# **Chapter 3**

# **Drinking Water Quality & Treatment**

#### **SOURCES:**

- Groundwater: Springs, Artesian wells
- Surface water: Rivers, Lakes, The sea (Desalination plants)

### **Groundwater**

- constant composition
- high mineral content
- low turbidity
- low color
- low or no D.O.
- high hardness
- high Fe, Mn



## **Surface water**

- variable composition
- low mineral content
- high turbidity
- colored
- D.O. present
- low hardness
- taste and odor



# **Objectives of Water Treatment**

- The goal of municipal water treatment is to provide water:
- 1. With quality that meets regulatory criteria or standards to be both
  - □ <u>potable</u>
    - Safe to drink protective of human health
    - Not necessarily esthetically pleasing, and

## □ palatable

- o esthetically pleasing
- presence of chemicals does not pose a threat to human health
- includes chloride, color, corrosivity, iron, manganese, taste and odor
- 2. In sufficient quantity that is continuous without interruption, and
- 3. At reasonable cost.

# **Water Quality**

- Defined in relation to intended use
  - Drinking water,
  - irrigation,
  - power generation, etc.
- Water Quality Parameters:
  - A. Physical (Aesthetic or Acceptability),
  - B. Chemical,
  - C. Radioactive,
  - D. Microbiological.

Contaminant	Maximum Contaminant Level (mg/l)
Inorganics	Y 14 Y 24 Y 25 Y 25 Y 25 Y 25 Y 25 Y 25 Y 2
Arsenic	0.05
Cadmium	0.01
Lead	0.015 action level <sup>a</sup>
Mercury	0.002
Selenium	0.01
Organic chemicals Pesticides	
Endrin	0.0002
Lindane	0.004
Methoxychlor	0.1
Herbicides	
2,4-D	0.1
2,4,S-TP	0.01
Silvex	0.01
Volatile organic chemicals	
Benzene	0.005
<b>Carbon tetrachloride</b>	0.005
Trichloroethylene	0.005
Vinyl chloride	0.002
Microbiological organisms	
Fecal coliform bacteria	1 cell/100 ml

**National Drinking Water Standards** 

**Table 22.2** 

<sup>&</sup>lt;sup>a</sup> Action level is related to the treatment of water to reduce lead to a safe level. There is no maximum contaminant level for lead. *Source:* U.S. Environmental Protection Agency.

# A. Physical Parameters

- 1. Suspended solids (big particles)
- 2. Turbidity (tiny particles)
- 3. Odor and Taste
- 4. Color
- 5. Temperature.



# 1. Suspended solids (big particles)

- Organic and inorganic particles in water are generally termed total suspended solids (TSS).
- Imhoff cones are used to measure for settleable solids, which are larger than TSS.
- TSS are measured by <u>filtering</u> a water sample, drying and weighing the filter paper.



# 2. Turbidity

# **Turbidity (NTU)**

Water Samples: 250 100 50 25 10

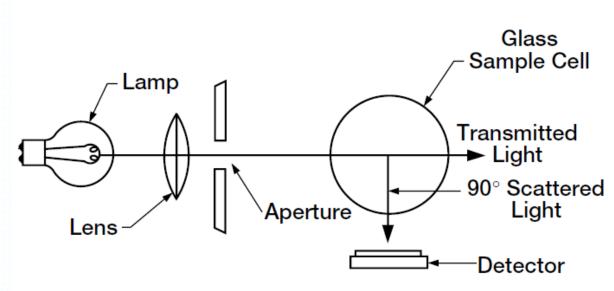


- 1. Used to assess the clarity of water
- 2. Due to *colloids*: particles that do not settle readily
- 3. When colloidal matter accumulates, light is scattered and the water appeared <u>turbid</u>.
- 4. Substances that cause turbidity include:
  - clay, silt,
  - tiny fragments of organic matter, and
  - microscopic organisms (e.g. bacteria, algae)

## **Turbidity measurement:**

- measured in units that relate the clarity of the water sample to that of standardized suspension of silica (SiO2).
- □ The interference in the passage of light caused by a suspension of 1 mg/l of silica is equivalent to one turbidity units (TU).
- ☐ Measurement: Nephelometers (NTU).









# <u>3. Odor</u>

- 1. A physical characteristics of drinking water that is important for *aesthetic reasons*.
- 2. It may be caused by dissolved or suspended colloidal particles of *organic nature*.
- 3. It is measured by human panel or **electronic nose (olfactory device)** expressed in terms of a <u>threshold odor number</u> (TON).

<u>TON:</u> It is the ratio by which the sample has to be diluted for the odor to become virtually unnoticeable.

Example: 50 ml is diluted to vol. of 200 ml.

The dilution # equals 200/50 = 4 = TON

# 3. Taste

- Taste is an aesthetic water quality parameter.
- 2. Caused by presence of organic and / or inorganic substances:
  - Organics cause both odor & Taste problems.
  - Inorganics (e.g. soil components) are usually odorless and may be responsible for taste.

## Odor & Taste:

most commonly caused by:

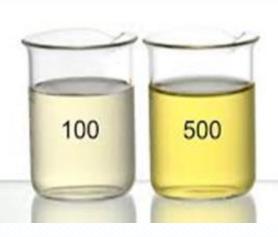
algae, decomposed organic matter, and dissolved gases.

## 4. Color

- is a physical characteristics of drinking water that are important for esthetic reasons.
- It may be caused by dissolved or suspended colloidal particles.
- One color unit is equivalent to the color produced by a 1 mg/L solution of platinum (Pt).







## 5. Temperature





- Warm water tastes flat. Cooling suppresses odors and tastes and makes water more palatable.
- Temperature effects the chlorination and purification of water.
- Disinfection takes longer when water is colder
- At lower T, purification capacity is reduced with reverse osmosis treatment equipment.
- Water having physical characteristics exceeding the limits or making it less palatable should not, as a general rule, be used for drinking.

# **B. Chemical Parameters**

# Total dissolved solids:

- Alkalinity
- Hardness
- Fluoride
- Metal ions
- Organics
- Nutrients

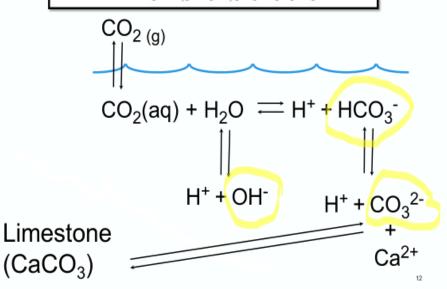
#### **ANALYSIS:**

- Gross: alkalinity, hardness.
- Specific ions: Ca<sup>2+</sup>, Mg <sup>2+</sup>, Na<sup>+</sup>, K<sup>+</sup>, Cl<sup>-</sup>, NO3<sup>-</sup>, pH (H<sup>+</sup> or OH<sup>-</sup>)



#### **Alkalinity**

The amount of alkalinity present is expressed in terms of CaCO<sub>3</sub>. Alk  $T = [HCO_3] + 2[CO_3] + [OH] - [H']$ 



## **Hardness**

- Multivalent cations, particularly magnesium and calcium, causes the hardness.
- These ions are easily precipitated and form scales especially in heat transfer equipment and a problem arise up.
- Correlated with TDS (as part of it), they represent total concentration of Ca and Mg, and is reported in equivalent CaCO3 (mg/L).
- Other ions (Fe2+, Al3+) may also contribute to hardness.

## Hard water problems:

- leaves solid deposits in boilers, hot water pipes, heaters, ...)
- They also react with soap and detergents requiring more soap or detergent and form a difficult—to-remove scum

Advantage: hard water is less corrosive than soft water.

# $Hardness, eq/m^3 = (Ca^{+2}) + (Mg^{+2})$

# **Chemical Parameters**

## Hardness in Water

Degree of Hardness	p.p.m. as CaCO <sub>3</sub>
Soft	0-75
Moderately Hard	75 - 150
Hard	150 - 300
Very hard	>300

# C. Radioactive Parameters (Primary MCLs)

Radionuclides

Radium 226	20 pCi/L	Beta particle and photon radioactivity	4 mrem/yr
Radium 228	20 pCi/L	Radon	300 pCi/L
Gross alpha particle activity	15 pCi/L	Uranium	$20~\mu \mathrm{g/L}$

1 Ci = decay rate of 1 g Radium-226 (= decay of 3.7x10 10 atoms/s)

# **D. Microbiological Parameters**

## **Microorganisms:**

- 1. Bacteria
- 2. Protozoa
- 3. Parasitic Worms
- 4. Viruses
- 5. Algae
- 6. Fungi
- 7. Pollution indicators: Coliform bacteria





- ☐ E. coli used as an indicator of water quality: normal inhabitant of intestines of humans and many animals.
  - √ Fecal Indicator Bacteria (FIB)
  - ✓ Indicator of presence of fecal matter
  - √ Total coliforms are typically reported

# **Microbiological Parameters**

- A fecal coliform bacteria test is used to indicate the likely presence of disease-causing bacteria in water.
- Results reported as Most Probable Number (MPN) per 100 mL.
- Incubation at moderate temperature (35°C) for 48 hr.

	MCL GOAL	MCL*
Total Coliforms (Including fecal coliforms & E. coli)	Zero	5%**
Viruses (Enteric)	Zero	99.99% killed or inactivated
Giardia lamblia	Zero	99.99% killed or inactivated

<sup>\*</sup>MCL - Maximum Concentration Level

Table 1. EPA National Drinking Water Regulations for Microorganisms.

<sup>&</sup>quot;No more than 5% of the water samples total coliform positive in a month. Every sample that has total coliforms must be analyzed for fecal coliforms. The presence of any fecal coliforms is unacceptable in drinking water.

# **Drinking Water Treatment**



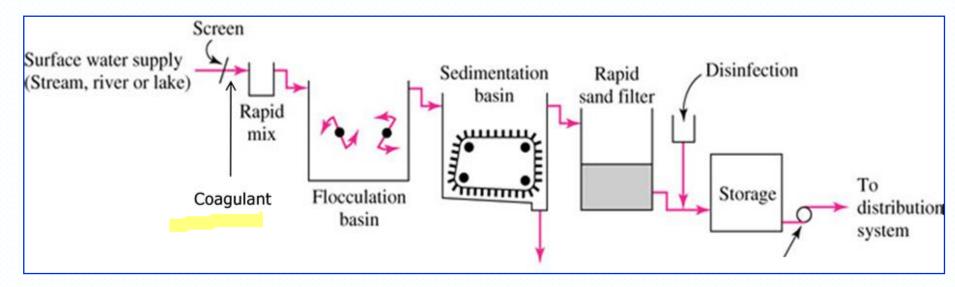
### **I. Surface Water Treatment**

- Primary objectives are to
  - 1) Remove suspended material (turbidity) and color
  - 2) Eliminate pathogenic organisms
- Treatment technologies largely based on <u>coagulation and</u> <u>flocculation</u>

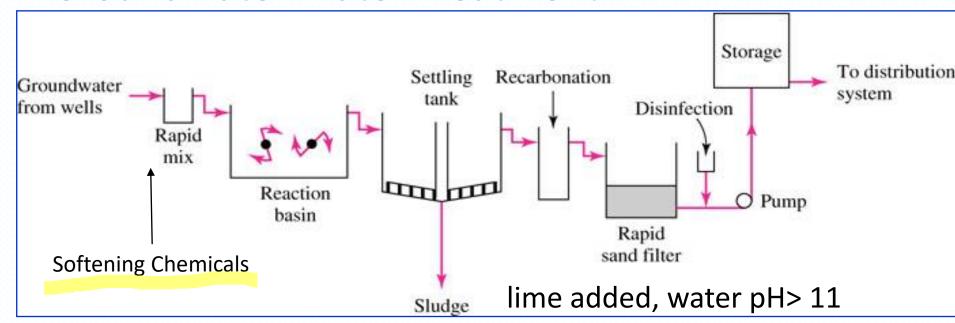
## **II. Groundwater Treatment**

- Primary objectives are to
  - 1) Remove hardness and other minerals
  - 2) Eliminate pathogenic organisms
- Treatment technologies largely based on <a href="checkbox">chemical precipitation</a>

## **Surface Water Treatment**



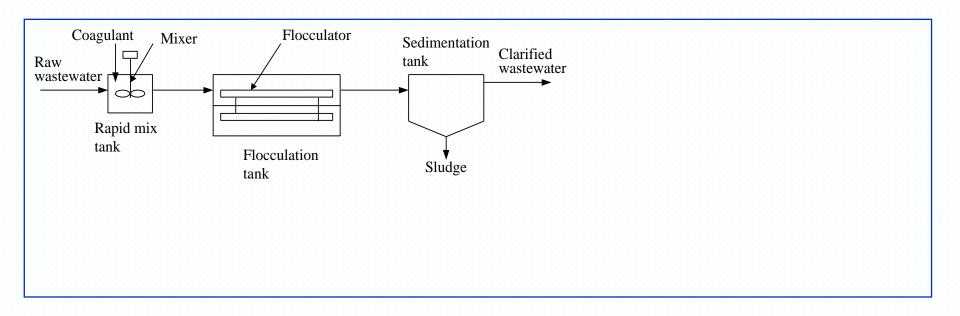
## **Groundwater Water Treatment**



## **Surface Water Treatment**

## Removal of turbidity (and suspended solids formed):

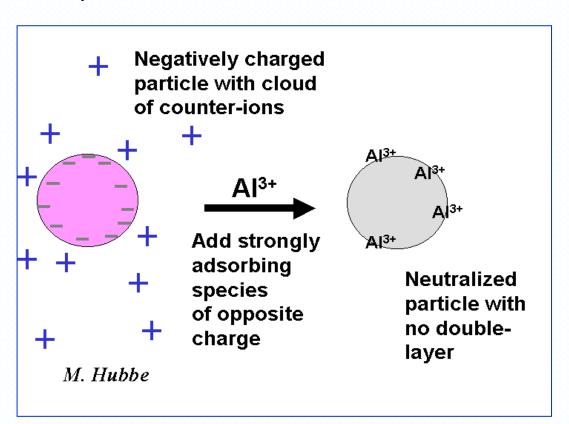
- 1. rapid mix (coagulation) tank
- 2. flocculation tanks
- 3. settling (sedimentation) tanks
- 4. Filtration

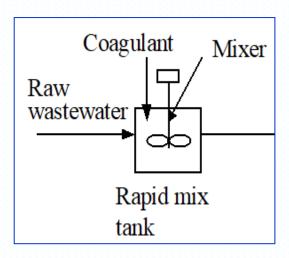


# Coagulation

## Addition and rapid mixing of a coagulant with the water to:

- 1) neutralize surface charges
- 2) collapse the (charged) surface layer around the particles
- 3) allow the particles to come together and agglomerate
- 4) allow the formation of floc that can readily settle





# **Good Coagulants**

- 1) Non-toxic and relatively inexpensive
- 2) Insoluble in neutral pH range
- 3) Do not leave high concentrations of metals in treated water
- 4) Trivalent cations are most effective in charge neutralization.

## **Common Coagulants**

#### Alum

- Hydrated aluminum sulfate [Al<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>·14H<sub>2</sub>O]
- Alum, when added to water, will be <a href="hydrolyzed">hydrolyzed</a> to form gelatinous hydroxide [Al(OH)3] precipitate.
- This will carry suspended solids as it settles by gravity.
- Optimum pH: 5.5 6.5.
- Anhydrous Fe<sup>3+</sup> (as FeCl<sub>3</sub>)
  - Forms Fe(OH)<sub>3</sub>(s) in a wide range of pH 4-11, optimum 4.5 5.5.

# **Aluminum Chemistry**

With alum addition, what happens to water pH?

$$Al_2(SO4)_3.14 H_2O \Leftrightarrow 2Al(OH)_3 \downarrow + 8H_2O + 3H_2SO_4$$

• 1 mole of alum consumes 6 moles of bicarbonate (HCO<sub>3</sub>-)

$$Al_2(SO4)_3.14 H_2O + 6HCO_3^- \Leftrightarrow 2Al(OH)_3 \downarrow + 6CO_2 + 14H_2O + 3SO_4^{-2}$$

• If alkalinity is not enough, pH will be reduced greatly

$$Al_2(SO_4)_3.14H_2O \leftrightarrow 2Al(OH)_3 \downarrow + 3H_2SO_4 + 14H_2O$$

• Lime or sodium carbonate may be needed to neutralize the acid.

## **Iron Chemistry**

$$FeCl_3 + 3HCO_3 \Leftrightarrow Fe(OH)_3 \downarrow + 3CO_2 + 3Cl^2$$

- With iron salt addition, what happens to water pH:
   1 mole of FeCl<sub>3</sub> consumes 3 moles of bicarbonate (HCO<sub>3</sub><sup>-</sup>)
- If alkalinity is not enough, <u>pH will reduce greatly due to hydrochloric acid HCl formation.</u>
- Lime or sodium carbonate may be needed to neutralize the acid.
   Lime is the cheapest.

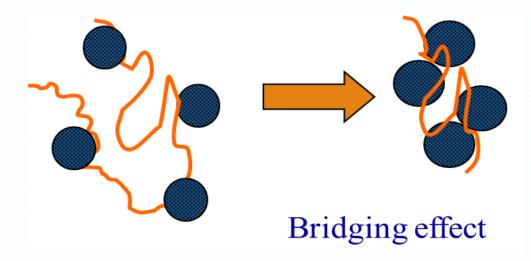
## **Flocculation**

- The process that leads to the formation of large voluminous flocs, which are loosely held aggregates of coagulated particles and solids.
- The slow mixing is one of the fundamentals necessary to promote collisions between particles to form flocs.

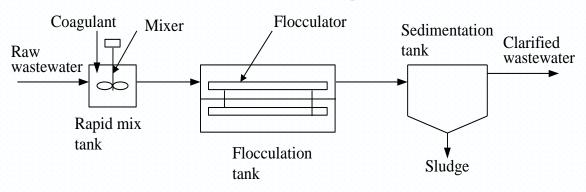
# Flocculation Chemical (Coagulant-Aids):

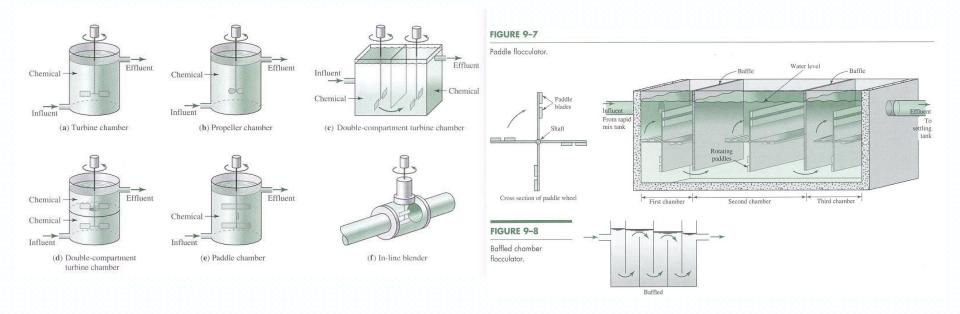
- 1. inorganic: polyaluminum chloride
- 2. Synthetic Polymers (Polyelectrolytes): polyacrylic acid,

polyacrylamide derivatives



# **Mixing & Flocculation Units**





# Jar Test Determining Coagulant Dose & Optimum pH

- ☐ The jar test a laboratory procedure to determine the optimum pH and the optimum coagulant dose
- ☐ A jar test simulates the coagulation and flocculation processes



Jar Test set-up

## Jar Test

## determining optimum pH

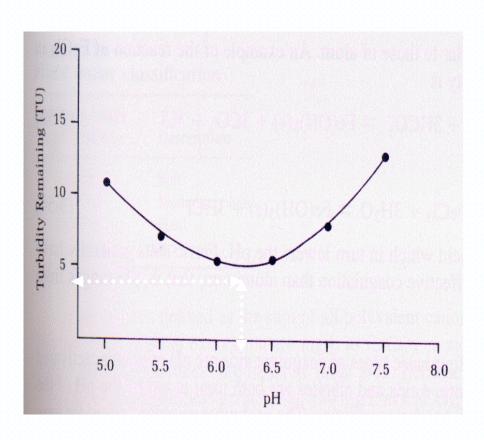
## ☐ Fill the jars with raw water sample – usually 6 jars ☐ Adjust pH while mixing using $H_2SO_4$ or NaOH (pH: 5.0; 5.5; 6.0; 6.5; 7.0; 7.5) ☐ Add same dose of selected coagulant to each jar (Coagulant dose: 5 or 10 mg/L) Rapid mix each jar at 100 to 150 rpm for 1 minute. ☐ Reduce the stirring speed to 25 to 30 rpm and continue mixing for 15 to 20 min. ☐ Turn off mixers and allow flocs to settle for 30 to 45 mins Measure the final residual turbidity in each jar Plot residual turbidity against pH.

## **Optimum coagulant dose**

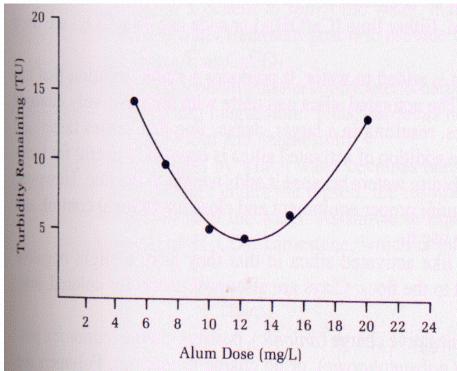
☐ Repeat all the previous steps
☐ This time adjust pH of all jars at
optimum found from first test while mixing
using H <sub>2</sub> SO <sub>4</sub> or NaOH
☐ Add different doses of the selected
coagulant (alum or iron) to each jar
(Coagulant dose: 5; 7; 10; 12; 15; 20 mg/L)
$oldsymbol{\square}$ Rapid mix each jar at 100 to 150 rpm for
1 minute.
☐ Reduce the stirring speed to 25 to 30
rpm for 15 to 20 mins
☐ Turn off the mixers and allow flocs to
settle for 30 to 45 mins
☐ Measure the final residual turbidity in
each jar
☐ Plot residual turbidity against coagulant
dose.

# **Coagulation Jar Tests**

# Jar Test – optimum pH



# Jar Test – optimum dose



# Coagulation / Flocculation Calculations

# Measuring the degree of mixing

Velocity gradient:

$$G = \sqrt{P/(\mu V)}$$

P = power input (W)

V = volume of basin(cu. m)

 $\mu$  = fluid viscosity (Pa.s)

#### G values for rapid mixing

## Gto values for flocculation

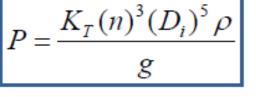
Detention	G (s <sup>-1</sup> )	Туре	G (s <sup>-1</sup> )	Gt <sub>0</sub>
time, t <sub>0</sub> (s)		Low turbidity, color	20 – 70	60,000 – 200,000
0.5	3500	removal coagulation		
10 – 20	1000	High turbidity, solids	50 - 150	90,000 - 180,000
20 – 30	900	removal coagulation		
30 – 40	800	Softening, 10% solids	130 - 200	200,000 – 250,000
Longer	700	Softening, 39% solids	150 - 300	390,000 – 400,000

# **Coagulation / Flocculation Calculations**

## Designing a flocculator

### Power input:





 $K_T$  = impeller constant n = rotational speed (rpm)Di = impeller dia (m)



Basic impeller styles, (Source: Courtesy of SPX Process Equipment,

## Values of impeller constant K<sub>T</sub>

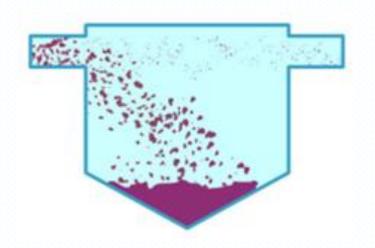
Type of impeller	K <sub>T</sub>
Propeller, pitch of 1, 3 blades	0.32
Propeller, pitch of 2, 3 blades	1.00
Turbine, 6 flat blades, vaned disc	6.30
Turbine, 6 curved blades	4.80
Fan turbine, 6 blades at 45°	1.65
Shrouded turbine, 6 curved blades	1.08
Shrouded turbine, with stator, no baffles	1.12

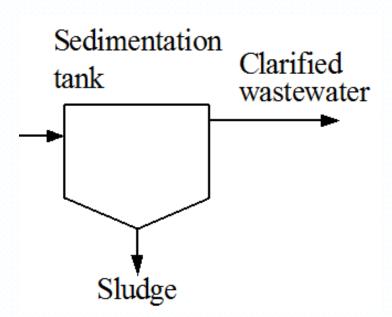
## Power and rotational speed of some standard mixers

Model	n (rpm)	P (kW)
JTQ50	30, 45	0.37
JTQ75	45, 70	0.56
JTQ100	45, 110	0.75
JTQ150	45, 110	1.12
JTQ200	70, 110	1.50
JTQ300	110, 175	2.24
JTQ500	175	3.74

JWI, Inc. of Holland

# 2. Sedimentation:





- ☐ Flocs settle out and is scraped and vacuumed off the bed of large sedimentation tanks.
- ☐ Clarified water drains out of the top of these tanks in a giant decanting process.

(For settling tank design and details, refer to Pre-requisite course notes (particulate separation).

## 3. Filtration

- □ Removal of those particles that are too small to be effectively removed during sedimentation.
- □ Sedimentation effluent: 1 10 JTU & Desired effluent level: <0.3 JTU
- □Separate non-settleable solids from water. Combined with coagulation/clarification, filtration can remove 84%-96% turbidity, over 97-99 Coliform bacteria and Giardia.

## Slow sand filters AND Rapid sand filters

Either slow or rapid filtration (depends on size of plant / volume of water considerations).

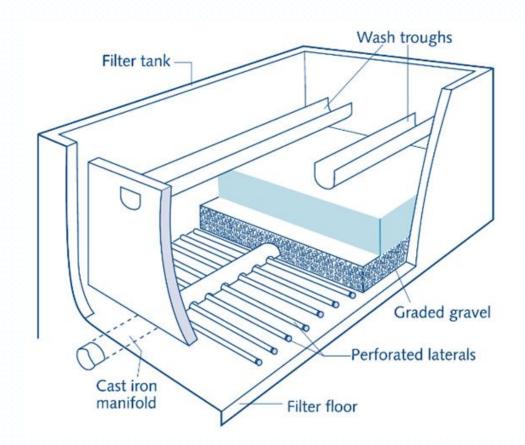
## 1) Rapid-sand filters

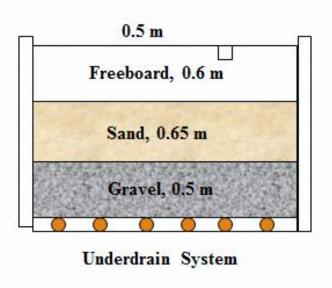
• force water through a 0.45-1 m layer of sand ( $d_p$ =0.4-1.2mm) and work faster, needing a smaller area. But they need frequent back-washing

## 3. Filtration

## 2) Slow-sand filters

 $(d_p=0.15-0.35mm)$  require a much larger area but reduce bacteriological and viral levels to a greater degree. The top 1 inch must be periodically scraped off and the filter occasionally back-washed





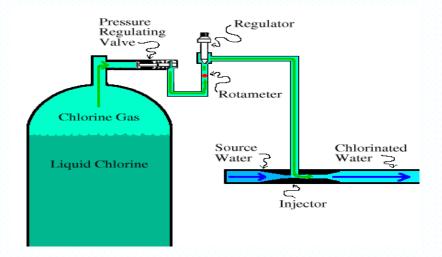
## 4. Disinfection:

- 1) Disinfection is typically the <u>last step</u> in a water (*and waste-water*) treatment system
- 2) Residual disinfectant is needed in distribution system after water (or wastewater) treatment
- 3) Water completely *free of suspended sediment* is treated with a powerful oxidizing agent usually chlorine gas.
- 4) A residual of chlorine disinfectant is left in the water to prevent reinfection. Chlorine can form *harmful byproducts* and has suspected links to stomach cancer and miscarriages. Many agencies now residually disinfect with Chloramine.
- 5) In addition to disinfection, <u>chlorine also</u> has the following functions:
  - taste and odor control as an oxidizing agent
  - b. oxidation of Fe<sup>2+</sup> and Mn<sup>2+</sup> in groundwater
  - c. ammonium removal in domestic waste treatment
  - d. slime, biofouling control

## **Disinfectants**

- 1. Gaseous Cl<sub>2</sub>
  - Most commonly used.
  - Advantage: provide residual chlorine for the protection from bacterial growth in distribution system
  - Disadvantage: The formation of disinfection by-products (trihalomethanes) presents a health risk
- 2. Hypochlorite: NaOCl or Ca(OCl)<sub>2</sub>
- 3. Ozone: generated on site
- 4. UV lamps: for small applications.



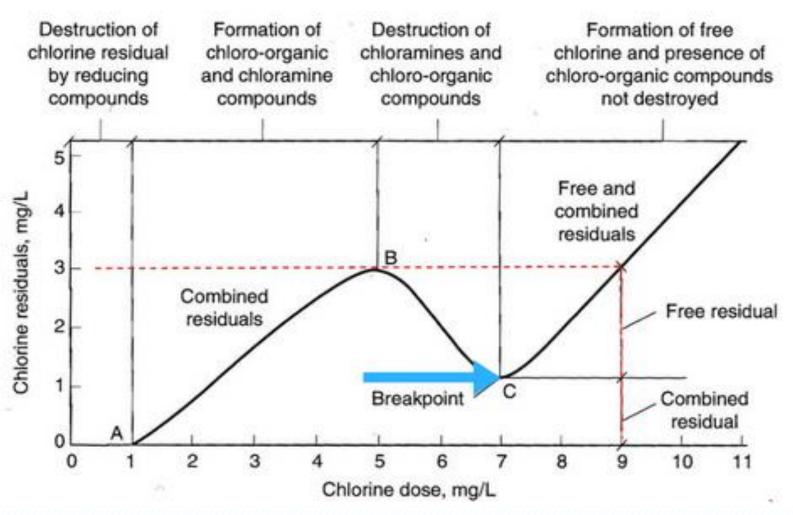


# **Chemistry of Chlorine in Water**

$$Cl_2 + H_2O \rightarrow H^+ + Cl^- + HOCl$$

- 1. HOCl is a weak acid (HOCl =  $H^+ + OCl^-$ ) with  $K_a = 4.5 \times 10^{-4}$
- 2. HOCl and OCl<sup>-</sup> are <u>free available chlorine</u> which are very effective in killing bacteria
- 3. Small amount of ammonium (NH<sub>4</sub><sup>+</sup>) in water is <u>desired</u> to form Chloramines: NH<sub>2</sub>Cl, NHCl<sub>2</sub>, NCl<sub>3</sub>
  - ☐ Chloramines (<u>combined available chlorine</u>) are weaker disinfectants than free available chlorine but are desired <u>residual chlorine</u> to be retained in water distribution system
- 4. Excessive amount of ammonium  $(NH_4^+)$  in water is <u>undesirable</u> because it consume excess demand of  $Cl_2$

# **Chlorine Demand or Breakpoint Chlorination**



- Combined chlorine is the proportion that combines with organic matter.
- Free chlorine is the amount that <u>remains</u> to kill microbes in water system.
- Total chlorine is the sum of Combined and Free chlorine.

# **Disinfection CT\* Concept**

CT = 0.9847 C \*\*0.1758 pH\*\*2.7519 temp\*\*- 0.1467

Inactivation is a function of <u>Contact time</u> <u>Concentration</u>, <u>pH</u>, and <u>Temperature</u>

## To get credit for 99.9% inactivation of Gardia microorganism:

# **Contact Time (min)**

chlorine	pH 6.5		pH 7.5	
(mg/L)	2°C	<b>10°C</b>	2°C	10°C
0.5	300	178	430	254
1.0	159	94	228	134

<sup>\*</sup>CT: Concentration (mg/L) x Contact Time (minutes)

# REFERENCES

- 1. Davis, M.L. and Cornwell, D.A. Introduction to Environmental Engineering, McGraw-Hill, 5<sup>th</sup> Edition, 2013.
- 2. Peavy, H.S.; D.R. Rowe and G. Tchobanoglous. Environmental Engineering, McGraw-Hill, 1985.
- 3. Parsons, S. and Jefferson, B. Introduction to Potable Water Treatment Processes, Wiley-Blackwell, 2006.
- 4. Spellman, F.R. and Drinan, J.E. The Drinking Water Handbook, 2<sup>nd</sup> Edition, CRC Press, 2012.



## Exercise-1

Example 6-2. Given the following analysis of a groundwater, construct a bar chart of the constituents, expressed as CaCO<sub>3</sub>.

Ion	mg/L as ion	EW CaCO <sub>3</sub> /EW ion	mg/L as CaCO <sub>3</sub>
Ca <sup>2+</sup> Mg <sup>2+</sup> Na <sup>+</sup>	103	2.50	258
Mg <sup>2+</sup>	5.5	4.12	23
Na <sup>+</sup>	16	2.18	35
$HCO_3$	255	0.82	209
$HCO_3$ $SO_4^2$	49	1.04	51
Cl-	37	1.41	52

Solution. The concentrations of the ions have been converted to CaCO<sub>3</sub> equivalents. The results are plotted in Figure 6-11.

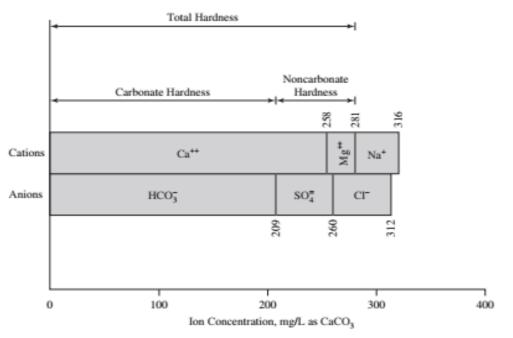


FIGURE 6-11 Bar graph of groundwater constituents.

#### Given a water having the following composition:

## Exercise-2

$$Ca^{++} - 40 \text{ mg/L} = 2 \text{ meq/L}$$

$$HCO_3 - 61 \text{ mg/L} = 1 \text{ meq/L}$$

$$Mg^{++} - 24 \text{ mg/L} = 2 \text{ meq/L}$$

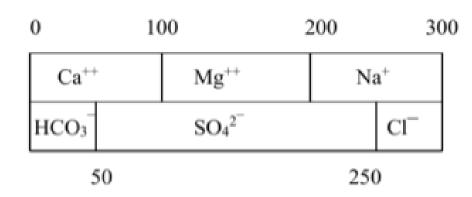
$$SO_4^{2^-}$$
 - 192 mg/L = 4 meq/L

$$Na^{+} - 46 \text{ mg/l} = 2 \text{ meg/L}$$

$$Cl^{-}$$
 - 35.5 mg/L = 1 meq/L

#### These totals must balance

a) Put these in the form of a bar diagram expressing all concentrations in terms of CaCO<sub>3</sub> (50 mg/L CaCO<sub>3</sub> = 1 meq/L)



#### From ion to CaCO3:

Ca: 50/20 (or 100/40)

Mg: 50/12 (or 100/24)

Na: 50/23

HCO3: 50/61

SO4: 50/48 (or 100/96)

CI: 50/35.5

#### From diagram you can see that

Mg hardness = 
$$100 \text{ mg/L}$$
 as  $CaCO_3$ 

$$(NCH) = TH - CH = 200-50 = 150 \text{ mg/L}$$
as CaCO3

Total alkalinity = bicarbonate alkalinity = 50 mg/L as CaCO<sub>3</sub> Why?

## Exercise-2

A surface water treatment plant is to process 30000 m $^3$ /d for a large city. A cubic rapid mixing tank will be used to blend 35 mg/L of alum solution with the flow during a detention time of 2 minutes. The solution temperature is 22°C with a viscosity of 9.6 x  $10^{-4}$  kg/m/s.

## Determine the following:

- a) Quantity of alum coagulant added (kg/day).
- b) Tank dimensions.
- c) Power input (kW) necessary for a velocity gradient (G) of 900 s<sup>-1</sup>.

G =
$$V(P/\mu V)$$
  
V= (Q).(t)