Operation of Sequencing Batch Reactors for Nutrient Removal

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Operation of Sequencing Batch Reactors for Nutrient Removal

7 Contact hours 9 CC10 hours

Aerobic Sequence Batch Reactors (SBR) are unique configured, activated sludge plants. Central to the SBR design is the use of a single tank for multiple aspects of wastewater treatment, e.g., BODs, TSS's and nutrient removal. The process operates with a single sludge in a single reactor basin to accomplish both biological treatment and solids-liquid separation. Operators will review the SBR processes and become familiar with process control tests and troubleshooting. Topics in this course will include the history of SBRs, process configurations, process operation and controls, aeration and mixing, performance expectations, troubleshooting, and instrumentation and automation control.

- 1. Discuss the history of Sequencing Batch Reactors.
- 2. Describe the process of Sequencing Batch Reactors.
- 3. Identify process control tests for Sequencing Batch Reactors.
- 4. Describe the instrumentation and automation control of Sequencing Batch Reactors.

Class Agenda:

- I History (30 minutes)
 - A. Origins 1915
 - B. Resurgence 1970's
 - C. Today
- II Process Description
 - A. Storage/equalization tank (30 minutes)
 - a. Minimum of two tanks
 - b. Conventional screening and grit removal provided as preliminary treatment
 - c. Primary sedimentation is not usually required unless influent suspended solids are excessive
 - B. Process configurations (30 minutes)
 - a. Basic/original
 - b. Cyclic Activated Sludge System (CASSTM)
 - c. Intermittent Cycle Aerated Extended System (ICEAS®)
 - C. Process Cycle Periods: (30 minutes)
 - a. Fill
 - b. React
 - c. Settle
 - d. Draw
 - e. Idle
 - f. Times per period

- D. Cycle options: (30 minutes)
 - a. Flow-paced
 - b. Time-paced
- III Process Operation and Control
 - A. Process control tests (60 minutes)
 - a. Organic loadings
 - b. Solids inventory
 - c. Sludge Quality SSV's and SVI's
 - d. Nutrients
 - i. NH₃
 - ii. NO_x
 - iii. PO₄
 - e. Trouble shooting analyses:
 - i. pH
 - ii. ORP
 - iii. Alkalinity
 - B. Aeration and Mixing (30 minutes)
 - C. Performance expectations: (30 minutes)
 - a. BOD5 and TSS removal
 - b. Nutrient Removal
 - D. Troubleshooting: (60 minutes)
 - a. Observations
 - b. Condition
 - c. Process control analysis
 - d. Possible causes
 - e. Control actions
- IV Instrumentation and Automation Control (I&CA) (60 minutes)
 - A. SCADA
 - B. PLC's and RTU's
 - C. Sensors and analyzers
 - D. Loop controls
- V. Summary and Conclusions (30 minutes)

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Sequencing Batch Reactors MCET | Fill | React | Settle | Draw | Idle |

Process Training Sessions

Before class starts, please:

ENVIRONMENTAL, HEALTH, AND SAFETY TRAINING

- Sign in on Attendance Sheet

During class, please:

- Asks questions
- Feel free to get up and leave the classroom at any time (e.g., rest rooms, phone calls, etc.)

At the end of class, please:

- Answer questions on Post Test
- Evaluate the class on form provided





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Housekeeping

- Start time 8:00 am
- Please mute/silence cell phones
- 10-minute Breaks every hour
- Lunch ~ 11:30 am 12:30pm
- End class ~ 3:30 4:00 pm

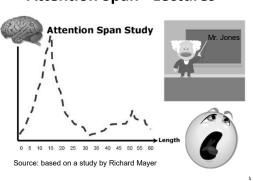


Instructor Expectations

- Begin and end class on time
- Be interactive
- Share experiences and needs
- Less lecture, more discussions
- Make this an enjoyable and informative experience!



Attention Span - Lectures



Ground Rules

- · Discussion is encouraged
- Participate at your own comfort level
- Use terms and examples, we all can understand
- Everyone is different, so please show respect for others in the room
- · Listen with an open mind
- Express opinions of things, not people
- Maintain confidences



Introduction

Class Objectives

Why are we Here?

- This class is intended to aid SBR plant operators improve operations through:
 - Monitoring and process control testing:
 - DO
 - NH₃; NO_X (NO₂, NO₃)
 - PO₄
 - ORP/pH
 - Automation using:
 - Probes
 - Analyzers

Learning Objectives

- To inform operators on equipment components; cycles and phases; and operational concepts of SBRs
- 2. To identify process control, monitoring strategies and process adjustments to enhance:
 - BOD and TSS removal
 - Nitrogen removal
 - Phosphorus removal



Agenda

- Overview
 - Introduction SBR characteristics
 - SBR cycles and phases
- ENR and phosphorus removal
 - Nitrogen removal
 - Phosphorus removal
- Troubleshooting
 - Nitrification/denitrification
 - High decant BOD and TSS
 - Changes in pH and alkalinity

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Ice Breaker and Discussion

- Let's introduce ourselves:
 - Name
 - Plant location
- Facilitated discussion:
 - How well is your SBR plant working?
 - Is your plant:
 - Automated?
 - Meeting nutrient removal standards?
 - Is there room for improvement?

(Allow 20 to 30 minutes for reflection and discussion)

SBR Overview

History, SBR Cycles and Phases

Open Sewers – 1800s



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Need for Wastewater Treatment



Epidemics (e.g., cholera) triggered interest in constructing sewage collection systems in large European cities:

Berlin: 1830London: 1830Hamburg: 1842

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History: Pre-Activated Sludge

- 1690 Sewers (Paris, France)
- 1860 Septic Tank (Louis Moureas)
- 1868 Trickling Sand Filter Process (Edward Frankland)
- 1882 Aeration of sewage (Argus Smith)
- 1911 Chlorination (London, England)
- 1914 Activated Sludge Process (Ardern and Lockett)

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Early History of Activated Sludge

- 1898 formation of the Royal Commission on Sewage Disposal in England
- In the early 1900's, England and the US were seeking small footprint wastewater treatment solutions
- 1912 Famous "30:20 + full nitrification" effluent standard was adopted in England
- New stringent BOD₅ effluent standard inspired creation of the activated sludge process

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Early History of Activated Sludge

 In 2012, in the US, Clark and Gage at the Lawrence Experimental Station began looking at aerating suspended solids in wastewater



 Lawrence Experimental Station became known as "the Mecca of sewage purification"

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Activated Sludge

- Growth and retention of suspended biological solids using oxygen to remove:
 - Soluble Organics (cBOD, COD)
 - Organic Solids (TSS, VSS)
 - Nutrients
 - Nitrogen
 - Phosphorus



Early History of Activated Sludge

- From 1913 1915, in England;
 - Lab-scale suspended biomass experiments were performed in England by Edward Ardern and his coworker, William Lockett under the direction of Dr. Gilbert Fowler



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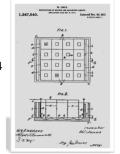
Early History of Activated Sludge

- In separate studies, Walter Jones of Jones and Attwood, Ltd. developed practical applications of the activated sludge process
- 1913 English patents by Jones and Attwood,
 - 25 cents/capita royalties paid by most WWTPs

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Early History of Activated Sludge

- Walter Jones' U.S. activated sludge patent 1,947,540 was issued in 1917 and expired in 1934
- The last of the Jones' US patents expired in 1935



Early History of Activated Sludge

- 1914/1915 Ardern and Lockett published their research findings on aeration of suspended solids (e.g., MLSS)
 - Added the concept of recycling sludge
 - First to use the term "activated sludge" $\,$
- 1915 First full-scale activated sludge plant in Salford, England
 - 80,000 gpd at fill-and-draw operation (SBR)
 - 12,000 gpd at continuous-flow operation (Plug-flow)

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Sequencing Batch Reactors

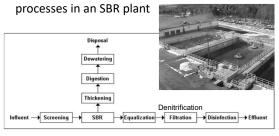




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SBR Unit Processes

Below is a flow chart of possible unit



SBR Systems

SBR systems can have either circular, square or rectangular tanks



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Sequencing Batch Reactor

- Fill-and-Draw activated sludge process
- Treatment cycle can be adjusted to allow:
 - Aerobic conditions BOD removal and nitrification
 - Anoxic conditions denitrification
 - Anaerobic conditions biological phosphorus release



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ENR Stages

- Anaerobic stage No oxygen nor NO₃-N;
 Phosphorus is released; enhances greater TP uptake in the aerobic stage
- Anoxic stage No oxygen; NO₃-N is converted to N₂ gas (Denitrification)
- <u>Aerobic stage</u> Plenty of oxygen; NH₃-N is converted to NO₃-N (Nitrification)

Microorganisms

- <u>Aerobic</u> (Oxic) Organisms requiring, or not destroyed, by the presence of free oxygen
- Anoxic: Organisms requiring, or not destroyed, by the absence of free oxygen; nitrates (NO₃) are present.
- Anaerobic Organisms requiring, or not destroyed, by the absence of free oxygen and NO₃
- <u>Facultative</u> Organisms able to function both in the presence or absence of free oxygen
- Heterotrophic Organisms that use organic materials as their source of cell carbon
- <u>Autotrophic</u> Organisms able to use carbon dioxide and other inorganic matter as their source of carbon
- Filamentous Bulking organisms that grow in thread or filamentous form

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SBR Systems

- Typically SBR plants consist of:
 - Equalization basins
 - Two or more treatment basins
 - Preliminary treatment, such as bar screens to remove rags, sticks and other debris





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SBR Systems

- It is recommended that SBR systems have an influent-flow equalization tank:
 - To handle wet-weather flow increases
 - To remove scum and grease from a single point prior to plant entry



SBR Systems

- Effluent from the SBR basins should also go into a flow equalization tank
 - Allows effluent to be metered
 - Reducing the adverse effects on disinfection or other downstream processes caused by surging flow





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SBR Systems – Process Control

- · Process control testing is essential
 - DO, pH and alkalinity testing
 - Nutrient analyses NH_3 , NO_X (NO_2 and NO_3), PO_4
 - Oxidation Reduction Potential (ORP)
- Location of the online probes should be placed to allow easy access
- Probes must be maintained and calibrated

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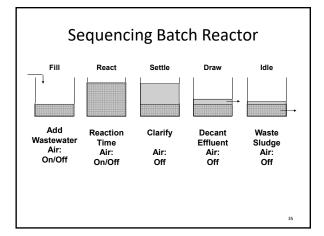
Sequencing Batch Reactor

- Total nitrogen limits of less than 5 mg/l can be achieved
 - Through nitrification in react phase and denitrification in settle and mix-fill phases
 - If more stringent TN discharges are required, then post denitrification filters should be considered
- Phosphorous limits of 2 mg/l or less can be met through biological uptake and chemical addition (aluminum or iron salts)

Sequencing Batch Reactor

- The SBR process cycle consists of five stages sequentially operating in the same tank:
 - Filling (Static, mixed or aerated)
 - Reaction (Air on or off; mixers on)
 - Clarify/Settle (Air and mixers off)
 - Decant/Effluent draw (Air and mixers off)
 - Idle (Sludge waste)

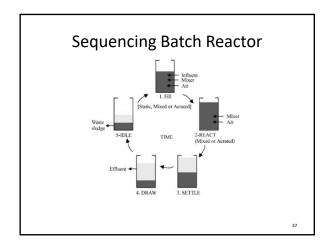
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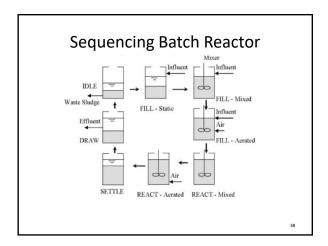


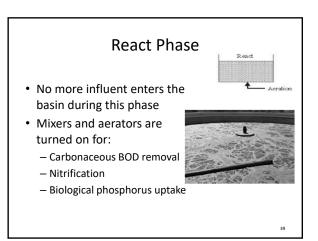
Fill Phase



- SBR receives influent wastewater
- Mixing and aeration can be varied:
 - $\underline{\textbf{Static Fill}}$ No mixing or aeration occurs during fill creating anaerobic conditions; e.g., PO_4 release
 - <u>Mixed Fill</u> Mixers are active, but aerators remain turned off to create anoxic conditions promoting denitrification
 - Aerated Fill Mixers and aerators are both on to convert anoxic conditions to aerobic conditions for PO₄ uptake and to reduce BOD loads before the React Phase





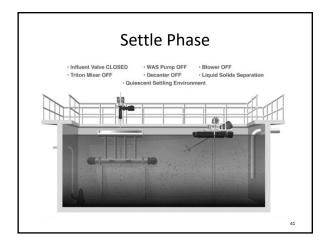


Settle Phase



- Aeration and mixing equipment are turned off allowing the activated-sludge to settle
- Sludge blanket must settle enough as to not interfere with the decant phase
- If the sludge settles too slowly, the biomass may be drawn off during the Decant Phase degrading effluent quality

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A decanter is used to remove the clear supernatant effluent; floating or fixed-arm decanters used: Floating decanters maintain the inlet pipe slightly below the water surface Fixed-armed decanters are cheaper can allow the operator to change the level of the

Decant Phase



decanter



Decant Phase



- Decanted volume ~ one-third of the basin volume and equal to influent fill-step volume
- No surface foam or scum is decanted



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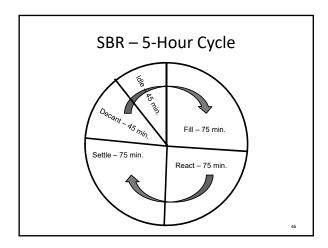
Idle Phase

- Occurs between the decant step and the fill step
- Operator should measure the sludge depth in the basin and waste sludge, if necessary
- Important to maintain adequate solids inventories to assure BOD and nutrient removal

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Sequencing Batch Reactor

Sequence	Volume taken up (as a % of capacity)	Sequence duration (as a % of cycle)	Cycle stage	Object of the sequence	Air
1	60 to 100	33	Feed	Substrate input (denitrification)	With or without (optional)
2	100	33	Reaction	Carbon removal (and nitrification)	With
3	100	16	Sottle	Clarification	Without
4	100 to 65	14	Decant V	Treated water removal	Without
5	65 to 60	4	Idlo 🗸	Excess sludge	Without



Diverse	SBR Cycle and Phase Times			
Phase	4 hours	5 hours	6 hours	
Fill	60 minutes	75 minutes	90 minutes	
React	60 minutes	75 minutes	90 minutes	
Settle	60 minutes	75 minutes	90 minutes	
Draw	30 minutes	45 minutes	60 minutes	
Idle	30 minutes	30 minutes	30 minutes	
Comments	High flows	Average Flows	Low Flows	

Nutrient Removal "Driver"

Chesapeake Bay & CWA¹ Regulations

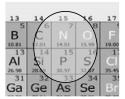
1. CWA – 1972 Clean Water Act

Nutrients

- TN Total Nitrogen (N_{org} + NH₃ + NO₃ + NO₂)
 TP Total Phosphorus (PO₄ + P_{org} + P_{poly})

Nutrients - Overview

Part of the Periodic Table



- $\checkmark\,$ Both Phosphorus and Nitrogen are considered essential for plant and animal life
- ✓ Both are called nutrients

How Too Much Nutrient Pollution Impacts the Chesapeake Bay Ecosystem Balanced Nutrient Diet Too Much Nutrients Chesapeake Bay Program - May

BNR Program

- EPA created the Chesapeake Bay Program in 1983; first Chesapeake Bay agreements signed in 1987
- BNR Programs initiated by Bay states
- For WWTPs greater than 0.5 mgd:
 - 95% of wastewater discharged into the Bay
 - Grant funding available for WWTP upgrades
- WWTP discharge goals:
 - Reduce TP from ~ 6 mg/l to < 3.0 mg/l</p>
 - Reduce TN from ~ 20 mg/l to < 8.0 mg/l

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BNR Program

- To reduce total phosphorus concentrations, most WWTPs began adding chemicals like FeCl₃ or alum
- To reduce total nitrogen concentrations, most WWTPs initiated a capital improvement project to add "Pre" and/or "Post" anoxic zones to already existing nitrification processes for denitrification

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Typical BNR Configurations Sequential BOD Removal Settling Nitrification then Denitrification Nitrates Nitrification SBR Nitrification/Denitrification

ENR Program

- New Chesapeake Bay Agreement enacted in 2000; ENR Program began in that same year
- For WWTPs greater than 0.5 mgd
 - 95% of wastewater discharged into the Chesapeake Bay
 - Grant funding available for upgrades
- WWTP discharge reduction goals:
 - Reduce TP from < 3.0 mg/l to < 0.3 mg/l
 - Reduce TN from < 8.0 mg/l to < 3.0 mg/l

Meeting Nutrient Discharge Limits **Process Strategies**

1. Multiple barriers for TN removal

- Pre-anoxic zone (first stage denitrification)
- Nitrification aerobic zone
- IFAS (enhanced nitrification, optional))
- Post anoxic zone (second stage denitrification)
 Denitrification filters (in lieu of post anoxic zone)

2. Multiple barriers for TP removal

- Biological uptake
 One (maybe two) chemical application points
- Filtration for TSS (particulate TP) removal

ENR Program

- To further reduce total phosphorus concentrations, most WWTPs began adding increased quantities of chemicals
- To further reduce total nitrogen concentrations, most WWTPs initiated a capital improvement project to add "Post" anoxic zones to already existing BNR facilities

Wastewater Discharge Limits

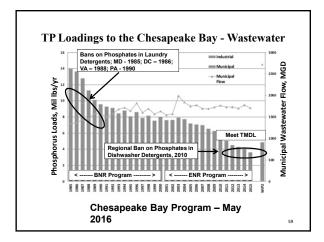
Typical **Total Phosphorus** Standards, mg/l

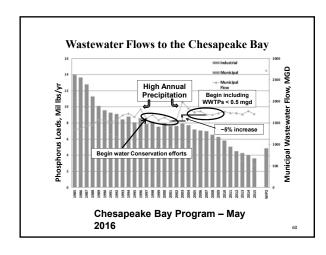
Moderate 0.5 - 1.0 (BNR)

Bay Target < 0.3 (ENR)

- Potomac River < 0.18 (ENR)
- Very Severe < 0.1</p>
- LOT/SOA(a) < 0.05

(a) Limit of Technology/State of the Art





Wastewater Discharge Limits

Typical **Total Nitrogen** Standards, mg/l

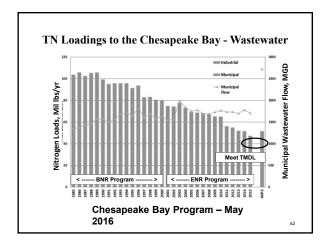
- Moderate 3.0-5.0 (BNR)

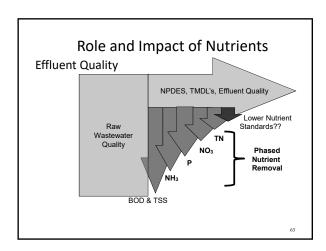
Bay Target < 3.0 (ENR)

Severe < 2.5Very Severe < 1.5

- LOT/SOA(a) < 1.0

(a) Limit of Technology/State of the Art





How will future regulations affect Nutrient Removal Requirements? Regulatory Challenges: - Clean Water Act (CWA) - Chesapeake Bay Program - State Ordinances • Nutrients • Sludge - Local Ordinances

ENR with SBRs

Nitrogen

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Forms of Nitrogen

FORM

REMOVAL PROCESS

- Organic-N
- Converts to ammonia; a small soluble portion is non-reactive (1.0 mg/l)
- Ammonia(um) (NH₃/NH₄+)
- Most abundant form; converts to nitrites/nitrates under aerobic conditions (nitrification)
- Nitrite (NO₂-)/Nitrate (NO₃-)
- Converts to N₂ under anoxic (no oxygen) conditions (denitrification)

Biological Nutrient Removal

- Removes total nitrogen (TN) and total phosphorus (TP) from wastewater
- BNR processes use microorganisms under different environmental conditions:
 - Anaerobic (w/o $\rm O_2$ and $\rm NO_3\text{-}N)$
 - Anoxic (w/o O₂)
 - Aerobic or oxic (with O₂)

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Nitrification Control Parameters

Temperature

- Nitrifiers lose about ½ their activity for each 10°C temperature drop
- In winter, put additional aeration basins online, or increase MLSS
- Either action will increase MCRT

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Nitrification Control Parameters

Dissolved Oxygen

• Maintain MLSS-DO at 2.0 - 4.0 mg/L

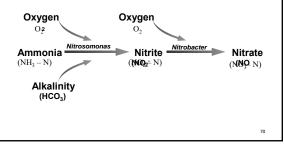
pH / Alkalinity

- Maintain MLSS-pH > 6.8
- Maintain alkalinity residual of at least 70 mg/L, preferably 100 mg/L

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Two-step Nitrification

• For 125 years, nitrification was believed to be solely a two-step process:



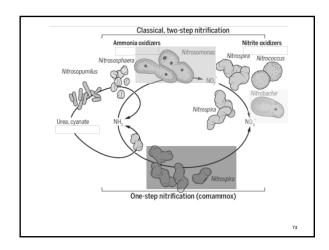
Two-step Nitrification

- Two-step nitrification depends on two organisms e.g., Nitrosomonas and Nitrobacter, which was the basis for hundreds of studies on wastewater nitrification
- A single microbe capable of catalyzing both nitrification steps may actually be a benefit by conserving more energy

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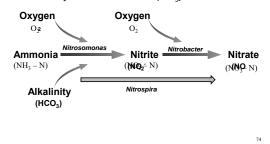
One-step Nitrification - Comammox

- Comammox (COMplete AMMonia Oxidixer) is the name for a single organism that can convert ammonia into nitrite then nitrate
- Existence of comammox organisms were first predicted in 2006
- In 2015, the presence of comammox organisms was confirmed within Nitrospira
- The Nitrogen cycle has since been updated



Nitrification

The oxidation (as by bacteria) of ammonia and organic nitrogen to nitrites (NO_2) and then further oxidation of nitrites to nitrates (NO_3).



Nitrification (Comammox)

NO2

Nitrification (Comammox)

NO2

Nitrification Nitrite oxidation

Nitrite Reduction

Environmental Conditions for Nitrification

- Nitrifying (Autotrophic) Bacteria
- CO₂ Carbon Source for Growth
- Sufficient SRT > 10 days
- Adequate Oxygen > 2.0 mg/l
- Adequate Alkalinity to prevent pH drop > 70 mg/l
- Process operating pH range 6.5 to 8.0
- No Toxics or inhibitory compounds
- Temperature has a significant impact on process

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Activated Sludge Environmental Factors

Parameter	Range		
Food	Proper amount of food to microorganisms (F:M)		
Hydraulic Flow Rate	Within plant design capacity. Excessive flows can result in suspended solids washout		
Oxygen	Many of the bacteria in wastewater require between 1 mg/L to 3 mg/L of dissolved oxygen		
Temperature	Most microorganisms in wastewater grow best between 10 and 25 degrees C. At >35 to 40 degrees C, thermophilic bacteria will take over		
Nutrients	Conventionally a cBOD: Nitrogen: Phosphorus ratio of 100:5:1 is recommended in addition to proper micronutrients such as iron and other trace minerals		
pH	Between 6.5-8.5 is recommended		
Alkalinity	There needs to be enough buffering capacity to maintain pH. Typically 60 mg/L or more of alkalinity at the end of treatment is desired		

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Let's Focus on Aeration





Aeration



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Jet Aeration Systems



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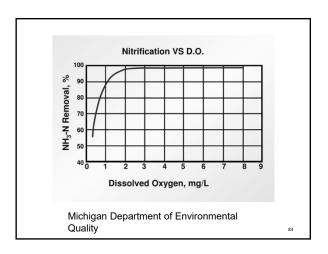
Nitrification- Oxygen Required

- Ammonia oxidation $-2NH_4++3O_2 \rightarrow 2NO_2-+4H++2H_2O$
- Nitrite oxidation $2NO_2$ + $O_2 \rightarrow 2NO_3$ -
- Overall: $NH_4+ + 2O_2 \rightarrow NO_3- + 2H+ + H_2O = 4.57$ g O2/g N for complete oxidation

Nitrification- Oxygen Required

- Ensure adequate D.O. during peak loads D.O. Profiling
- Increase D.O. as needed Blower Capacity?
- Ideal to implement automated, real-time D.O. control to maintain minimum D.O.'s in react phase
 - Air delivery to match D.O. demand (load) variations
 - Reduce energy costs
 - Avoid D.O. bleed through to downstream anoxic stages

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Air Demand Requirements, Ibs/day Treatment Equation | Ib O2/Ib oxidized | Organic Removal BODoxidized = BODinf - BODeff 1.0 - 1.2 | TKNoxidizable = TKNinf - TKNassimilated Nitrification | TKNoxidizable = TKNoxidizable - TKNeff | NPDES Effluent Requirement | OTR Equation | BODs Limit | 1.2 * BODoxidized | BODs + NH3-N Limit | 1.2 * BODoxidized + 4.6 * TKNoxidized |

Importance of Dissolved Oxygen

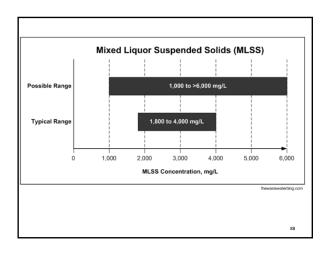
- Oxygen is sparingly soluble in water
- DO is a growth-limiting substrate
- Critical oxygen concentration is about 10% to 50% of DO saturation in water
 - $-\,$ 10% minimum saturation (~ 1.0 mg/l DO) for BOD removal to less than 15 mg/L
 - 20% minimum saturation (~ 2.0 mg/l DO) for complete nitrification

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D.O. - Percent Saturation in Water Water Temperature in Degrees Celsius Oxygen in mg/L Source: Department of Fisheries and Aquatic Sciences, University of Florida

Activated Sludge Processes

Observations – Aeration Tank



Aeration Tank Observations

- Foam:
 - Low MCRT off white, small amounts
 - High MCRT dark brown, larger amounts
- Color of mixed liquor:
 - Low MCRT chocolate brown
 - High MCRT dark chocolate brown



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Process Variations and F/M Ratio Young Sludge" 100 0.20 0.40 0.60 0.80 1.00 1.20 F:M Ratio

Young Sludge

- Start-up or High BOD Load
- Few Established Cells
- Log Growth
- High F:M
- Low MCRT



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Young Sludge



Poor Flocculation Poor Settleability Turbid Effluent

White Billowing Foam

High O₂ Uptake Rate



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Old Sludge

- Slow Metabolism
- Decreased Food Intake
- Low Cell Production
- Oxidation of Stored Food
- · Endogenous Respiration
- Low F:M
- High CRT
- High MLSS



Old Sludge

Dense, Compact Floc Fast Settling Straggler Floc







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Spray Wash for Foam Control



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Nitrification Process Controls

- Temperature
- Flow
- Wasting rate
- SRT
- DO in aeration zone
- pH/Alkalinity in aeration zone
- NH₃-N and NO_x-N probes:
 - End of aerobic stage
 - Plant effluent
 - At end of anoxic stage

Nitrification-Related Process Instruments and Parameters

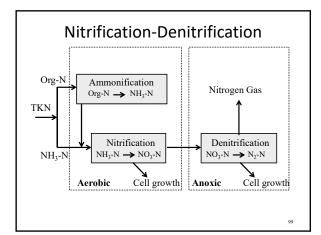
- Temperature
- Flow meters
- Flow rates:
 - Influent/Effluent
 - WAS
- Solids ret. time (SRT)
- pH/alkalinity
- ORP

- Airflow distribution
- DO probe(s)
- DO conc., mg/L
- Ammonia probe(s)
- Ammonia conc., mg/L
- Nitrate probe(s)
- Nitrate conc., mg/L

...

ENR with SBRs

Denitrification



Denitrification Reduction of nitrates or nitrites commonly by bacteria usually resulting in the escape of nitrogen in the air. Carbon Source Carbon Dioxide (CO₂) Nitrate (NO₃) Alkalinity

Carbon for Denitrification

- · Influent WW Carbon
 - Utilized in first anoxic zone
 - EBPR can compete for carbon
 - Limited carbon available for secondary anoxic zones and effective denite
- Endogenous Carbon
 - Slow kinetics limited denite in post-anoxic zones
- Supplemental Carbon
 - Methanol typically used
 - But requires methylotrophic population!
 - Alternatives to methanol ethanol, acetic acid, glycerin, sugars, mono-propylene glycol, proprietary products

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Other Carbon Sources

- Alcohols
 - Methanol
 - Ethanol
- Glycerol/glycerin (Biodiesel by-products)
- Acetates (Acetic acid, sodium acetate)
- Carbohydrates (Sucrose, sugar water, corn syrup)
- MicroCTM Carbohydrate (1000), glycerin (2000), alcohol based blends (3000)

SBR Nitrification/Denitrificatioin

- TKN enters the SBR during the "Fill-mix" stage
- TKN is converted to nitrates during the "aerated react" stage
- Most nitrates formed remain in the "low-water" remaining after clarification
- · Nitrates in the "low-water" are converted to nitrogen gas during the "Fill-mix" stage
- ORP plus nitrate probes would be helpful to determine absence of nitrates

Denitrification-Related Process Instruments and Parameters

- Temperature
- Flow meters
- Flow rates
 - Inflows
 - Internal Recycle
- pH/alkalinity
- ORP

- DO probe(s)
- DO conc., mg/L
- Nitrate probe(s)
- Nitrate conc., mg/L

Oxidation-Reduction Potential

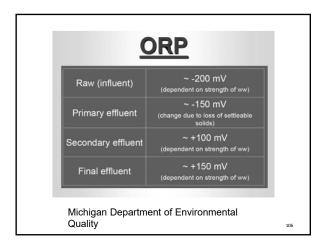


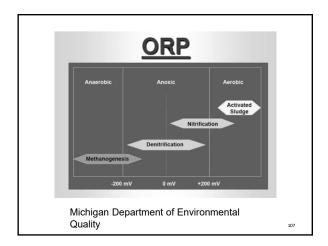
The electrical potential (mv) required to transfer electrons from one compound to another.

> Used as a qualitative measure of the state of oxidation.

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ORP Ranges for Processes		
Biochemical Reaction	ORP, mV	
Wastewater Treatment:		
Nitrification	+100 to +350	
cBOD degradation with air (O ₂)	+50 to +250	
Denitrification	+50 to -50	
Anaerobic Digestion:		
Acid formation (fermentation)	-100 to -225	
Methane production	-175 to -400	
	108	

SBR Denitrification

- Nitrates are converted to nitrogen gas during the "Fill-mix" stage in the absence of oxygen
- Percent nitrate removal in the "Fill-mix" stage is a function of low water volume:
 - = <u>Low-water volume</u> (Low-water + cycle fill Volumes) – % Nitrate Removed =
- Optimum nitrate removal occurs with multiple SBR tanks in service, e.g., high low-water volumes compared to low influent volumes

ENR Phosphorus

Forms of Phosphorus

FOD84

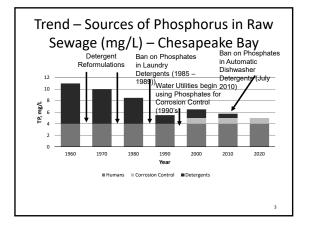
Organic-P

REMOVAL PROCESS

- Converts to orthophosphate form; a small soluble portion is non-reactive (e.g., 0.05 mg/l)
- Condensed Phosphates
- -----

Orthophosphate

- Converts to orthophosphate form
- Most abundant form; chemically reactive and consumed by biological growth



Biological Uptake

- Conventional Biological Uptake
 - To satisfy biological needs (2.0 to 3.0% by weight)
- Enhanced Biological uptake (5 to 7% by weight)
 - Stress induced
 - Release of phosphorus under anaerobic conditions
 - Uptake of phosphorus under aerobic conditions

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Biological Uptake

 <u>Assimilation</u> - Phosphorus removal from wastewater has long been achieved through incorporation of P as an essential element in the biomass

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Assimilation

- Microorganisms are 2-3% P (dry weight)
- Removing biological sludge removes P
- Why doesn't P go to 0?
 - Because carbon is limiting, not phosphorus

Phosphorus-Related Process Instruments and Parameters

- · Flow meters
- Flow rates:
 - Influent/Effluent
 - WAS
- pH/alkalinity
- ORP

- Phosphate analyzers
- Phosphate conc., mg/L



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Biological Uptake

- Enhanced Biological Phosphorus Removal (EBPR) phosphate accumulating organisms (PAOs) store polyphosphate as an energy reserve in intracellular granules
 - Under anaerobic conditions, PAOs release orthophosphate, utilizing the energy to accumulate simple organics and store them as polyhydroxyalkanoates (PHAs) such as poly-β-hydroxybutyrate (PHB)
 - Under aerobic conditions, the PAOs then grow on the stored organic material, using some of the energy to take up orthophosphate and store it as polyphosphate.

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Enhanced Biological P Removal (EBPR)

- Enhanced bio-P removal depends on:
 - Anaerobic conditions (zero dissolved oxygen and zero nitrate)
 - Volatile fatty acids (VFA, rbCOD)
 - Solids management (SRT, WAS, and side streams)

PAO - Phosphate Accumulating Organisms

PAO Able to <u>store soluble organics</u> as Polyhydroxybutyrate (PHB)





EBPR Organisms

- Originally identified as *Acinetobacter sp.* (later turned out this was incorrect)
- Now generically called "phosphate accumulating organisms" (PAOs)
- Some competing organisms accumulate glycogen (GAOs) instead of P
- PAOs require oxygen for growth (aerobic)

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Enhanced Biological P Removal (EBPR)

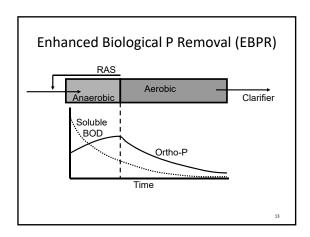
- For EBPR: must be anaerobic, then aerobic
 - P released under anaerobic conditions
 - P then taken up under aerobic conditions
 - 5 mg/L (inf.) \rightarrow 15 mg/L (anaerobic) \rightarrow < 1 mg/L (aerobic)
 - Biological removal or biologically mediated chemical precipitation?
 - − ~1980 largely agreed it was biological

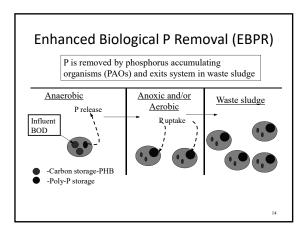
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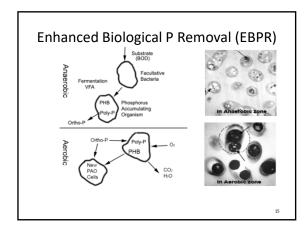
Enhanced Biological P Removal (EBPR)

- Step 1: Anaerobic Phase
 - BOD removal
 - Phosphorus release
- Step 2: Aerobic Phase
 - Phosphorus uptake and creation of new PAOs
 - Phosphorus removal by sludge wasting









Enhanced Biological P Removal (EBPR)

Anaerobic Conditions

Heterotrophic Bacteria Break Down **Organics**

Fermentation

Volatile Fatty Acids (VFAs) Acetate (Acetic Acid)



Selection of <u>PAO - Phosphate Accumulating Organisms</u>
(Able to Out-Compete Other Aerobic Heterotrophic Bacteria for Food When Ana

Enhanced Biological P Removal (EBPR)

Anaerobic Conditions

PAO Take Up VFAs and Covert them to Polyhydroxybutyrate (PHB)



PAO Able to <u>store soluble organics</u> as Polyhydroxybutyrate (PHB)

Ortho-P is Released Into Solution

Enhanced Biological P Removal (EBPR)

Aerobic Conditions

Rapid Aerobic Metabolism of Stored Food (PHB) **Producing New Cells**

> $\frac{PO_4}{}$ Used in Cell Production Excess Stored as Polyphosphate ("Luxury Uptake")



Enhanced Biological P Removal (EBPR)

Aerobic Conditions

PO₄ Used in Cell Production Excess Stored as Polyphosphate

Biomass 5 to 7% P by Weight

(Normal 2 to 3 %)



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Phosphorus Profile July Anaerobic Anoxic Aerobic Anaerobic zone P - Release P - Uptake

Advantages

- Less chemical costs
- Less chemical storage and handling
- Less chemical sludge disposal

Disadvantages

- Not as reliable (initially maybe it is now)
- Limitation need enough BOD
- May be difficult to get < 0.1 mg/L consistently
- Digestion (especially anaerobic) releases P

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Phosphorus Removal with Chemicals

- Precipitation and adsorption with chemical addition:
 - Ferric chloride
 - Aluminum sulfate
 - Polyaluminum chlorides (PACI)
- With effluent filtration, TP concentrations can be reduced to $\sim 0.05 \ mg/l$

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Chemical Precipitation

- Phosphate is an anion: PO₄3-
- Cations can be added to bind with phosphate:
 - Ca²⁺
 - Al³⁺
 - Fe³⁺
- Each forms an insoluble precipitant with alkalinity

Reaction with Lime

• Reaction with lime:

 $5Ca^{2+} + 3PO4^{3-} + OH^{-} \longleftrightarrow Ca_{5}(PO_{4})_{3}(OH)(s)$ hydroxyapatite

• But when lime is added to water:

 $Ca(OH)_2 \leftrightarrow Ca^{2+} + 2OH^ OH^- + HCO_3^- \leftrightarrow H_2O + CO_3^{2-}$ $Ca^{2+} + CO_3^{2-} \leftrightarrow CaCO_3(s)$

• So required dose of lime depends on alkalinity

- Once carbonate is used up, P will be removed

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Co-Precipitation with Metal Salts

Ortho Phosphates
React with
Metal Salts and Alkalinity
To form
Insoluble Phosphorus Compounds

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Phosphorus Removal with Chemicals

- Precipitation of PO₄:
 - FePO₄ and AlPO₄ only exist at very low pH's (< 5.0)
- Co-precipitation: Fe and Al <u>along with</u> alkalinity form metal-phosphate-hydroxide floor
- Adsorption of soluble (PO₄-3) phosphate onto metal hydroxide flocs
- As floc is formed and settles, it also entraps particulate phosphorus

Phosphorus Removal with Chemicals

Chemical Reactions – two mechanisms:

- $\underline{\text{\textbf{Co-precipitation}}}$ (Remove TP to $^{\sim}$ 0.5 mg/l)
 - <u>2</u>Al + 3OH + PO₄ ----> <u>2</u>Al(OH)₃PO₄
- Adsorption (Remove TP < 0.5 mg/l to \sim 0.05)
 - x (AI + 3OH) ---> x (AIOH₃) ↓
 - $x (AIOH_3) + PO_4 ----> x (AI(OH)_3) \cdot PO_4$
 - x > 2; more chemical required as PO₄ levels drop
- Both reactions form Metal (Al or Fe)-Phosphate-Hydroxide floc

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Co-Precipitation Iron Reactions

- FeCl₃ + PO₄⁻³ ---> FePO₄ + 3Cl⁻¹
- FeCl₃ + 3HCO₃⁻¹ ----> Fe(OH)₃ + 3CO₂ + 3Cl⁻¹
- Simplified: Fe + PO_4 -----> $FePO_4$ Fe + 3OH -----> $Fe(OH)_3$
- Combined:

 $2Fe + PO_4 + 3OH ----> 2FePO_4(OH)_3$ Complex (Mole Ratio = 2.0)

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Co-Precipitation Aluminum Reactions

- $Al_2(SO_4)_3 \cdot 14H_2O$ + $2PO_4^{-3}$ -----> $2AIPO_4$ + $3SO_4^{-2}$ + $14H_2O$
- $Al_2(SO_4)_3 \cdot 14H_2O + 6HCO_3^{-1} -----> 2Al(OH)_3 + 6CO_2 + 14H_2$
- Simplified: Al + PO_4 ----> $AIPO_4$
 - Al + 3OH -----> Al(OH)₃
- Combined:

 $\underline{\underline{2}}$ Al + PO₄ + 3OH ----> 2AlPO₄(OH)₃ Complex \ \]



Chemical Addition – Dosages

- 1. Determine incoming TP and biological uptake
- 2. Assume initial chemical mole ratio of 2:1:
- 2. Assume mitted circimited fillowed at 1.1.

 2Al + PO₄ + 3OH = AlPO₄ + Al(OH)₃

 1.7 mg/l of Al per mg/l P

 3.3 mg/l of Al₂O₃ per mg/l P

 1.21 GPH of Alum (@8.3% Al₂O₃ or 0.95 lbs Al₂O₃/Gallon) to remove 1 mg/l P in 1 mgd of flow
- 2Fe + PO₄ + 3OH = FePO₄ + Fe(OH)₃

 3.6 mg/l of Fe per mg/l P

 10.4 mg/l of FeCl₃ per mg/l P

 0.9 GPH of FeCl₃ QB3% FeCl₃ or 1.45 lbs Fe/Gallon) to remove 1 mg/l P in 1 mgd of flow
- 3. Adjust dose to meet discharge standard

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Chemical Addition - Alkalinity/pH

Chemical addition consumes alkalinity and can depress pH:

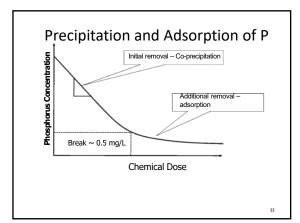
- Al₂(SO₄)₃ x 14H₂O + 6HCO₃⁻¹ ---> 2Al(OH)₃ + 6CO₂ + 3SO₄⁻² + 14H₂O

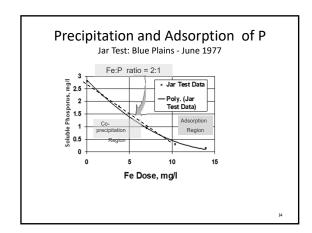
 Al + 3OH -----> Al(OH)₃

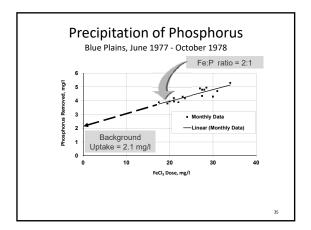
 5.6 mg/l of CaCO₃/mg Al

 3.0 mg/l of CaCO₃/mg Al₂O₃
- FeCl₃ + 3HCO₃-1 ---> Fe(OH)₃ + 3CO₂ + 3Cl⁻¹
 Fe + 3OH -----> Fe(OH)₃

 - 2.7 mg/l of CaCO₃/mg Fe
 0.9 mg/l of CaCO₃/mg FeCl₃







Chemicals used for Phosphorus Precipitation

Chemical	Formula	Removal mechanism	Effect on pH
Ferric Chloride	FeCl ₃ M.W. = 162.3	Metal hydroxides	Removes alkalinity
Aluminum Sulfate (Alum)	Al ₂ (SO4) ₃ .14.3(H ₂ O) M.W. = 599.4	Metal hydroxides	Removes alkalinity
Ferrous sulfate (pickle liquor)	Fe ₂ SO ₄	Metal hydroxides	Removes alkalinity
Poly Aluminum Chloride	AlnCl _(3n-m) (OH) _m Al ₁₂ Cl ₁₂ (OH) ₂₄	Metal hydroxides	none
Lime	CaO, Ca(OH) ₂	Insoluble precipitate	Raises pH above 10

Chemical Addition – Effects on pH

- Alum or iron salts will decrease alkalinity and pH, especially at higher dosages
- Lime raises pH
- PACI will not lower alkalinity or pH

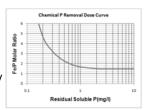
Chemical Addition Rates

- · Dependent on:
 - TP Discharge Limitations
 - Influx TP Loading
 - Biological P Removal Rates
 - Chemical to P Molar Ratios:
 - Al/Fe Salts, Range: 1.6-2.1 to reach 0.5 mg/l P
 > 3.0 to reach < 0.25 mg/l P
 > 5.0 to reach < 0.2 mg/l P

 - >10 to reach < 0.15 mg/l P
 - Dependent on Alkalinity

Phosphorus Removal w/Chemicals

- Add chemical to precipitate phosphorus
- Alum & ferric chloride
- Consumes alkalinity
- Increases sludge production



Chemical Addition - Advantages

- Can be used in every single current wastewater application from Lagoons to EBPR (Enhanced Biological Phosphorous Removal)
- Easy plant trials chemical and feed pumps are the only requirements
- · Low capital costs
- Easy to adjust to changing influent concentrations and flows
- Relatively less complicated method of removal

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Chemical Addition - Disadvantages

- Operation and Maintenance Costs can be higher due to chemical usage
- Handling and Storage of different chemicals and freeze protection
- Iron Products are not recommended in front of UV disinfection due to staining
- If fed at wrong area, it can reduce nutrient levels to beneficial bacteria and cause die off.
- Possible changes to dewatering chemistries
- Increased sludge production

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Metering Pump Definition

- Convey (like any pump)
- Measure (repeated displacement of defined volume)
- Adjust
 - Volume per displacement
 - Frequency of displacements

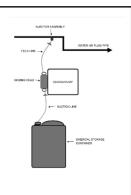




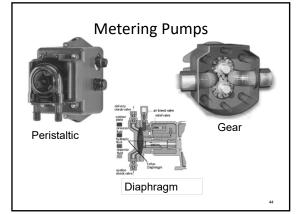


Metering Pumps

- Reciprocating piston, diaphragm, or plunger
- Rotary gear, screw, lobe
- Peristaltic series of rollers to push through tubing or hoses



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Effluent Filtration Applications

- Gravity filters are needed to reduce effluent particulate phosphorus to less than 0.3 mg/L
- Membranes may be needed to reduce effluent particulate phosphorus to less than 0.1 mg/L

Effluent Filtration Application

- Assuming that 2-3% of organic solids is P, then an effluent total suspended solids (TSS) of 10 mg/L represents 0.2-0.3 mg/L of effluent P.
- In plants with EBPR the P content is even higher
- Sand filtration or other method of TSS removal (e.g., membrane) is likely necessary for plants with low effluent TP permits

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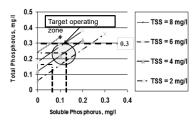
TSS Removal Requirements

TP Limit, mg/L	Max TSS, mg/L
0.1	3.0
0.2	5.0
0.3	7.0
0.4	9.0
0.5	11

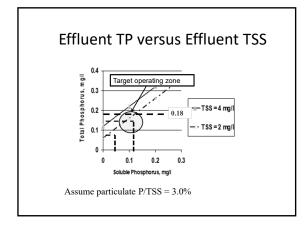
Assume soluble P = 0.05 mg/L; particulate P/TSS = 3.0%

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Effluent TP versus Effluent TSS



Assume particulate P/TSS = 3.0%



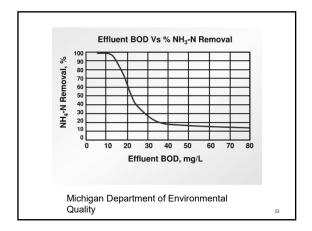
Troubleshooting

Nitrification/Denitrification

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Optimizing Nitrification

- Remove $BOD_5 \le 20 \text{ mg/L}$
- Optimize dissolved oxygen in aerobic zones
- Last step: add alkalinity only if needed



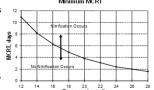
Obstacles to Achieving Nitrification

- Inadequate aeration capability
- Inadequate biomass quantity(MCRT)
- Poor hydraulics limiting MLSS in tanks
- Poor sludge settling/excessive filamentous bacteria
- · Insufficient alkalinity
- Inhibitory chemicals

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Nitrification Process Monitoring

- Key Factors:
 - Slow growth requires adequate aerobic
 SRT or MCRT or CRT
 - **DO** typically >2mg/L
 - pH 6.5-7.5
 - Target effluent alkalinity of 50 to 75 mg/L as CaCO₃



- Overall Reaction:
 - NH₄⁺ + 2 O₂ → NO₃⁻ + 2H⁺ + H₂O

Operational Controls for Nitrification

Nitrifiers Grow Slowly

CRT Must Be Long Enough
> 5 Days (minimum)
Best > 8 Days

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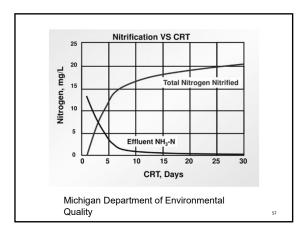
Operational Controls for Nitrification

Aerobic React Time Must Be Long Enough (> 5 hrs.)

(BOD Removal Must Occur First)

F:M Ratio Must Be Low Enough (< 0.25)

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Effect of Temperature on Nitrification

As temperature increases, nitrifier growth rate increases (within the range of 4° C to 35° C). T \uparrow μ \uparrow

As nitrifier growth rate increases, required MCRT decreases. MCRT ↓ μ1



Rule of Thumb: For every 10°C increase in temperature, nitrifier growth rate doubles, required MCRT is cut in half and required MLSS concentration is also reduced.

Effect of Temperature on Nitrification

Lower Temperatures Cause Slower **Nitrifier Growth Rate**

Minimum of 59 Degrees F. (15 °C) for 90 % Nitrification

Below 50 Degrees F. (10 °C) Expect Maximum of 50 % Nitrification

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Nitrification – Temperature Impacts

- Need to plan for temperature transitions especially in areas with significant seasonal changes.
- Build solids inventory, decrease F:M ratio, and increase MCRT/SRT as seasonal (cold weather) changes approach.

Wastewater Temperature Range (°C)	Target Food to Mass Ratio	Target Average Aeration MLSS (mg/L)	Typical MCRT/SRT, Days	NH ₃ , mg/l
14-18	0.05	3000	15	1 - 2
18-22	0.075	2500	10	0.5 - 1
22-25	0.1	2000	7	0.5 - 1

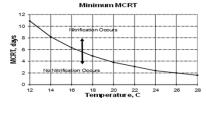
Nitrification Process Monitoring

- Key Factors:
 - 1. MCRT solids inventory
 - 2. DO concentration
 - 3. Alkalinity

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Nitrification Process Monitoring

- Key Factor 1
 - Slow growth requires adequate aerobic SRT
 - MAINTAIN ADEQUATE SOLIDS INVENTORY

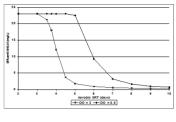


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Nitrification Process Monitoring

■ Key Factor 2

Maintain target DO concentration



Effect of Dissolved Oxygen Concentration on Nitrification

As dissolved oxygen increases, nitrifier growth rate increases up to DO levels of about 5 mg/L.

DO ↑ u



Rule of Thumb:

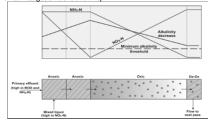
Maintain dissolved oxygen concentration at 2.0 mg/l or higher for complete nitrification.

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Nitrification Process Monitoring

Key Factor 3

Maintain target effluent alkalinity

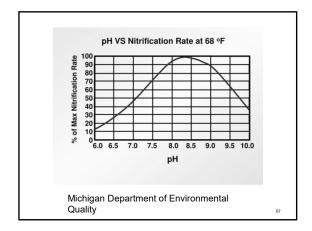


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Nitrification- Alkalinity/pH

Nitrifiers utilize inorganic carbon/consume alkalinity:

- Aerobic chemoautotrophs
 - $NH_4^+ + 2HCO_3^- + 2O_2 \rightarrow NO_3^- + 2CO_2 + 3H_2O$
- Theoretically, 7.14 g alkalinity as CaCO_3 consumed for each g of ammonia nitrogen oxidized to nitrate.
- Typically, lime or caustic soda is added to make up for alkalinity loss.



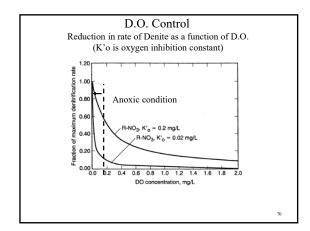
Inhibition of Denite Process

- Environmental Conditions
 - pH
 - Temperature
 - Aerobic Conditions! Keep DO< 0.2 mg/L
- Insufficient amount of rbCOD (Carbon Substrate).
- Presence of Chemical Inhibitors:
 - Substrates, intermediates, and products of denitrification
 - Synthetic organic chemicals
 - Heavy metals Hg, Ni, Pb, etc.

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High DO in the Anoxic Zones

- High DO in the anoxic zones may be more of a problem during the winter because more DO can be absorbed by colder water and biological kinetics are reduced.
- Lower the nitrate recycle rate in the winter if necessary



Denitrification Problems

Possible Causes	Solution
Not enough nitrates being returned to anoxic zone	Increase nitrate recycle pump speed
Not enough BOD entering anoxic zone	 Bypass primary clarifiers, or Add supplemental carbon (for example, methanol) to anoxic zone
BOD entering the anoxic zone breaks down too slowly	Add readily available carbon source such as methanol to anoxic zone or increase the anoxic zone hydraulic retention time
High DO in the anoxic zone	Try to limit backmixing of air from the aerobic zones or decreasing the DO in the AT influent. Decrease nitrate recycle rate if necessary.

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Troubleshooting

High/Low Dissolve Oxygen

Typical Aeration Basin Control Strategy - DO DO Setpoint DO setpoint chosen to minimize historical NH₃ breakthrough

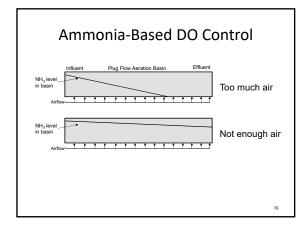
New Aeration Basin Control Strategies

- Ammonia-based DO control
- Nitrate-based DO control

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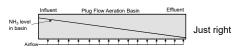
Objective of Ammonia-Based Aeration Control

- Aeration options:
 - Full nitrification
 - Incomplete nitrification
 - Reduce effluent ammonia peaks
- Potential benefits of incomplete nitrification include:
 - Decreased energy expenses (for aeration)
 - Possibly increased denitrification with less supplemental carbon addition
 - Possibly improved Bio-P removal



Ammonia-Based DO Control

- Operator selects effluent ammonia setpoint
 - Complete nitrification, NH $_3$ -N $^{\sim}$ 0.1 mg/L
 - Incomplete nitrification, NH₃-N ≤ 1.0 to 2.0 mg/L



- When effluent ammonia is greater than setpoint, controller increases DO
- When effluent ammonia is below setpoint, controller decreases DO

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Objective of Ammonia based Aeration control

- Aeration is limited to:
 - Prevent complete nitrification
 - Reduce effluent ammonia peaks
- Potential benefits include:
 - Decreased energy expenses (for aeration)
 - Possibly increased denitrification with less supplemental carbon addition
 - Possibly improved Bio-P removal

Ammonia Feed Forward – Feedback Control Not, Septemblish Not Septemblish

Troubleshooting

High Decant BOD and TSS

Loss of Solids due to High Blankets

- Old sludge (low F/M) decrease MCRT
- Young sludge (high F/M) increase MCRT
- · Filamentous bulking
- Slime bulking add nutrients
- Foam trapping pretreatment for oil & greases

Rapid Sludge Settling

- Monitor with SSVs, SVI, F/M
 - Pin floc
 - Straggler floc
- Probable cause, low F/M Increase F/M by decreasing MLVSS

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High Effluent BOD, TSS, NH₄-N

- Low MLSS increase
- Low D.O increase
- High organic loading increase MLSS and aeration cycle
- High nitrogen loadings and effluent NH₄
 - Increase MLSS and aeration cycle
 - Low D.O.? Nitrifiers? Alkalinity?
 - Short aeration times?

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Troubleshooting

Changes in pH and Alkalinity

pH Alkalinity Changes

- H₂S production
- Respiratory release of CO₂
- Nitrification inhibition
- Nitrification
- Denitrification
- Metal salt addition

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IC&A

Overview

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Before You Can Control a Process Variable, You Must be able to Monitor It with Reliable Sensors







IC&A Drivers

- Instrumentation, control, and automation (IC&A)
 - Initiated in the 1970s
 - Major push in the 1980s
 - Meet nutrient regulatory requirements
- Improve process performance and cost efficiency
- BNR/ENR processes require effective DO control (enough but not too much)

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Process Control

- Operators use a combination of probes and analyzers to monitor <u>and control</u> BNR/ENR processes:
 - DO
 - Nitrification (Ammonia profile)
 - Denitrification (Nitrate profile)
 - Phosphorus removal

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Process Control Loops

- Monitoring and control of a process
- WWTPs rely on four building blocks:
 - A process model concept
 - Monitoring and control strategies
 - <u>Sensors that provide accurate and precise on-line</u> <u>data to controllers</u>
 - Actuators or control elements that implement controller output

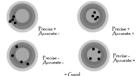
Process IC&A - What's Needed?

- Sensors Probes and/or analyzers:
 - Accurate
 - Precise
- Communications network Analog, digital, radio, and/or telemetric
- Controllers:
 - Modules or Remote Terminal Units (RTUs)
 - Programmable Logic Controllers (PLCs)
- Actuators, e.g., valves, pumps, blowers

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Accuracy and Precision

- Accuracy: How close is it to the actual reading?
- Precision/Repeatability: Does it provide the same answer each time?



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Why Automate?

- Monitor basic information about the processes
- Accurate and precise instrumentation is now available to automate system processes
- Eliminate manual measurements, e.g. dependency on delayed lab measurements
- Save time, save money, and increase efficiency
- Allow facilities to operate at limits of technology

Process Control

- WWTPs are never in a steady state; subject to "disturbances"
- A major incentive for automated process control is to minimize impacts of "disturbances" on plant processes:
 - Wastewater influent constituents, their concentrations, and flow rates
 - Discrete events such as rainstorms, peak loadings, and spills

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Process Control

- Key factors in automated process control systems:
 - Responsiveness
 - Ability to deal with disturbances
- "A responsive control system" means the controlled variable responds quickly to adjustments in the manipulated variable
- Frequency and magnitude of disturbances should be minimum

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Definitions

- Input Variable This variable shows the effect of the surroundings on a process and normally refers to factors that influence the process:
 - Manipulated inputs: variables in the surroundings that can be controlled by an operator or the control system
 - Disturbances: inputs that can not be controlled by an operator or control system
- Output variables- Also known as control variables; these are variables that are outputs of the process.

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Process Control

- Three physical properties are typically monitored in wastewater:
 - Liquid flow: Influent/effluent, recirculation, return activated sludge (RAS), sludge wasting quantities, chemical addition
 - 2. <u>Constituent Concentrations:</u> DO, MLSS, BOD₅, TSS, nutrients, sludge solids
 - 3. Gas volumes: air, digester gas

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Common Controlled Variables

- Aeration
- BOD₅ and TSS loadings
- Ammonia, Nitrate, and Phosphate loadings
- · Chemical Addition
- Internal Recycles (MLE processes)
- Low water level and fill level during fill stage (SBR)
- Sludge Wasting Rates

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Common Controlled Variables

- Aeration
 - Set DO levels in different sections of process
 - Control aeration time (cyclic aeration)
- BOD₅ and TSS loadings
 - Maximize removal of BOD₅ and TSS before nitrification/denitrification
- Ammonia, Nitrate, and Phosphate mass loadings
 - Avoid overloading unit processes

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Common Controlled Variables

- Chemical Addition
 - Methanol, Ferric/Alum, alkalinity feed rates
- Internal Recycles (MLE processes)
 - Set recycle flow rates based on process conditions
- Low water level to fill level (SBRs)
- Sludge Wasting Rate
 - Control Solids Retention Time One of the most important parameters for advanced BNR

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Nitrification-Related Process Instruments and Parameters

- Temperature
- Flow meters
- Flow rates:
- Influent/Effluent
- WAS
- Solids ret. time (SRT)
- pH/alkalinity
- ORP

- Airflow distribution
- DO probe(s)
- DO conc., mg/L
- Ammonia probe(s)
- Ammonia conc., mg/L
- Nitrate probe(s)
- · Nitrate conc., mg/L

Denitrification-Related Process Instruments and Parameters

- Temperature
- Flow meters
- Flow rates
 - Inflows
 - Internal Recycle
- pH/alkalinity
- ORP

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- DO conc., mg/L
- Nitrate probe(s)
- · Nitrate conc., mg/L

Phosphorus-Related Process Instruments and Parameters

- Flow meters
- Phosphate analyzersPhosphate conc., mg/L
- Flow rates:
 - Influent/Effluent
 - WAS
- pH/alkalinity
- ORP

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What are sensors?

- Devices which measure a target variable
- Two components
 - Sensing Element
 - Tracks the variable being measured
 - Sends signal to transmitter
 - Transmitter
 - Converts signal for use on local display
 - Sends signal to controller/SCADA





Sensor image from: http://www.capitolacientific.com/core/media/media.ni?id=51573&c=1290437&h=767aff1ece59e1f3e

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Sensors - Calibration and Validation

- Example: side-by-side grab sampling with immediate filtration/analysis and comparing grab value with instrument value
- Typically conducted three times per week dependent on plant and sensor



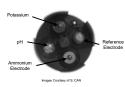
Three Main Sensor Types

- Ion Selective Electrode (ISE) probes
- Wet Chemistry (Colorimetric) analyzers
- Optical (UV) probes

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Ion Selective Electrode (ISE)

Probe-type sensors that use an ISE probe and reference electrode



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Detection by ISE Probe

- Specific ions adhere to membrane on measurement electrode
- Those ions do not affect reference electrode
- Measure potential (voltage) difference
- Replace cartridge ~ 6-12 months





Nutrient ISE Probes

Brand	Model
YSI	VARiON (IQ SensorNet)
Hach	AN ISE sc
Endress & Hauser	ISEmax

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Wet Chemistry (Colorimetric) Analyzers

- Utilize a colorimetric method for measuring a constituent in a sample
- Withdraw a sample from the wastewater flow and pump it to a nearby analyzer





ACHE

www.capitalh2o.com/image/page_photos/che n_front.jpg

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Optical (UV) Probes

- Utilize an ultraviolet (UV) light source to measure an absorbance and/or transmittance of UV light waves passing through a sample
- Similar to UV light absorbance spectrophotometers in a lab

UV Transmittance Path



Optical (UV) DO Probes

Brand	Model
YSI	FDO (IQ SensorNet)
Hach	LDO Model 2
Endress & Hauser	Oxymax
Insight IG	Model 1000
ATI	Q45D

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Why are Sensors important?

- Automate data collection
- Monitor performance when no one is looking
- Optimize process performance
- Minimize energy use and chemical consumption
- Sensors can be paired with calibrated process models to enhance operations

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IC&A Drivers

- Collect and record data for creating reports, profiling process performance, and storing data
- Reduce costs:
 - Operating costs, e.g. chemicals, energy (for aeration), labor
 - Capital costs, e.g., increase nutrient removal capacity by 10% to 30%; possibly reduce future system investments by another 20% to 50%

SBR Systems – Process Control

 Any online instrument should be tied into the plant SCADA system, allowing operators to verify and make changes as needed to the SBR control parameters.

Supervisory Control

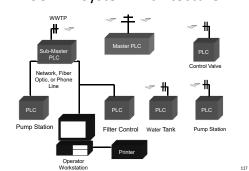
Denotes a control system that manages the activities of several integrated unit operations

A control system that directs and coordinates the activities of several interacting pieces of equipment

- SCADA
- CMMS Avantis



SCADA System Architecture



Programmable Logic Controller (PLC)

- Introduced around 1970 to replace electromechanical relay controllers
- Microprocessor-based
- Executes instructions/algorithms that implement logic, sequencing, counting, and arithmetic functions for controlling equipment and processes

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SCADA System Architecture

- Goals:
 - Graphical Representation of Entire System
 - Continuous Information at Operator Workstations
 - Automatic Control
 - Remote Control
 - Alarming/Paging
 - Trending/Reporting
 - Easily Expandable

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Typical Hardware – PLC's

 Programmable Logic Controllers (PLC's) -Standardized on Allen-Bradley and Micrologix controllers





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Typical Hardware – Radio Communication

• MAS/MDS radios and modems for Radio Communication





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Typical Hardware – Phone Line Communication

• Mille Research modems for Phone Line Communication



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Typical Software

- Standardized on Rockwell Software:
 - RSLogix for programming
 - RSLinx for communications
 - RSView for Operator Interface
 - RSMessenger for Alarm Paging
- LapLink Software allows connection to operator workstation from a remote computer through a dialup modem

SCADA Economic Advantages

- Implementation of "On-Call" shifts
- Reduced Operator Travel Time





SCADA Economic Advantages

- Greater Speed, Accuracy, and efficiency
- Increased Reliability
- Reduced Maintenance
- Safety and Security





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Questions?



Thank You

Maryland Center for Environmental Training College of Southern Maryland La Plata, MD