TKN N02+N03 Filt. TKN NH4-N TKN N02+N03 Filt. TKN 5.7 5.7 5.4 0 0 0 O 0 O O O .0003287 O O O** 

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**Source: AMES NIT.TOWER INFL/EFFL Date: 032686 Avg. Total Flow, 1/ sq. m x si .58 Recycle, percent total flow: 27**  Periodic Dosing Cycle, sec: **Distribution Time, sec/cycle: 18**  Avg. Distributed Flow, 1/sq. m x s: 1.9 **Media Spec. Surf. Area, aq.m/cu. mi 137.8 Onda % Wetted Areat 31 Water Temp., Ci 12 DIO/DT: .942 Infl. COD, mg/li Infl. Filt. COD, mg/li Infl. B0D5, mg/ls Infl. Filt. BODS, mg/li SS, mg/ls VSS, mg/ls pHi Infl. Alk, mg CaC03/ls Infl. D. O., mg/1: Oxidized Media Influent**  10 c **Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. aurface area 0-2.4 NH4-N 9.7**   $\bullet$ **TKN**   $\bullet$ **N02+N03**   $\mathbf{o}$ **Filt. TKN**   $\mathbf{o}$ **NH4-N 2.4-4.9**   $\mathbf{o}$ **TKN**   $\bullet$ **N02+N03**   $\mathbf{o}$ **Filt. TKN**   $\bullet$ **0-4.88**  .000351 **NH4-N 9.7**  5 **TKN**   $\mathbf{o}$ **N02+N03**   $\bullet$ **Filt. TKN**   $\bullet$ 

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**Source: AMES NIT.TOWER INFL/EFFL Date: 032786 Avg. Total Flow, 1/ sq. m x si .581 Recycle, percent total flow: 27**  Periodic Dosing Cycle, sec: **59 a. 59 a. 59**<br>Distribution Time, sec/cycle: 59 a. 59 Distribution Time, sec/cycle: **Avg. Distributed Flow, 1/sq. m x s: 1.9 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: Water Temp., C: 11.7 DIO/DT: .95 Infl. COD, mg/1: Infl. Flit. COD, mg/1# Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SB, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C pth Concentration Nitr** $\mathfrak{m}$  **Mitr** $\mathfrak{m}$  **/1 m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 10.7 0 TKN O N02+N03 0 Filt. TKN O 2.4-4.9 NH4-N 0 TKN O**  TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 **Filt. TKN<br>NH4-N 10.7 6.3 0-4.88 NH4-N 10.7 6.3 .0004468 TKN** 0 **N02+N03** 0<br> **Filt.** TKN 0 **Filt. TKN 0** 

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**Source: AMES NIT.TOWER INFL/EFFL Date: 033186 Avg. Total Flow, 1/ sq. m x si .577 Recycle, percent total flow: 26 Periodic Dosing Cycle, sec: 59**  Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.9 **Media Spec. Surf. Area, aq.m/cu. m: 137.8 Onda % Wetted Area:** 31<br> **Water Temp., C:** 15.7 **D10/DT: .84** Water Temp., C: **Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. BOD5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C**  Concentration **m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 7.3 O TKN O N02+N03 0 Filt. TKN O 2.4-4.9 NH4-N 0 TKN O N02+N03 O Filt. TKN** 0<br>
NH4-N 7.3 4.1 .0002553 **0-4.88 NH4-N 7.3 4.1 .0002553 TKN O N02+N03 O Filt. TKN O** 

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Source: AMES NIT.TOWER INFL/EFFL. Date: 040186<br>Avg. Total Flow, 1/ sq. m x s: 0.583 **Avg. Total Flow, 1/ sq. m x st .583 Recycle, percent total flows 26**  Periodic Dosing Cycle, sec: **Distribution Time, sec/eyelet 18**  Avg. Distributed Flow, 1/sq. m x s: 1.9 **Media Spec. Surf. Area, sq.m/cu. mi 137.8 Onda % Wetted Area: Water Temp., Ci 13.5 DlO/DTt .899 Infl. COD, mg/li Infl. Filt. COD, mg/lt Infl. B0D5, mg/lt Infl. Filt. BODS, mg/li SS, mg/lt VSS, mg/lt pHt Infl. Alk, rag CaC03/li Infl. 0. O., mg/lt Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 11.8 6 TKN** 0 TKN 0<br>NO2+NO3 0<br>Filt. TKN 0 **Filt. TKN O 2.4-4.9 NH4-N 0 TKN O N02+N03 0 Filt. TKN** 0<br> **NH4-N** 11.8 4.5 .000303 **0-4.88 NH4-N 11.8 4.5 .000303 TKN O N02+N03 O** 

**Filt. TKN 0** 

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**Source: AMES NIT.TOWER INFL/EFFL Date: 040286 Avg. Total Flow, 1/ sq. m x s: .584 Recycle, percent total flow: 26 Periodic Dosing Cycle, sec: 59 Distribution Time, sec/cycle: 18**  Avg. Distributed Flow, 1/sq. m x s: 1.9 **Media Spec. Surf. Area, sq.ra/cu. m: 137.8 Onda % Wetted Area: 31 Water Temp., Ci 13 DIO/DT: .913 Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 16.7 O TKN O N02+N03 O Filt. TKN O 2.4-4.9 NH4-N 0 TKN O N02+N03 0 Filt. TKN O 0-4.88 NH4-N 16.7 6.6 .0004521 TKN** 0 **N02+N03 0 Filt. TKN O** 

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Source: AMES NIT.TOWER INFL/EFFL Date: 040386<br>Avg. Total Flow, 1/ sq. m x s: 0100386 Avg. Total Flow, 1/ sq. m x s: **.377 .377 .388 .377 .388 .377**<br>Recycle, percent total flow: 25 **Recycle, percent total flow: 25 Periodic Dosing Cycle, sec: 59 Distribution Time, sec/cycle: 18 Avg. Distributed Flow, 1/sq. m x s: 1.9 Media Spec. Surf. Area, sq.m/cu. mi 137.8 Onda % Wetted Areas 31**  Water Temp., C: 12.2 **DIO/DT: Infl. COD, mg/1: Infl. Filt. COD, mg/1; Infl. B0D5-, mg/13 Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/li Infl. D. O., mg/1: Media Influent Oxidized 10 C. Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 9.3 0 TKN O W02\*N03 0 Filt. TKN O 2.4-4.9 NH4-N 0 TKN O**  TKN 0<br>
NO2+NO3 0<br>
Filt. TKN 0 **Filt. TKN O 0-4.88 NH4-N 9.3 4.7 .0003261 TKN** 0 **N02+N03 O Filt. TKN O** 

 $\sim 10^{11}$  MeV  $\sim 10^{11}$ 

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Source: AMES NIT.TOWER INFL/EFFL Date: 040386 **Avg. Total Flow, 1/ sq. m x s: .581 Recycle, percent total flow: 24 Periodic Dosing Cycle, sec: 59 Distribution Time, sec/cycle: 18**  Avg. Distributed Flow, 1/sq. m x s: 1.9 **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 31 Water Temp., C: 12.6 DIO/DT: .925 Infl. COD, mg/1 : Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1\* VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 11.7 O TKN O N02+N03 O Filt. TKN O 2.4-4.9 NH4-**<br>TKN **TKN O N02+N03 O Filt. TKN O 0-4.88 NH4-N 11.7 6 .0004143 TKN O N02+N03 O**  Filt. TKN

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 $\sqrt{3}$  g  $\sim$ 

**273** 

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**274** 

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 $\label{eq:2.1} \frac{1}{\sqrt{2\pi}}\int_{0}^{\infty}\frac{1}{\sqrt{2\pi}}\left(\frac{1}{\sqrt{2\pi}}\right)^{2\alpha} \frac{1}{\sqrt{2\pi}}\,d\mu.$ 

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**276** 

 $\ddot{\phantom{a}}$ 

 $\ddot{\phantom{a}}$ 

**Source: AMES NIT.TOWER INFL/EFFL Date: 040486**  Avg. Total Flow, 1/ sq. m x s: **Recycle, percent total flow: 25 Periodic Dosing Cycle, secr Distribution Time, sec/cycle:** 18<br> **Avg. Distributed Flow, 1/sq. m × s: 1.9** Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. mi 137.8 Onda 7. Wetted Areai 31**  Water Temp., C: **Infl. COD, mg/ls Infl. Filt. COD, mg/li Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SB, mg/1I VSS, mg/1: pH: Infl. Alk, mg CaC03/li Infl. D. O., mg/1: Media Influent Dxidized 10 C**<br>Depth Concentration Nitrogen Nit.Rate **pth Concentration Nitrogen**<br>m mg/l **m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 5.7 O TKN O N02+NQ3 0 Filt. TKN O 2.4-4.9 NH4<br>TKN TKN O N02+N03 O Filt. TKN O 0-4.88 NH4-N 5.7 4.8 .0003443 TKN O N02+N03 0 Filt. TKN O** 

 $\omega \propto \omega_{\rm{max}}$ 

 $\sum_{i=1}^{n}$ 



**Source: AMES NIT.TOWER INFL/EFFL Date: 040586 Avg. Total Flow, 1/ sq. m x s: .584 Recycle, percent total flow: 25**  Periodic Dosing Cycle, sec: **59 approximately 1998**<br>Distribution Time. sec/cycle: **59 approximately** 18 Distribution Time, sec/cycle: 18<br>Avg. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 31 Water Temp., C: 11.2 DIO/DT: .965 Infl. COD, mg/1: 73 Infl. Filt. COD, mg/1: Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 9.5 O TKN 11.7 0**  TKN 11.7 0<br> **N02+N03** 10.3 0<br>
Filt. TKN 0 **Filt. TKN O 2.4-4.9 NH4-N 0 TKN O N02+N03 O Filt. TKN** 9.5 5.8 00042 **0-4.88 NH4-N 9.5 5.8 .00042**  TKN 11.7 2.2 .0001593<br>**NO**2+NO3 10.3 2.5 .000181 **N02+N03 Filt. TKN O** 

 $\sim 10^{11}$  MeV  $\sim 10^{11}$ 

**279** 

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Source: AMES NIT.TOWER INFL/EFFL Date: 040686 **Avg. Total Flow, 1/ sq.** M **x si .568 Recycle, percent total flow: 24 Periodic Dosing Cycle, sec: 59 Distribution Time, sec/cycle: 18 Avg. Distributed Flow, 1/sq. n x s: 1.9 Media Spec. Surf. Area, sq.m/cu. mi 137.8 Onda % Wetted Areas 31**  Water Temp., C: **Infl. COD, mg/li 98 Infl. Filt. COD, mg/1: Infl. B0D5, mg/1I Infl. Filt. BODS, mg/1: SS, mg/18 VSS, mg/1: pH: Infl. Alk, mg CaC03/li Infl. D. O., mg/1I Media Influent Oxidized 10 C**  Concentration<br>mg/1 **m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 7.6 O TKN 11.3 O N02+N03 O Filt. TKN O 2.4-4.9 NH4-N O TKN O N02+N03 0 Filt. TKN O 0-4.88 NH4-N 7.6 5.1 .0003465**   $-.0001155$ **N02+N03 O**  Filt. TKN

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**Source» AMES NIT.TOMER INFL/EFFL Date: 041086**  Avg. Total Flow, 1/ sq. **m** x s: **.589 Recycle, percent total flow: 28 Periodic Dosing Cycle, sec: 59 Distribution Time, sec/cycle: 18**  Avg. Distributed Flow, 1/sq. m x s: 1.9 **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 31 Water Temp., C: 13 DIO/DT: .913 Infl. COD, mg/l**: 108 **108 108 108 Infl. Filt. COD, mg/1: Infl. B0D5, mg/1I Infl. Filt. BODS, mg/1: SS, mg/1: 54 VSS, mg/1: 44 pH: Infl. Alk, mg CaC03/l: 170 Infl. D. O., mg/1 : Media Influent Oxidized 10 C pth Concentration Nitr** $\frac{1}{2}$  **Mitr** $\frac{1}{2}$  **Mitr** $\frac{1}{2}$  **Mitr** $\frac{1}{2}$  **Mitr** $\frac{1}{2}$  **Mitr** $\frac{1}{2}$  **Mitr** $\frac{1}{2}$  **Mitr m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 20.1 O TKN 22.8 O TKN 22.8** 0<br>102+N03 8.7 0 **Filt. TKN O 2.4-4.9 NH4-N 0 TKN** 0 **N02+N03 0 MO2+NO3** 0<br> **Filt. TKN** 0<br> **MH4-N** 20.1 4.4 .000304 0-4.88 NH4-N 20.1 4.4 .000304 **TKN 22.8 3.8 .0002626 N02+N03 8.7 4.8 .0003316 Filt. TKN O** 

 $\sim 1000$  MeV  $\sim 100$ 

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**Source: AMES NIT.TOWER INFL/EFFL Date: 041086 Av0. Total Flow, 1/ sq. m x si .571 Recycle, percent total flow:** 26<br> **Periodic Dosing Cycle, sec: 29** Periodic Doming Cycle, sec:<br>
Distribution Time, sec/cycle:<br>
18 **Distribution Time, sec/cycle:** 18<br> **Avg. Distributed Flow, l/sq. m x s: 1.9** Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. #: 137.8 Onda % Wetted Area: 31 Water Temp., C: 12.3 DIO/DT: .934 Infl. COD, mg/li 122 Infl. Filt. COD, rag/1: Infl. BOOS, mg/lf Infl. Filt. BODS, mg/lt SB, mg/lr VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/lt Media Influent Oxidized lO 1 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 13.5 0 TKN 17.6 0**  TKN 17.6 0<br>NO2+NO3 8.8 0<br>Filt. TKN 0 **Filt. TKN 0 2.4-4.9 NH4-N 0 TKN** 0 **N02+N03 0 Filt. TKN**<br>NH4-N 13.5 4 0−4.88 NH4−N 13.5 4 .0002741<br>TKN 17.6 −.3 −.0000206 **TKN 17.6 -.3 -.0000206 N02+N03 8.8 5.4** • **00037** 

 $\omega \propto \omega_{\rm{max}}$ 

**Filt. TKN 0** 

E.

**Source: AMES NIT.TOWER INFL/EFFL Dates 041186 Avg. Total Flow, 1/ sq. m x s: .578 Recycle, percent total flow: 27**  Periodic Dosing Cycle, sec: **Distribution Time, sec/cycle: 18 Avg, Distributed Flow, 1/sq. m x s: 1.9 Media Spec. Surf. Area, sq.m/cu. it: 137.8 Onda % Wetted Area: 31 Water Temp., C: Infl. COD, mg/1: 95 Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: 80 VSS, mg/1: 72 pH: Infl. Alk, mg CaC03/l: 184 Infl. D. O., mg/1:**  Media Influent **Oxidized 10 C**<br>Depth Concentration Nitrogen **Nit.Rate Concentration**<br>mg/l **m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 11.8 0 TKN 13.7 O N02+N03 9.8 O Filt. TKN O 2.4-4.9 NH4-N 0 TKN O N02+N03 O Filt. TKN O 0-4.88 NH4-N 11.8 6.2 .0004153** 

**TKN 13.7 3.3 .0002211** 

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**Filt. TKN 0** 

**N02+N03 9.8 5.2 .0003483** 

**Source: AMES NIT.TOMER INFL/EFFL Dates 041186 Avg. Total Flow, 1/ eq. m x si - .578 Recycle, percent total flow: 27**  Periodic Dosing Cycle, sec: 39<br>Distribution Time, sec/cycle: 39<br>18 **Distribution Time, sec/cycle:** 18<br> **Avg. Distributed Flow, 1/sq. m x s: 1.9 Avg. Distributed Flow, 1/sq. m x si Media Spec. Surf. Area, sq^m/cu. mi 137.8**  Onda % Wetted Area: 31<br> **Water Temp., C:** 33 **Dio/DT:** 313 **Water Temp., Ci 13 DIO/DT: .913 Infl. COD, mg/li 96 Infl. Filt. COD, mg/li Infl. B0D5, mg/li Infl. Filt. BODS, mg/li 88, mg/ls VSS, mg/1: pH: Infl. Alk, mg CaC03/1: Infl. D. 0., mg/1: Media Influent Oxidized pth Concentration**<br>m mg/l **m mg/1 mg/1 10 C Nit. Rate kg/d sq.m tot. surface area 0-2.4 NH4-N 8.7 TKN 12**<br>**NO2+NO3** 9.2  $N02+NO3$ **Filt. TKN**   $2.4 - 4.9$ **TKN N02+N03 Filt. TKN 0-4.88 NH4-N 8.7 TKN 12**<br>**NO2+NO3** 9.2  $NQ2+ND3$ **Filt. TKN 4 1.6 4.7 0**  O **0 0 0**  O **o o .0002712 .0001085 .0003187**  O

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**Source: AMES NIT.TOWER INFL/EFFL Date: 041786 Avg. Total Flow, 1/ sq. m x s: .563 Recycle, percent total flow: 29**  Periodic Dosing Cycle, sec: 53<br>Distribution Time, sec/cycle: 53<br>17 Distribution Time, sec/cycle: 17<br>Avg. Distributed Flow, 1/sq. m x s: 1.8 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda % Wetted Area: 31 Water Temp., C: 15.1 DIO/DT: .853 Infl. COD, mg/1: 154 Infl. Filt. COD, mg/1; Infl. B0D5, mg/1: Infl. Filt. B0D5, mg/1\* SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. 0., mg/1: Media Influent Oxidized 10 C Concentration m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 21.2 0 TKN 26.6 O**  TKN 26.6 0<br> **N02+N03** 9.2 0<br>
Filt. TKN 0 **Filt. TKN O 2.4-4.9 NH4-N 0 TKN O N02+N03 O Filt. TKN 0 0-4.88 NH4-N 21.2 4.7 .00029**  TKN 26.6 −11.8 −.0007281<br>NO2+NO3 9.2 6.7 .0004134 **N02+N03 9.2 6.7 .0004134**  Filt. TKN

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**Source: AMES NIT.TOMER INFL/EFFL Date: 121686 Avg. Total Flow, 1/ sq. m x s: 1.17 Recycle, percent total flow: 38 Periodic Dosing Cycle, sec: 35<br>Distribution Time. sec/cycle: 39<br>18** Distribution Time, sec/cycle: 18<br>Avq. Distributed Flow. 1/sq. m x s: 2.3 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda 7. Wetted Area: 33 Water Temp., C: Infl. COD, mg/1: Infl. Filt. COD, mg/1: Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1s Media Influent Oxidized 10 C Concentration** Nitrogen **m mg/1 mg/1 kg/d sq.m tot. surface area 0-2.4 NH4-N 11.3 .2 .0000529 TKN O TKN** 0<br>
NO2+NO3 0<br> **Filt.** TKN 0 **Filt. TKN**<br>NH4-N 11.1 1 **2.4-4.9 NH4-N 11.1 1 .0002643 TKN O N02+N03 0 NO2+NO3** 0<br> **Filt. TKN** 0<br>
NH4-N 11.3 1.2 .0001586 **0-4.88 NH4-N 11.3 1.2 .0001586 TKN O N02+N03 0** 

 $\sim 10^{11}$  m  $^{-1}$ 

**Filt. TKN 0** 

**Source: AMES NIT.TOMER INFL/EFFL Date: 121786 Avg. Total Flow, 1/ sq. m x si 1.16 Recycle, percent total flow# 38 Periodic Dosing Cycle, sec: 36**  Distribution Time, sec/cycle: 18<br>Avg. Distributed Flow, 1/sq. m x s: 2.3 Avg. Distributed Flow, 1/sq. m x s: **Media Spec. Surf. Area, sq.m/cu. mi 137.8 Onda % Wetted Area: 33 Water Temp., Ci 14 DIO/DT: .885 Infl. COD, mg/li 73 Infl. Filt. COD, mg/li 44 Infl. BODS, mg/li Infl. Filt. BODS, rog/1: SS, mg/1: VSS, mg/ls pHi Infl. Alk, mg CaC03/lt Infl. D. O., mg/ls Media Influent Oxidized Depth** Concentration **m mg/1 mg/1 10 C Nit. Rate**  kg/d sq.ni tot. -surface area **0-2.4 2.4-4.9 0-4.88 NH4-N TKN N02+N03 Filt. TKN NH4-N TKN N02+N03 Filt. TKN NH4-N TKN 9.7 13.9 11.6 11.4 9.4 13.3 11.5 9.7 13.9 .3 .6 -.1 .9 .3 1.5 1.2 .9**  .0000791 **.0001583 -.0000264 0 .0002374 .0000791 .0003957**  O **.0001583 .0001187** 

> **11.6 11.4**

**1.4 1.8** 

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**N02+N03 Filt. TKN**  **296** 

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**Source: AMES NIT.TOWER INFL/EFFL Date: 012787 Avg. Total Flow, 1/ sq. m x s: 1.243 Recycle, percent total flow: O Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.243 Media Spec. Surf. Area, sq.m/cu. m: 137.8 Onda '/. Wetted Area: 26 Water Temp., C: 11.3 DIO/DT: .962 Infl. COD, mg/l:** 104<br> **Infl. Filt. COD, mg/l:** 50 **Infl. Filt. COD, mg/1: 50 Infl. BODS, mg/1: Infl. Filt. BODS, mg/1: 88, mg/1: 42 V8S, mg/1: pH: Infl. Alk, mg CaC03/l: 159 Infl. D. O., mg/1I Media Influent Oxidized**  Concentration<br>mg/1 **m mg/1 mg/1 10 C Nit. Rate kg/d sq.m tot. surface area 0-2.4 NH4-N 10.8 1 16.6** 3<br>7.2 **1.5 N02+N03 7.2 1.5 Filt. TKN 12.8 .9 2.4-4.9 NH4-N 9.8 2.5 13.6** 2.2<br>**8.7** 2.3 **N02+N03** 8.7 2.3<br>
Filt. TKN 11.9 2.2 **Filt. TKN 11.9 2.2 0-4.88 NH4-N 10.8 3.5 16.6 5.2**<br>7.2 **3.8 N02+N03 7.2 3.8 Filt. TKN 12.8 3.1 .0003073 .0009218 .0004609 .0002765 .0007682 .000676 .0007067 .000676 .0005377 .0007989 .0005838 .0004763** 

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Pretreatment Tower Data



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**TKN** 0 **N02+N03** 0<br>
N02+N03 0<br>
Filt. TKN 0 Filt. TKN 0<br>
NH4-N 5.3 1.6 .0005726

**TKN** 0 **N02+N03 O** 

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**0.0-2.44 NH4-N 5.3 1.6 .0005726** 

Filt. TKN

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**Source: AMES PRETREATMENT TOWER Date: 072186 Avg. Total Flow, 1/ sq. m x st 1.41 Recycle, percent total flow:**  O **Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.41 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 33 Water Temp., C: 22.5 DIO/DT: ,695 Infl. COD, mg/ls Infl. Filt. COD, mg/lt Infl. B0D5, mg/1: 12 Infl. Filt. BODS, mg/1: VSS, mg/1 : SS, mg/18 pH: Infl. Alk, mg CaC03/l: Infl. D. O., mg/1: 5 Media Influent Oxidized 10 C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N**  5 **1.1 .0007788 TKN**  O **N02+N03 0 Filt. TKN**  O  $1.2 - 2.4$ **3.9 .9 .0006372 TKN**  O **N02+N03 0 Filt. TKN**  O 5  $\overline{2}$ **.0007082 0.0-2.44 NH4-N TKN**  O **N02+N03 0** 

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**Filt. TKN** 

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Source: AMES PRETREATMENT TOWER Date: 072386<br>Avg. Total Flow, 1/ sq. m x s: 1.37 Avg. Total Flow, 1/ sq. m x s: **Recycle, percent total flows 0 Periodic Dosing Cycle, sect Distribution Time, sec/cycle:**  Avg. Distributed Flow, 1/sq. m x s: 1.37 **Media Spec. Surf. Area, sq.m/cu. mi 98 Onda % Wetted Area: 33**  Water Temp., C: **Infl. COD, mg/ls Infl. Filt. COD, mg/ls Infl. B0D5, mg/1:** 15 **15 Infl. Filt. BODS, mg/ls SS, mg/1: VSS, mg/1: pHs Infl. Alk, mg CaC03/lt Infl. D. O., mg/ls 4.6 Media Influent Oxidized 10 C Concentration Nitrogen m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 6.4 .6 .0003908 TKN O N02+N03 0 Filt. TKN O 1.2-2.4 NH4-N 5.8 .7 .0004559 TKN** 0 **N02+N03 O Filt. TKN O 0.0-2.44 NH4-N 6.4 1.3 .0004234 TKN** 0 **N02+N03 0 Filt. TKN O** 

 $\sim$  2008/07/18

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**Source: AMES PRETREATMENT TOWER Date: Avg. Total Flow, 1/ sq. m x s: Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x sr 073086 1.91 0 1.91 Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: Infl. COD, mg/ls Infl. Filt. COD, mg/li Infl. B0D5, mg/1: Infl. Filt. BODS, mg/1: 24.2 SS, mg/1: pH: Infl. D. 0., mg/lr VSS, mg/1 : Infl. Alk, mg CaC03/l: 4.7 DIO/DT: 98 36 .663 18 Media Influent Oxidized lO C Depth Concentration Nitrogen Nit. Rate m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 6.9 1.2 .0010978 TKN O N02+N03 0**  Filt. TKN<br>NH4-N 5.7 1.1 **1.2-2.4 NH4-N 5.7 1.1 .0010063 TKN O N02+N03 O Filt. TKN** 6.9 2.3 .0010524 **0.0-2.44 NH4-N 6.9 2.3 .0010524 TKN** 0

**N02+N03 0** 

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**Filt. TKN 0** 

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**Filt. TKN** 

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Source: AMES PRETREATMENT TOWER Date: 111186 **Avg. Total Flow, 1/ sq. m x st 1.72 Recycle, percent total flow: 0 Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.72 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 34 Water Temp., C: 16.3 DIO/DT: .827 Infl. COD, mg/ls 92 Infl. Filt. COD, mg/1: Infl. BODS, mg/1: 35 Infl. Filt. BODS, mg/1: SS, mg/1: VSS, mg/1: pH: Infl. Alk, mg CaC03/lt Infl. D. O., mg/1: 5.1 Media Influent Oxidized Concentration**<br>mg/1 **m mg/1 mg/1 10 C Nit. Rate kg/d aq.m tot. surface area 0-1.2 NH4-N TKN N02+N03 Filt. TKN**   $1.2 - 2.4$ **TKN N02+N03 Filt. TKN 0.0-2.44 NH4-N TKN N02+N03 Filt. TKN 19.7 22.3 18.9 19.7 22.3 .8**  1.8 **2.6 -1.1**  .0008221 **0 0**  O **.0018498**  O O O **.0013363 —.000S654 0**  O

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Source: AMES PRETREATMENT TOWER Date: 012587 Avg. Total Flow, 1/ sq. m x s: **1.75**<br>Recycle. percent total flow: **1.75 Recycle, percent total flow: Periodic Dosing Cycle, sect Distribution Time, sec/cycle:**  Avg. Distributed Flow, 1/sq. m x s: 1.75 **Media Spec. Surf. Area, sq.m/cu. rai 98 Onda % Wetted Area: 34 Water Temp., C: 12.5 DIO/DT: .928**  166<br>
Infl. COD, mg/l: 166<br>
Infl. Filt. COD, mg/l: 66 **Infl. Filt. COD, mg/1: 66 Infl. BOD5, mg/li Infl. Filt. BODS, mg/1; SS, mg/lt VSS, mg/1: finfl. Alk, mg CaCOS/l:** 5.7 **Infl. D. O., mg/1: 5.7 Media Influent Oxidized 10 C**  Concentration<br>mg/l **m mg/1 mg/1 kg/d sq.m tot. surface area 0-1.2 NH4-N 14.2 0 TKN 22.2 O N02+N03 O Filt. TKN O 1.2-2.4 NH4-N O TKN O N02+N03 O Filt. TKN** 0<br>
NH4-N 14.2 2.2 .0012909 **0.0-2.44 NH4-N** 14.2 2.2<br>TKN 22.2 3.2 **TKN 22.2 3.2 .0018777 N02+N03 0**  Filt. TKN

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Source: AMES PRETREATMENT TOWER Date: 012787<br>Avg. Total Flow, 1/ sq. m x s: 1.76 Avg. Total Flow, 1/ sq. m x s: **1.76 1.76 1.76 1.76 Recycle, percent total flow: 0 Periodic Dosing Cycle, sect Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x si 1.76 Media Spec. Surf. Area, sq.m/cu. m: 98**  Onda % Wetted Area: <sup>37</sup> **Canadian Conduct 2 Area: 34**<br>
Water Temp., C: 342 **D**10/DT: 342 **Water Temp., Ci 12 DlO/DTi .942 Infl. COD, mg/li** 164<br> **164 Infl. Filt. COD, mg/li** 26 **Infl. Filt. COD, mg/1: 56 Infl. B0D5^ mg/1: 47 Infl. Filt. BODS, mg/1: SS, mg/1I VSS, mg/1: pH: Infl. Alk, mg CaC03/l: Inf1. D. O., mg/1: 5.7 Media Influent Oxidized Concentration m mg/1 mg/1 10 C Nit. Rate kg/d sq.m tot. surface area**   $0 - 1.2$ **1.2-2.4 NH4-N 11.7 0 TKN** 20 0 **N02+N03 4.4 0 Filt. TKN 14.5 O NM4-N** 0 **TKN** 0 TKN 0<br>**NO2+NO3** 0<br>Filt. TKN 0 **Filt. TKN**<br>0.0-2.44 NH4-N 11.7 2.3 **NH4-N 11.7 2.3 .0013778 TKN 20 4.5 .0026957 4.4** 3.3 .0019769<br>14.5 2.2 .0013179 Filt. TKN

 $\sim 1000$  MeV  $\sim 100$ 

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**Brazil** 

Huxley WPCP Oxygen-Limited Data, Winter of 1986

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 $\sum_{k=1}^{n}$ 

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Source: HUXLEY WPCP INFL/EFFL Date: 022686 **Avg. Total Flow, 1/ sq. m x ai ,249 Recycle, percent total flowt O** 

**Media Spec. Surf. Area, sq m/cu. m: 98**  Water Temp., C: 11.5 **Infl. COD, mg/ls Infl. Fil t. COD, mg/lt Infl. B0D5, mg/ls Infl. Filt. BODS, mg/li SS, mg/1: 317 pH# 7.9 Infl. Alk, mg CaC03/l: Infl. D. O., mg/lc**  <sup>I</sup>**DlO/DTi .956 133 86 VSS, mg/lt Media Influent Oxidized Depth Concentration Nitrogen m mg/1 mg/1 10 C Ni t. Rate kg/d eq.m** 



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 $\sum_{i=1}^{n}$ 

**325** 

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**Source, HUXLEY WPCP INFL/EFFL Avg. Total Flow, 1/ sq. m x a: Recycle, percent total flows Date: 031886 .401**   $\bullet$ 

**Media Spec. Surf. Area, sq.m/cu. ms Water Temp., Ci 13.2 DlO/DTt Infl. COD, mg/lt Infl. Filt. COD, mg/lt Infl. B0D5, mg/li Infl. Filt. BODS, mg/ls SS, mg/1( 35 VSS, mg/1i pHs 7.8 Infl. Alk, mg CaC03/lt Infl. D. O., mg/lt 98 .907 25** 

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 $\sim 10^{11}$  km s  $^{-1}$ 

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Data From

Duddles and Richardson (1973)

and

Baxter and Woodman, Inc. (1973)

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## Data From

## Parker and Richards (1985),

## Richards (1984),

## and

Richards and Reinhart (1986)

**Source: Parker and Richards** (1985) **Date: 112482 Avg Total Plow, 1/ sq. m \* s: .394 Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avgv Distributed Plow, 1/sq. ass: .394 Media Depths m: 2.44 Media Spec. Surf. Area, sq.m/cu. •: 98 Onda % Wetted Area:** 23<br>Mater Temp.. C: 25 256<br>Mater Temp.. C: 25 256 **Water Temp., C: COD, mg/1: 0.45 micron PIIt. €00, mg/1: B0D5, mg/1: Pilt. BOD5; mg/l:** 50<br>SS, mg/l: 50<br>SS, mg/l: 50<br>SS, mg/l: 50 **SS, mg/1: VSS, mg/1: pH: Alk, mg CaC0371: Total Applied Oridized 10 C Nitrification<br>
Influent Conc. Nitrogen, Rate, kg/d/sq. m** Influent Conc. Nitrogen, Rate, kg/d/sq. m **and Nitrogen**<br> **Rate, kg/d/sq. m and ng/l cotal surface area mg N/1 mg/1 total surface area NH4 18.9 1.8 .000219 N02+N03 0 0 0.000000** 

**TKN** 

**0.000000** 

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**Source: Parker and Richards (1985) Date:** Avg. Total **Plow**, 1/ sq. a x s: Recycle, percent total flow: **Periodic Dosing Cycle, sec: Distribution Time, sec/cycle;**  Avg. Distributed Flow, 1/sq. m x s: **Media Depth» a: Media Spec. Surf. Area, sq.a/cu. a: Onda % Wetted Area: Water Teap., C: 14 COD, ag/lr a.45 alcron Mit. COO, ag/1: BODS, ag/1: Flit. BODS, ag/1: SS, ag/1: pH: DIO/DT: VSS, ag/l: Alk, ag CaC03/l:**  iK2482 **.394 34 .394 2.44 98 23 .885 Total Applied Influent Cone, ag N/1 Orldized Nitrogen, ag/1 10 C Nitrification Rate, kg/d/sq. a total surface area NH4**  16.7 **.000630 N0ZfN03 4.9 .000617 TKN o.oooooa** 

 $\sim 100$  MeV  $\sim 100$ 

**0.45 alcron Pilt. TKN** 

**Source: Parker and^ Richards** (1985) **Date; 072585**  Avg. Total **Flow**, 1/ sq. n x s: **.767 Recycle, percent total flow: 0**  Periodic Dosing Cycle, sec: **Distribution Time, sec/cycle:**  Avg. Distributed Flow, 1/sq. a x s:  $\blacktriangleright$ **.767 Media Depth, m: 2.44 98 Media Spec. Serf. Area, sq.a/cu. a: 25 Onda % Wetted Area: Water Temp., C: 25 DIO/OT: .649 COD, mg/1: 0.45 micron Pllt. COO, mg/1: B0D9, mg/1: Pllt. BOOS, mg/1: V3S, mg/1: S8, mg/1: pH: AIk, mg CaC03/l: Total Applied Oxidized 10 C Nitrification Rate, kg/d/sq. m Influent Cone, Nitrogen, mg N/1 mg/1 total surface area NH4**  10 **5.7 .001025 N02+N03 .6 4.1 .000737 TKN 0.000000** 

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**0.45 micron Pllt. TWt** 

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 $1.44$ 

**Sources Richards (1984) Period 1 Avg. Total Flow, 1/ sq. m x s» .482 Recycle, percent total flow: 0**  Periodic Dosing Cycle, sec: 12<br>Distribution Time. sec/cycle: 5 **Distribution Time, sec/cycle: 6 Avg. Distributed Flow, l/sq. m x s: .96 Media Depth, m: 6.1 Media Spec. Surf. Area, sq.m/cu. m: 137.8**<br>Onda % Wetted Area: 26 **Onda % Wetted Area: 26**  Water Temp., C: **COD, mg/1: Effluent 1.5 micron filt. COD, mg/1: 43 Effluent B0D5, mg/1: 11 Effluent 1.5 micron filt. B0D5, mg/1: 6 SS, mg/1: 11 VSS, mg/1: 7 pH: 7 Alk, mg CaC03/l: 21 Total Applied Oxidized 10 C Nitrification Influent Cone. Nitrogen, Rate, kg/d/sq. m rag N/1 mg/1 total surface area NH4**  8.6  $\mathbf{B}$  $.000257$ **N02+N03**   $7.7$  $.000248$ **TKN 0.000000** 

**0.45 micron Filt. TKN** 

**Sources Richards (1984) Period 1**  Avg. Total Flow, 1/ sq. m x s: 475 **Recycle, percent total flows O**  Periodic Dosing Cycle, sec: **Distribution Time, sec/cycles 6 Avg. Distributed Flow, 1/sq. m x s: .96 Media Depth, ms 6.1 Media Spec. Surf. Area, sq.m/cu. m: 144.4 •nda 7. Wetted Areas 26**  Water Temp., C: **COD, mg/1s Effluent 1.5 micron filt. COD, mg/1s 41 Effluent BODS, mg/1s 15 Effluent 1.5 micron filt. B0D5, mg/1s 7 SS, mg/1s 17 VSS, mg/1: 11 pH: 7.2 Alk, mg CaC03/l: 36 Total Applied Oxidized 10 C Nitrification Influent Cone. Nitrogen, Rate, kg/d/sq. m mg N/1 mg/1 total surface area NH4**   $9.6$ 8  $.000242$ **N02+N03 0.000000 TKN 0.000000** 

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**0.45 micron Filt. TKN** 

**Sources Richards and Reinhart (1986) Period 3 Avg. Total Flow, 1/ sq. m x si .472 Recycle, percent total flow: O Periodic Dosing Cycle, sec: 12**<br>Distribution Time. sec/cycle: 6 **Distribution Time, sec/eyele: 6 Avg. Distributed Flow, 1/sq. m x s: .95 Media Depth, n: 6.1 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 28**  Water Temp., C: **COD, mg/1 : Effluent 1.5 micron filt. COD, mg/i: 43 Effluent B0D5, mg/li Effluent 1.5 micron filt. BODS, mg/ls 8 S8, mg/lx 49 VSS, mg/lt 36**  Alk, mg CaC03/lr **Total Applied Oxidized 10 C Nitri fication Influent Cone. Nitrogen, Rate, kg/d/sq. m mg N/1 mg/1 total surface area NH4 11.8 5 .000247 N02+N03 .09 5.2 .000256 TKN**   $3.9$ .000192  $15.1$ 

 $\sim 10^{11}$  MeV  $\sim 10^{11}$ 

**0.45 micron Filt. TKN**
Sources Richards and Reinhart (1986) Period 3 **Avg. Total Flow, 1/ sq. m x as .472 Recycle, percent total flows O Periodic Doming Cycle, secs 12**  Distribution Time, sec/cycles<br>Ava. Distributed Flow. 1/sa. m x s: 195 **Avg. Distributed Flow, 1/aq. m x ss .95 Media Depth, ms 6.1 Media Spec. Surf. Area, sq.m/cu. mt 101 Onda % Wetted Areas 27**  Water Temp., C: **COD, mg/1s Effluent 1.5 micron filt. COD, mg/1: 46 Effluent BODS, mg/1s Effluent 1.5 micron filt. B0D5, mg/1s 13 SS, mg/1s 49 VSS, mg/1s 36 pH: 7.2 Alk, mg CaC03/ls 92 Total Applied Oxidized 10 C Nitrification Influent Cone. Nitrogen, Rate, kg/d/sq. m mg N/1 mg/1 total surface area NH4**   $11.8$  $3.4$  $.000163$ **N02+N03**   $3.1$  $.000148$ **TKN**   $15.1$  $2.4$  $.000115$ 

 $\sim 10^{11}$  MeV  $_\odot$ 

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**0.45 micron Filt. TKN** 

**Source: Richarde and Reinhart (1986) Period 1 Avg. Total Flow, 1/ sq. m x s: 1 Recycle, percent total flow: 0 Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1 Media Depth, m: 6.1 Media Spec. Surf. Area, sq.m/cu. m: 101 Onda % Wetted Area: 27**  Water Temp., C: **COD, mg/1: Effluent 1.9 micron filt. COD, mg/1\* 80 Effluent BODS, mg/1: 38 Effluent 1.9 micron filt. B0D9, mg/1: 23 SS, mg/1: 59 VSS, mg/1: 44**<br> **pH: 7.2** Alk, mg CaCO3/1: 102 **pH: 7.2 Alk, mg CaC03/li 102 Total Applied Oxidized 10 C Nitrification**<br>
Influent Conc. Nitrogen, Rate, kg/d/sq. m **Influent Cone. Nitrogen, Rate, kg/d/sq. m mg N/1 mg/1 total surface area NH4**  10.5  $1.1$ .  $.000121$ **N02+N03**  .000088  $\cdot$  8  $.03$ **TKN**   $4.3$ **18.1**  .000475 **0.49 micron** 

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**Filt. TKN** 

**Source: Richards and Reinhart (1986) Period 1 Avg. Total Flow, 1/ sq# m x s: .389 Recycle, percent total flow: O Periodic Dosing Cycle, sec: 12** 12<br>Distribution Time. sec/cycle: 15 **Distribution Time, sec/cycle: 6 Avg. Distributed Flow, 1/sq. m x s: .778 Media Depth, m: 3 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 26**  Water Temp., C: **COD, mg/l: Effluent 1.5 micron filt. COD, mg/l: 48 Effluent B0D5, mg/l: Effluent 1.5 micron filt. BODS, mg/l: 17 SS, mg/l: 92 VSS, mg/l: 41**  Alk, mg CaC03/l: **Total Applied Oxidized 10 C Nitrification Influent Cone. Nitrogen, Rate, kg/d/sq. m mg N/1 mg/l total surface area NH4 11.4 2.7 .000200 N02+N03 .04 1 .000074 TKN**   $17.1$  $\blacktriangleleft$ .  $.000297$ **0.45 micron** 

**Filt. TKN** 





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**0.45 micron Filt. TKN** 

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Source: Richard and Reinhart (1986) Period 2 **Avg. Total Flow, 1/ sq. m x s: .25 Recycle, percent total flow: O**  Periodic Dosing Cycle, sec: **Distribution Time, sec/cycle: 6 Avg. Distributed Flow, 1/sq. m x s: .5 Media Depth, m: 3 Media Spec. Surf. Area, sq.ra/cu. m: 98 Onda 7. Wetted Area: 23**  Water Temp., C: **COD, mg/1: Effluent 1.5 micron filt. COD-, mg/1: 44 Effluent B0D5, mg/1: Effluent 1.5 micron filt. BOD5, mg/1: 12 SS, mg/1: 49 VSS, mg/1: 36 pH: 7.2 Alk, mg CaC03/l: 92 Total Applied Oxidized 10 0 Nitrification Influent Cone. Nitrogen, Rate, kg/d/sq. \* mg N/1 mg/1 total surface area NH4**  11.8 6 .000319 **N02+N03**   $5.7$ .000303  $.09$ **TKN**   $5.7$  $15.1$ .000303

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**0.45 micron Filt. TKN** 

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**Sources Richards and Reinhart (1986) Avg. Total Flow, 1/ sq. m x a: Recycle, percent total flow: Periodic Dosing Cycle, sect**  Distribution Time, sec/cycle: **Avg. Distributed Flow, 1/sq. m x s: Media Depth, mi Media Spec. Surf. Area, sq.m/cu. ms Onda % Wetted Area: Period 5 Water Temp.,** Ci **COD, mg/1s Effluent 1.5 micron filt COD^ mg/13 Effluent 60DS, mg/1: Effluent 1.5 micron filt BOD5, mg/11 SS, mg/1I 45 pHi 7 23.5 D10/DT» VSS, mg/11 Alk, mg CaC03/l: .972**  O **.972**  6.1 **98 28 .677 32 29**  10 **94 Total Applied Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m total surface area NH4 8.1 1.2 .OOOl 14 N02+N03 .5 1.3 ,000124 TKN 13.1 13.2** .000304

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**0.45 micron Filt. TKN** 

**Source: Richards and Reinhart (1986) Period 6 Avg. Total Flow, 1/ sq. m x s: Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: 20 DlO/DTi COD, mg/1: Effluent 1.5 micron filt. COD, mg/1: Effluent BODS, mg/1: Effluent 1.5 micron filt. BODS, mg/1: SS, mg/1: 45 VSS, mg/1: pH: 7 Alk, mg CaCOS/l: ,667**  O **,667 6.1 98 24 .742 32 25 9 99 Total Applied Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m total surface area NH4 12 3.1 .000222 N02+N03 .2 1.9 .000136 TKN 19 1.2 .000086** 

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**0.45 micron Filt. TKN** 

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**Source: Richards and Reinhart (1986) Period 4 Avg. Total Flow, 1/ aq. m x s: .25 Recycle, percent total flow: O**  Periodic Dosing Cycle, sec: 12<br>Distribution Time. sec/cycle: 12<br>6 Distribution Time, sec/cycle:<br>Avg. Distributed Flow, 1/sq. m x s: **6** 6 **Avg. Distributed Flow, 1/sq. m x s: .5 Media Depth, m: 3 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda y. Wetted Area: 23**  Water Temp., C: **COD, mg/1: Effluent 1.5 micron fiIt. COD, mg/1: 29 Effluent BODS, mg/1: 32 Effluent 1.5'micron filt. B0D5, mg/1: 9 S3, mg/1: 50 VSS, mg/1: pHr 7 Alk, mg CaC03/l: 96 Total Applied Oxidized 10 C Nitrification Influent Cone. Nitrogen, Rate, kg/d/sq. m mg N/1 mg/1 total surface area NH4 8.6 6.2 .000312 N02+N03**   $4.1$ .000207 .05 **TKN**   $7.7$ .000388 12.9

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**0.45 micron Filt. TKN** 

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Data From

Sampayo (1973)

and

Sampayo and Metcalf (1984)

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**Source: Sampayo (1973) Date: 060472 Avg. Total Plow# 1/ sq. a s s: Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Tlae» sec/cycle: Avg. Distributed Plow, 1/sq. m % s: Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: 19.5 DlO/DT: COD, mg/1: 0.45 micron PIIt. COD, mg/1: B0D5, mg/1: Pilt. B0D5, mg/l:**<br>**98, mg/l:** 9 **SS, mg/1: 9 VSS, mg/1: pH: 7.7 AlK, mg CaG03/l: .679 50 .679 6.55 88.6 24 .845 7 2 Total Applied Influent Cone, mg N/1 NH4 5.5 Oxidized Nitrogen, mg/1 5.2 10 C Nitrification Rate, kg/d/sq. m total surface area .000444 N02+N03 8.9 7.7 .000658 TKN 8 6.5 .000555** 

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**0.45 micron Pilt. TKN** 

Source: Sampayo (1973) Avg. Total **Flow**, 1/ sq. **a** x s: **Recycle» percent total flow: Periodic Dosing Cycle» sec: Distribution Time» sec/cycle: Avg. Distributed Flow» 1/sq. m % s: Media Depth» m: Media Spec. Surf. Area» sq.m/cu. m: Onda % Wetted Area: Date Water Te\*p.»** 0: **15.9 COD» mg/1: 0.45 micron Pilt. C8D» mg/1: B0D5» mg/1: F lit. B0D5» mg/1: SS» mg/1: pH: DtO/DT: 060572 .679 50 .679 6.55 88.6 24 .845 17 7.6 VS8» mg/1: Alk» mg CaC03/l: 7 3 Total Applied Influent Gone, mg N/1 Oxld'lzed^ Nitrogen» mg/1 10 0 nitrification Rate» kg/d/sq. m total surface area NH4 3.2 2.7 .000231 N02+W03**  10.1 **7.9 .000675 TKN 9.2 6.8 .000581** 

**0.45 micron Fllt. TKI»** 

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Source: Sampayo (1973) **Date:**<br> **Avg. Total Flow, 1/ sq. ax s: Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle:**  Avg. Distributed Plow, l/sq. m x s: **Media Depth, a: Media Spec. Surf. Area, sq.a/cu. a: Onda % Wetted Area: Water Temp., C: 15.5 DlO/DT: COD, ag/1: 0.45 micron Pllt. COD, mg/1: B0D5, mg/1: put. B005, mg/1: SS, mg/1: 5- VSS, mg/1: pH: 7.5 Alk, mg CaC03/l: 060672 .679 50 .679 6.55 88.6 24 .845 6 3 Total Applied Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, Kg/d/sq. m total surface area NH4 5.5 5.3 .000453 N02+N03 7.1 6.3 ,000538 TKN**  7.4 6 .000513

**0.45 micron put. TKN"** 

**Source: Sampayo (I973> Avg. Total Plow, 1/ aq. a s s: Recycle, percent total flowt Periodic Dosing Cycle, sec: Distribution Time, sec/cycle:**  Avg. Distributed **Flow**, 1/sq. m x s: **Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Date: Water Temp.,** 0: **15.5 COD, ag/l: 0.45 micron PIIt. COD, mg/1: B0D5, mg/1: Pilt. B0D5, mg/l:**<br>88. mg/l: 7 88, mg/l:<br>pH: **pH: 7.6 DIO/DT: 060772 .679 50 .679 6.55 88.6 24 .845 39 31 6**  1 **VS3, mg/1: Aik, mg CaCO3/l: Total Applied Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate# kg/d/sq. m total surface area NH4 6.7 6.5 .000555 N02»N03 9.1** 8.1 **.000692 TKN 7.4 .000513** 

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**0.45 micron put. TKN** 

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**Source: Sampayo (1973> Date:**  Avg. Total **Flow**, 1/ sq. n x s: **Recycle» percent total flow: Periodic Dosing Cycle» sec: Distribution Time» sec/cycle; Avg. Distributed Plow» 1/sq. m : s: Media Depth» m: Media Spec. Surf. Area» sq.m/cu. m: Onda % Wetted Area: Water Temp.» C: 19.5 DIO/DT: COD» mg/1: 0/45 micron Flit. COD» mg/1: B0D5» mg/lr flit. B009» mg/1: S3» mg/1: pH: 17 7.5 VS8» mg/1: Aik» mg CaC03/l: 0608»2 .679 50 .679 6.55 88.6 24 .845 43 31 7 Total Applied Influent Cone, mg N/1**  Oxidized<sup>.</sup> **Nitrogen» mg/1 to C Nidification Rate, Kg/d/sq. m total surface area NH4 7.2 7.1 .000606 N02\*N03 7.7 6.8 .0005»! TKti**  10.2 **9.8 .000752** 

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**0.45 micron Flit. TKN** 

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Source: Sampayo and Metcalf (1984) Date: 060782 **Avg. Total Plow, 1/ sq. m \* s: .679 Recycle, percent total flow: 33 Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m % s: Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: 19.2 COD, mg/1: 0.45 micron Flit. COD, mg/1: B0D5, mg/1: Mit. B0D5, mg/1 : S3, mg/1: pH: DIO/DT: 20 7.6 VSS, mg/1: Alk, mg CaC03/l: .679 7.315 98 24 .761 230 Total Applied Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m total surface area NH4 6.7 4.6 .000286 N02+NO3 7.2 4.2 .000262 TKN 10.5 5.2 .000324** 

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**0.45 micron Flit. TKN** 

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Source: Sampayo and Metcalf (1984) Date: Avg. Total **Flow**, 1/ sq. **n** x s: **Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Plow, 1/sq. m % s: Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: 20.3 DlO/DT: COD, mg/1: 0.45 micron Pilt. COD, mg/1: B0D5, mg/1: Pilt. B0D5, mg/1: SS, mg/1: 29 VSS, mg/1: pH: 7.6 Alk, mg CaC03/l: 060882 .679 33 .679 7.315 98 24 .736 Total Applied Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m total surface area NH4 7.3 .000361 N02+N03 11.1 .000241 TKN 9.8 4.7 000283** 

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**0.45 micron Pilt. TKN** 

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Source: Sampayo and Metcalf (1984) Date: **060982 Avg. Total Plow, 1/ sq. m x s: .679 Recycle, percent total flow: 33 Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: .679 Avg. Distributed Flow, 1/sq. m % s: Media Depth, m: 7.315 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 24 Water Temp., C: 19.9 DIO/DT: .744 COD, mg/1: 0.45 micron Flit. COD, mg/1: B0D5, mg/1: Flit. B0D5, mg/1: SS, mg/1: 14 VSS, mg/1: pH: 7.8 Alk, mg CaC03/l: 226 Total Applied Oxidized 10 C Nitrification Nitrogen, Influent Cone, Rate, kg/d/sq. m mg tt/1 mg/1 total surface area NH4 4.1 2.9 .000177 N02+N03 5.2 1.9** 000116 **TKN 2.2 000134**  6

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**0.45 micron**  Filt. TKN

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Source: Sampayo and **Metcalf** (1984) Date: **080282**  Avg. Total **Plow, 1/ sq. a x s:** 1.398 **25 Recycle, percent total fl^ow:**  Periodic Dosing Cycle, sec: **Distribution Time, sec/cycle: Avg. DIstribated Plow, 1/sq. m % s: 1.358 Media Depth,** a: **7.315 Media Spec. Surf. Area, sq.a/cu. m: 98 Onda % Wetted Area: 31 Water Teap., Ct 22.5 DiO/OT: .695**  COD, mg/l: **a.45 micron Pilt. €00, mg/1:**  5 **B0D9, ag/l: Pilt. B0D5, mg/1: SB, mg/1: II VSS, mg/1: pH: 7 A>lk, mg CaC03/l: 10 C Nitrification Total Applied Oxidized Influent Cone, Nitrogen, Rate, kg/d/sq. m total surface area mg H/l**  mg/t NH4 **1.5 .000171**  5 M02»N03 **7 2.1 .000239 TKN 8.5 2.5 .000284** 

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**0;45 alcron Pilt. TKN** 

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**Source: Sampayo and Metcalf (1984) Date: 080682 Avg. Total Flow, 1/ sq. m x s: 1.358 25 Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.358 7.315 Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 31 Water Temp., C: 24 DIO/DT: .667 COD, mg/1: 0.45 micron Flit. COD, mg/1: B0D5, mg/1:**  5 **Flit. B0D5, mg/1: vss, mg/1 : SS, mg/1: 13 pH: 7 Alk, mg CaC03/l: Total Applied 10 C Nitrification Oxidized Influent Cone, Nitrogen, Rate, kg/d/sq. m mg N/1 mg/1 total surface area NH4 5.6 3.7 .000404 N02+N03 11.2 2.8 . 000306 TKN 8.7 2.3 .000251** 

 $\sim 1000$  km s  $^{-1}$ 

**0.45 micron Flit. TKN** 

**Source: Sampayo and Metcalf (1984) Date: 080982 1.358 Avg. Total Plow, 1/ sq. m z s: 25 Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.358 7.315 Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 31 Water Temp., C: 22.5 DIO/DT: .695 COD, mg/1: 0.45 micron Flit. COD, mg/1:**  5 **B0D5, mg/1: Flit. B0D5, mg/1: .SS, mg/l:**<br>pH: **VSS, mg/1: pH: 6.8 Alk, mg CaC03/l: 10 C Nitrification Total Applied Oxidized Nitrogen, Rate, kg/d/sq. m Influent Cone, total surface area mg N/1 mg/1 NH4 3.7 .000421**  6 **N02+N03 7.9 4.2 .000478 TKN**  8.1 **4.1 .000466** 

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**0.45 micron Flit. TKN** 

**Source: Sampayo and Metcalf <1984) Date:**  081082 **1.358 Avg. Total Flow, 1/ sq. m x s: Recycle, percent total flow: 25 Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.358 Media Depth, m: 7.315 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 31 Water Temp., C: 22.1 DIO/DT: .703 COD, mg/1: 0.45 micron Flit. COD, mg/1: B0D5, mg/1:**   $\overline{\mathbf{z}}$ **Filt. BOD5, mg/l:**<br>**SS, mg/l:** 18 **SS, mg/1: 18 VSS, mg/1: pH: 6.9 Alk, mg CaC03/l:**  188 **Total Applied Oxidized 10 C Nitrification Influent Cone, Rate, kg/d/sq. m Nitrogen, mg N/1 total surface area mg/1 NH4**  5.1 **3.9 .000449**  N02+N03 **5.3 4.5 000518 TKN 6.7 2.4 .000276** 

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**0.45 micron Flit. TKN** 

**Source: Sampayo and Metcalf <1984) Date:**  Avg. Total Flow, 1/ sq. m x s: **Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m % s: Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Netted Area:**  081182 **Nater Temp., C: 22.1 COD, mg/1: 0.45 micron Filt. COD, mg/1: B0D5, mg/1: Filt. B0D5, mg/1: DIO/DT: SS, mg/1: pH:**  18 **6.9 VSS, mg/1: Alk, mg CaC03/l: 1.358 25 1.358 7.315 98 31 .703 8 Total Applied Influent Conc. mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m total surface area NH4 5.6 4.2 .000483 N02+N03**  10.1 **5.4** 000621 **TKN 8.6 4.8 000552 0.45 micron** 

 $\sim 10^{11}$  km s  $^{-1}$ 

 $\sqrt{\mathcal{E}_{\mathbf{p}_1}}$  (  $\mathcal{E}_{\mathbf{p}_2}$ 

**Filt. TKN** 

**Source: Sampayo and Metcalf <1984) Date: Avg. Total Flow, 1/ sq. m x s: Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: 22.7 DIO/DT: COD, mg/1: 0.45 micron Flit. COD, mg/1: B0D5, mg/1: Flit. B0D5, mg/1: SS, mg/1: 8 VSS, mg/1: pH: 7 Alk, mg CaC03/l:**  081282 **1.358 25 1.358 7.315 98 31 .691 5 232 Total Applied Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m total surface area NH4 6.2 4.6 .000520 N02+N03 4.8 5.9 .000667 TKN 8.9 5.9 000667** 

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**0.45 micron Flit. TKN** 

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**081382 Source: Sampayo and Metcalf (1984) Date: Avg. Total Flow, 1/ sq. m z s: 1.358 Recycle, percent total flow: 25 Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.358 Media Depth, m: 7.315 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 31 Water Temp., C: 23.2 DIO/DT: .682 COD, mg/1: 0.45 micron Flit. COD, mg/1:**  5 **B0D5, mg/1: Filt. BOD5, mg/l:**<br>**SS, mg/l:** 11 **SS, mg/1: 11 VSS, mg/1: pH: 6.9 Alk, mg CaC03/l: Total Applied Oxidized 10 C Nitrification Rate, kg/d/sq. m Influent Cone, Nitrogen, mg N/1 total surface area mg/1 NH4 5.8 .000446 N02+N03 6.3 4.4 .000491 TKN 9.5 4.4 .000491** 

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**0.45 micron Flit. TKN** 

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**Source: Sampayo and Metcalf (1984) Date: 093082 Avg. Total Flow, 1/ sq. m z s: 1.358 Recycle, percent total flow: 25 Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m z s: 1.358 Media Depth, m: 7.315 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 31 .701 Water Temp., C: 22.2 DIO/DT: COD, mg/1: 0.45 micron Filt. COD, mg/1: B0D5, mg/1:**   $\mathbf{9}$ **Filt. BOD5, mg/l:**<br>**SS, mg/l:** 36 **SS, mg/1: 36 VSS, mg/1: pH: 7.1 Alk, mg CaCOS/l: Oxidized 10 G Nitrification Total Applied Nitrogen, Influent Cone, Rate, kg/d/sq. m total surface area mg N/1 mg/1 NH4**  6.1 **3.3 .000379 N02+N03 000459**   $\blacktriangleleft$ **TKN 9.5** -1.6 **-.000184** 

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**0.45 micron Filt. TKN** 

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**Source: Sanpayo and Metcalf (1984) Date:**  100182 **1.358 Avg. Total Flow, 1/ sq. m x s: 25 Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.358 Media Depth, m: 7.315 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 31 Water Temp., C: 22.3 DIO/DT: .699 COD, mg/1: 0.45 micron Flit. COD, mg/1: B0D5, mg/1:**  8 **Filt. BOD5, mg/l:**<br>**SS, mg/l:** 37 **SS, mg/1: 37 VSS, mg/1: pH: 7.1 Alk, mg CaC03/l: 258 10 C Nitrification Total Applied Oxidized Influent Cone, Nitrogen, Rate, kg/d/sq. m mg N/1 mg/1 total surface area NH4 6.4 3.4 .000389 N02+N03 4.7 5.9 .000675 TKN 9.1 2.6 .000297** 

 $\sim 100$  MeV  $\sim 100$ 

**0.45 micron Flit. TKN** 

**Source: Saapayo and Metcalf (1984) Date: Avg. Total Plow, 1/ sq. a z s: Recycle# percent total flow: Periodic Dosing Cycle, sec: Distribution Tiae, sec/cycle: Avg. Distributed Plow, 1/sq. a x s: Media Depth, a: Media Spec. Surf. Area, sq.a/cu. a: Onda % Wetted Area: Water Teap., C: 21.7 DIO/DT: COD, ag/1: 0.45 aicron Filt. COD, ag/1: B0D5, ag/1: Pilt. B0D5, ag/1: SS, ag/1: 39 VSS, ag/1: pH: 7.5 Alk, ag CaC03/l: 100482 1.358 25 1.358 7.315 98 31 .71 15 Total Applied Influent Conc. ag N/1 Oxidized Nitrogen, ag/1 10 C Nitrification Rate, kg/d/sq. a total surface area NH4**  9.6 **4.5 000523 N02+N03 3.6 6.3 .000732 TKN 13.4 4.6 .000535** 

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**0.45 aicron Pilt. TKN** 

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**Source: Saapayo and Metcalf (1984) Date: 100582 1.358**  Avg. Total Flow, 1/ sq. **n** x s:  $\ddot{\phantom{a}}$ **Recycle, percent total flow: 25 Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m % s: 1.358 Media Depth, m: 7.315**  Media Spec. Surf. Area, sq.m/cu. m: **98 31 Onda % Wetted Area: Water Temp., C: 22.3 DIO/DT: .699 COD, mg/1: 0.45 micron Filt. COD, mg/1: 13 BODS, mg/1: Filt. B0D5, mg/1: VSS, mg/1: SS, mg/l:**<br>pH: **pH: 7.7 Alk, mg CaC03/l: 234 Total Applied 10 C Nitrification Oxidized Rate, kg/d/sq. m Influent Cone, Nitrogen, mg N/1 mg/1 total surface area NH4 11.5 8.2 .000938 N02+N03 7.2 .000824 TKN 15.7 6.8 000778** 

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**0.45 micron Filt. TKN** 

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**Source: Sampayo and Metcalf <1984> Date:**  100682 **Avg. Total Plow, 1/ sq. m % s: 1.358 25 Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Flow, 1/sq. m x s: 1.358 Media Depth, m: 7.315 Media Spec. Surf. Area, sq.m/cu. m: 98 Onda % Wetted Area: 31 Water Temp., C: 22.9 DIO/OT: .688 COD, mg/1: 0.45 micron Pilt. COD, mg/1: B0D5, mg/1:**  4 **Pilt. BOOS, mg/1: SS, mg/1: 17 VSS, mg/1: pH: 7.6 Alk, mg CaC03/l: Total Applied Oxidized 10 C Nitrification Rate, kg/d/sq. m Influent Cone, Nitrogen, mg N/1 total surface area mg/1 NH4 7.9 5.6 .000631 N02+N03 7.4 5.1 .000574 TKN 9.2 6.6 .000743 0.45 micron** 

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**Source: Sampayo and Metcalf (1984) Date: Avg. Total Plow, 1/ sq.** m x **s: Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Plow, 1/sq. m s s: Media Depth, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: 22.2 DIO/DT: COD, mg/1: 0.45 micron Pllt. COD, mg/1: B0D5, mg/1: Pllt. BODS, mg/1: SS, mg/1: 17 VSS, mg/1: pH: 7.5 Alk, mg CaC03/l: 100782 1.358 25 1.358 7.315 98 31 .701**  12 **Total Applied Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m total surface area NH4 10.4 7.9 .000906 N02+N03 3.8 8.3 000952 TKN 11.1 7.1 .000815** 

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**0.45 micron Flit. TKN** 

100882 Source: Sampayo and Metcalf (1984) Date: **Avg. Total Plow, 1/ sq. a x s: 1.358**  25 **Recycle» percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle:**  Avg. Distributed **Plow**, l/sq. **a** x s: **1.358 7.315 Media Depth, a: Media Spec. Surf. Area, sq.a/cu. a: 98 Onda % Wetted Area: 31 Water teap., C: 22.4 DiO/DT: .697 COD, ag/1: 0.45 aicron Pilt. COD, ag/1: B005, ag/1:**  6 **Pilt. BOD5, mg/l:**<br>SS, mg/l: 23 **33, ag/1: 23 VS3, ag/1: 202 pH: 7.4 Alk, ag CaC03/l: Total Applied Oxidized 10 C Nitrification Influent Cone, Nitrogen, Rate, kg/d/sq. a ag N/1 ag/1 total surface area NH4 6.2 4.6 .000525 N02+N03 5.2 ,000684 TKN 9.4 5.7 .000650** 

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**0.45 aicron Pilt. TKN** 

**369** 

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**Source: Sampayo and Metcalf (1984> Date: Avg. Total Plow\* 1/ sq. m % s; Recycle, percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Plow, 1/sq. m % s: Media Deptli, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Temp., C: 20.8 DiO/DT: COD, mg/1: 0.45 micron Pilt. COO, mg/1: B0D5, mg/1: Pilt. B0D5, mg/1: SS, mg/1: 19 VSS, mg/1: pH: 7.5 Alk, mg CaC03/l:**  101182 **1.358 25 1.358 7.315 98 31 .727 8 Total Applied Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m total surface area NH4 4.8 3.6 .000428 N02+N03 10.7 5.9 ,000702 TKN 7.4 .000714** 

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**0.45 micron Pilt. TKN** 

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Source: Sampayo and Metcalf (1984) Date: Avg. Total **Plow**, 1/ sq. m x s: **Recycle, percent total flow:**  Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed **Plow**, 1/sq. m x s: **Media Depth, »: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area: Water Tenp., C: 20.6 DIO/DT: COD, \*g/l: 0.45 micron Pilt. COD, mg/1: BODS, mg/l: Pilt. BOD5, mg/l:**<br>**SS, mg/l:** 44 **SS, mg/1: 44 VSS, mg/1: pH: 7.6 Alk, mg CaC03/l:**  101282 **1.3S8 25 1.358 7.315 98 31 .731 15**  216 Total Applied **Influent Cone, mg N/1 Oxidized Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m total surface area NH4 4.3 1.9 .000227 N02+N03 6.9 3.2 .00\*383 TKN 13.6 10.2** .001220

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**0.45 micron Pilt. TKN** 

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Source: Sampayo and Metcalf (1984) Date: Avg. Total **Plow**, 1/ sq. **a** x s: **Recycle» percent total flow: Periodic Dosing Cycle, sec: Distribution Time, sec/cycle: Avg. Distributed Plow, 1/sq. m % s: Media- Depth, m: Media Spec. Surf. Area, sq.m/cu. m: Onda % Wetted Area:**  Water Temp., C: 21.1 010/DT: **COD, mg/1: 0.4S micron Pllt. COD, mg/1: B0D5, mg/1: Pllt. BODS, mg/1: SS, mg/1: 41 VSS, mg/1: pH: 7.8 Alk, mg CaC03/l: 101382 1.398 25 1.398 7.319 98 31 .721 13 Total Applied Influent Cone, mg N/1 Ox id 1 zed-Nitrogen, mg/1 10 C Nitrification Rate, kg/d/sq. m tota-1 surface area NH4 6.6 3.4 .000401 N02+N03**  10 **2.2** .000260 **TKN 8.7 2.7 .000319** 

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**0.49 micron Pllt. TKN**