# fundamental Water treatment – Basic Track Cooling water system Prepared by: Eng. Islam Batta







#### Content:

Types of cooling towers

Heat exchanger

Cooling water treatment

Cooling tower controller



# **Cooling Towers**



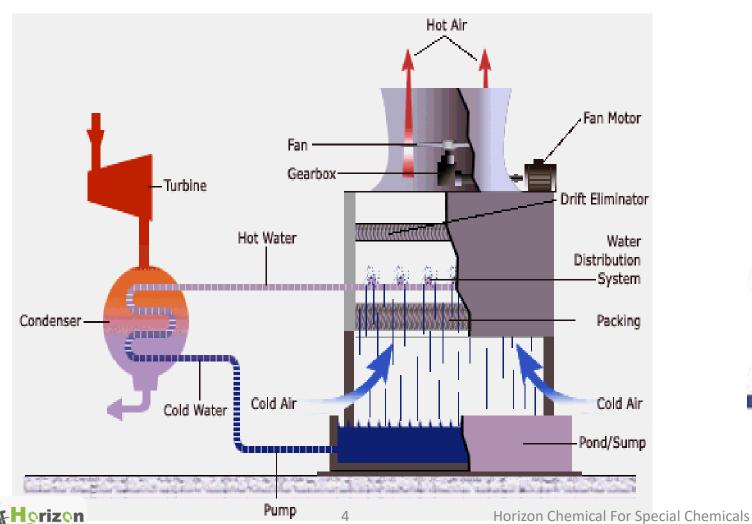




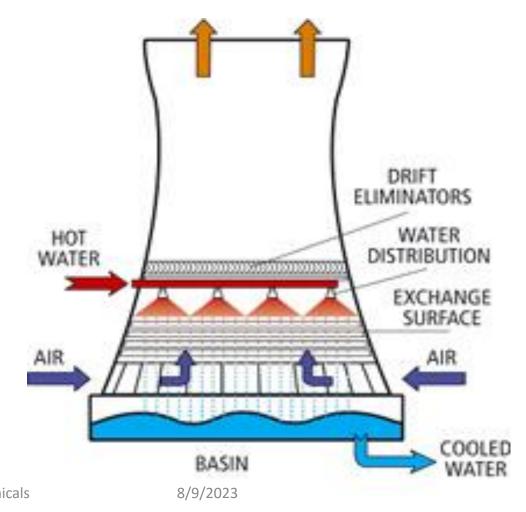


### **Types Of Cooling Towers**

#### **Induced Draft**





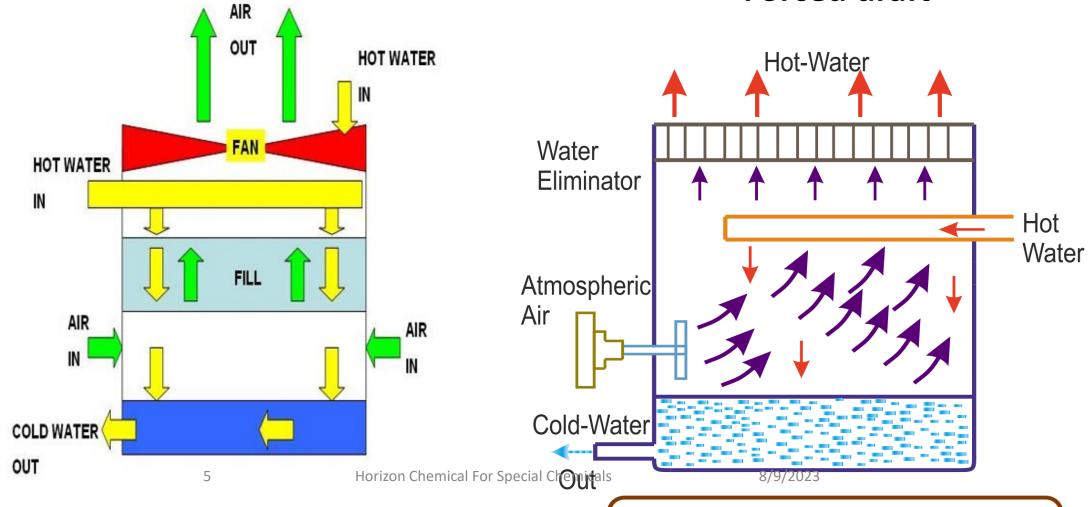


#### Non-natural draft

#### Induced draft •

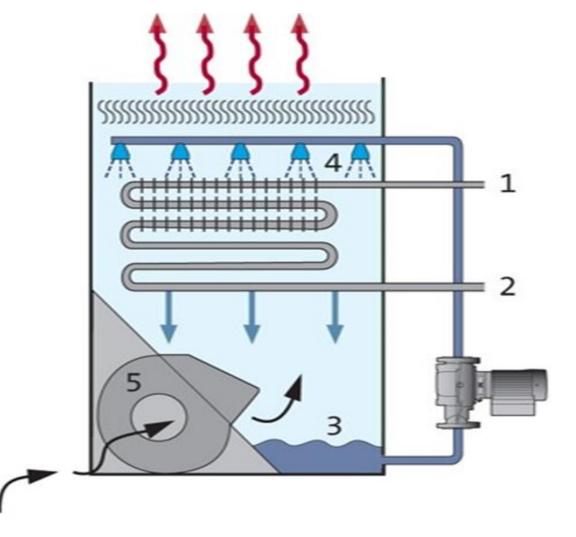








### **Evaporation Condenser**

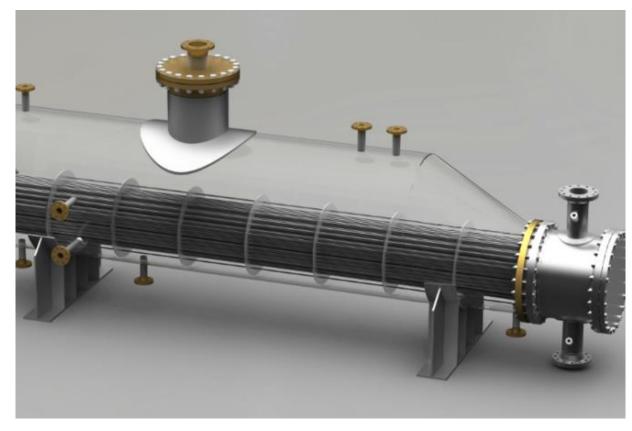






# Heat Exchanger

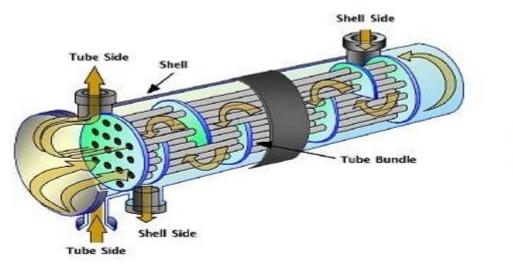






### Heat exchanger







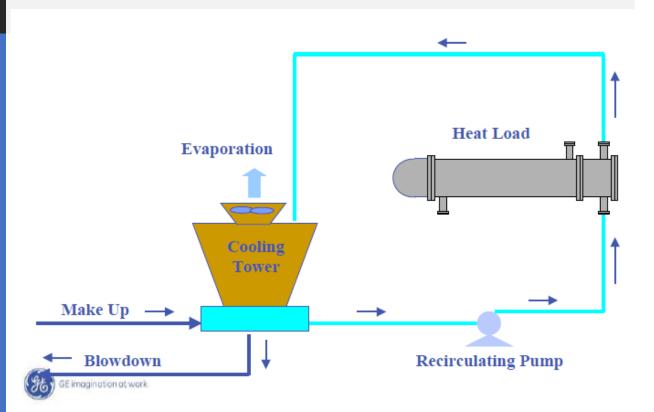
Shell & Tube

Plate



# Horizon Chemical For Special Chemicals

## **Cooling Water Treatment**





1. Scale

2. Corrosion

- 3. Fouling
- 4. Micro growth





# 1. Scale deposition

Due to the increase in the cycle Of concentration, the concentration of the dissolved ions in cooling water increases, and the solubility of one or more of the constituents is exceeded the solubility product. the compound precipitates and forms a deposit

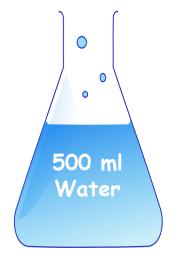
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### Cycles of Concentration

If 5 grams salt are dissolved in each flask, then TDS is...



5 g Salt TDS = 5,000 ppm



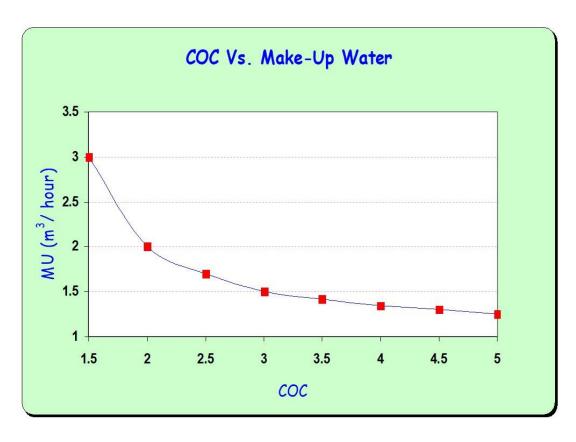
5 g Salt TDS = 10,000 ppm



5 g Salt TDS = 20,000 ppm

# Scale treatment (Adjust water chemistry –first defense line) why shall we operate at high COC?

Makeup water volumes based on an Evaporation rate of 50,000 gal/day. (Bleed rate assumes no Windage loss)				
COC	Bleed (gal/day)	Makeup (gal/day)		
2x	50,000	100,000		
3x	25,000	75,000		
4x	16,666	66,666		
5x	12,500	62,500		
6x	10,000	60,000		
7x	8,333	58,333		
8x	7,143	57,143		
9x	6,250	56,250		
10×	5,555	55,555		





# Scale treatment (Adjust water chemistry –first defense line) LANGELIER SATURATION INDEX

"Conditions at which a given water is in equilibrium with calcium carbonate".

$$pHs = (pK_2 - pK_s) + pCa + pAlk$$

 $pK_2 = 2^{\circ}$  dissociation constant of CaCO<sub>3</sub>

pK<sub>s</sub> = solubility product of CaCO<sub>3</sub> pCa = -log calcium concentration

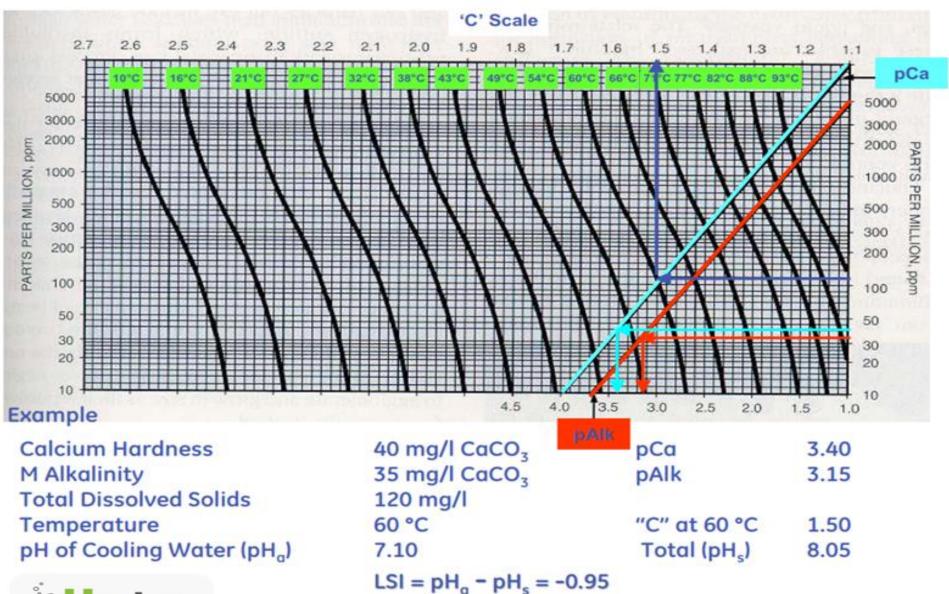
pAlk = -log alkalinity

$$LSI = pH_a - pH_s$$

LSI -ve : water tends to dissolve calcium carbonate and hence the water has corrosive tendency

LSI +ve: water has a calcium carbonate scaling tendency







## Scale Treatment (Adjust Water Chemistry –First Defense Line) Ryznar Saturation Index(RSI)



Ryznar studied the actual operating results obtained with waters of various Saturation Indices

$$RSI = 2pH_s - pH_a$$

#### Ryznar Stability Index

- > 7.5 or 8 probability of corrosion increases
- scaling may not occur > 7
- scaling tendency increases, < 6 probability of corrosion decreases



#### RYZNAR STABILITY INDEX



RSI = 2	pH <sub>s</sub> -	$pH_a$
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Ryznar Stability Index

> 7.5 or 8 probability of corrosion increases

> 7 scaling may not occur

< 6 scaling tendency increases

probability of corrosion decreases

#### **Examples:**

Calcium Hardness	200 mg/l CaCO <sub>3</sub>	pCa	2.70	
M Alkalinity	160 mg/l CaCO <sub>3</sub>	pAlk	2.50	
Total Dissolved Solids	400 mg/l	-		

Total Dissolved Solids 400 mg/l
Temperature 60 °C "C" at 60 °C 1.56

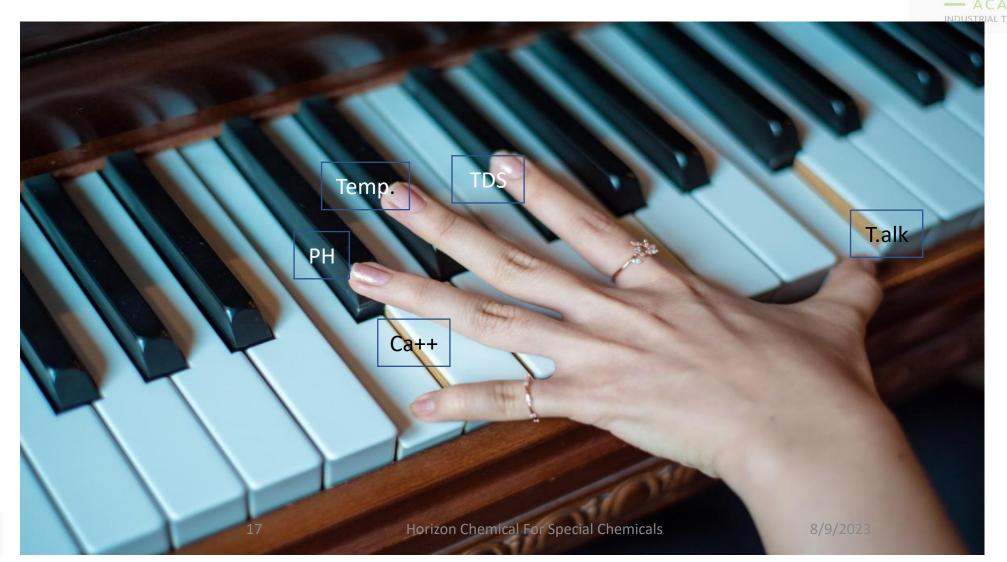
Total (pH<sub>s</sub>) 6.76

pH of Cooling Water (pH $_a$ ) 7.80 LSI = pH $_a$  - pH $_s$  = +1.04 SCALING

 $RSI = 2pH_s - pH_a = 5.72$  SCALING



# Scale treatment (Adjust water chemistry –first defense line)

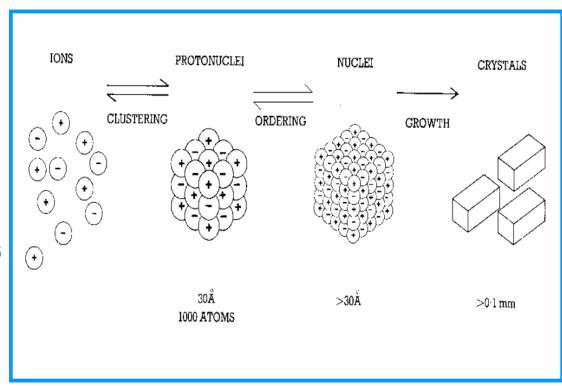




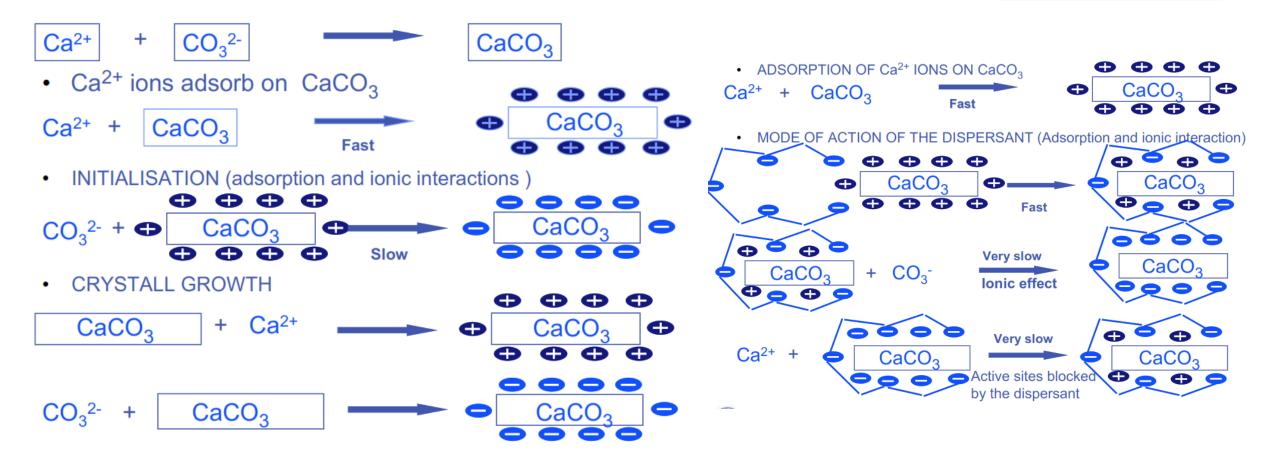


#### CaCO<sub>3</sub> SCALE INHIBITION CHEMISTRIES

**EDTA**  INHIBITION **CHELATING (COMPLEXING) PHOSPHONATES AEC ADSORPTION** CRYSTALL GROWTH INHIBITION DECREASE OF CRYSTALL GROWTH CYNETICS DISTORSION OF THE CRYSTALL < **DISPERSION OF PARTICLES TETRAPHOSPHONATES** DISPERSION **POLYACRYLATES POLYCARBOXYLATES POLYMALEATES** 









- ACADEMY -

#### **Polyphosphates**

Sensitive to hydrolysis (into oPO4 ) depending on : temperature (50 - 60°C) residence time pH in the system

#### **Phosphonates**

Chelating effect

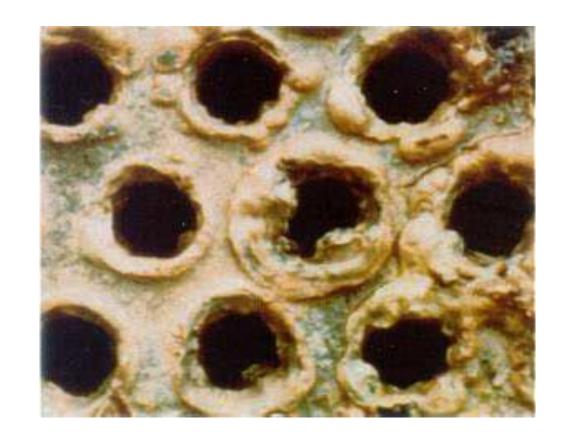
Blocks the generation of CaCO3 crystals

Reduce speed of growth

Depending on the type of phosphonate

Sensitivity to chlorination (oxidation)

Little to no risk of Ca phosphonate precipitation





Polyacrylates \_\_\_\_\_

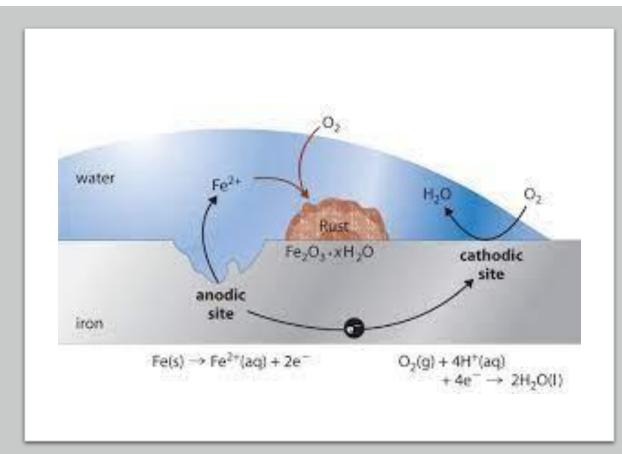




**Polymethacrylates** 

#### 2. Corrosion





$$Fe \to Fe^{2+} + 2e^{-}$$
 (Anodic)  
 $2H_2O + O_2 + 4e^{-} \to 4OH^{-}$  (Cathodic)  
 $Fe^{2+} + 2OH^{-} \to Fe(OH)_2$  (Chemical)

### Corrosion in cooling towers



- There are many reasons for corrosion, and here are some of them
- Scales (under deposit corrosion)
- Erosion
- Micro growth
- Low Isi
- Absence of anodic and /or cathodic inhibitors
- Higher conc. Of aggressive anions (So4, cl-)





# (chemical treatment –second defense line) Classification of corrosion inhibitors

Table 3.9 Classification of corrosion inhibitors

Type of corrosion inhibitor		Example of corrosion inhibitor	Characteristics of protective film
	Oxide film type (passivation film type)	Chromates, Nitrites, Molybdates	Fine and thin film (30–200Å), closely adherence to metal, excellent corrosion protection
Precipitation film type	Chemicals forming the insoluble salts with calcium ions, etc., in water	Polyphosphates, Orthophosphates, Phosphonates, Zinc salts	Relatively porous and relatively thick film, slightly poor adherence to metal, fairly good corrosion protection
	Chemicals forming water insoluble salts with the protected metal ion	Mercaptobenzothiazole, Benzotriazole, Tolyltriazole	Relatively close and relatively thin film, fairly good corrosion protection
	Adsorption film type	Amines, Surfactants	Formation of perfect adsorption layer on clean metal surface in acid or non-aqueous solution and formation of non-uniform adsorption layer on unclean surface in fresh water

### Cathodic Inhibitors (Precipitating film)



- They can either:
- Slow down the rate of cathodic reactions.
- Selectively precipitate on cathodic areas to impede or limit diffusion of reducible species (H<sub>2</sub>, O<sub>2</sub>) to these areas.
- Applied Mechanisms:
- 1.Cathodic Poisoning: compounds of Arsenic (As) and Antimony (Sb) make the recombination and discharge of hydrogen make difficult.
- 2. Precipitation: Calcium, Zinc, and Magnesium Oxides precipitate to form protective films on the cathode.



# Anodic (Passivating) Inhibitors HORIZON HORIZON

- They cause a large anodic "shift" of the corrosion potential forcing the metal surface to the "passivation range".
- Two main types are known:
- 1) Oxidizing anions: such as Chromate, Nitrate, and Nitrite. They can passivate steel in the absence of oxygen.
- 2) Nonoxidizing ions: such as Molybdate, Phosphate, and Tungstate. They passivate steel in the presence of Oxygen.
- Most effective and very widely used.
- Less Expensive.



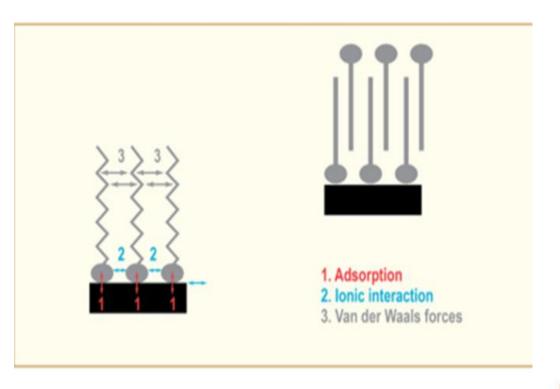
# Adsorption film (film forming)



- They effect the entire surface of corroding metal in equal concentration.
- Film-forming: They form a hydrophobic film on metal surface which provides a barrier to dissolution of metal in the electrolyte.
- Adsorbed on surfaces depending on their charge and surface charge.
- Can be Cationic (Amines) or Anionic as (Sulfonates).
- Example...
- Ethanol amine (used to stop Ethylene glycol corrosion in cooling systems).







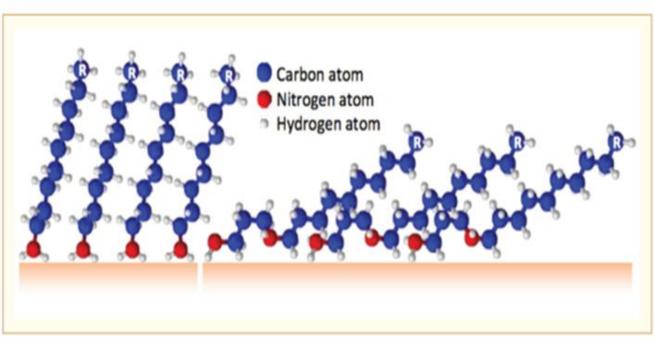


Figure 3-2: Schematic representations of FFA films: monoamine (left), polyamine (right) [Jack et al., 2015]



### General Effect - Precipitation Inhibitors

- HORIZON directly EMY —
- Film forming: they block both anodic and cathodic sites "indirectly" EMY
- They cause the formation of precipitates leading to the formation of protective film.
- Depend much on the hardness value in water (high hardness is less corrosive because of more salt tendency to precipitate).
- High pH dependence.
- Common inhibitors are Silicates and Phosphates.
- Sodium silicate is used in domestic waters to prevent "rust water".
- Phosphates require oxygen for effective inhibition.



#### Classification of Corrosion Inhibitors second defense line



Most Widely Classified as: -

1) Cathodic (Precipitating film)

2) Anodic (Passivating).

3) Adsorption film (film forming)

CaCO<sub>3</sub>  $Ca_3 (PO_4)_2$ **POLY PO4** ZINC

and Azoles

CLOT -MoO<sub>4</sub><sup>2-</sup> hc NO2 PO<sub>4</sub>3- hc Na<sub>3</sub>SiO<sub>4</sub> Poly PO<sub>4</sub> hc

### Corrosion



**Pitting** Tubercule







## Microbiological Control

- Active action •
- Sulfate reducing bacteria
- Nitrifying bacteria
- De-nitrifying bacteria
- Iron reducing Bacteria



- Slime forming
- Algae colonies
- bio film





# Microbiological Control (chemical treatment –second defense line)



- Oxidizing Biocides
- "Indiscriminate oxidation"
  - non-specific
  - react also with suspended solids and organic matter
  - halogenated by-products may be formed

- Non-Oxidizing Biocides
  - Specific Reactions
- specific mode of action
- organisms may acclimatize to the biocide
- more expensive



# Microbiological Control (chemical treatment –second defense line)

- Oxidizing Biocides
- "Indiscriminate oxidation"
  - Chlorine gas
  - Sodium hypochlorite
  - Calcium hypochlorite
  - Chlorine dioxide
  - Solid halogen donors
  - Activated Bromide
  - Peracetic Acid
  - Ozone

- Non-Oxidizing Biocides
  - Specific Reactions

Dbnpa

**Bronopol** 

**Quaternary Ammonium Salts** 

Glutaraldehyde

Iso-thiazolinone

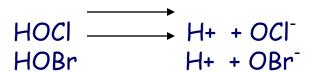
Triazines

Blends



34

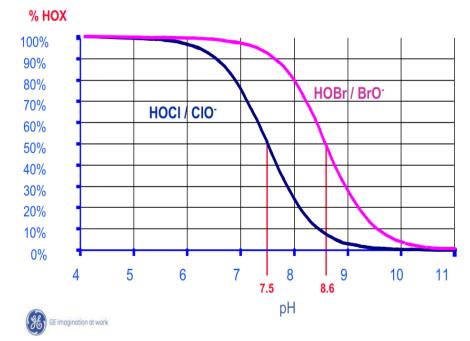
# Oxidizing Biocides & pH Effect



→  $H+ + OCl^{-}$  (Hypochlorite ion, less effective)  $H+ + OBr^{-}$  (Hypobromite ion, less effective)



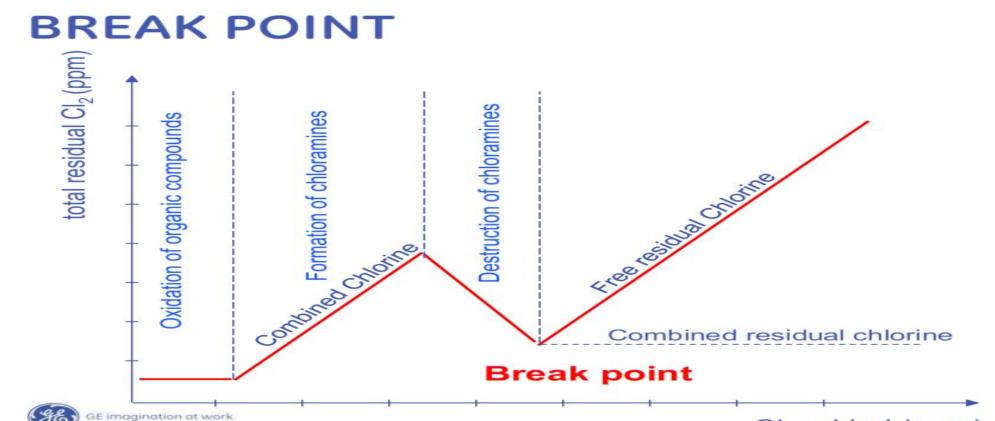
	pH value in Cooling Water					
	6.0	7.0	7.5	8.0	8.5	9.0
% Available HOCl	97	76	50	24	9	3
% Available HOBr	100	98	94	83	60	33





# Chlorine dosing (break point )







Cl<sub>2</sub> added (ppm)

## 3.Fouling



Definition: fouling is the accumulation of unwanted material on solid surfaces, most often in aquatic environment.

Fouling occurs on a surface of a component, system or plant performing a useful function causing disturbance or interference with this function.

#### Types:

- 1- Colloidal.
- 2- Biofouling.
- 3- Organic.
- 4- Scaling.



# Fouling first defense line



- •Dispersant polymers (polyphosphates, polyacrylates...etc)..
- Side-stream sand filtration.
- Polyelectrolytic precipitation (Coagulants and Flocculants).



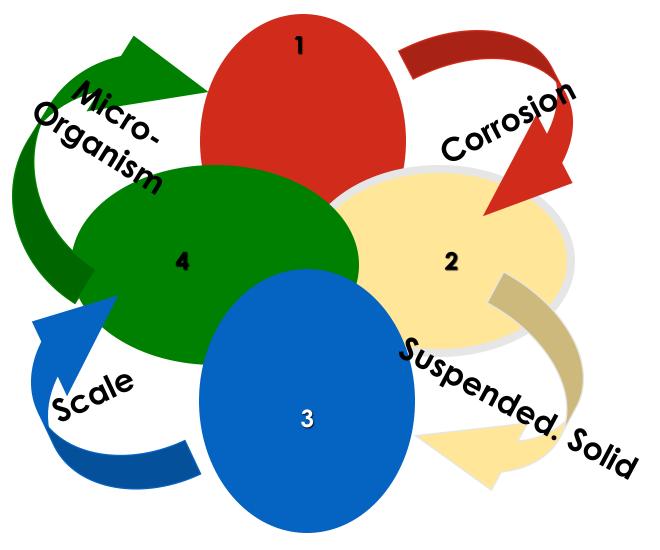
#### The Main Problems In Water

 Water is the strongest solvent on the earth

we find in water:

- (1) suspended solids
- (2) dissolved solids
- (3) dissolved gases

all these components dissolved in water cause problems in the operation of boilers, which must be addressed





## Cooling tower controller









#### Why cooling tower controller?



#### **Definition:**

- a cooling tower controller is used to automate the chemical feed of your corrosion and scale prevention chemical and biocide.
- it also controls the (conductivity blowdown set points-adjust PH and chlorine injection ) to maximize water reuse and to ensure a successful water treatment program.

#### Why controller?

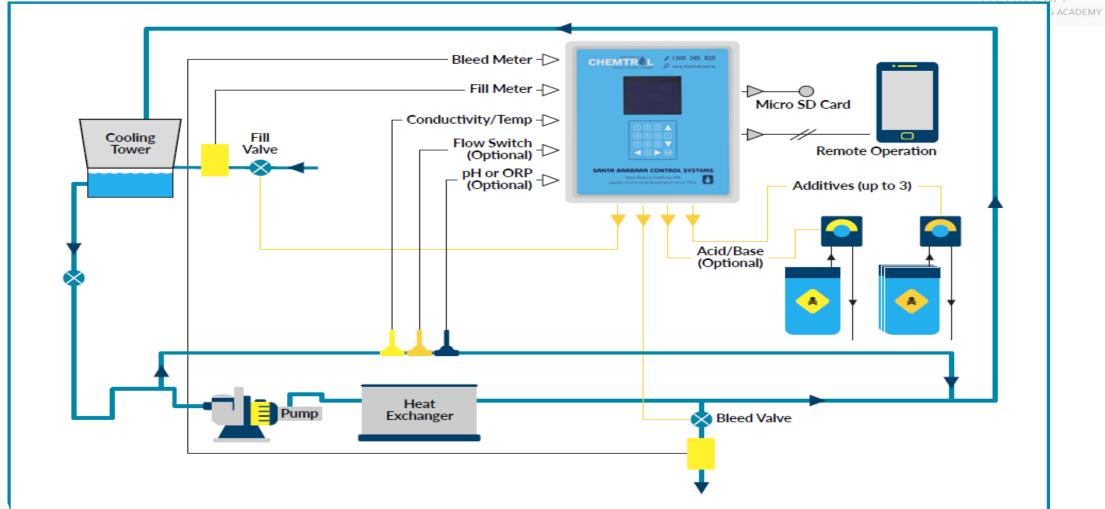
- Avoid water waste
- Avoid chemical waste
- Improve system performance
- Minimize corrosion, scaling, and micro growth problems
- Record and trace operational problems
- Easy way for reporting water treatment parameters
- Labor reduction, eliminating failure due to human factor



41

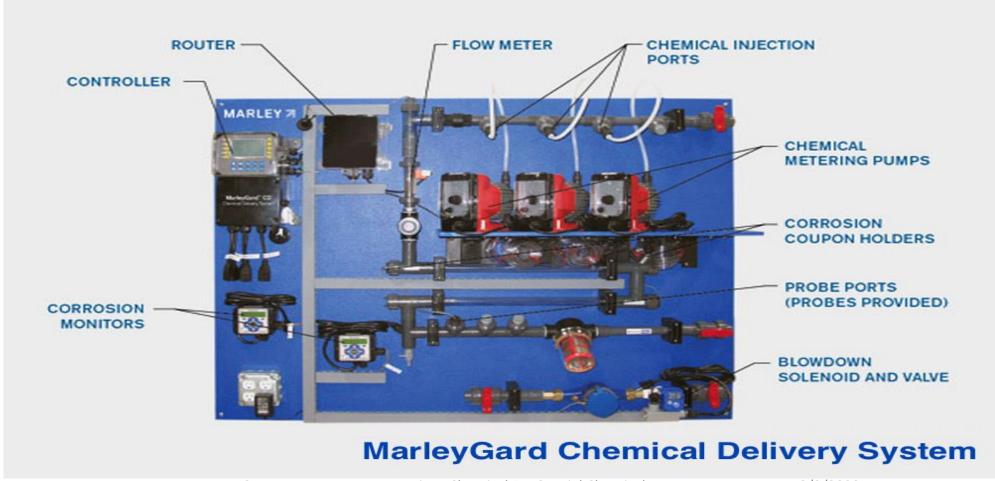
#### Cooling tower controller





# Cooling tower controller (Rac component)

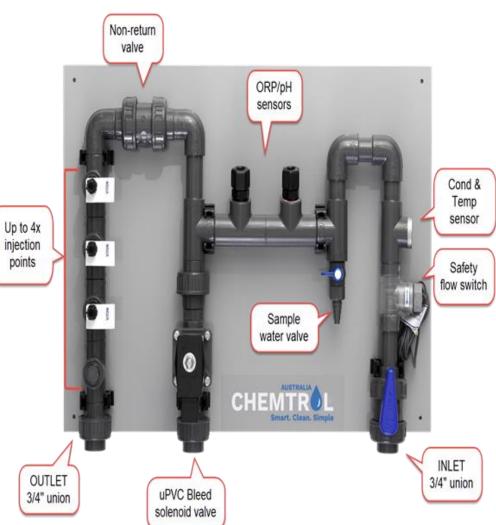














## Cooling tower controller input



Input

Water meter

Chemical levels

Conductivity or induced conductivity

Standby

Flow

conductivit y probe

Temperature probe

PH

Stand by

Flow

PH level

PH probe

Temperature probe

ORP

Or

free chlorine

Stand by

Flow

ORP level

ORP probe

Temperature probe

Corrosion

Stand by

Flow

PH level

PH probe

Temperature probe

Tracer

Stand by

Flow

TRC level

TRC probe

Temperature probe

MA

0-20 mA input

Stand by

Flow

level

Temperature probe



### Cooling tower controller input











Corrosion



Tracer





#### Cooling tower controller in put VS out put



Conductivity or induced conductivity

Bleed



alarm

PH **Propotional** PH dosing system alarm

**ORP** Or free chlorine Proportional ORP dosing system alarm

Corrosion Corrosion inhibitor dosing system alarm

Tracer TRC dosing system alarm

# Communication" remote management system



INSTRUMENT CONFIGURATION	PLUS	WHEN	REQUIREME NTS	FUNCTION
USB	USB output	You do not need a PC on your plant: you can download data log on a USB device	/	RS485 output to link other EMECinstruments Data Log on USB device
ETHERNET	LAN network between instrument and web	Remote control via WEB ERMES	LAN wiring (RJ- 45)	RS485 output to link other EMECinstruments Web ERMES remote control (PC, smartphone or tablet) Alarm messages via email
MODBUS	Connection to other devices (PLC) via RS485	PLC plant management	/	PLC connection output for reading and modifying parameters





GSM/GPRS 1	3G/4G modem between instrument and web	Remote control via WEB ERMES	Network coverage	RS485 output to link other EMEC instruments Web ERMES remote control (PC, smartphone or tablet) Alarm messages via email Alarm messages via SMS
WIFI 1	WIFI network between instrument and web	Remote control via WEB ERMES	Network coverage	RS485 output to link other EMECinstruments Web ERMES remote control (PC, smartphone or tablet) Alarm messages via email



#### Brain storming



Could you treat cooling tower without using chemicals ??



