to Optimize Process Control How to Utilize Your Lab Data

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Why Do Lab Results Matter?

- Valuable record of plant operations
- plant is running Tell you how efficiently your
- Required by regulatory agencies
- Help troubleshoot your process Help determine when an expansion is necessary

should be performed carefully For these reasons, lab tests and consistently!

What Can Impact the Quality of Your Data?

- Representative Sampling
- Sample Handling
- Sample Contamination
- Equipment Contamination
- Sample Hold Time
- **Biological Changes**
- Chemical Changes
- Physical Changes



Know What to Expect...

What is my wastewater comprised of?

- Domestic Only Flow
- Significant or Categorical Industries
- Inflow or Infiltration
- Combined Sewers

why? significantly, you should be If your influent changes asking

Know Your Expected Load...

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TKN
                                TSS
                                                       BOD - 0.18 to 0.22 ppd / capita
                                                                                       Flow
- 0.029 to 0.032 ppd / capita
                             - 0.20 to 0.25 ppd / capita
                                                                                    - 75 to 130 gpd / capita
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Loading = 6042 ppd / 30,000 people
                                                                 is 1.5MGD = 6042 ppd BOD
                                                                                           If your CBOD is 483 mg/l and flow
        0.20 ppd / capita
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PERFECT!

But What If?

in your What if there town? are only 7,000 people

Loading = 6042 ppd / 7000 people 0.86 ppd / capita

processing plant or dairy etc. Now that's a problem.. Extreme example unless you have a food

Correlation to Process Operations Laboratory Analysis and the





- рH sample. the acidity or basicity of a **1**S മ numerical expression O f
- Acidic 0 to 7
- Basic 7 to 14
- рH 10 Neutral times more is a log - 7 is considered scale acidic than I so pH neutral pH of 5 of 4
- and 100 times more acidic of 6. than pH

- preferred; ω standard starting with 7. calibration is ALWAYS
- Should run duplicates
- manufacturer than calibration standards. Check standard from another

- Ensure slope is within equipment
- manufacturer standards. coating the probe. interfere with the analysis by Oils, greases and fats can



pH as it Relates to Process Control

Preliminary and Primary Treatment

- to through to other processes. less than 7 can cause greases stay in suspension and pass
- Consistently low pH at the cleaning should be performed. user. System sampling and/or decomposition - or an industrial collection system due to anaerobic conditions influent can be a sign of in the

pH as it Relates to Process Contro

Fixed Film Systems

- Applies to RBCs, trickling filters О **К** roughing towers
- pH outside the range of 6.5 will prohibit nitrification. carbonaceous compounds; above 8.5 can cause reduced treatment of († 0
- pH >10 or <5 can cause excessive sloughing.
- necessary Chemical adjustment may be

pH as it Relates to Process Control

Activated Sludge Processes

- will occur at pH less between 7 and 8.3 and inhibition Optimum growth for nitrifiers is than 7.
- pH consistently less than 6.5 can settleability problems. create bulking and MLSS support filamentous growth and

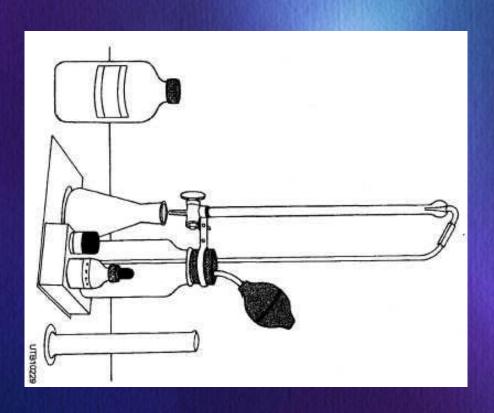
pH as it Relates to Process Control

Sludge Digestion

- Aerobic digestion is best done around pH of 7; if less than 6 nitrification or carbon dioxide look for causes - low alkalinity,
- become toxic to methane formers. will cause dissolved ammonia to between 6.8 and 7.2. pH less than Anaerobic digestion is best done to methane formers and above 8 6 will cause VFAs to become toxic

clue as to what might be causing the of impact it could be having on your numbers and ask yourself what kind Don't just record your influent and indication of a problem and also a effluent pH — pay attention to the process. pH is often our first problem!

Alkalinity



Alkalinity

- Alkalinity is the water absorb before there is a water or how much acid can the substantial change in pH neutralizing capability of acid the
- method so turbid or colored water Typically a color titration can cause interferences.
- Great process control parameter!

Alkalinity

- or plastic. Samples can **9** collected in glass
- Technically there is a 14 day sample. biological processes occurring in slow the biological action of the practical or cooled to 0-6°C to available alkalinity, the sample hold time; however, because should be analyzed as soon as the sample can change the

- measured and reported in terms In equivalent calcium carbonate (CaCO3). wastewater alkalinity y LS
- measured to a pH of 4.5 SU. is total alkalinity which is For wastewater, the measurement
- indicator of biological activity. Alkalinity is utilized as an

- NO3 and sulfate SO4. In available - dissolved 02, nitrate of oxygen that are typically wastewater there are 3 types
- dissolved oxygen to convert food Aerobic microorganisms use to energy.
- Nitrifiers ammonia to nitrate use 02 to convert

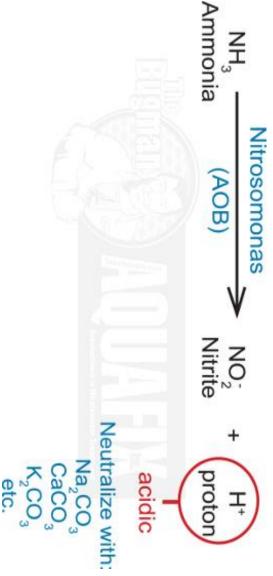
- nitrogen gas. In bacteria use the oxygen from the nitrate and it is converted to an anoxic environment,
- compounds. will use the oxygen from sulfate sulfide and other sulfur which is converted to hydrogen are no longer available, microbes conditions when DO and Nitrate And finally, in anaerobic

- Alkalinity is lost in an nitrification. activated sludge system during
- oxidized. for every mg of ammonium that is 7.14 mg of alkalinity is utilized
- M nitrification. can lack of carbonate alkalinity effectively stop

- თ . დ In significantly at pH values below sensitive and will decline addition, nitrification Z L PH
- Maintaining an appropriate level Double carbon the pH and provide inorganic of alkalinity will help buffer Benefit! for the nitrifiers

producing. Influencing factors are: Basically aerobic wastewaters are net acid

- Biological nitrification in aeration tanks or aerobic digesters
- Acid forming stage of anaerobic digestion
- Gas chlorination for disinfection
- Chemical addition of aluminum or iron



So how much alkalinity is enough?

- Alkalinity levels should be at least 8 times the concentration of influent ammonia.
- ability. Denitrification does add back some alkalinity if your plant has that
- Rule of thumb is to ensure at least completion to push nitrification to 75 to 150 mg/l residual alkalinity

How Much Alkalinity Do I Need?

Example:

Influent ammonia - 30 mg/l

So, $30 \text{ mg/l} \times 7.14 \text{ mg/l}$ alkalinity = 214.2 mg/l alkalinity required

nitrify 30 mg/l ammonia. alkalinity required to fully 214.2 mg/l is the minimum amount of

How Much Alkalinity Do I Need?

And alkalinity is 145 mg/l then, if the actual influent

214.2 mg/l needed - 145 mg/l actual= 69.2 mg/l alkalinity deficiency

or chemical addition of calcium Supplemental alkalinity will be carbonate etc. necessary through denitrification

How Much Alkalinity Do I Need?

mg/1. In alkalinity to be between 75-150 an alkalinity residual (effluent addition, most experts recommend

- Assists in stabilizing the pH neutral range. range in the process in the near
- microorganism balance is best at neutral pH (7 to 7.4) removal efficiency and proper
- Leads to better settleability.

Operations! Maintaining a Sufficient Amount of Alkalinity can Provide for Alkalinity is the Key to Steady State Expanded Treatment Capacity! Improved Performance and

Total Suspended Solids



Liquor Suspended Solids Total Suspended Solids / Mixed

- One operator / analyst can run of the most important test an
- Routinely utilized for:
- **Efficiencies** Process Loading / Removal
- Time (MCRT) and Sludge Age Calculate the Mean Cell Residence
- Determine Sludge Wasting Rate
- Calculate (SVI) the Sludge Volume Index

Iotal Suspended Solids

- dried and weighed Wastewater put through a that has been pre-washed and then filter
- being utilized micron or less; and varies in size depending on what type of funnel is Filter has a pore size of 2.0
- Vacuum applied and liquid pulled filter to be dried at 104°C through leaving residue on the
- ml multiplied by 1000 Weight of filter and residue filter divided by sample volume in

Iotal Suspended Solids

Interferences with analysis:

- Trying to filter too large a sample volume can create a water tight mat on the filter paper
- Water trapping can occur in the result 105°C resulting in a falsely high that will not evaporate at 103-
- disperse oil prior to analysis Samples with visible oil and grease should be mixed in blender to

Iroubleshooting TSS

paper (stuck to drying pan) mg/l between weighing concentrated in bottle paper not dried long enough or if Standard result too high - Filter not mixed or poured properly papers not pre-washed, balance not used previously, standard could be Standard result too low - Sample calibrated properly, loss of filter Filter blank changed more than 0.5 filter

ISS as it Relates to Process Control

- mg/1Average Raw Wastewater 120 to 400
- between 50 and 70% of influent TSS Primary clarification should remove
- Average MLSS is typically between systems 1000 and 4000 mg/l; however some (MBRs) run considerably higher

TSS as it Relates to Process Control

- your Also utilized in the solids refer to design criteria for loading rate calculations; facility to check capacity
- Key use for your facility. conjunction with SVI etc. to loadings... these can be used in develop a key operating window removal efficiencies, is MCRT calculations, 0/0 0/0



- Requires 3 pieces of equipment:
- Settleometer
- Paddle
- Timer
- activated sludge facilities Should be performed daily at
- Information that can be noted:
- Settled volume at 5 minutes
- Settled volume at 30 minutes
- Time "pops" μ. Η and when the blanket

- basin into the **Ensure the sample is** clarifier liquor flowing representative of the mixed secondary from aeration
- Always sample after basins any significant differences) sample from each basin and note combine (but can occasionally

- Test conditions and sampling influences on the test results! technique can have strong
- Always collect in a wide-mouth container
- minutes of collection Testing should begin within 10
- Sample should not be shaken or agitated during transport to testing site

Settleability - The Procedure

- Mix the MISS thoroughly sample carefully but
- aeration into the settleometer of additional turbulence or Pour rapidly with the least amount
- paddle Stir the settleometer gently with
- Stop all movement of the mixed 5 minute intervals remove and immediately set timer at liquor with the paddle and slowly

Settleability — The Observations

- sludge During the the visual characteristics first 5 minutes of the watch
- How do they agglomerate?
- What do the particles look like?
- What does the "clear" like? water look

by the sludge? mark - what is Then take a reading at the 5 minute the volume occupied

Settleability – The Readings

- minutes for the first 30 minutes Readings should be taken every 5
- minutes of the 60 minute test minutes for the remaining 30 Readings should be taken every 10
- Make note of when / if the sludge After the test is complete, allow sample to sit for surface. finally swells and floats to the several hours.

Settleability as it Relates to Process Control

- secondary clarifier give the operator an idea of how The main purpose of the test is to the sludge is settling in the
- measure depth of sludge in clarifier and sludge detention time quickly (less than 90 minutes), If the sludge rises or "pops"
- The lower the overflow rate, the more opportunity for denitrification to occur higher the detention time the

Settleability as it Relates to Process Control

milliliters occupied by 1 gram of 30 minutes activated sludge after settling for Calculation Index (SVI) - volume of sludge in of the Sludge Volume

over time. activated sludge as the conc. of SVI will help the operator evaluate the solids in the system change the settling characteristics of the

$$SVI = \frac{SSV_{30} \times 1000 \text{ mg/g}}{MLSS \times \text{mg/l}}$$

- every plant is different the SVIs between 75 and 150 ml/g settling sludge; however, since to plant. optimum or best SVI varies plant typically indicate a good
- Set your benchmark when plant is running well!

because of rapid settling. leaves pin-floc in the supernatant over-oxidized sludge. Sometimes settles rapidly. 80 ml/g or less usually dense and Denotes old and

Sometimes accompanied by higher than allowable TSS in the effluent.

pin floc. blanket that does not leave behind produce range. 100 to 200 ml/g - Most clear effluent in this SVI Slightly slower settling facilities

with small amount of filaments Typically irregular shaped floc colonize on. forming a backbone for bacteria to

dense. 250 ml/g or higher - very slow to settle with poor compaction. MLSS looks light and fluffy and not

may be necessary to determine bulking. Microscopic examination growth. actual cause. Also check pH (less also indicate filamentous sludge Sign of a young sludge or could than 6.5 can encourage filamentous

Control Parameters So Important? Why Are All of These Process

Conclusions

- NPDES limits. output and routinely meet our help us optimize our facility Process control parameters can
- You have to know what to expect something has changed! to be able to recognize when
- perform the tests. Biology and chemistry within your optimized if you take the time to systems can be controlled and

Questions?

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