



WATER TREATMENT SKAA 2912

WATER QUALITY PARAMETERS (PHYSICAL)

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Solids

- Sources (mixture of inorganic and organic compounds)
 - Soil erosion
 - Urban runoff
 - Decaying plants and animals

OPENCOURSEWARE

- Wastewater and septic effluent
- Effects
 - Aesthetic and health aspects
 - Adsorption point/centre for chemicals and microorganisms
 - Stain textiles
 - May contain elevated levels of ions such as nitrate, arsenic, lead, copper, calcium, magnesium etc
 - Corrosive to plumbing fixtures/Scale formation due to dissolved ions



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- Turbidity the amount of cloudiness in water.
- Measure light-transmitting properties of water with respect to particles/colloidal matter



- Unit: nephelometric turbidity unit (NTU)
- Measurement apparatus: Secchi disk, turbidity tube, turbidity meter
- For drinking water, turbidity < 5 NTU.
- For chlorination to be effective, it is best to achieve < 1 NTU







- Color is closely related to turbidity.
- Colloidal matter in the water gives color to surface waters
- Generally, pH ↑, color intensity ↑
- Source:
 - Decomposed organic debris in the form of humic acids, tannins, lignins
 - Dye wastes
 - Wastewater (domestic and industrial)
 - Minerals
- Effects:
 - Aesthetics
 - Some are toxic for human consumption
 - Stain textiles
 - High consumption of energy in industrial boilers, equipment and tools due to the insulation caused by minerals – reduce efficiency and life of an equipment





Color

- Two types:
 - Apparent color: Color due to both dissolved and suspended matters.
 - True color: Color of a water sample that consists only dissolved matter (suspended material is already removed)
- Measurements:
 - Visual comparison method (dilution)
 - Spectrometric method
 (light transmission)
 Most commonly used





Principles of Spectrometric Method

• Beers-Lambert Law

$$A = \log \frac{I_0}{I} = \varepsilon c \ell$$



where

A = absorbance, unit/cm

- I_0 = intensity of light entering the solution
- I = intensity of light leaving the solution
- ϵ = molar absorptivity, mol⁻¹ L cm⁻¹
- $c = concentration, mol L^{-1}$
- ℓ = length of absorbing layer, cm





Taste & Odor

- Taste reflects inorganic constituents and odor reflects organic constituents
- Temperature and pH can affect taste
- Sources:
 - Specific metals (Fe, Cu, Mn, Zn) Taste
 - MgSO₄ and CaCO₃ Taste Inorganic compounds
 - Chlorine (when reacts with organic compounds especially phenols), monochloramine – Taste & Odor
 - Microorganisms (Geosmin, 2-methylisoborneol (MIB), hydrogen sulfide normally happen in distribution system) – Taste & Odor
 - Organic compounds from petroleum and/or degradation of organic matter
- Effects:
 - Aesthetics
 - Some are toxic at elevated concentrations (e.g. H₂S, reaction from sources and other chemicals such as chlorine)
 - Produce psychological stress (poor appetite for food, lower water consumption, nausea and vomitting)
 - Impaired respiration







- The most common physical assessment of a water quality
- Sources:
 - Warmer water discharge into the environment → thermal pollution (normally by industrial boilers)
 - Absorption of heat from sunlight by sediments in water (Erosion)
- Effects:
 - Affect chemical reactions and properties (e.g. gas solubility, density and viscosity)
 - Disturb biological (microorganisms and aquatic life) activities (temperature, biological activity)







• The relationship between rate constant and temperature is represented by Arrhenius equation



 Rate constant, k, will change with temperature but activation energy does not. The increase of 10°C will double the k.





Conductivity

- Measurement of the ability of a solution to conduct an electrical current
- Dependent on the concentration and mobility of ions (1 ion concentration, 1 conductivity)
- Can be used as a surrogate measure of total dissolved solids (TDS) concentration.

TDS \cong EC (µmho/cm)×(0.55-0.90)

 Unit: millisiemens per meter (mS/m), micromhos per centimeter (µmho/cm)





Conductivity

- Electrical conductivity can be estimated if the concentration and mobility of the ions in a solution are known
- For this estimation, we need
 - concentration of the ions
 - equivalence conductance of the ions
 - ionic strength of a solution
 - activity coefficient of the ions in a solution





Ionic Strength

- Quantification of the effect of all ions in a solution
- Chemical interactions between the ions depend strongly on the ionic charge
- **†** ionic charge, **†** ionic strength

$$\mathbf{I} = \frac{1}{2} \sum_{\text{all ions}} \mathbf{c}_i \mathbf{z}_i^2$$

where

 c_i = the concentration of i in mol/L z_i = the charge on species i (diamensionless)





Activity Coefficient

• A factor that considers the deviation of a chemical reaction from its ideal behavior.

Actual reactivity of a substance

 $\gamma = \frac{\gamma}{\text{Reactivity of a substance in the standard state}}$

- fionic charge, f attraction and shielding effect,
 the activity coefficient
- To predict the activity coefficients, three equations used are:
 - Debye-Huckel limiting law
 - Extended Debye-Huckel
 - Davies





Activity Coefficient

• For water at 25°C,

Debye-Huckel limiting law (I<0.005 M)	$\log \gamma_{D-H} = -0.51 z^2 I^{\frac{1}{2}}$
Extended Debye-Huckel (I<0.1 M) Appropriate in solution where only one salt dominates the ionic strength	$log \gamma_{Ext.D-H} = -0.51z^2 \frac{I^{1/2}}{1+0.33aI^{1/2}}$ a = ion size, Ångstroms
Davies (I<0.5 M)	$\log \gamma_{\text{Davies}} = -0.51 z^2 \left(\frac{I^{1/2}}{1 + I^{1/2}} - 0.3I \right)$





Ionic Strength and Activity Coefficient

• Example:

Part 1

A solution contains 10^{-3} M NaCl, 10^{-3} of NaHCO₃ and 10^{-4} of CaCl₂. Calculate the

- a. ionic strength
- b. activity coefficient (Debye-Huckle)

of the solution.

Part 2

If NaCl concentration becomes 0.1 M, what changes do you observe in the ionic strength and activity coefficient?







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