

Environmental



**Summer
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Water Quality Parameters

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7.3
10/10/19

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Water Quality

Q: When is dirty water?

Answer: of course depends on what we mean by dirty.

مبشر اننا نقسم جودة المياه فديكون ذاتيا ولعينة المياه والعايير المنظمة.

Water Quality Parameters

- Physical أي عوامل يمكن
تصورها بالرائحة
 - Temperature
 - Color
 - Turbidity
 - Odor and Taste
- Chemical مواد كيميائية
 - pH
 - Dissolved Oxygen
 - Hardness
 - Chlorine
- Biological مواد من مصدر حيوي
 - Bacteria
 - Fungi
 - Algae
 - Protozoa
- Radiological اشعاع
 - Radioactive materials

* هذه التصنيفات تستخدم لتقييم سلامة المياه
للاستخدام البشري، الزراعة أو صناعة بناء
كما أنها مختلفة

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* هذه المعايير تستخدم لتحليل الخصائص الفيزيائية للمياه والتي تكون

Physical Parameters

are the first step in water quality assessment before chemical/biological analysis.

1. Suspended solids (big particles) جسيمات عالقة كبيرة الحجم that remain undissolved in water

2. Turbidity (tiny particles) تurbidity تسبب تعكر Causing cloudiness
attracting water clarity

3. Odor and Taste رائحة ذوق Indicators of chemical/organic contamination

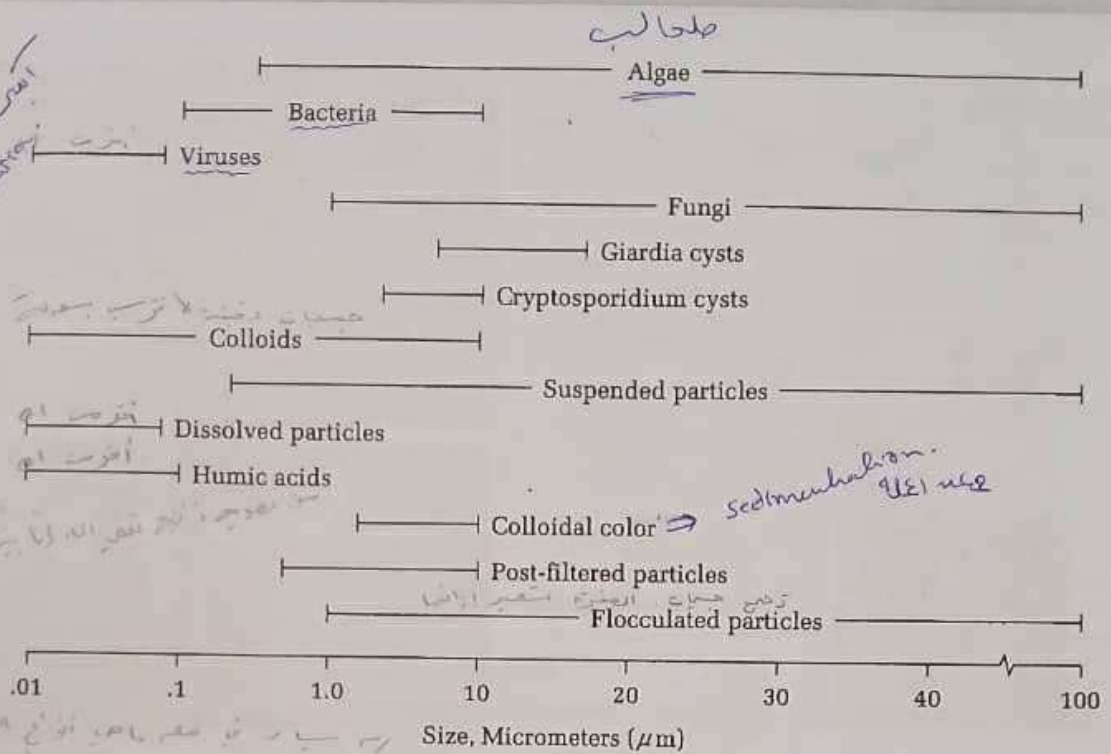
4. Color لون May signal impurities or organic matter

5. Temperature درجة حرارة Influences oxygen solubility and
microbial growth

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Sizes of Particles in Water



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Solids by Size ←

Classification	Diameter (mm)
Dissolved	< 0.000 001
Colloidal	0.000 001 – 0.001
Suspended	0.001 - 0.1
Particulate	> 0.1

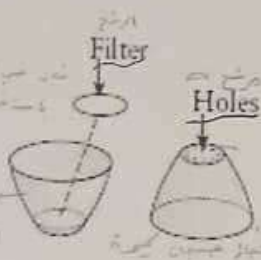
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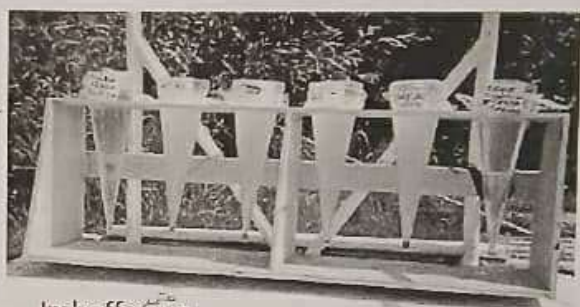
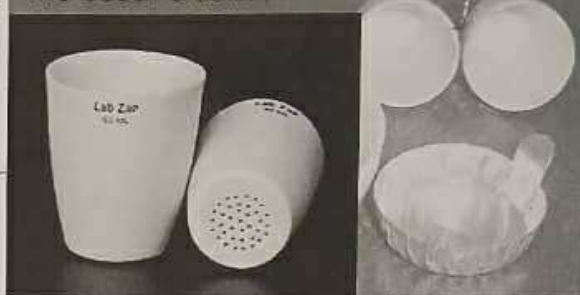
SS Separation from Water

Filtration
Suspension
Dissolved

Filtration
Distillation



The Gooch crucible



Imhoff cone

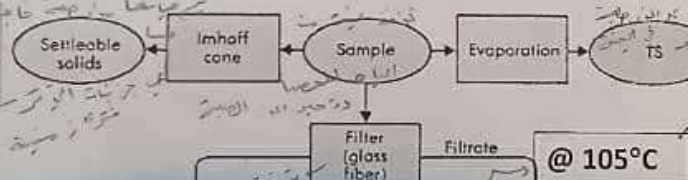
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1" 1/2
check

Suspension
dissolve

Fixed
Filtration
Filtrate



- TS = total solids
- SS = suspended solids
- VSS = volatile suspended solids
- FSS = fixed suspended solids
- TVS = total volatile solids
- TFS = total fixed solids
- DS = Dissolved solids

$TS + TDS = TSS$
Total Solids + Dissolved Solids = Total Suspended Solids

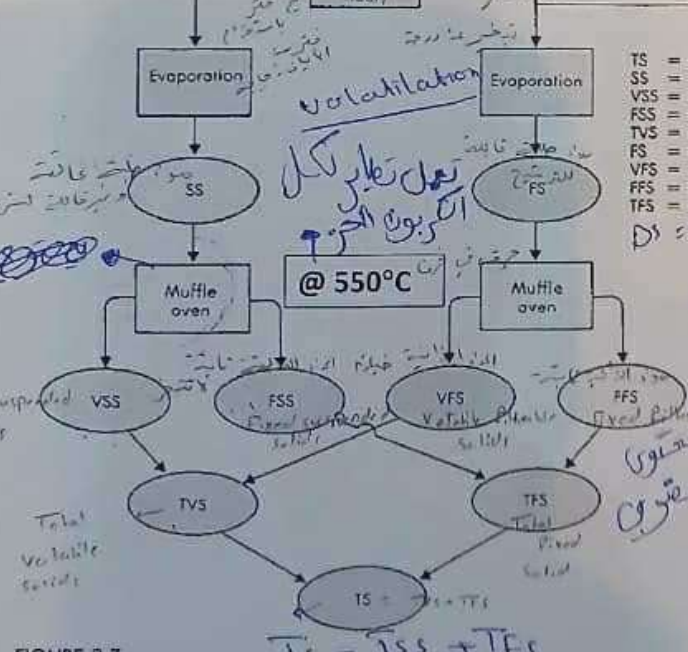


FIGURE 2.7

Interrelationships of solids found in water and wastewater. In much of the water quality literature, the solids passing through the filter are called dissolved solids.

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TS & SS

$$TS = \frac{W_{ds} - W_d}{V} \times 10^6$$

where,

TS = total solids, mg/L

W_{ds} = weight of dish plus the dry solids after evaporation, g

W_d = weight of the clean dish, g

V = volume of sample, mL

$$SS = \frac{W_{df} - W_d}{V} \times 10^6$$

where,

SS = suspended solids, mg/L

W_{df} = weight of dish plus dry filtered solids, g

W_d = weight of clean crucible and filter, g

V = volume of sample, mL

TS

DS

SS

➤ Total solids = dissolved solids + suspended solids.

▪ A **Gooch crucible** is used to separate suspended solids from dissolved solids.

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Solids: FS & VS

$$FS = \frac{W_{du} - W_d}{V} \times 10^6$$

Where,

FS = fixed solids, mg/L

W_{du} = weight of dish plus unburned solids, g

W_d = weight of clean crucible and filter, g

V = volume of sample, mL.

$$VS = TS - FS$$

Where,

VS = volatile solids, mg/L.

Although volatile solids are considered to be organic, some of the inorganics are decomposed and volatilized as well at the 550°C - 600°C used for the test. However, this is not considered a serious drawback.

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Turbidity

It is the muddy or cloudy appearance of clay or such other particles that presents hindrance to the path of light.

The turbidity is measured by a turbidity rod or a turbidity meter with physical observations and is expressed as the suspended matter in mg/l or ppm (part per million).

Colloids, clay, silt, tiny fragments of organic matter and microscopic organisms (bacteria, algae) are some of the substances that cause turbidity.

Units of turbidity

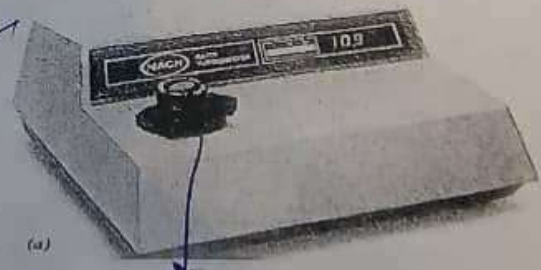
The standard unit of turbidity is that which is produced by 1 mg of finely divided silica in one liter of distilled water

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 methods:

Nephelometers (NTU)

Jackson candle turbidimeter (JTU)



Taste and Odor

- A physical characteristics of drinking water for aesthetic reasons.
- The dissolved inorganic salts or organic gases may impart taste and odor to the water.
- Inorganics (e.g. soil components) are usually only be responsible for taste.

- T & O are most commonly caused by: algae, decomposed organic matter, and dissolved gases.
- Oxidants (Chlorine, potassium permanganate) are used to oxidize the materials causing the taste and odor. Or Activated carbon filters can be used.

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T & O measurement

- The extent of taste or odor is measured by a term called odor intensity which is related with threshold odor, which represents the dilution ratio at which the odor is hardly detectable.

➤ The water to be tested is gradually diluted with odor free water and the mixture at which the detection of taste and odor is *just lost* is determined.

➤ The odor is measured (by human nose) and expressed in terms of a threshold odor number (TON) = $(V_t + V_d) / V_t$.

V_t : volume tested, V_d : volume added.

TON=1 is the lowest value possible.

Example: 50 ml is diluted to vol. of 200 ml. The dilution no. equals $200/50 = 4 = \text{TON}$

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Color

- It is a physical characteristic of drinking water that are important for aesthetic reasons.
- It may be caused by dissolved (true color) or suspended colloidal particles (apparent color) or by dissolved organic matters from decaying vegetation.
- It is measured in the labs by Nessler's tubes by comparing the sample with the known color intensities. More precisely tintometer measures it.
- One color unit is equivalent to the color produced by a 1 mg/l solution of platinum.

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Temperature

- Warm water tastes flat.
- Cooling water suppresses odors and tastes and makes it more palatable.
- Temperature also affects the chlorination and purification of water.
- Disinfection takes longer when water is colder, and purification capacity is reduced with reverse osmosis treatment equipment.
- An average temperature of 15°C is generally suitable.

1. Total dissolved solids:

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

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Hardness

- Hardness is caused by multivalent cations (or minerals)—such as calcium, magnesium and iron—that dissolve from soil and rocks (particularly limestone).

➤ While hardness does not cause health problems, it does reduce the effectiveness of soaps, and cause scale formation especially in heat transfer equipment.

- Total hardness (TH) is defined as the sum of the multivalent cations in the water. Mainly, Calcium (Ca^{2+}) and Magnesium (Mg^{2+}).

$$\text{TH} = \sum (\text{Multivalent cations}) \cong \text{Ca}^{2+} + \text{Mg}^{2+}$$

- Typical units for hardness are mg/L as CaCO_3 and meq/L

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Hardness Classification

Hardness

- Soft < 50 mg/L
- Moderate = 50 - 150 mg/L
- Hard = 150 - 300 mg/L
- Very Hard > 300 mg/L

هناك 3 أنواع

quantitative

Quantitatively
بالمقاييس والقياسات

Qualitatively
نوعياً
على أساس قطرة تاشبه اللون
فهيمن يعرف الماء Hard or soft
وهو شغل بالمصانع

➤ Treatment usually left to consumer (domestic, industrial, etc.) depending on needs

ما ينبغي للستهلاك

حلال

Hardness Classification

Classification	Hardness	
	meq/L	mg/L as CaCO_3
Extremely soft to soft	0-0.9	0-45
Soft to moderately hard	0.9-1.8	46-90
Moderately hard to hard	1.8-2.6	91-130
Hard to very hard	2.6-3.4	131-170
Very hard to excessively hard	3.4-5	171-250
Too hard for ordinary domestic use	>5	>250

لازم
آنت
unit
هنا

Source: L.A. Lipe and M. D. Curry, "Ion Exchange Water Softening," a discussion for water treatment plant operators, 1974-75 seminar series sponsored by Illinois Environmental Protection Agency.

Hardness Calculations

مهم جداً هذا القانون

$C_q = \frac{C}{EW} \rightarrow (mg/L)$
 where C_q = concentration in meq/L
 C = concentration in mg/L
 EW = equivalent weight in g/eq or mg/meq $\rightarrow$ molecular weight - atomic weight

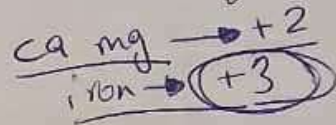
mg/L $\rightarrow$ Concentration
 meq/L

$C_q = \frac{C}{EW}$

$$EW = \frac{AW \text{ or } MW}{n} \rightarrow \text{cationic}$$

where, atomic weight (AW) and molecular weight (MW), valence or ionic charge (n, which is always positive):

$$C_q \times 50$$



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To convert to the standard unit mg/L as $CaCO_3$, the meq/L concentration is multiplied by the equivalent weight of $CaCO_3$, which is 50.0 mg/meq (100 g/mol / 2 eq/mol):

$$C_{CaCO_3} = C_q \times 50.0$$

$EW = \frac{40 + 12 + 16 \times 3}{2} \rightarrow [Ca^{2+}]$

where C_{CaCO_3} = concentration in mg/L as $CaCO_3$
 C_q = concentration in meq/L

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✓ Example 10.4

Problem The concentration of calcium in a water sample is 100 mg/L. What is the concentration in (a) meq/L and (b) mg/L as CaCO₃?

Solution The valence or ionic charge of calcium is +2, so n is 2 eq/mol. Calcium's atomic weight is 40.1 g/mol. Therefore, its equivalent weight is (Equation 10.5):

$$C_1 = \frac{C}{EW}$$

$$EW = \frac{AW}{n} = \frac{40.1 \text{ g/mol}}{2 \text{ eq/mol}} = 20.0 \text{ g/eq} = 20.0 \text{ mg/meq}$$

- a. The concentration in meq/L is then simply obtained through unit conversion (Equation 10.4):

$$C_q = \frac{C}{EW} = \frac{100 \text{ mg/L}}{20.0 \text{ mg/meq}} = 5.0 \text{ meq/L}$$

- b. Again, the concentration in mg/L as CaCO₃ is simply obtained through unit conversion (Equation 10.6):

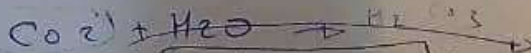
$$\begin{aligned} C_{\text{CaCO}_3} &= C_q \times 50 \\ &= (5.0 \text{ meq/L})(50 \text{ mg/meq}) \\ &= 250 \text{ mg/L as CaCO}_3 \end{aligned}$$

Note that the *correct* unit includes "as CaCO₃."

Alkalinity

7.2-8.4 pH → **Alkalinity** is a measure of the buffering capacity of water (or the capacity of the water to neutralize acid, or H⁺).

- It is not the same as pH; water does not have to be basic to have high alkalinity.
- The natural sources of alkalinity in water are the atmosphere and limestone.



Atmospheric carbon dioxide (dissolved in water) forms carbonic acid (H₂CO₃), which can dissociate into bicarbonate (HCO₃⁻) and carbonate (CO₃⁻²).

Limestone formations (CaCO₃) can also dissolve in water and produce HCO₃⁻ and CO₃⁻². Both HCO₃⁻ and CO₃⁻² can "grab" (or neutralize) H⁺ that is added to the water. Notice that carbonate can grab two H⁺ while bicarbonate can grab only one.

Alkalinity Calculations (mol/L)

$$\begin{aligned} \text{Alkalinity (mol/L)} = & \left(\frac{\text{mol HCO}_3^-}{\text{L}} \right) \left(\frac{1 \text{ mol ALK}}{\text{mol HCO}_3^-} \right) \\ & + \left(\frac{\text{mol CO}_3^{2-}}{\text{L}} \right) \left(\frac{2 \text{ mol ALK}}{\text{mol CO}_3^{2-}} \right) \\ & + \left(\frac{\text{mol OH}^-}{\text{L}} \right) \left(\frac{1 \text{ mol ALK}}{\text{mol OH}^-} \right) \\ & - \left(\frac{\text{mol H}^+}{\text{L}} \right) \left(\frac{1 \text{ mol ALK}}{\text{mol H}^+} \right) \end{aligned}$$

or simply

$$\text{Alkalinity (mol/L)} = [\text{HCO}_3^-] + 2[\text{CO}_3^{2-}] + [\text{OH}^-] - [\text{H}^+]$$

(1 H)
2 H
↓
 $\frac{\text{mol}}{\text{L}}$
Alkalinity

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Alkalinity Calculations (meq/L)

2H : 2 eq / 1 mol $\text{CO}_3^{2-} \text{ (eq/L)} = \left(\frac{\text{mol CO}_3^{2-}}{\text{L}} \right) \left(\frac{2 \text{ eq}}{\text{mol}} \right)$

$$\begin{aligned} \text{Alkalinity (meq/L)} = & \left(\frac{1 \text{ meq HCO}_3^-}{\text{L}} \right) \left(\frac{1 \text{ meq ALK}}{\text{meq HCO}_3^-} \right) \\ & + \left(\frac{1 \text{ meq CO}_3^{2-}}{\text{L}} \right) \left(\frac{1 \text{ meq ALK}}{\text{meq CO}_3^{2-}} \right) \\ & + \left(\frac{1 \text{ meq OH}^-}{\text{L}} \right) \left(\frac{1 \text{ meq ALK}}{\text{meq OH}^-} \right) \\ & - \left(\frac{1 \text{ meq H}^+}{\text{L}} \right) \left(\frac{1 \text{ meq ALK}}{\text{meq H}^+} \right) \end{aligned}$$

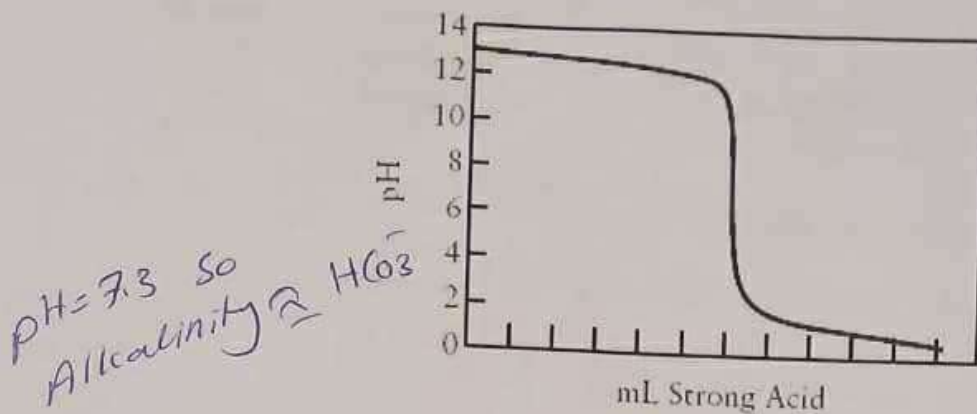
or simply

$$\text{Alkalinity (meq/L)} = [\text{HCO}_3^-] + [\text{CO}_3^{2-}] + [\text{OH}^-] - [\text{H}^+]$$

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Alkalinity: Titration curve



At pH less than 7.3, the carbonate concentration is much less than the bicarbonate concentration, and around neutral pH, $[\text{H}^+] \approx [\text{OH}^-]$. Therefore, for most potable water sources:

$$\text{Alkalinity} \approx (\text{HCO}_3^-)$$

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Carbonate hardness & Noncarbonate hardness

$$\text{TH} = \text{carbonate hardness (CH)} + \text{noncarbonate hardness (NCH)}$$

- Carbonate hardness is the component of total hardness associated with the anions carbonate (CO_3^{2-}) and bicarbonate (HCO_3^-);
- Carbonate hardness (total alkalinity) is a temporary hardness that is removed by heat;
- Noncarbonate hardness is equal to the difference between total hardness and carbonate hardness;
- If the alkalinity is equal to or greater than the total hardness, then the noncarbonate hardness is zero.

$$\text{NCH} = \text{TH} - (\text{CH})$$

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Example

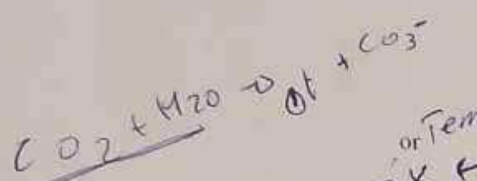
حای زیره بارقسان

10.6

Problem From the following water analysis, determine the total hardness, carbonate hardness, and noncarbonate hardness in (a) milliequivalents per liter (meq/L) and (b) milligrams per liter (mg/L) as CaCO_3 .

Component	Concentration (mg/L)
CO_2	6.0
Ca^{2+}	50.0
Mg^{2+}	20.0
Na^+	5.0
Alkalinity	120 as CaCO_3
SO_4^{2-}	94.0
pH	7.3

Component	EW = $\frac{\text{Mol}}{\text{X}}$	Concentration
Ca^{2+}	20.0	50 mg/L
Mg^{2+}	12.2	20 mg/L
Alkalinity	50.0	120 meq/L



24.31

TH = (2.5 meq/L) + (1.6 meq/L) = 4.1 meq/L

TH = (125 mg/L as CaCO_3) + (82 mg/L as CaCO_3) = 207 mg/L as CaCO_3

207 >> Alkalinity

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Examples

- Ex 10.4 ✓
- Ex 10.5 ✓
- Ex 10.6 ✗

INTRODUCTION TO ENVIRONMENTAL ENGINEERING

THIRD EDITION

P. AARNE VESILIND Bocknall University
SUSAN M. MORGAN Southern Illinois University Edwardsville
LAUREN G. HEINE Clean Production Action

WATER AND WASTEWATER ENGINEERING

Design Principles and Practice

Mackenzie L. Davis, Ph.D., P.E., RCEE
Michigan State University

- All Examples in Chapter 7

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8
expt 321

7.2
7.3

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Microorganisms:

▪ Bacteria

الميكروبات ▪ Protozoa

الديدان طفيلية ▪ Parasitic Worms

▪ Viruses

الطحالب ▪ Algae

الفطريات ▪ Fungi

مؤشرات تلوث ▪ Pollution indicators: Coliform bacteria

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Microorganisms

البحث عن مسببات الأمراض على السطح من المياه العذبة في الوقت نفسه (أو غير ذلك)

✓ Looking for pathogens in most surface waters is a perfect example of the proverbial needle in a haystack!

➤ E. coli used as an indicator of water quality: normal inhabitant of intestines of many animals

➤ Indicator organisms: the indicator most often used is a group of microbes called coliforms (which includes the 150 strains of E. coli).

Coliform test:

- Membrane filtration and 24 or 48 hours of incubation.
- Results reported as Most Probable Number (MPN) per 100 mL
- Incubation at moderate temperature (35°C) for 48 hr

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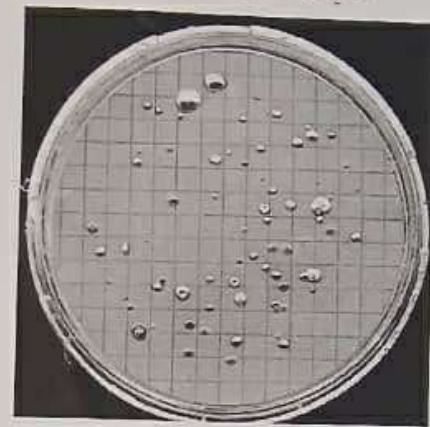
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A fecal coliform bacteria test is used to indicate the likely presence of disease-causing bacteria in water.

يستخدم هذا الاختبار للكشف وجود
بكتيريا سببية للأمراض في الماء

لغرض
تقدير
تركيز البكتيريا

عينة ماء
مختلطة
بماء جاف



تظهر
مستعمرات
بكتيريا
في
صفحة

✓ a high coliform count is suspicious, and although the water may in fact be perfectly safe to drink, it should not be consumed.
✓ The opposite is also true!

Radiological Parameters

Radioactive substances

Disinfection By-Products (interim)			Primary MCLs
Total trihalomethanes	0.10		
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 µg/L

وحدة قياس الإشعاع البيولوجي

تقبل
الاشعة
بمعدل
محدد
على
الإنسان

- The rem, or roentgen equivalent man, takes into account the biological effect of absorbed nuclear radiation, so it measures the extent of biological injury.
- One Sv is numerically equal to 100 rem.
- pCi/L (pico Curies of radon per liter of air)

غاز تنفسي الهواء

Water Quality Management

إدارة جودة المياه

Dr. Motasem Saidan

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فهم مفهوم 1 = رتبة بين مقياس المياه وقاعدتها

Alkalinity

تأثير الماء = قدرة الماء على معادلة الأحماض وهي ليست مقياساً مباشراً
بل مقياس للمحتوى الكيميائي من القواعد، القابلية للذوبان في الماء
قاعدية الماء = درجة حموضته الماء (pH) عندما تكون $pH > 7.7$

Univ. of Jordan/ Chem. Eng. Dept.

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Water Quality Management

Water quality management is the science that predicts how much waste is too much for a body of water

Assimilated-amount of waste that can be tolerated by a body of water

assimilated capacity

Determined by knowing the type of pollutants discharged and their effect on water quality

13/8 Midterm

كمية القابلية
بالماء
المحتمل

Measures of Water Quality

What are some parameters that determine water quality?

The parameters that have been covered in the previous lectures

Biochemical Oxygen Demand (BOD)

Chemical Oxygen Demand (COD)

Dissolved oxygen (DO)

water quality

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Biochemical Oxygen Demand (BOD)

BOD

Measurement of the rate at which microbes degrading organic matter use oxygen

One can calculate the theoretical oxygen demand

If you know the chemical formula – for example $1.67 \times 10^{-3} \text{ M}$ glucose



$$6 \times 1.67 \text{ moles/L} \times 32 \text{ g/mole} \times 1000 \text{ mg/g} = 321 \text{ mg/L}$$

BOD is used because wastewater isn't one organic molecule

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Five-day BOD

BOD₅ is the amount of dissolved oxygen consumed by microorganisms during the biochemical oxidation of organic matter to carbon dioxide in 5-day standard test at 20°C.

➤ A very low rate of use would indicate

- /1) contamination is absent,
- /2) the available microorganisms are uninterested in consuming the available organics, or
- /3) the microorganisms are dead or dying.

or unavailable or inactive.

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The standard BOD test

The standard BOD test is run:

- in the dark at 20 °C for five days
- At specified temperature because the rate of oxygen consumption is temperature dependent.
- In a reaction that must occur in the dark because algae may be present.
- universally using a standard BOD bottle (about 300mL volume),
- In a special nonreactive glass and supplied with a ground-glass stopper



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BOD test procedures:

1. A special 300 mL BOD bottle is filled with a sample of water that has been appropriately diluted and inoculated with microorganisms.
2. Blank samples containing only the dilution water are also placed in BOD bottles and sealed.
3. The sealed BOD bottles containing diluted samples and blanks are incubated in the dark at 20°C for the desired number of days.
4. After five days has elapsed, the samples and blanks are removed from the incubator and the dissolved oxygen concentration in each bottle is measured.

Fresh Water at 1 atm*

Temperature (°C)	Dissolved Oxygen (mg/L)
0	14.60
1	14.19
2	13.81
3	13.44
4	13.09
5	12.75
6	12.43
7	12.12
8	11.83
9	11.55
10	11.27
11	11.01
12	10.76
13	10.52
14	10.29
15	10.07
16	9.85
17	9.65
18	9.45
19	9.26
20	9.07
21	8.90
22	8.72

Source: *For atmospheric pressures of less than 1 atm, the dissolved oxygen concentration has to be reduced because there is less pressure to push the oxygen into the water. Published correction factors are available.

- النهاية
- Ultimate BOD (BOD_u or L): which is the O_2 demand after a very long time—when the microorganisms have oxidized as much of the organics as they can. The ultimate BOD is usually run for 30 days.

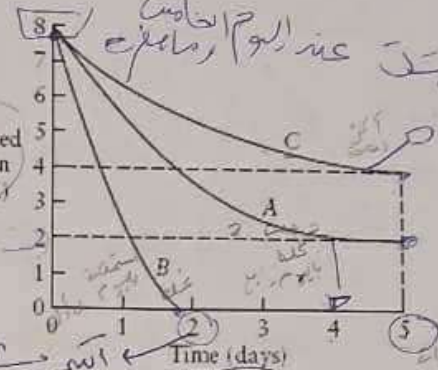
$$BOD = I - F$$

Where;

I = initial DO, mg/L

F = final DO, mg/L

Dissolved oxygen (mg/L)



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dilution

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Dilution

What if the F is =0?

$DO \rightarrow$ بعد اليوم الرابع

منزل لا يصغر $\rightarrow$ هاهي مشكلة

Dilution of the sample is required when $F = 0$. If $F = 0$ we don't know how much DO would have been used.

Initial - Final < 2

DO

DO

هاهي مشكلة لانهم

Dilution اقل

$$BOD = (I - F)D$$

where, D = dilution factor.

total volume of bottle

$D =$

$$\frac{[\text{total volume of bottle}]}{[\text{volume of dilution water}]}$$

$$D = \frac{\text{Total volume of bottle}}{\text{Volume of sample}}$$

$$\frac{300}{300}$$

300 ml

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10

$$D = \frac{300}{15}$$

Example

$$BOD_t = (1-f)D$$

$$D = \frac{300}{15}$$

Volume up to 300

Determine the BOD₅ for a 15 ml sample that is diluted with dilution water to a total volume of 300 ml when the initial DO concentration is 8 mg/l and after 5 days, has been reduced to 2 mg/l

$$D = 6 \quad F = ?$$

$$BOD_5 = (8 - F) \times 6 = 120$$

$$I = 8$$

$$F = 2$$

$$D = 300/15 = 20$$

$$BOD_5 = (8 - 2) \times 20 = 120$$

Dilution Factor

Remark: The assumption in the dilution method is that the results from each dilution of a single sample will yield the same BOD value (**No Sliding Scale**)

Examples: 9.4 9.5

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Sample	Dilution
3 ml	291
1.5	298.5
0.75	299.25

$$D_1 = 99$$

$$D_2 = 199$$

$$D_3 = 399$$

$$BOD = (1 - f) D$$

How much dilution?

not a fact

The BOD of the sample is seldom known before the BOD test is conducted, so the dilution must be estimated.

The test is not very precise if the drop in DO during the five days of incubation is less than about 2 mg/L or if the DO remaining in the bottle is less than 2 mg/L.

Generally, a final BOD is estimated from previous tests or similar waste and at least two dilutions used, with one expected to have at least 2 mg/L DO remaining and one expected to use at least 2 mg/L of DO.

The dilution is then calculated as

$$D = \frac{\text{expected BOD}}{\Delta DO}$$

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$$D = \frac{DO_{\text{initial}} - DO_{\text{final}}}{\Delta DO}$$

2 cases

very important

بما ان معدل كدورة السبب يتبع درجات الحرارة، اذا ما اعطى حرارة المختبر

room temp

Example: The five-day BOD of an influent to an industrial wastewater treatment plant is expected to be about 800 mg/L based on similar wastewaters. What dilutions should be used in a five-day BOD test?

$D = \frac{\text{expected BOD}}{I - f}$
 $I = 10$
 $\text{BOD} = 800$

$$D = \frac{\text{expected}}{(I - f)} = \frac{800}{(10 - f)}$$

assume $D \geq 2$
 $f \geq 2$

two dilution must set

Case 2 = $\frac{800}{8} = 100$

Case 1 = $\frac{800}{2} = 400$

$2 = 10 - f$
 $f = 8$

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also flow rate

دليل على dilution

Seeding: the addition of active microorganisms that take up oxygen

- May be required in samples that do not have their own
- If seeding is necessary, any BOD that is contributed by the seed must be subtracted

$$\text{BOD}_t = \left[(I - F) - (I' - F') \left(\frac{X}{Y} \right) \right] D$$

where BOD_t = biochemical oxygen demand measured at some time, t , mg/L

I = initial DO of bottle with sample and seeded dilution water, mg/L

F = final DO of bottle with sample and seeded dilution water, mg/L

I' = initial DO of bottle with only seeded dilution water, mg/L

F' = final DO of bottle with only seeded dilution water, mg/L

X = seeded dilution water in sample bottle, mL

Y = volume of BOD bottle, mL

D = dilution of sample

Dilution factor

seeding

300 mL

Total Volume 300 mL

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بانتقال هو مني افراف
 Same = initial

العينه 300
 10 → 300
 290 → 300
 10

Example

Standard BOD test with a 1:30 dilution with seeded dilution water is run. Both bottles begin at saturation, 9.2 mg/L. After five days, the bottle with waste has a DO of 2 mg/L, while the DO of the seed = 8 mg/L. Find the BOD₅.

BOD =

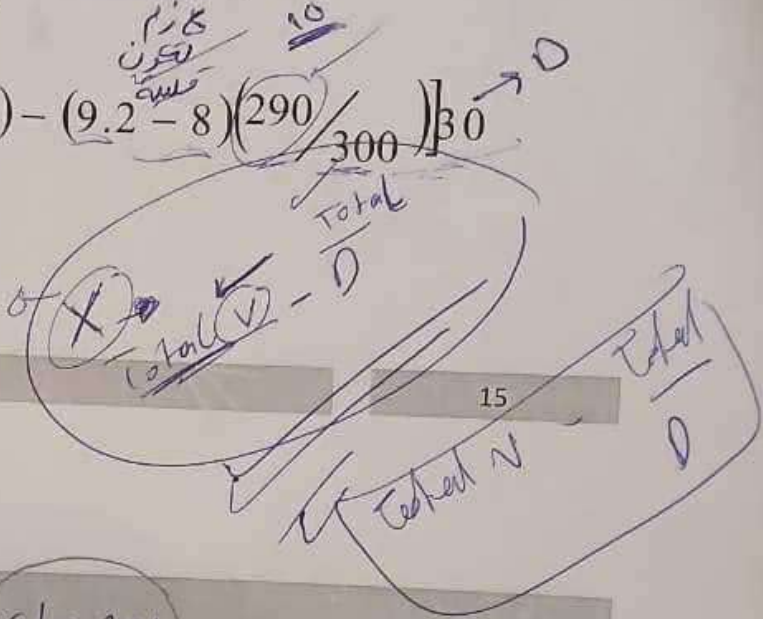
$(L - f) - \frac{b}{9.2}$
 6
 2

BOD₅ = $[(9.2 - 2) - (9.2 - 8)(290/300)] \times 30$
 = 181 mg/L

300 - 290/30

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Volume
 Seed



$X = \text{total } V - \frac{\text{total } v}{D}$

300
 30

BOD Kinetics

rate of

Thus the mass balance is:

Rate of DO Accum. = Rate of DO consumed

$z = D_0$

$\frac{dz}{dt} = -r$

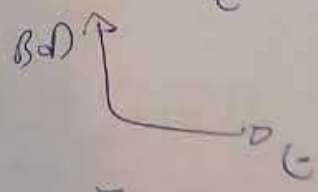
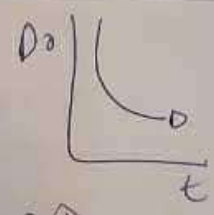
z = dissolved oxygen, mg/L

Assume that it is a first-order reaction

$\frac{dz}{dt} = -k_1 z \Rightarrow \frac{dz}{z} = -k_1 dt \Rightarrow z = z_0 e^{-k_1 t}$

$-r_A = k_1 C_A$
 $-r_A = k_1 z$

$\frac{dz}{dt}$



$\int_{z_0}^z \frac{dz}{z} = \int_0^t -k_1 dt$
 $\ln \frac{z}{z_0} = -k_1 t$

بانتقال هو مني افراف
 Same = initial

العينه 300
 10 → 300
 290 → 300
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Example

Standard BOD test with a 1:30 dilution with seeded dilution water is run. Both bottles begin at saturation, 9.2 mg/L. After five days, the bottle with waste has a DO of 2 mg/L, while the DO of the seed = 8 mg/L. Find the BOD₅.

BOD =

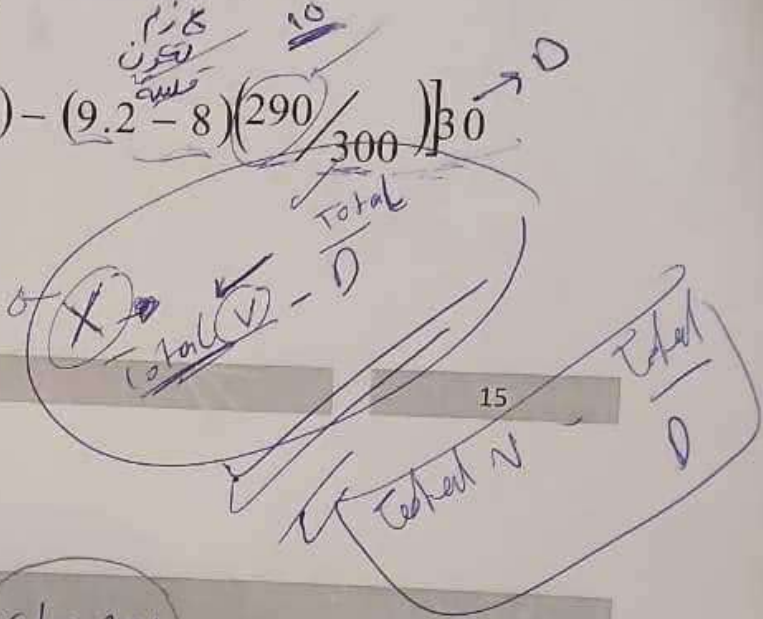
$(L - f) - \frac{b}{9.2}$
 6
 2

$BOD_5 = [(9.2 - 2) - (9.2 - 8)(\frac{290}{300})] 30$
 = 181 mg/L

$\frac{300 - 290}{30}$

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Volume
 Seed



$X = \text{total } V - \frac{\text{total } v}{D}$

$\frac{300}{30}$

BOD Kinetics

rate of

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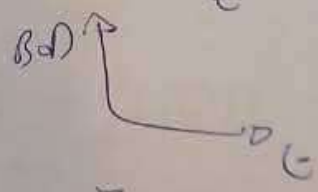
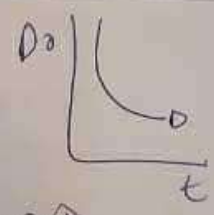
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Assume that it is a first-order reaction

$\frac{dz}{dt} = -k_1 z \Rightarrow \frac{dz}{z} = -k_1 dt \Rightarrow z = z_0 e^{-k_1 t}$

$-r_A = k_1 C_A$
 $-r_A = k_1 z$

$\frac{dz}{dt}$



$\int_{z_0}^z \frac{dz}{z} = \int_0^t -k_1 dt$
 $\ln \frac{z}{z_0} = -k_1 t$

BODs = consumption y $\leftarrow y$
 availability z $\leftarrow z$
 to be consumed

As O_2 is used, the amount still to be used is z , the amount already used is y

$y = BOD_t$
 $L = y + z$ @ $t=0, y=0, L=z_{t=0}$

Where, L = ultimate demand

$z = L - y$

$L - y = z_0 e^{-k_1 t}, z_0 = L$

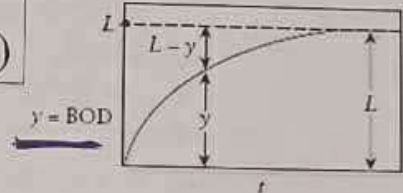
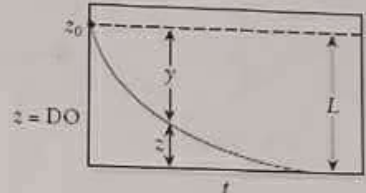
$y = L - L e^{-k_1 t} = L(1 - e^{-k_1 t})$

Where;

y = BOD at any time t

L = ultimate BOD

k_1 = deoxygenation constant, day^{-1}



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Average k

TABLE 1.4

Average BOD rate constants at $20^\circ C$

Substance	k_{10}, day^{-1}
Untreated wastewater	0.15-0.28
High-rate filters and anaerobic contact	0.12-0.22
High-degree biotreatment effluent	0.06-0.10
Rivers with low pollution	0.04-0.08

Numerical value of the rate constant k depends on:

- Nature of waste
- Ability of organisms in the system to use the waste
- Temperature

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Temperature Effect on k

- The BOD rate constant is adjusted to the temperature of receiving water using this:

Correction for Temp $k_T = k_{20} (\theta)^{T-20}$ "C"

- Where;

- k_t = BOD rate constant at the temperature of interest
- k_{20} = BOD rate constant determined at 20°C
- T = temperature in °C
- θ = Temperature coefficient (For domestic WW, 1.135 and 1.024 for re-aeration)

1.135 domestic

1.024 re-aeration

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تبعاً إلى BODs
التي تم
الملاحظة
في
الوقت
الذي
يكون
الوقت
الذي
يكون
الوقت
الذي
يكون

time BOD = y

Example 9.8: Find the BOD₅ for a waste with an ultimate BOD = 282 mg/L and a $k_1 = 0.348$ /day

$$y = L(1 - e^{-k_1 t})$$

$$= 282 \text{ mg/L} (1 - e^{-0.348 / \text{day} \times 5})$$

$$= 232.5 \text{ mg/L}$$

$$y = L - Le^{-k_1 t}$$

✓ Example 9.8

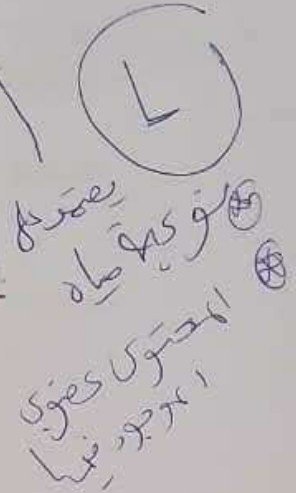
9.9

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تتبع BOD_5 مختلف باختلاف نوعية المياه

البيوت - هو الامونيا الذي تسببها الفول
ومنه من البكتيريا (بكتيريا الفم)
الاسود الحمرية العنقية العنقية
(مثل الامونيا) وتكون في الفم
البيوت - منه امد الحمرية
من المواد العنقية الحمرية - بعد اليوم 5
البيوت - مرتبان الفترات
 NO_3
 NH_3

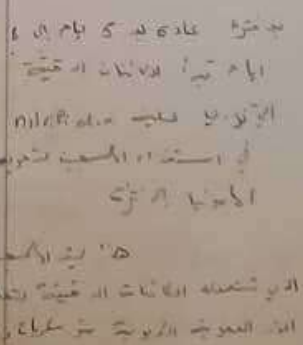


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Same Temp
 Same K
 Same Rod

21

& Nitrogenous Oxygen Demand

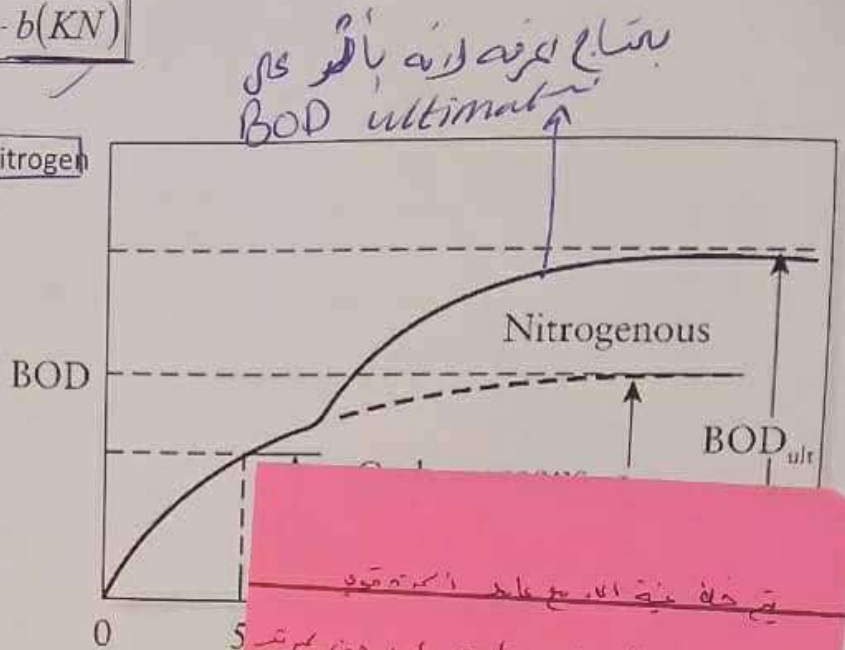
[illegible]

Nitrogenous
 Content

BOD Ultimate

$$BOD_{ultimate} = a(BOD_5) + b(KN)$$

Where $\frac{KN}{a, b}$ Kjeldahl nitrogen constants



BOD
Biological

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Chemical Oxygen Demand

Procedure:

① A sample is mixed with a strong chemical oxidizing agent (potassium dichromate, $K_2Cr_2O_7$) plus a strong acid (sulfuric, H_2SO_4) and then heated. The COD is determined by measuring the consumption of $K_2Cr_2O_7$. ③ The chemicals oxidize all the organic matter—biodegradable and non-biodegradable.

The COD for a sample will be greater than the BOD unless the sample contains primarily easily biodegradable wastes. For example, DDT, a refractory organic, would yield a high COD but a zero BOD.

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noway have COD
COD << BOD

So why is COD used?

تسمى هذه (COD) صاعدة ١.٥
← من standard ومواد مستخدمة
وذلك ما ان المرات

- COD can be obtained in about 3 hr while BOD takes 5 days at least.
- It also provides more consistent results because it is chemically, rather than biologically based.
- Sometimes the COD is also used to estimate BOD_u and to indicate the presence of biologically resistant organics.

الوقت الذي يستغرقه الكيمياء والبيولوجيا COD لتقدير الطلب البيولوجي النهائي BOD_u والوقت الذي يستغرقه الكيمياء والبيولوجيا BOD_u لتقدير الطلب البيولوجي النهائي COD

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HW

9.1 ✓
9.4 ✓
9.7 ✓
9.16 ✓
9.18 ✓
9.21 ✓
exmp 9.9
exm 9.4

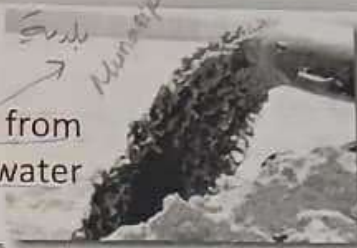
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Sources of Water Pollutants

Point source:

- Point source pollution comes mainly from industrial facilities and municipal wastewater treatment plants.
- The range of pollutants is vast, depending only on what gets "thrown down the drain."
- collected by a network of pipes or channels and conveyed to a single point of discharge in receiving water.
- Point sources are the easiest to identify and control
- Point source pollution can be controlled by waste minimization and proper wastewater treatment



easy to control:-

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* اي اشي قديم مرتبط بمصادر غير مسجل عليه .

سور فوهة

الناس جميعهم عندهم (مصادر غير مسجلة) ← لا مطار، لكن ينقل سيول القمامات ← اكلت غير المسجلة

Non-Point source:

- Urban and agricultural runoff that are characterized by overland discharge
- This type of pollution occurs during rainstorms and spring snowmelt



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Non-point Source

- The term non-point source encompasses a large range of sources such as:

- when rain or snow moves through the ground and picks up pollutants as it moves towards a major body of water
- the runoff of fertilizers from farm animals and crop land
- air pollutants getting washed or deposited to earth
- storm water drainage from lawns, parking lots, and streets

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Runoff →

المجرى (جري)

للمعونة فقط

Sediment

Soil particles transported from their source

Biochemical Oxygen Demand (BOD)

- Oxygen depleting material
- Leaves
- Organic material

Toxics

- Pesticides
 - Herbicides
 - Fungicides
 - Insecticides
- Metals (naturally occurring in soil, automotive emissions/tires)
 - Lead
 - Zinc
 - Mercury
- Petroleum Hydrocarbons (automotive exhaust and fuel/oil)

Nutrients

- Various types of materials that become dissolved and suspended in water (commonly found in fertilizer and plant material):
 - Nitrogen (N)
 - Phosphorus (P)

Bacteria/ Pathogens

Originating from:

- Pets
- Waterfowl
- Failing septic systems

Thermal Stress

Heated runoff, removal of streamside vegetation

Debris

Litter and illegal dumping

Pollutants Found in Runoff

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Water quality management in rivers

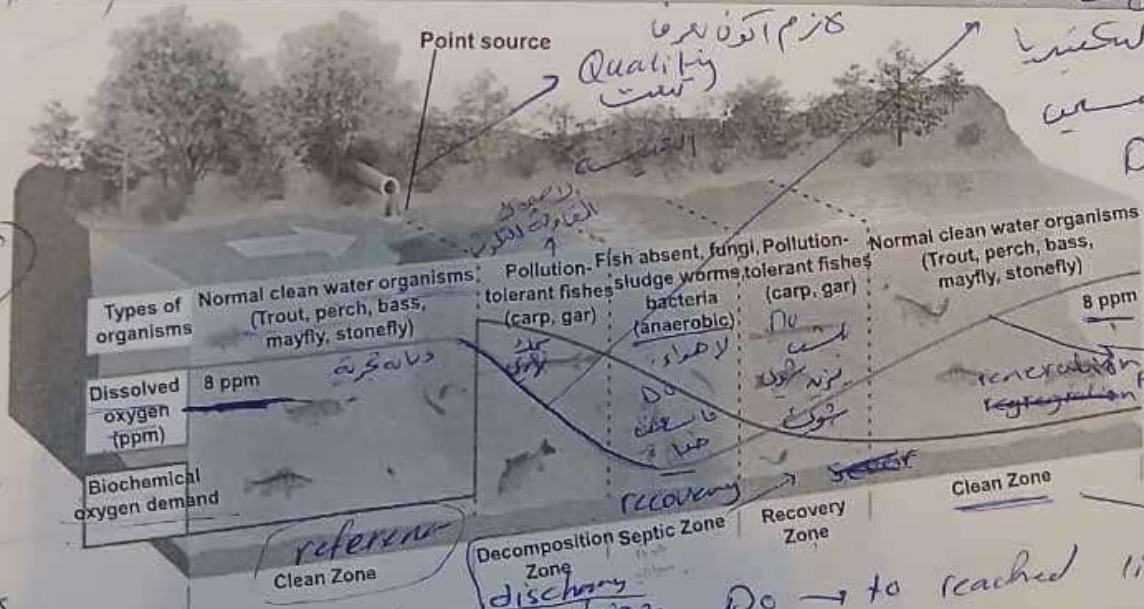
- Main goal is to control the discharge of pollutants so that water quality is not degraded above the natural background level

Controlling waste involves:

- Measuring pollutants levels (x, z, t)
- Predicting their effect on the water quality
- Determining background water quality that would be present without human intervention
- Evaluate the levels acceptable for intended uses of the water

Trout →
 Perch →

distance → 20-40 km



4 → still survive
 < 4 →

BOD →

heater → DO كافت po / Soln + 520
 anoxic → DO بتقدير / شديتة باكونها، لستة

مختلطة
 بيطفئ بين الحما
 8 وقت تسف
 DO

DO (ppm) at 20° C

Good		8-9
Slightly polluted		6.7-8
Moderately polluted		4.5-6.7
Heavily polluted	4-4.5	
Gravely polluted	Below 4	

لازم يكون
 6

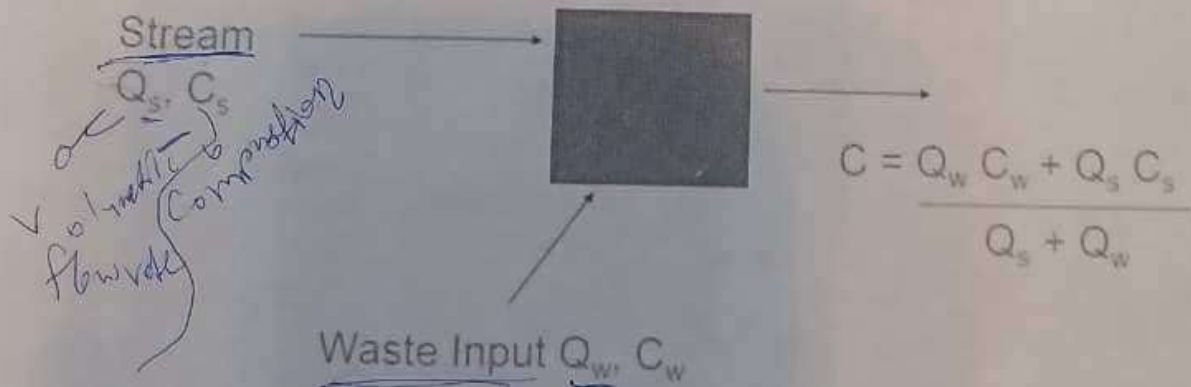
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below
 غاز فستان بلس بزنه
 anaerobic

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عدل الاصفلا
 Input rate - Output rate - decay rate =
 Accumulation rate

Steady state conservative system



Transport characteristics that affect concentration

- Velocity ↓
- Dilution (mixing) ↓
- Dispersion
- Degradation (mass loss)
- Adsorption (to soils)
- Sedimentation (to bottom)
- Aquatic Life (attached)

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Dissolved Oxygen

- If the discharge of oxygen-demanding wastes is within the self-purification capacity, the DO is high

- If the amount of waste increases, it can result in detrimental changes in plant and animal life

- Objective of water quality management is to assess the capability of a stream to absorb waste

- Oxygen use speeds up as the temperature increases and slows down as the temperature decreases

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Do Sag Curve

- DO concentration dips as oxygen-demanding materials are oxidized and then rises as oxygen is replenished from atmosphere and photosynthesis

Major sources of oxygen:

- Reaeration from the atmosphere
- Photosynthesis of aquatic plants

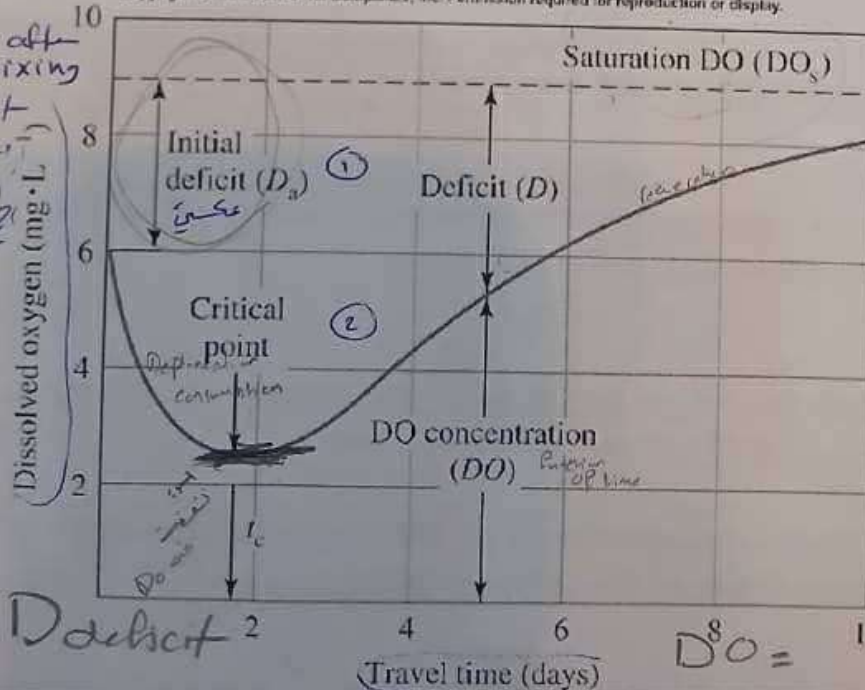
Factors of oxygen depletion:

- BOD of waste discharge
- DO in waste discharge is less than that in the river
- Nonpoint source pollution
- Respiration of organisms and aquatic plants

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(reference)

$$DO = \text{Satu} - D_{\text{deficit}}$$

$$DO_c = \text{Sat} - D_c$$

$$DO = \text{Sat} - D$$

$$DO_c = DO_s - D_c$$

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Quality of water ← لازم أنون لحرف mixing ← ما قبل

Steps in Developing the DO Sag Curve

1. Determine the initial conditions
2. Determine the re-aeration rate from stream geometry
3. Determine the de-oxygenation rate from BOD test and stream geometry
4. Calculate the DO deficit as a function of time
5. Calculate the time and deficit at the critical point

two values

at critical

تناقص يوتي للاكسجين

الدراسة اشياء BOD

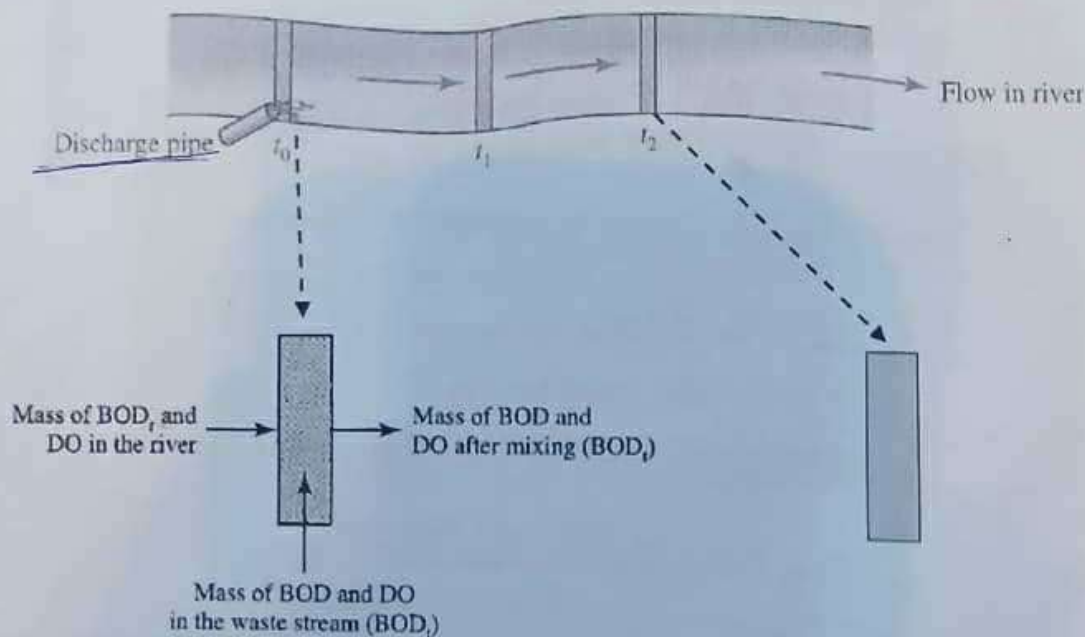
بتسبب اكسجين
تتوزع للاكسجين
أهم الامور

Measure - saturation = deficit

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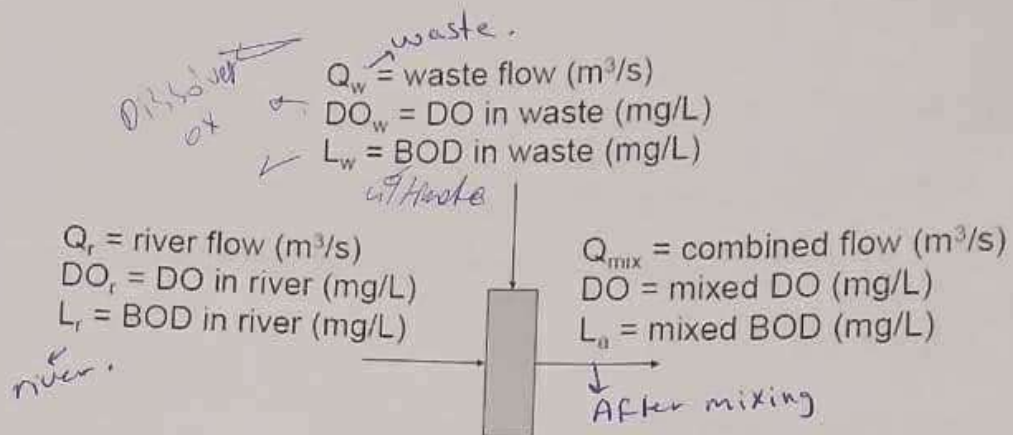
Selecting System Boundaries



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Initial Mixing



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DO

1. Determine Initial Conditions

A. Initial dissolved oxygen concentration

$$DO = \frac{Q_w DO_w + Q_r DO_r}{Q_w + Q_r}$$

B. Initial dissolved oxygen deficit

$$D = DO_s - DO$$

where D = DO deficit (mg/L)

DO_s = saturation DO conc. (mg/L)

critical point - saturation

$$DO = DO_s - \frac{Q_w DO_w + Q_r DO_r}{Q_{mix}}$$

After mixing

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DO → and 1 DO_s
Theoretical DO_s is in
table 2.10

$D_{o, sat}$ is a function of temperature. Values can be found tabulated in books.

C. Initial ultimate BOD concentration

$$L_o = \frac{Q_u L_u + Q_r L_r}{Q_u + Q_r}$$

Reaeration $\rightarrow$ function of geometry and flow

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- Reaeration $\rightarrow$ D_o يرفع $\uparrow$
- de-oxygenation $\rightarrow$ D_o ينزل $\downarrow$

2. Determine Reaeration Rate

A. O'Connor-Dobbins correlation:

$$k_r = \frac{3.9u^{1/2}}{h^{3/2}}$$

where k_r = reaeration coefficient @ 20°C (d⁻¹)

u = average stream velocity (m/s)

h = average stream depth (m)

أقصى عمق،
نطاق معين

B. Correct rate coefficient for stream temperature

$$k_r = k_{r,20} \Theta^{T-20}$$

where $\Theta = 1.024$

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Table 4-1. Reaeration Constants

Type of watercourse	k'_2 at 20°C ^a (days ⁻¹)
Small ponds or backwaters	0.10–0.23
Sluggish streams	0.23–0.35
Large streams, low velocity	0.35–0.46
Large streams, normal velocity	0.46–0.69
Swift streams	0.69–1.15
Rapids	>1.15

^aFor temperatures other than 20°C, $k'_2(T) = k'_2(20^\circ\text{C})(1.024)^{T-20}$.

$$k'_2 = \frac{3.9v^{1/2} \sqrt{(1.037)^{(T-20)}}}{H^{3/2}}$$

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element here play the kinetic rate test.

3. Determine De-oxygenation Rate BOD_{test}

- a. Rate of de-oxygenation = $k_d L_t$ calculate k_d by BOD test
 where k_d = deoxygenation rate coefficient (d⁻¹)
 L_t = ultimate BOD remaining at time (of travel downstream) t .

b. If $k_d(\text{stream}) = k(\text{BOD test})$

order of reaction.
 zero order very fast.

and

$$L_t = L_0 e^{-k_d t}$$

rate of deoxygenation = $k_d L_0 e^{-k_d t}$

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c. However, $k = k_d$ only for deep, slow moving streams. For others, $k_d = k + \frac{u}{h} \eta$

geometry play
rule here
also.

So D well
test.

$$k_d = k + \frac{u}{h} \eta$$

where η = bed activity coefficient (0.1 - 0.6)

Soil -
up
Civ 100 1000

d. Correct for temperature

$$k_r = k_{r,20} \Theta^{T-20}$$

where $\Theta = 1.135$ (from 4 to 20 °C) or

$\Theta = 1.056$ (from 20 to 30 °C)

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we have two mechanism
→ de-oxygenation,
- reoxygenation.

4. DO as function of time

- Mass balance on moving element

$$\frac{dD}{dt} = k_d L_t - k_r D$$

oxygenation. → reoxygenation

- Solution of differential equation is

Numerical
Soln.

$$D_t = \frac{k_d L_a}{k_r - k_d} (e^{-k_d t} - e^{-k_r t}) + D_a (e^{-k_r t})$$

deficit

Value
After.

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5. Calculate Critical time and Deficit

$$t_c = \frac{1}{k_r - k_d} \ln \left[\frac{k_r}{k_d} \left(1 - D_a \frac{k_r - k_d}{k_d L_a} \right) \right]$$

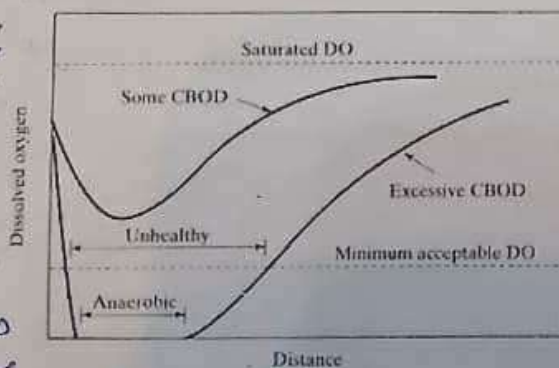
$$D_c = \frac{k_d L_a}{k_r - k_d} (e^{-k_d t_c} - e^{-k_r t_c}) + D_a e^{-k_r t_c}$$

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← لا نصل minimum
← بعد هذا وقت طويل حتى يتم recovery

More Oxygen Sag Curves

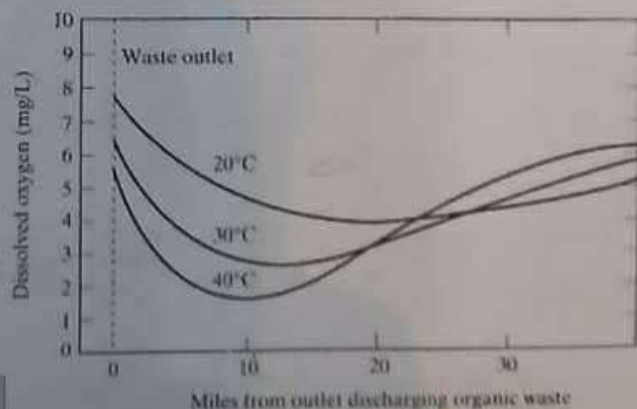


Effect of BOD Level:

- sag becomes more severe
- longer distance (or time) at unhealthy DO levels

Effect of temperature:-

- sag deepens and shortens
- may cause a portion of river to have unhealthy DO levels



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وقت التنبؤ

الوقت، المسافة

time, unit, Temperature

Example

Calculate the DO at a point 1.609 km downstream from a waste discharge point for the following conditions. Report answers to two decimal places. Rate constants are to be temperature adjusted.

velocity
+
distance
= time

Parameters	Stream
K_d (at 20°C)	1.911 (d ⁻¹)
K_r (at 20°C)	4.49 d ⁻¹
Flow	2.4 m ³ /s
Speed	0.100 m/s
DO _a (after mixing)	0.00
Temperature (°C)	17
BOD _L (after mixing)	1100 kg/d

time = distance / velocity

سؤال مسن

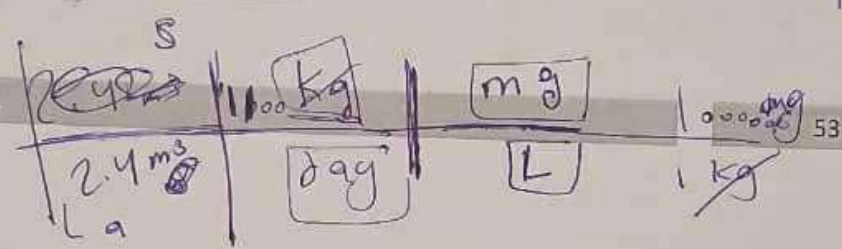
mg/L

La

1 mg · 10⁻³ g

criticle.
 $DO_c = \frac{K_r}{K_d} L_a e^{-k_d t}$; $k = k_{20} \theta^{(T-20)}$; $\theta = 1.047$

الوقت
20
24 x 3600 M.saidan



Example

24 hours
3600 s
1 hour

سؤال
مس
مس
مس

a. Calculate travel time

distance / velocity

per day $t = \frac{(1.609 \text{ km}) (1000 \text{ m/km})}{(0.100 \text{ m/s}) (86400 \text{ s/d})} = 0.186 \text{ d}$

$\frac{1.609 \text{ Km}}{0.1 \text{ m/s}} \times \frac{1000 \text{ m}}{1 \text{ km}} \times \frac{1 \text{ s}}{1 \text{ day}}$

b. Convert kg/d to mg/L

BOD_L

$L_a = \frac{(1100 \text{ kg/d}) (10^6 \text{ mg/kg})}{(2.40 \text{ m}^3/\text{s}) (86400 \text{ s/d})} = 5.31 \text{ mg/L}$

$\frac{16090 \text{ s}}{24 \times 3600 \text{ s}}$

c. Calculate deficit

-kt
Dae

$D = \frac{(1.911)(5.31)}{4.49 - 1.911} (e^{-(1.911)(0.186)} - e^{-(4.49)(0.186)}) + 0$

$D = 3.93(0.7009 - 0.4338) = 1.046 \text{ mg/L}$

مس
مس
مس

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deficit

Saturation - DO
= Deficit

DOE $\rightarrow$ critical dissolve oxygen : is the actual amount of oxygen remaining in the water at its lowest point.

Example

DOc $\rightarrow$ critical dissolve oxygen
 $\rightarrow$ is the actual amount of oxygen remaining in the water at its lowest point

Dc $\rightarrow$ critical oxygen deficit

$\rightarrow$ is the amount of oxygen that has been consumed at the same point

$$DO = DO_s - D$$

Appendix at 17 °C find $DO_s = 9.74 \text{ mg/L}$

$$0.46 = 8.69 \text{ mg/L}$$

نقص
الأكسجين
في
الماء

نقص الأكسجين في الماء
نقص الأكسجين في الماء
نقص الأكسجين في الماء
نقص الأكسجين في الماء

نقص الأكسجين في الماء

Appendix ??

$$DO_c = \frac{kD}{kD} L_n e^{-kD}$$

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$$= 8.7 \text{ mg/L}$$

HW

The town of ABC has filed a complaint that the City of XYZ is restricting its use of the Alpha River because of the discharge of raw sewage. The water quality criterion for the Alpha River is 5.00 mg/L of DO. The ABC Town is 15.55 km downstream from XYZ City.

- What is the DO at City ABC?
- What is the critical DO and
- Where (at what distance) downstream does the critical DO occur?
- Is the assimilative capacity of the river restricted?

The following data pertain to the 7-day, 10-day low flow at XYZ.

Parameter	XYZ Wastewater	Alpha River	Units
Flow	0.1507	1.08	m ³ /s
BOD ₅	128		mg/L
BOD _L		11.4	mg/L
DO	1.00	7.95	mg/L
k at 20°C	0.4375		day ⁻¹
Velocity		0.390	m/s
Depth		2.8	m
Temperature	16	16	°C
Bed-activity coeff		0.200	

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سؤال امتحان

نقص

deficit
saturation = 0

نقص الأكسجين في الماء
نقص الأكسجين في الماء
نقص الأكسجين في الماء
نقص الأكسجين في الماء

نقص الأكسجين في الماء
نقص الأكسجين في الماء
نقص الأكسجين في الماء
نقص الأكسجين في الماء

سببه وجود أشجار في الماء

NPK
Nitrogen → Nitrate
phosphorus → phosphate

Lake Eutrophication

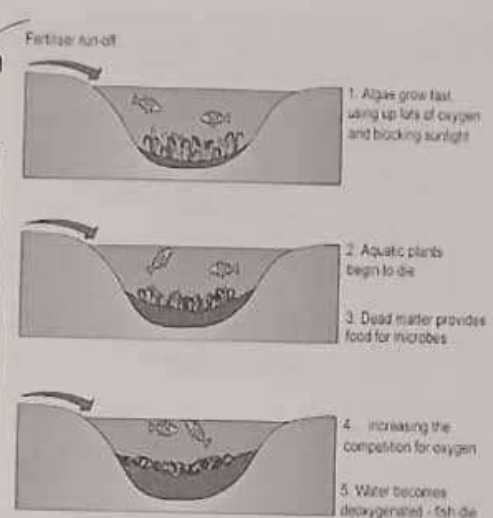
بحيرة التخت

- As lakes age, they become more Productive وقان
- Natural processes: natural Eutrophication
- Pollutant loading: cultural Eutrophication

تزيد البكتيريا
بالفقدان
خلال جريان
الفقدان
موت البكتيريا
يؤدي إلى انخفاض
مستوى الأكسجين
الذائب

Eutrophication: nutrient enrichment of lakes
mostly from runoff of plant nutrients (nitrates
and phosphates)
During hot dry weather can lead to algae
blooms
Decrease of photosynthesis
Dying algae then drops DO levels
Fish kills, bad odor

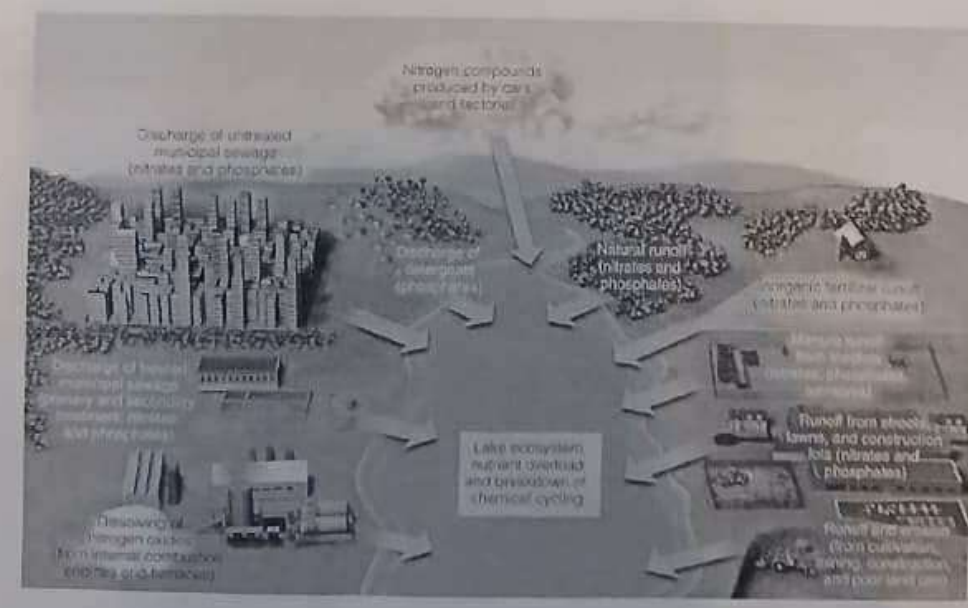
الزراعة والحدائق



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من مصادر مياه العادمة

Nutrient loading and Eutrophication

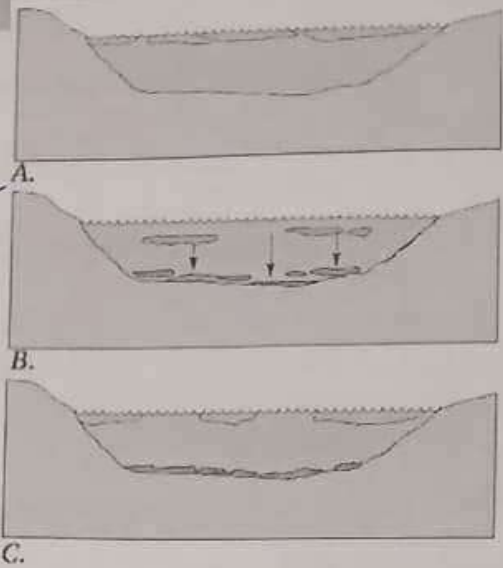


© 2005 Brooks/Cole - Thomson

Eutrophication in Lakes

Solutions:

- 4. Advanced sewage treatment (N, P)
- Household detergents ✓
- Soil conservation ✓
- Remove excess weed build up
- Pump in oxygen or freshwater

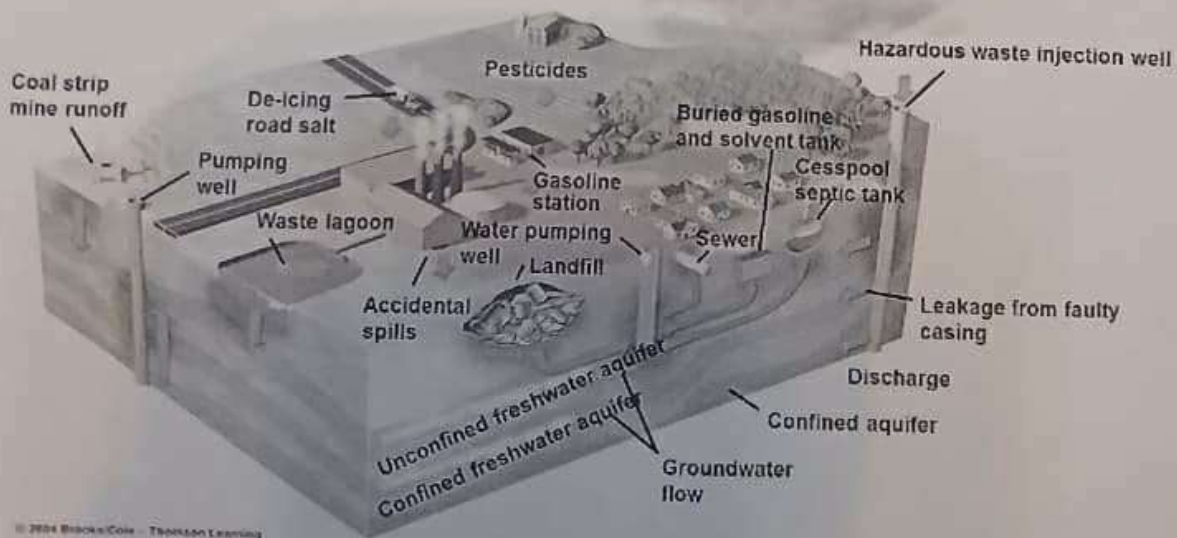


Schematic Diagram showing effects of algal bloom on water quality. (A) Abundant growth of algae in sunlit shallow water when nutrients are abundant. (B) In colder weather, algae die and sink to the lake bottom. (C) The next growing season, more algae thrive at the surface while older material decays at the bottom, increasing BOD and releasing more nutrients to fuel new growth.

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Groundwater Pollution



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1cc 5

Water Treatment

wast. treat. met.
water

صيانة، المعالجة

315

Dr. Motasem Saidan

m.saidan@gmail.com

Univ. of Jordan/ Chem. Eng. Dept.

1

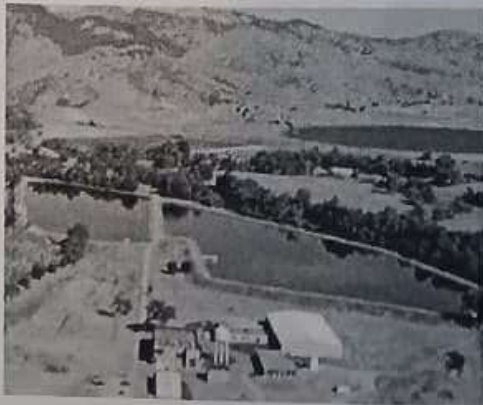
This means that the outcome, quality or characteristics of water can change depending on where the water comes from - either surface water or ground water.

surface water: water that is found above ground in lake, river and reservoir. It is often exposed to pollution from the environment.

Drinking Water Treatment

ground water: water located below the earth's surface, stored in soil or rock layers (aquifers). It is usually cleaner but may contain minerals or be harder to access.

- It depends on the source: Surface or Ground Water



Coagulation : تتم فيه إضافة مادة كيميائية (مثل البصم) أو لوريد الحديد (يك) إلى الماء (مضان) لاجل إزالة الجسيمات العالقة في الماء. مما يؤدي إلى عدم اعتقادها بغيرها في كتلات صغيرة microflocs.

Flocculation : تكون كتلات صغيرة يتم تعريضها لمزيج بطيء من الكتلات لتتجمع مع بعضها البعض فتتكون كتل أكبر وكثافة أكبر بالمثل لتتجمع في قاع الخزان لتعمل كما ذببت في الماء ويتم تجميعها بسهولة

Coagulation
Coagulation
mat

Treatment technologies largely based on coagulation and flocculation



floc or flocc
كتل أو كتل

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هو تجميع الجسيمات العالقة في الماء وتجميعها في كتل أكبر وتسمى flocc

mixing rate → معدل الخلط
Coag. Flocc. chemical
وحدة
الخلط
تكونوا مع
نوعها

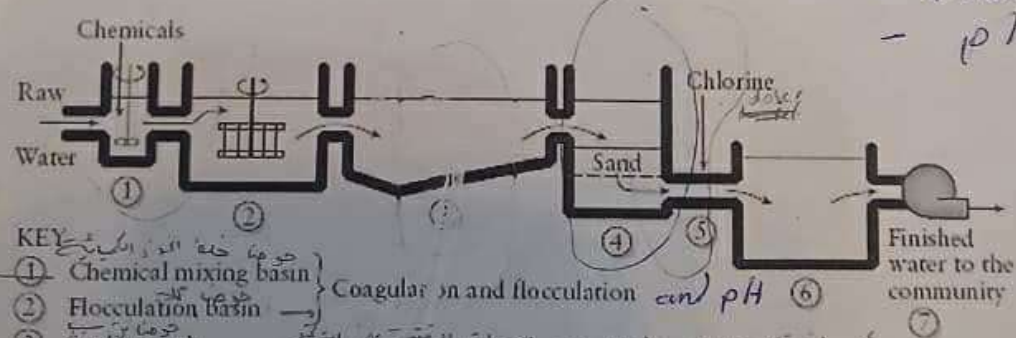
Chemical Addition
- Coagulation
- flocculation
- pH

Coagulation
flocculation

بصم الجسيمات
أكبر الجسيمات العالقة
floc
كتل

Alum

بصم بالماء
لأنه يكون بهن زبد

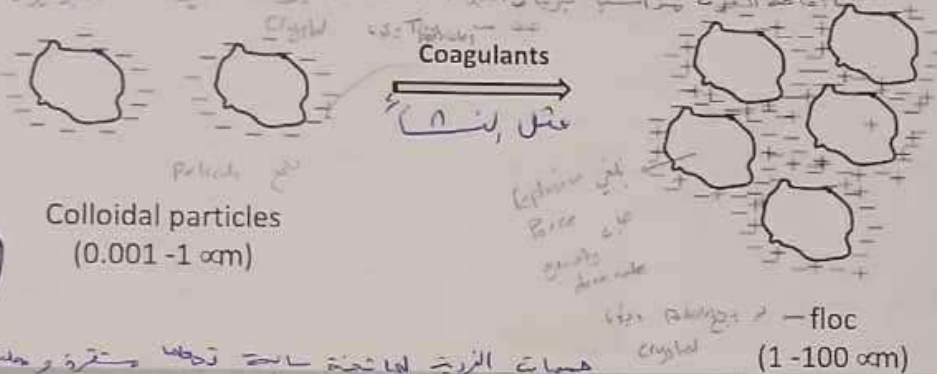


- KEY
- ① Chemical mixing basin
 - ② Flocculation basin
 - ③ Settling tank
 - ④ Rapid sand filter
 - ⑤ Disinfection with chlorine
 - ⑥ Clean water storage basin (clear well)
 - ⑦ Pump

Figure 10.11 A typical water treatment plant.

Coagulation and Flocculation

- Coagulation is the chemical alteration of the colloidal particles to make them stick together to form larger particles called **flocs** that readily settle.
- Tiny (colloidal) clay and silt particles: negative surface charge
- Particles repel so suspension is considered stable
- Coagulants (alum (aluminum sulfate), and coagulant aids, such as lime and polymers) are added to the water to neutralize the charge on the particles.



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polymer
- bio polymer
الاصطناعي

الهدف من عملية Coagulation هو كسر الشحنة على الجزيئات
 ② المساعدة على جعل الجزيئات الصغيرة "لزجة" من تلقاء نفسها، لتكون جزيئات أكبر
 ③ making the tiny particles "sticky" so they can coalesce and form large, quick settling particles
 والسبب انه تنجس الماء من الملوثات العالقة

Particulate size (meters)

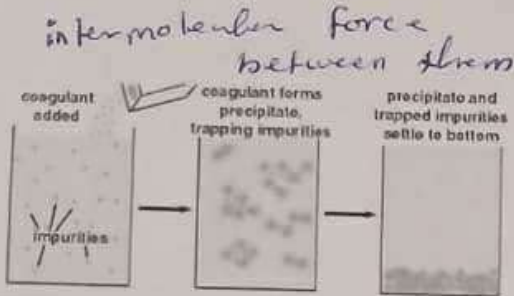
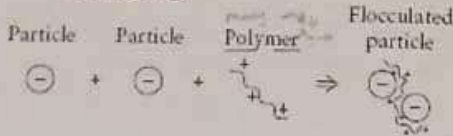
	10 ⁻¹⁰	10 ⁻⁹	10 ⁻⁸	10 ⁻⁷	10 ⁻⁶	10 ⁻⁵	10 ⁻⁴	10 ⁻³
Electromag Waves	X-rays	UV	Visible	Near Infrared	Far Infrared	Micro waves		
Technical Definitions	Molecules	Colloids			Suspended Solids			
Typical Particulates in Water		Viruses	Humic Acids	Bacteria	Algae	Sand		
Methods for Size Analysis		Electron Microscope		Optical Microscope	Coulter Counter	Sieving		
Potential Removal Systems	Reverse Osmosis	Ultra Filtration		Coagulation/Granular Media Filtration	Gravity Separation			

Coagulation: charge neutralization and bridging

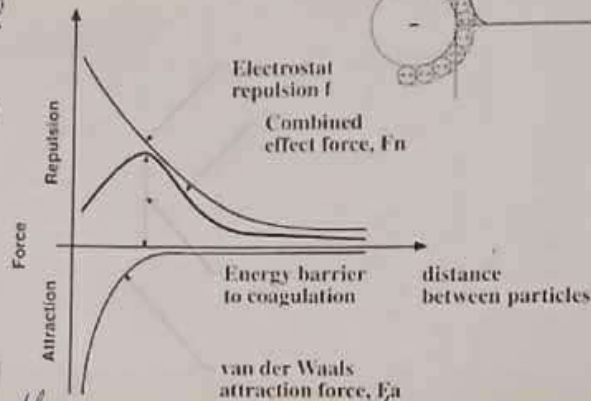
Charge neutralization:

will cause instability and collision

Bridging



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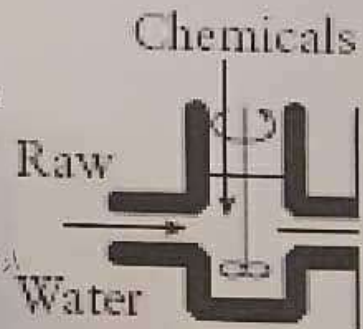


The specific combination of these compound is dependent on:

- pH of the water
- The temperature
- method of mixing

Rapid mixing

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



✓ neutralize surface charges

✓ collapse the surface layer around the particles

✓ allow the particles to come together and agglomerate

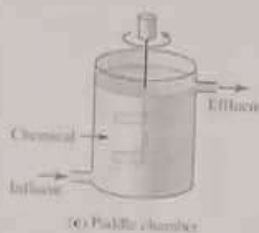
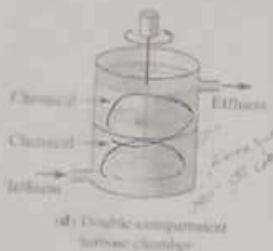
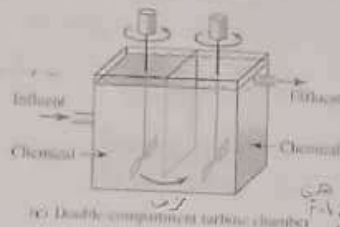
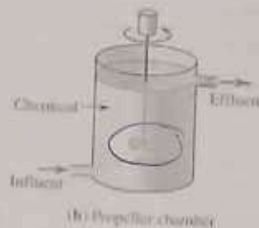
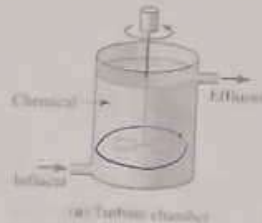
✓ allow formation of floc that can readily settle

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The Alkalinity also measured (Alam) because these salts react with the Alkalinity in the water and reducing its buffering capacity

الماء مشان اخله تغير لونه
 Outflow → (زقزقة)
 Flow → يا هو وقت اهلون كس يطلع

Resist
 resistance time
 geometry
 الهم من تغير



ما برسيب
 انا بكتيها
 و بعد الفلتر
 مخرج
 number

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Coagulants: Non-toxic and relatively inexpensive

Alum

✓ Hydrated aluminum sulfate $[Al_2(SO_4)_3 \cdot 18H_2O]$

✓ Alum, when added to water, will be hydrolyzed to form gelatinous hydroxide $[Al(OH)_3]$ precipitate. This will carry suspended solids as it settles by gravity.

Anhydrous Fe^{3+}

➤ Forms $Fe(OH)_3(s)$ in a wide range of pH 4-11

Anhydrous Fe^{2+} ($FeSO_4 \cdot 7H_2O$)

➤ Must be oxidized to Fe^{3+} first at pH higher than 8.5

Coagulant Aids (Polyelectrolytes)

➤ Natural: Starch, cellulose derivatives, and gums (polysaccharides)

➤ Synthetic polymers

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Flocculation

- It is the agglomeration of destabilized particles into a large size particles known as flocs which can be effectively removed by sedimentation or flotation.
- It needs gentle mixing.
- To enhance the agglomeration, organic polymers (flocculent) are added.

The chain is long enough to allow active groups to bond to multiple colloids

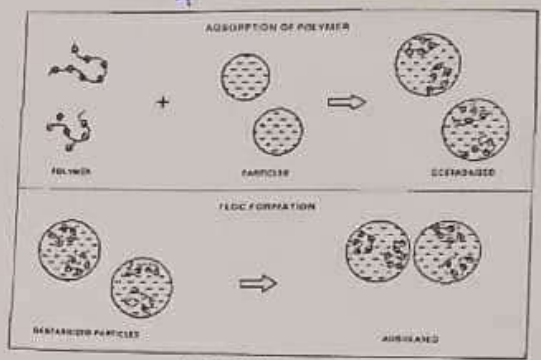
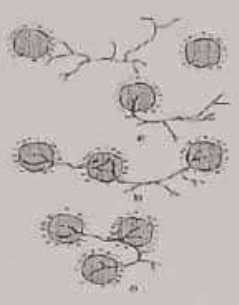
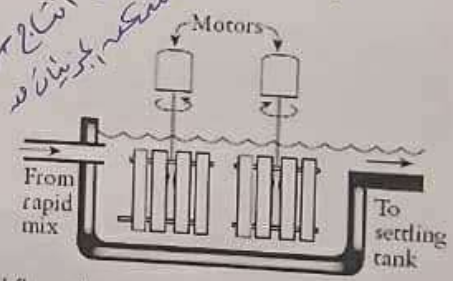


Figure 1.9 Flocculation process

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- The intent of the process of flocculation is to produce differential velocities within the water so that the particles can come into contact.

الهدف من العملية هو إنتاج اختلافات في السرعات داخل الماء حتى تتمكن الجسيمات من التماسك مع بعضها البعض.

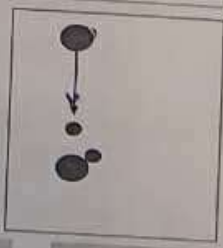
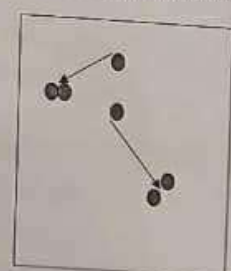


Typical flocculator used in water treatment.

Transport Mechanisms:

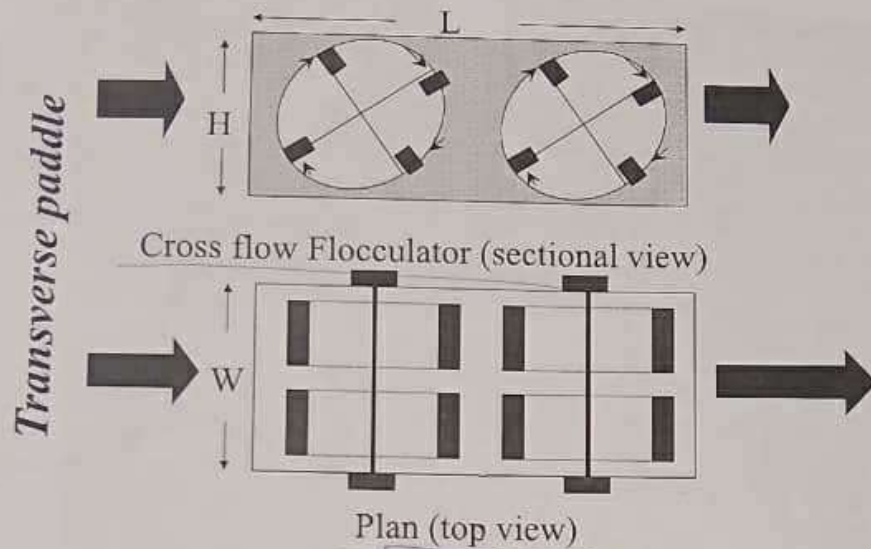
Brownian motion: for relatively small particles which follow random motion and collide with other particles (perikinetic motion)

Differential settling: Particles with different settling velocities in the (vertical alignment) collide when one overtakes the other (orthokinetic motion)



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Mechanical Flocculator



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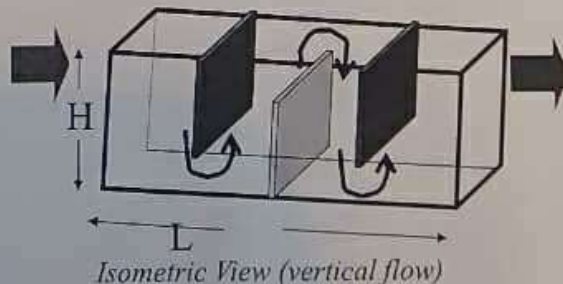
Hydraulic Flocculation

Small Volume
resists fine
particles
settling down

- **Horizontally baffled tank**
The water flows horizontally. The baffle walls help to create turbulence and thus facilitate mixing

- **Vertically baffled tank**

The water flows vertically. The baffle walls help to create turbulence and thus facilitate mixing



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Flocculator perforated wall

← المياه تدخل عبر الهاي الشقوق →



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للسؤال صيد

Determining Coagulant Dose

حساسية مخبرية لتدوير

Jar Tests

pH اختبار درجة لونية

Coagulant Dose

اختبار جرعة

تستخدم جرعة الماء بحدود 100-200 mg/L. الاختبار يتم في مختبر لتحديد الجرعة المناسبة للمياه الخام. ☐ The jar test – a laboratory procedure to determine the optimum pH and the optimum coagulant dose

مع تثبيت parameters وتعديل الثاني
درجتين يتدرج نعل، لكس

☐ A jar test simulates the coagulation and flocculation processes

Determination of optimum pH

الحال particles
مغلقة او ذائبة
لازم نفس
Volume

- Fill the jars with raw water sample (500 or 1000 mL) – usually 6 jars
- Adjust pH of the jars while mixing using H_2SO_4 or NaOH/lime (pH: 5.0; 5.5; 6.0; 6.5; 7.0; 7.5) (5-8.5)
- Add same dose of the selected coagulant (alum or iron) to each jar (Coagulant dose: 5 or 10 mg/L)



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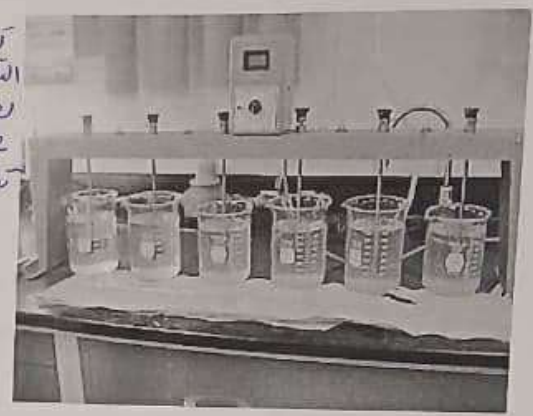
هذا على كل 1000 ml
5-10 mg/L
لازم انفس
تغير لوني

chemicals

لحمية
Sample

Jar Tests – determining optimum pH

- Rapid mix each jar at 100 to 150 rpm for 1 minute. The rapid mix helps to disperse the coagulant throughout each container
- Reduce the stirring speed to 25 to 30 rpm and continue mixing for 15 to 20 mins
This slower mixing speed helps promote floc formation by enhancing particle collisions, which lead to larger flocs
- 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 pH



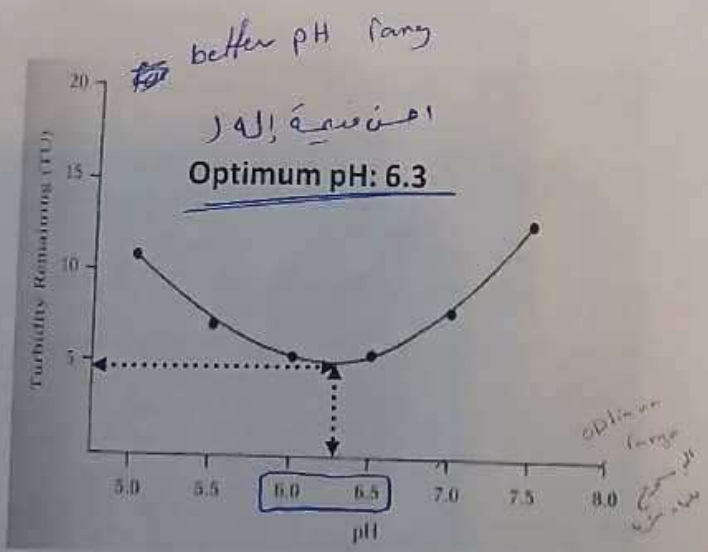
turbidity ↓
effect
less turbidity
so coagulation
floc
بالتالي شغل

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Jar Tests – optimum pH

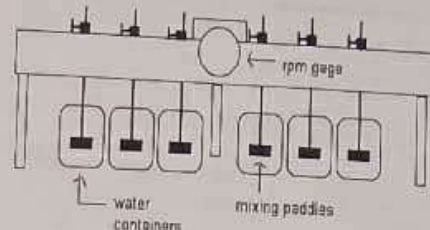
➤ The pH with the lowest residual turbidity will be the optimum pH

إذا قل يكون فإله تأثير
وإذا زاد يكون إله تأثير



Jar Test: Optimum coagulant dose

- ☐ Repeat all the previous steps
- ☐ This time adjust pH of all jars at optimum (6.3) found from first test while mixing using H_2SO_4 or NaOH/lime
- ☐ Add different doses of the selected coagulant (alum or iron) to each jar (Coagulant dose: 5; 7; 10; 12; 15; 20 mg/L)



- ☐ Rapid mix each jar at 100 to 150 rpm for 1 minute. The rapid mix helps to disperse the coagulant throughout each container
- ☐ Reduce the stirring speed to 25 to 30 rpm for 15 to 20 mins

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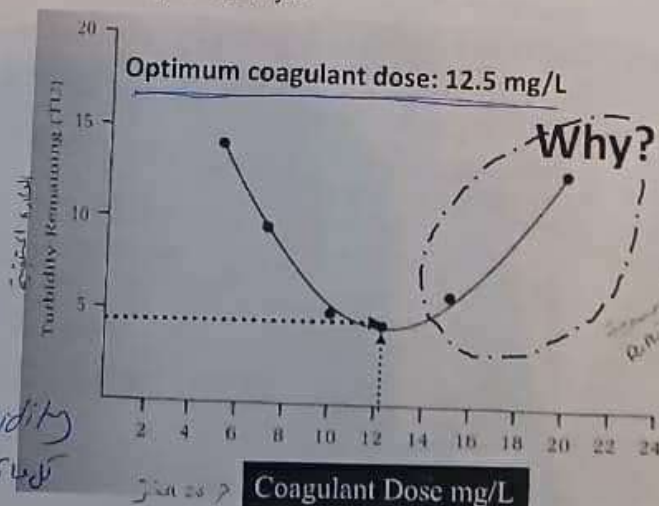
الته coagulation على يكون ~~مستقر~~ ليحل توازن لا
 particle surface charge
 + كاتيونات - صالبة
 + كاتيونات - صالبة

Optimum coagulant dose

لماذا غير

- ☐ Turn off the mixers and allow flocs to settle for 30 to 45 mins
- ☐ Then measure the final residual turbidity in each jar

- ☐ Plot residual turbidity against coagulant dose



➤ The coagulant dose with the lowest residual turbidity will be the optimum coagulant dose
 lowest point is the best because they less turbidity
 كاتيونات - صالبة

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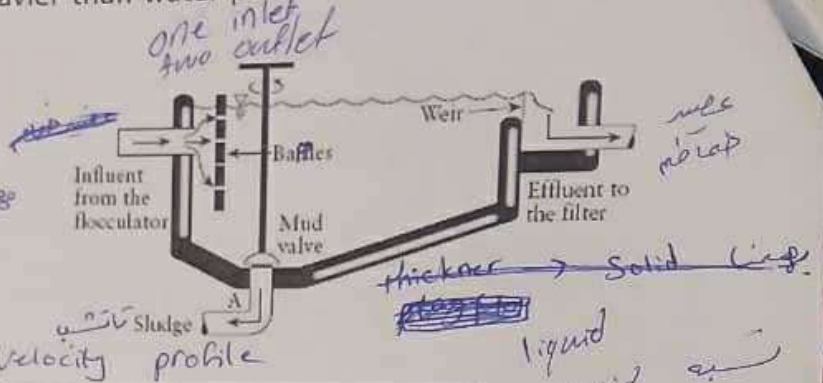
separation between liquid (carrier fluid) and suspension inside.

Sedimentation

- Separation of formed flocs from water in gravity settling tanks that simply allow the heavier-than-water particles to settle to the bottom.

Settling depends on:

- particle size (volume)
- particle shape
- particle density
- fluid density
- fluid viscosity.



Particle Diameter (mm)	Typical Particle	Settling Velocity (m/s)
1.0	Sand	2×10^{-1}
0.1	Fine sand	1×10^{-2}
0.01	Silt	1×10^{-4}
0.001	Clay	1×10^{-6}

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Ideal settling tank

- visualized hydraulically as perfect plug flow reactors and the outlet should to deal with it

Overflow rate:

horizontal velocity

overflow velocity

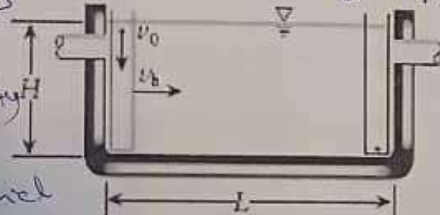
$v_0 = \frac{Q}{A_s} = \frac{m^3/s}{m^2}$

$v_0 = \frac{H}{t}$

$t = \frac{H}{v_0}$

depth

velocity of particle to reach the bottom



two velocity

horizontal

with flow

Examples: 10.16 10.17 10.18

retention time

$$U = HLW$$

$$\bar{t} = \frac{U}{Q} = \frac{HLW}{(HW)v} = \frac{L}{v}$$

$$U = HWL$$

$$A_s = WL$$

$$U = A_s H$$

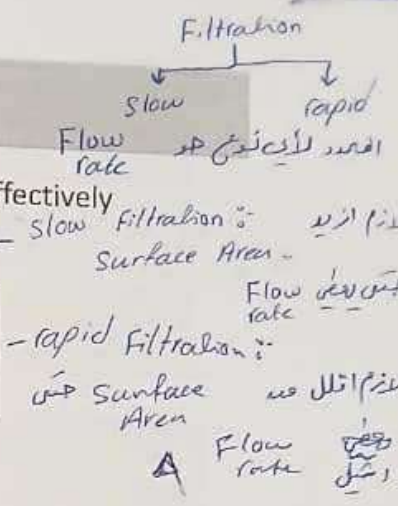
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porosity = هي التي تتغلغل tiny particles
التي ماضية

Filtration

انواع خلاص رديف و ملاصق



is a most Removal of those particles that are too small to be effectively removed during sedimentation

لنستخرج الزاوية
شكل وصال است
الشعر حسب

- > Sedimentation effluent: 1-10 JTU
- > Desired effluent level: <0.3 JTU

- Slow sand filters
- Rapid sand filters

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الاسم هو... نبتة في البركة... من تنظيف سطح المياه... افاعة... بجمع الماء... سطح البركة... النظام... (skimmer)

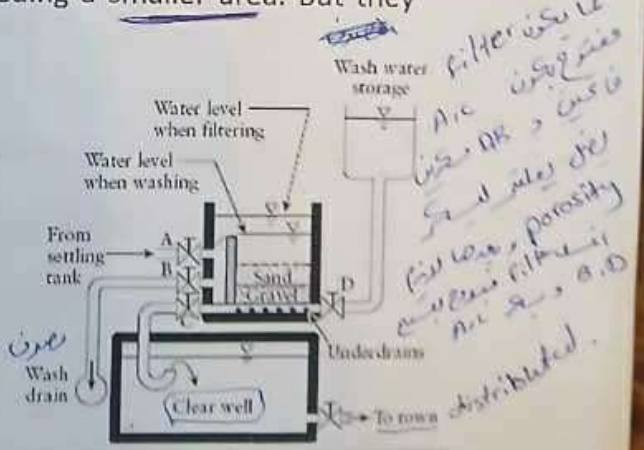
Rapid sand filter

Filter
stand by
Filter
porosity
dp
flow
particles
main purpose

The operation of a rapid sand filter involves two phases: filtration and washing.

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

✓ rapid sand filter becomes clogged, resulting greater head loss through the filter, so it must be cleaned. This cleaning is performed hydraulically by a process called backwashing (10 - 15min).



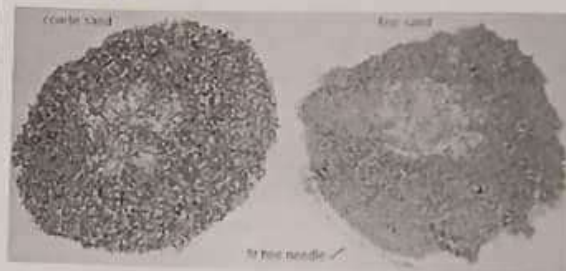
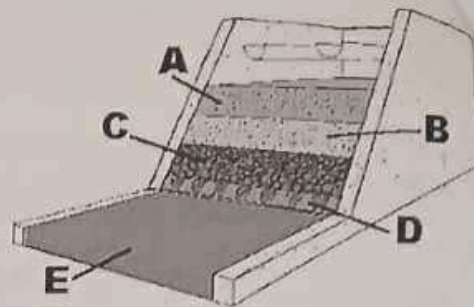
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الاسم هو... bed expansion... turbulent... sand & gravel... medium... (porosity, liquid, A, B, C, D)

Rapid Filter Media

- Single media: sand
- Dual media: anthracite coal and sand
- Multimedia: anthracite coal, sand and garnet



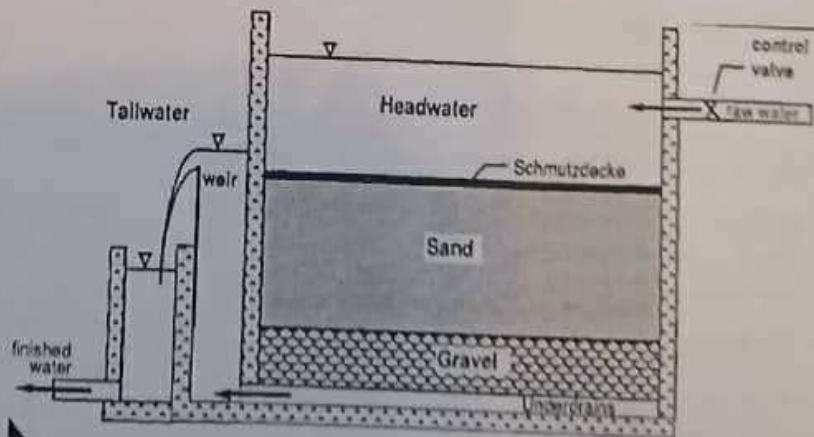
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Slow sand filter

Filter
Surface area
much larger than
rapid filter.

- Slow-sand filters:** ($d_p=0.15-0.35\text{mm}$) require a much larger area. The top 1 inch must be periodically scraped off and the filter occasionally back-washed.

طريقة الترشيح البطيء
البيكتيريا او الغالبية
die/m → لا تتكاثر
في الطبقة السطحية



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Biological Layer: The Schmutzdecke

- Due to the organic and biological matter in the water to be filtered, a bio-layer grows on top of the sand layer.
- The bio-layer contributes to water treatment by consuming organic contaminants including bacteria and viruses.
- The majority of biological activity occurs in the top 20 cm of the filter.



Biofilm Development



Layer of Schmutzdecke

29

oxidizing agent

كثير من
الميكروبات
والبكتيريا
Boundary
طبقة هلامية

Disinfection

- water completely free of suspended sediment is treated with a powerful oxidizing agent, usually chlorine, chlorine then ammonia (chloramine), or ozone. $O_3 \rightarrow$ very oxidizing agent but the life time is short.
- Disinfection is typically the last step in a water / wastewater treatment system.
- A residual 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.

Disinfectants

Gaseous Cl_2

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

Chlorine dioxide (ClO_2):

- No disinfection by-products such as trihalomethanes

• $\text{Ca}(\text{ClO})_2$:

- Safer than Cl_2

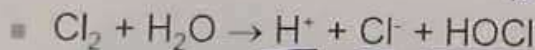
• Ozone: generated on site

• UV lamps

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Chemistry of Chlorine in Water



HOCl = Hypochlorous acid

OCl^- = Hypochloride ion

- HOCl is a weak acid with $K_a = 4.5 \times 10^{-4}$ ($\text{HOCl} \rightleftharpoons \text{H}^+ + \text{OCl}^-$)

- HOCl and OCl^- are free available chlorine which are very effective in killing bacteria

- Small amount of ammonium (NH_4^+) in water is desired

- Chloramines: NH_2Cl , NHCl_2 , NCl_3

- Chloramines (combined available chlorine) are weaker disinfectants than free available chlorine but are desired residual chlorine to be retained in water distribution system

- Excessive amount of ammonium (NH_4^+) in water is undesirable because it consume excess demand of Cl_2

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Disinfectant	Minimum Residual* (mg/L)		MRDL** (mg/L)	MRDLG*** (mg/L)
	Leaving Plant	In Distribution System		
Chloramines (as Cl_2)	2.0		4.0	4.0
Combined residual		0.5		
Chlorine (as Cl_2)	2.0		4.0	4.0
Chlorine dioxide (as ClO_2)	2.0		0.8	0.8
Free chlorine residual		0.2		

* From Illinois and Missouri regulations.

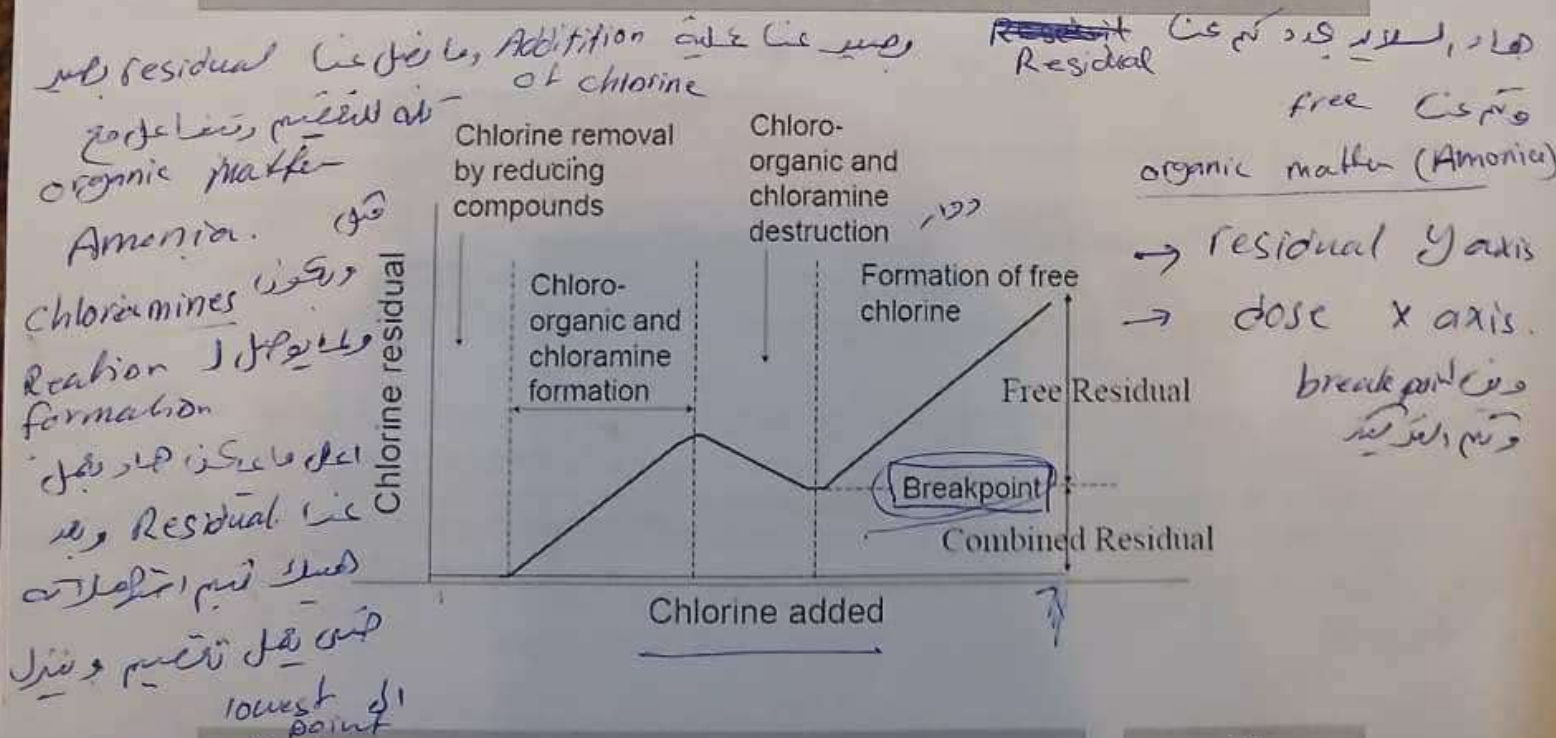
** MRDL = maximum residual disinfectant level.

*** MRDLG = maximum residual disinfectant level goal.

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Chlorine Demand or Breakpoint Chlorination

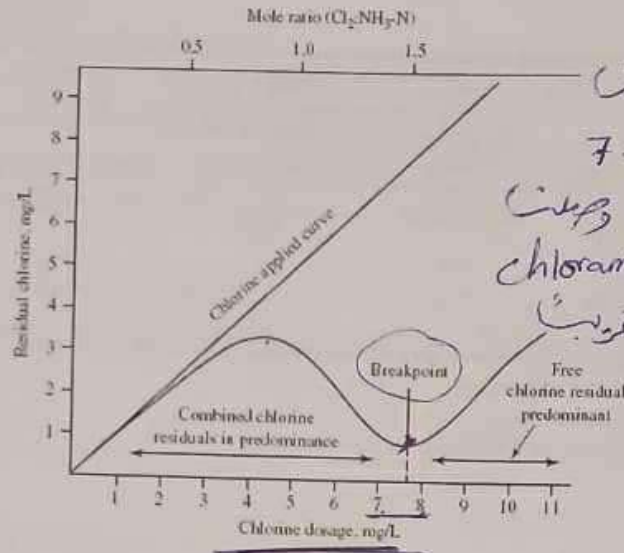


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Final Residual = Free concentration + Combined concentration

~~Free~~ total chloramines → Combined
free → فائز اي واه



* على اي dose يكثر
7-8 ← breakpoint
* على اي dose يكثر
chloramine ← maximum
على 4 و هو تقريبا

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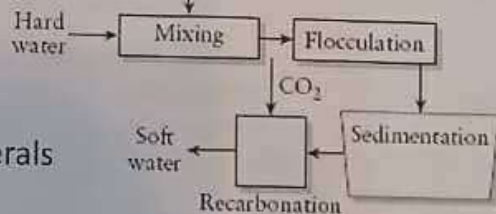
35

Groundwater Treatment

منظور هون كيمياء
TDS ذائبه
Lime and/or soda ash

➤ Primary objectives are to

- ➔ 1. Remove hardness and other minerals
ازالة الصلابة والمعادن الذائبة
- ➔ 2. Eliminate pathogenic organisms
القضاء على الكائنات المرضية



- Treatment technologies largely based on chemical precipitation

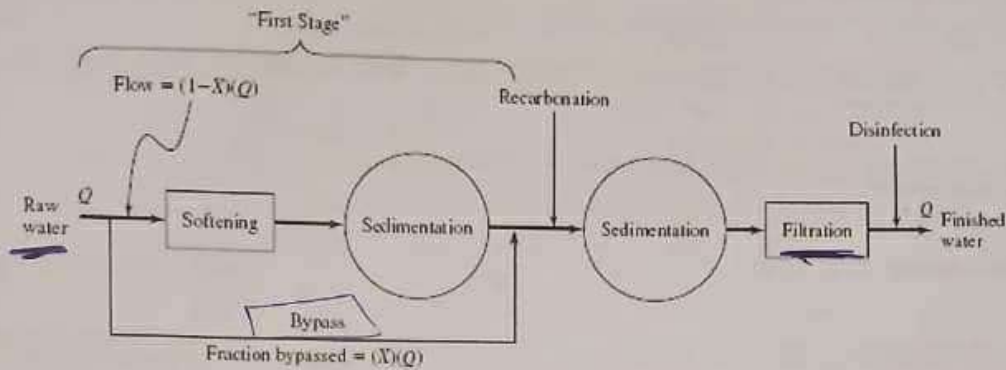
الترسيب الكيميائي

we need chemical reaction

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امنه طريقة نقل فينا PH هو نضج CO₂ inside



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Water Softening

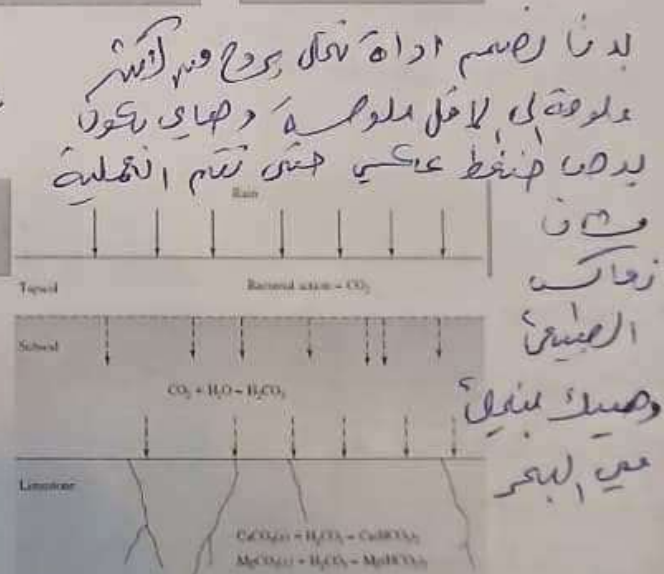
- Groundwater is generally harder than surface water.
- Hard waters produce scale in hot-water pipers, heaters and boilers.
- The removal of ions that cause hardness is called softening.

- Softening can be accomplished by the lime-soda process, ion exchange, nanofiltration, or reverse osmosis.

- Principal cations causing hardness and the major anions associated with them (in decreasing order of abundance in natural waters)

Cations: Ca^{2+} , Mg^{2+} , Sr^{2+} , Fe^{2+} , Mn^{2+}

Anions: HCO_3^- , SO_4^{2-} , Cl^- , NO_3^- , SiO_3^{2-}



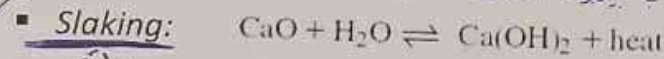
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Water Softening Methods

1. LIME-SODA SOFTENING

عند وجود عسار بكاربونات استخدام الجير فقط
 اللين يسو لفين على
 Quicklime (CaO), commonly called lime, and hydrated lime [Ca(OH)₂]:
 عند غياب بكاربونات استخدام الجير مع صودا كاربونات



Softening Methods;

lime-only process: when Ca^{2+} is present primarily as "bicarbonate hardness"

lime-soda [Ca(OH)₂-Na₂CO₃] process: when bicarbonate is not present at substantial level

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Hardness من الكالسيوم و المغنسيوم
 Calcium Carbonate
 والمغنسيوم هيدروكسيد
 magnesium hydroxide

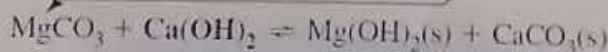
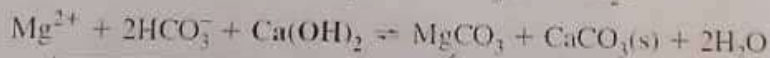
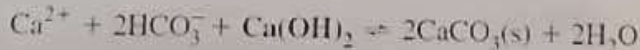
RXNs

كيفية برسم تفاعل كيميائي

Neutralization of carbonic acid



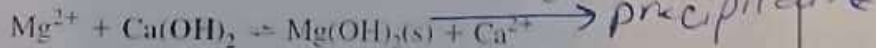
Precipitation of carbonate hardness



Precipitation of noncarbonate hardness due to calcium



Precipitation of noncarbonate hardness due to magnesium



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كربونات و هيدروكسيد
 لينة و صلبة
 في الماء
 لا نرمض في كربونات
 الكالسيوم

كربونات و هيدروكسيد
 الالهة صلبة
 CO₃ + Mg

هناك الطريقة
 كبيرة و صلبة
 large scale
 و صلبة

Ion exchange

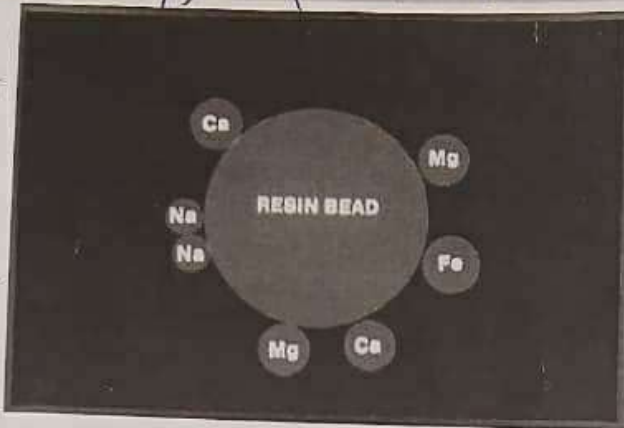
polymer thermoset
مطهرات صلبة مثل البلاستيك
التي لا تتغير عند التسخين
Resin

Thermoset
يصلب إذا سخنت
تصلب
Thermoplastic
التي تليق
اصقة

قادرة على أخذ صلب وتصلب
فإنها لا تتغير عند التسخين
يعني صلابة
fixed bed

المواد بعد softening
سواء صلبة أو سائلة بالصدور
تغير صلبة إلى سائلة
Hardener

وتغير مكانها
موزون
عن صلب المواد
carrier

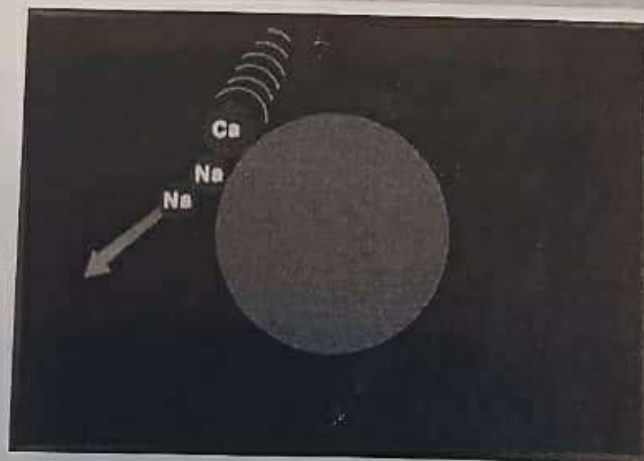


Cation resin bead attract and hold positively charged ions.

Those ions remain on the bead until the bead encounters other ions for which it has a greater affinity.

بفضل هذه
لقد عالجنا الماء
الساخن
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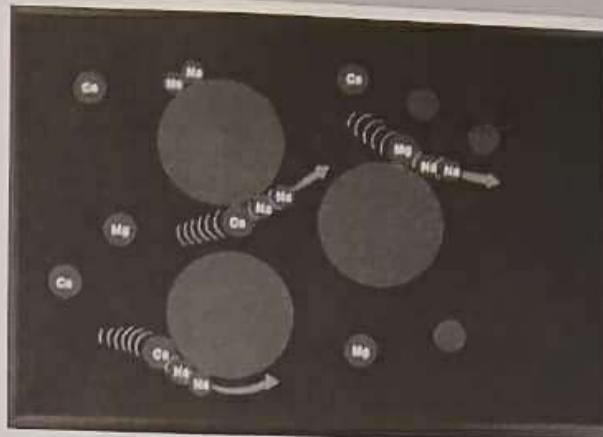


- Resin bead releases Na^+ and absorbs Ca^{+2} and Mg^{+2} ions for which it has a greater attraction.
- Ions are not destroyed or changed chemically; they are simply replaced on the resin bead.
- This process is known as ion exchange.

(Softener)

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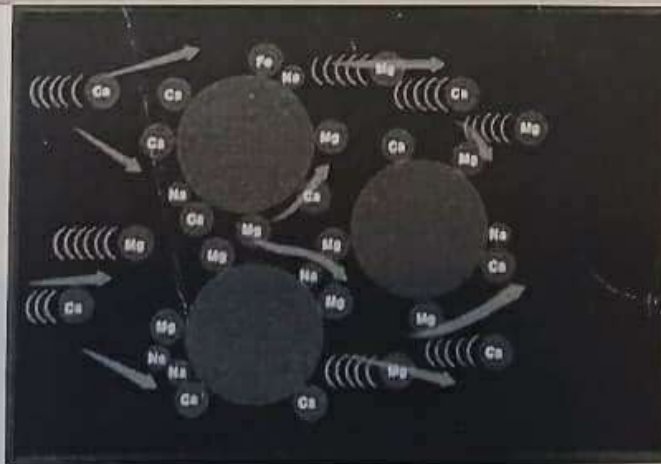
44



- Resin beads attract Ca^{+2} and Mg^{+2} ions and release Na^+
- Water has been softened because the Ca^{+2} and Mg^{+2} concentrations, which cause water hardness have been reduced.

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Resin
← كيم ريف
قادر على تليين
الماء

After a vast number of Ca^{+2} and Mg^{+2} ions have become attached to the resin beads, and most of the Na^+ ions have been released, the resin can no longer soften the water.

- If no new chemical reaction is set, the incoming Ca^{+2} and Mg^{+2} ions flow untouched through the unit because there is no room for them on the resin beads.

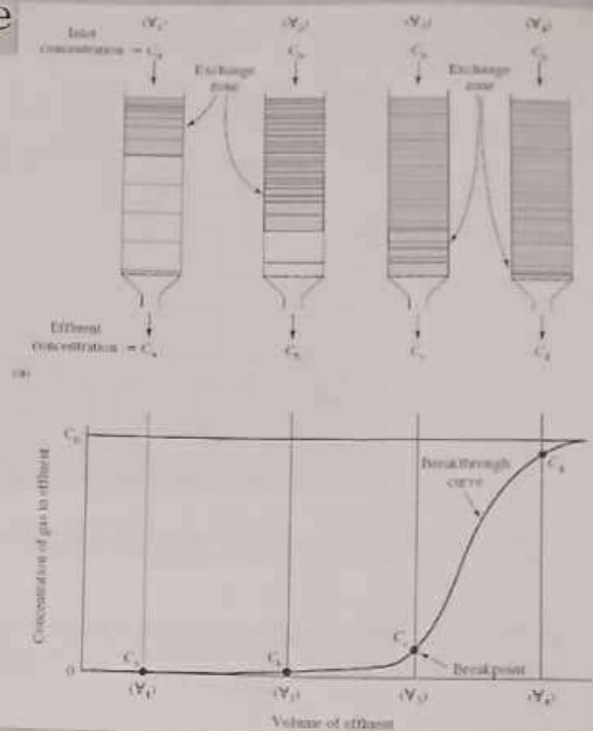
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تدفق على سطح
بالرأس

Breakthrough Curve

- It is more typical at municipal plants for the breakthrough criterion to be set much lower, for example, at 5% to 10% of the influent concentration.



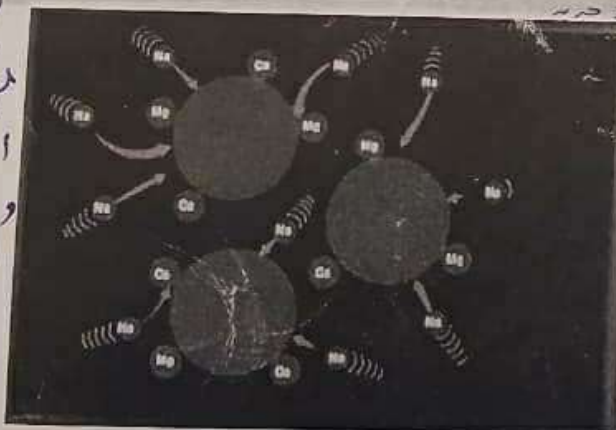
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اي Softner في جنبه tank
full with Sodium (ملح)
و بيكون عباره عن مياه اقل من كبريتات الصوديوم
Saturation

Resin Regeneration

الحاد مشع
(مرد يوم)
مبتدئين
انكاليوم
و هفتيد



- The reaction can be reversed by greatly increasing the concentration of sodium in the solution.
- Reverse process drive the Ca^{+2} and Mg^{+2} ions off the resin beads and replace them with Na^{+} ions.
- Although the resin beads prefer Ca^{+2} and Mg^{+2} ions, the excessive concentration of Na^{+} ions overcomes this affinity.

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← إذا كان قناري قاري فله المياه

والقارة (المياه) القارية

التي هي من مياه البحر

السكني في كثير من الأحيان

المياه في كثير من الأحيان

قارة فقط (البحر)

Wastewater Treatment

Dr. Motasem Saidan

m.saidan@gmail.com

Univ. of Jordan/ Chem. Eng. Dept.

1

Wastewater Treatment

Wastewater:

➤ Usually refer to sewage treatment, or domestic wastewater treatment

➤ process of removing contaminants from wastewater, both runoff and domestic

Wastewater comes from:

- Residences (kitchen, bathroom)
- Commercial institution
- Industrial institution (usually require specialized treatment process)

WW is collected and transported via a network of pipes and pump stations to a municipal treatment plant

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2

عزل Quality لمياه مياه القمامة Typical Wastewater Characteristics

الخصائص
التي
توجد
في
BOD
COD
نقص

أي مادة عادمة
داخلة على محطة
تكون تتسبب في
تلوث مياه الشرب

Parameter	Concentration (mg/L)
Biochemical Oxygen Demand (BOD)	250
Total suspended solids (TSS)	250
Chemical oxygen demand (COD)	500
Nitrogen, Total	40
Ammonia	30
Organic	10
Nitrate	0
Phosphorus, Total	10
Ortho	6
Organic	4
Total organic carbon (TOC)	150
Chloride	50

250-300

500-800

لا يتم فرز النفايات

An estimate of the flow rate : 600L/capita-day

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3

توليد مياه الصرف مع مياه القمامة وتلوث المياه الجوفية، لذلك يجب معالجة مياه القمامة قبل استخدامها.

Purpose:

To control

- To manage water discharged from homes, businesses, and industries to reduce the threat of water pollution.

لتقليل تلوث المياه

- To reduce the amount of pollutants in the water to such a level that the water can be returned to the environment without causing stress on aquatic life and be of sufficient quality for subsequent users.

(مستوى جودة المياه)

كمية الملوثات

تقليل كمية الملوثات

في مياه الشرب

دون أن يتسبب في الإجهاد للحياة المائية ويكون من جودة كافية لاستخدامها

التي يمكن استخدامها

للمياه الجوفية

التي تستخدم في الزراعة

والتي تكون ذات جودة

كافية للاستخدام

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ممكن تكون

Utility level

صالحة للاستخدام

BOD, COD, TSS

4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100

التي تكون ذات جودة كافية لاستخدامها

Types of Treatments

Types of Treatments

- (d) Mechanical treatment primary $\rightarrow$ Remove all of suspended

- Influx (Influent) 320

- Removal of large objects

- Removal of sand and grit

- Primary Sedimentation

- (2) ▪ Biological treatment

- Trickling bed filter

- Activated sludge

- (3) ▪ Chemical treatment Tertiary

- Disinfection

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←، فیهما المی زرع مع فحولة عادم مکنون BOD والی اقل من 25 وحصی مکنون قبل 250

* Activated sludge → المخمرات النشطة
تستخدم في التخلص من المواد العضوية
الذائبة في مياه الصرف الصحي.

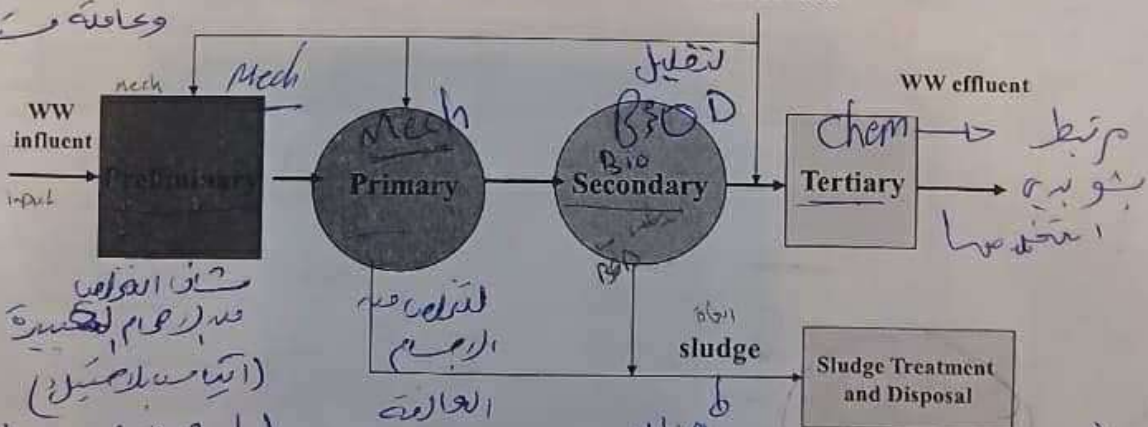
Wastewater treatment stages

Wastewater treatment stages

Bottom

و يتفقون لاكثر من اربعة و شموله m_{max}

Disinfectant



settling
down
for TSS

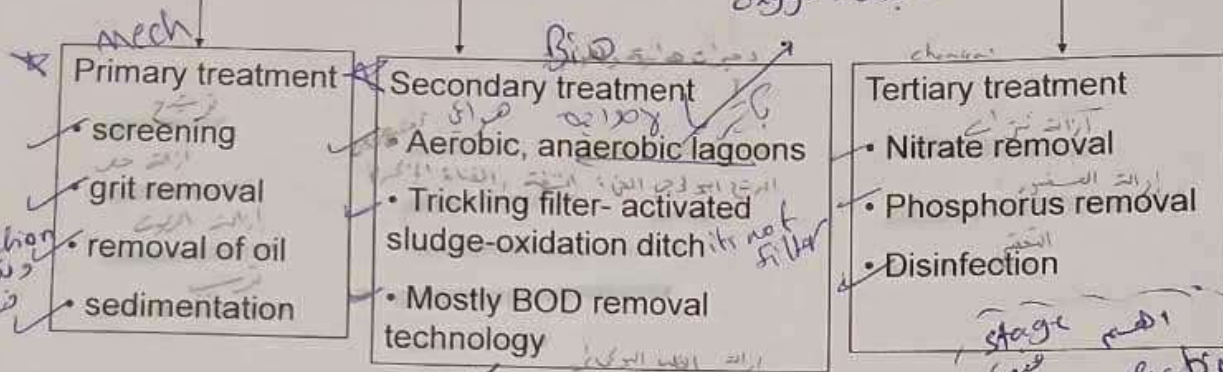
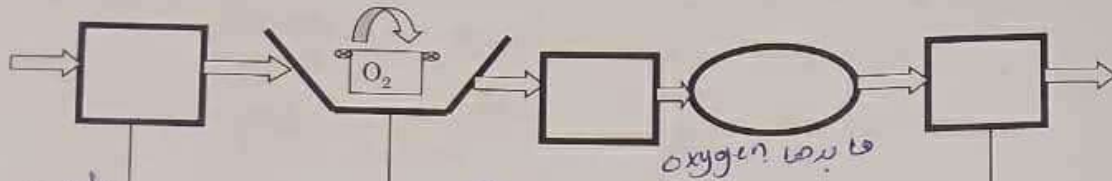
clarifier le μ

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Soln
partially (1/2) study e
20% - 10%
وار

6
مكونه اميد

Wastewater Treatment Processes



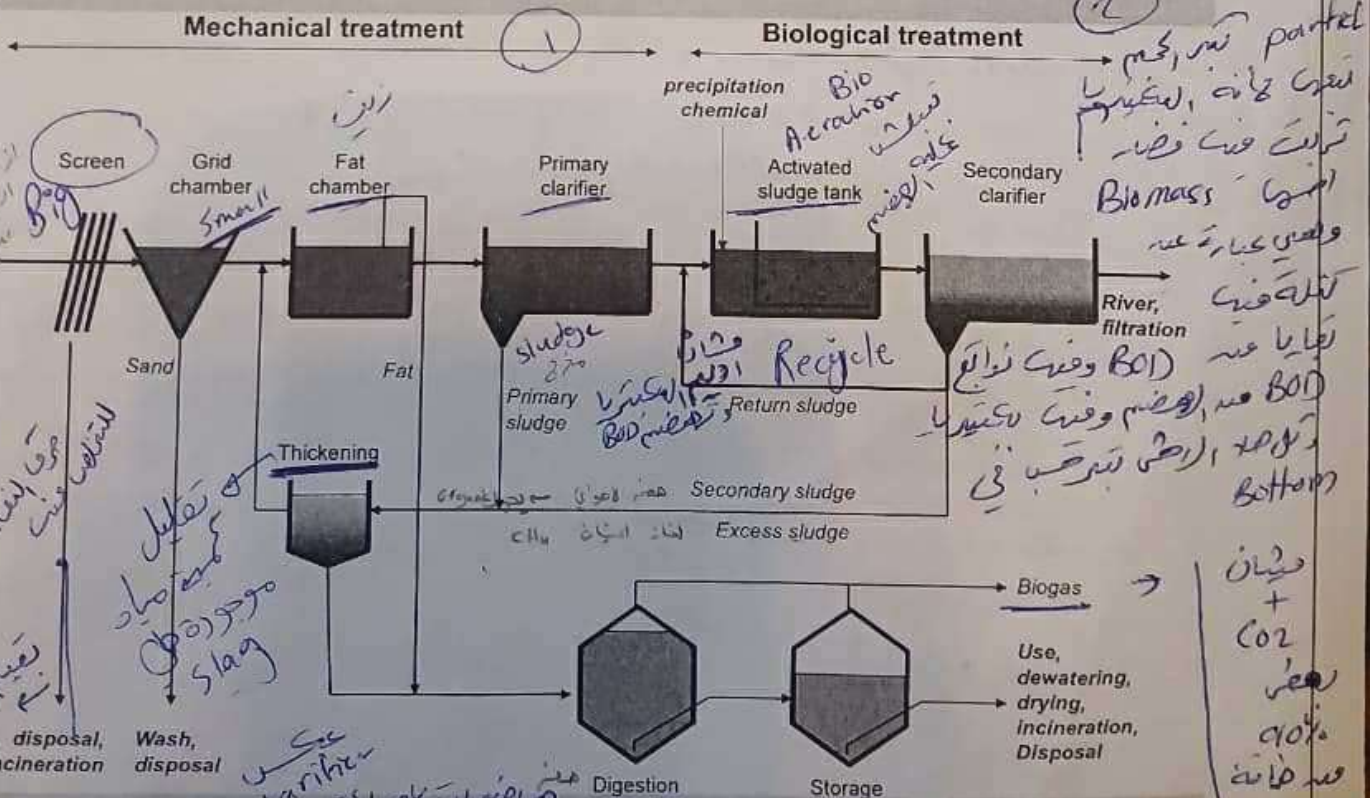
Aerobic $\Rightarrow$ CO_2 H_2O
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Anaerobic $\Rightarrow$ biogas $\Rightarrow$ غاز فيثان
 CO_2

farther treatment Digestion في مرحلة هضم الكتل sludge وادارة

Flowsheet
رسم تخطيطي
مياه

Layout of a WWTP



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Thickener

solid particle في مرحلة الترسيب

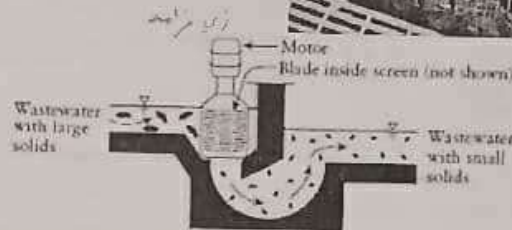
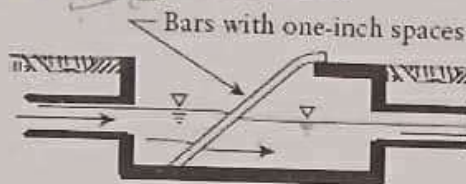
فيثان + CO_2
90%
في فائض
الطاقة

Preliminary Treatment

- removes large objects and non-degradable materials
- protects pumps and equipment from damage
- bar screen and grit chamber

Bar Screen

catches large objects that have gotten into sewer system such as bricks, bottles, pieces of wood, etc



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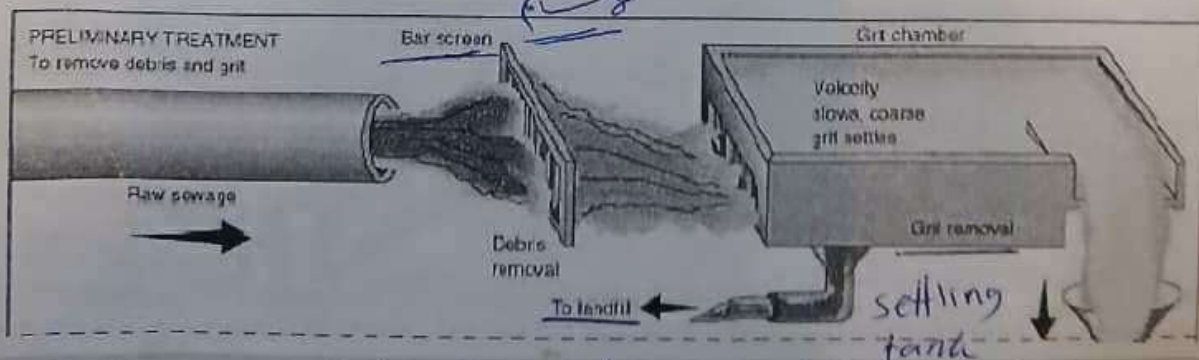
9

Surface hardnesses → (احتمالاً دهریان و طبعی) → این هائی

Preliminary Treatment

- Grit removal chamber : to removes rocks, gravel, broken glass, etc.

- Grit is composed primarily of sand, cinders, and gravel
- Grit causes excessive wear and abrasion in pipes and pumps
- Grit accumulates in downstream tanks where flow velocities are insufficient to keep it in suspension. As grit accumulates, it reduces the effective tank volumes and thus treatment effectiveness
- Grit removal is done by gravity settling (the high specific gravity of grit)



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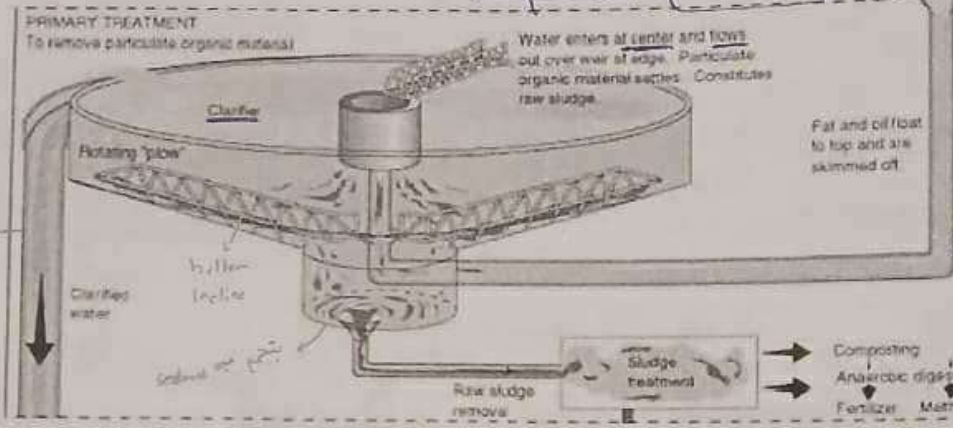
10

Primary Treatment clarifier

- a physical process

- wastewater flow is slowed down and suspended solids settle to the bottom by gravity

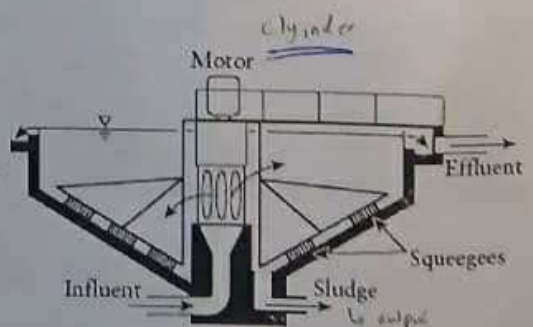
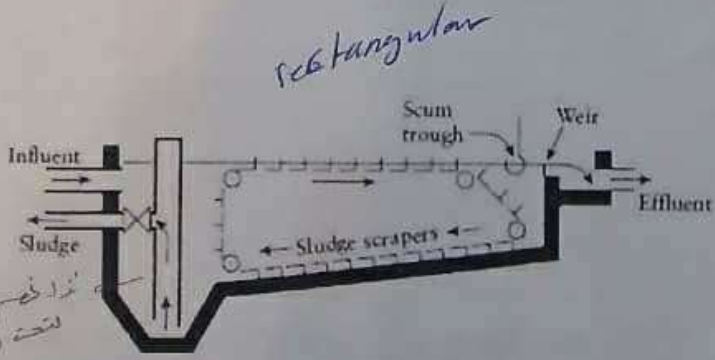
- the material that settles is called sludge or biosolids



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Sedimentation tanks and clarifiers

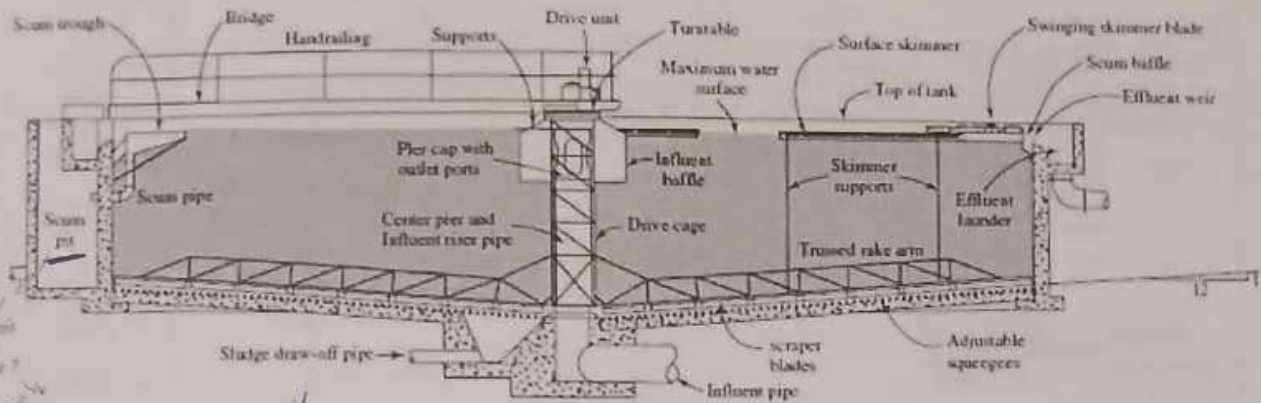
- The settling tank that follows preliminary treatment, such as screening and grit removal, is known as the primary clarifier.
- Primary treatment, in addition to removing about 60% of the solids, removes about 30% of the demand for oxygen and perhaps 20% of the phosphorus (both as a consequence of the removal of raw sludge).



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cylindrical

بالنوع في زي قاشطان مكان لا تظهر لمزيت او المواد او الغشوة



scum pit → آتت فيه الرغوة لا يكون مليان
فلواتت بالاصفر يكونان، له نسبة وزنية

rectangular

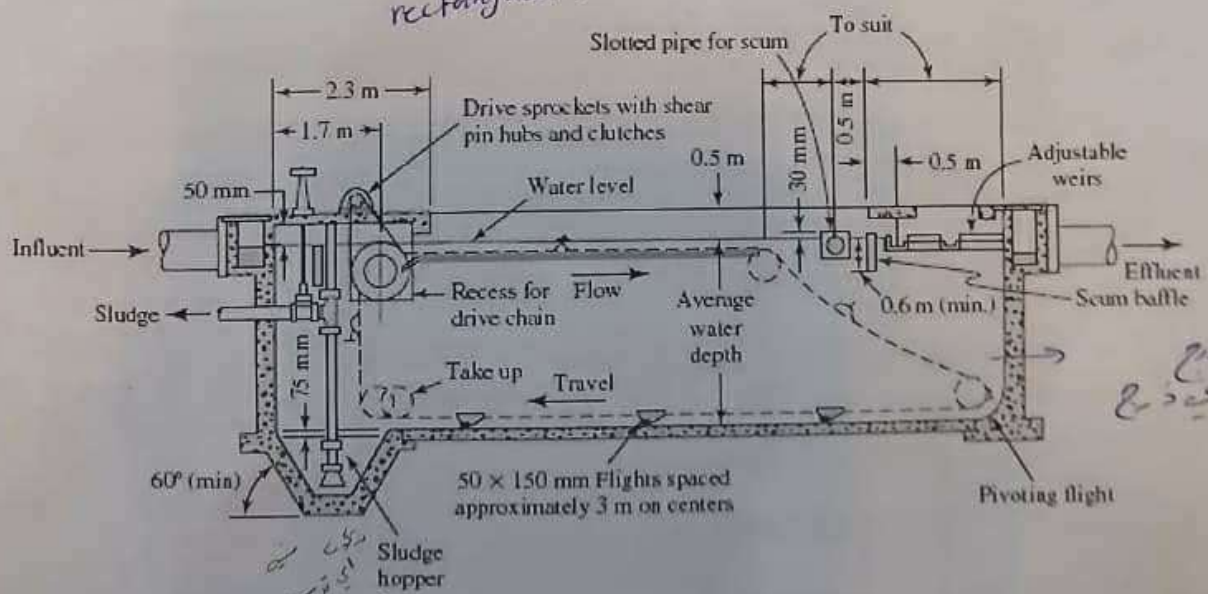
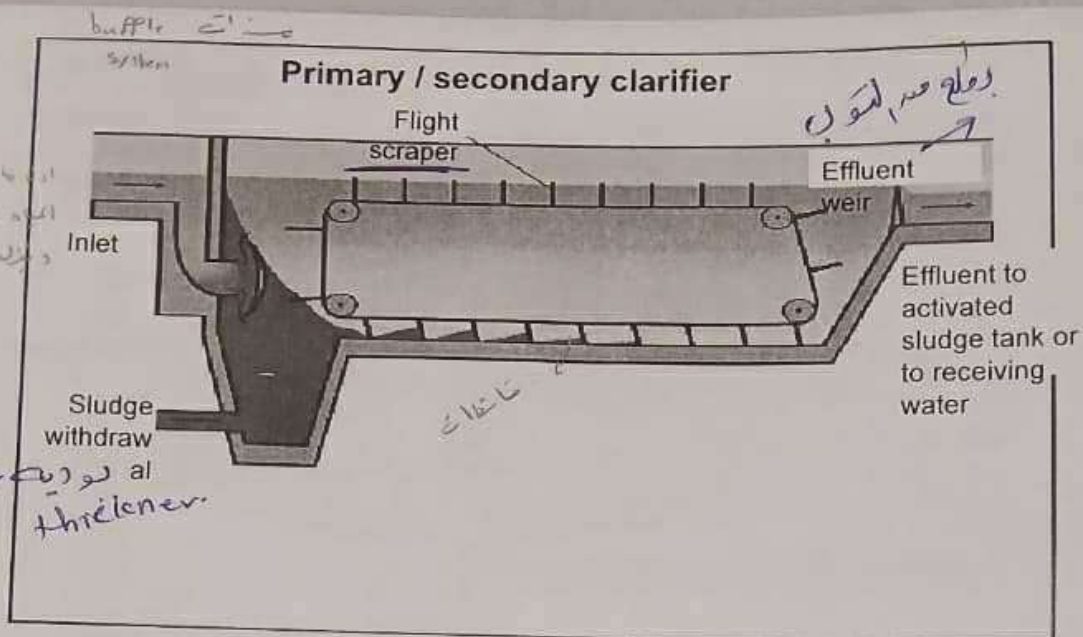


FIGURE 21-3 Rectangular primary settling tank. (Source: Davis and Cornwell, 2008.)



13 Umweltbereich Wasser

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Primary Settling Tank Design

- Size:
 - rectangular: 3-24 m wide x 15-100 m long
 - circular: 3-90 m diameter

- Detention time: 1.5-2.5 hours

- Overflow rate: 25-60 m³/m²·day

- Typical removal efficiencies:

solids: 50-60%

BOD₅: 30-35%

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Effects of primary clarifier on wastewater

Compound	Unit	Inlet	Outlet*	$\eta = \frac{C_{in} - C_{out}}{C_{in}}$
TSS	g TSS / m ³	360	180	0.5
BOD ₅	g O ₂ / m ³	300	230	0.23
COD	g O ₂ / m ³	600	450	0.25
TKN	g N / m ³	60	56	0.067
NH ₄ -N	g N / m ³	40	40	0
NO ₂ -N	g N / m ³	0	0	0
NO ₃ -N	g N / m ³	1	1	0
P _{tot}	g P / m ³	10	9	0.1
Alkalinity	mol HCO ₃ ⁻ / m ³	= f(Drinking water) + NH ₄ -N		

* Short residence time

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Secondary Treatment

A Biological Process

- The objective of secondary treatment is to remove/reduce BOD using

microbial action from soluble to suspended solids.

Substrate

Organic Matter + Bacteria + O₂ → New Cells (Biomass) + H₂O, CO₂, NH₃



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organic Matter → substrate

① organic Matter

② Biomass

③ Sludge

Aerobic + Anaerobic = بدون أكسجين
 BOD بالأكسجين ينتجها عليه وينتجها
 Aerobic = CO_2 H_2O ← BOD
 Anaerobic = ← BOD غازات كريهة

Secondary clarification
 - يتم فيه تسخين قاع الحمأة ويطرد على
 - يتم فيه تسخين قاع الحمأة ويطرد على
 - يتم فيه تسخين قاع الحمأة ويطرد على
 - يتم فيه تسخين قاع الحمأة ويطرد على

Trickling filter -
 Fixed bed
 هو عبارة عن
 حوائط في
 media وها
 أو media من
 أو الرمال أو الزجاج أو البلاستيك
 عبارة عن طرقات، حوائط
 موزعة على سطحها
 يكون زبد جل ولا يمر داخل
 الماء عند قاعه
 porosity
 media و BOD موجود في
 ولا يطفئ فيه طبقة رقيقة
 رقيقة شفافة

methods

aerobic and anaerobic

biological treatment is a
 which the pollutants in
 water (organic matter)
 and by microorganisms

in the presence of molecular
 oxygen

Anaerobic: biological treatment is
 a process in which the pollutants
 in the waste water (organic
 matter) are stabilized by
 microorganisms in the absence of
 molecular oxygen

suspension while converting
 matter to gases and cell tissue
 (Activated sludge)

Attached growth is a biological w.w.t
 in which microorganisms responsible
 for the conversion of organic matter
 to gases and cell tissue are attached
 to some material such as rocks,
 sand, or plastic (Trickling filter).

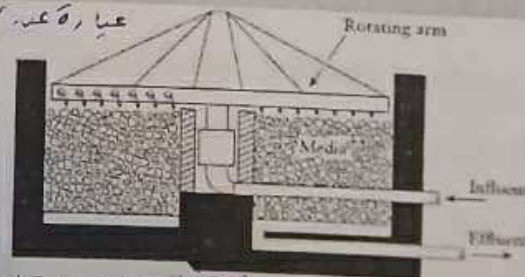
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Secondary Treatment Method

The Trickling filter

porosity
 اللام فيه
 media
 الصخر أو الزجاج
 البلاستيك



- It does not "filter" the water.
- It consists of a bed of media (such as fist-sized rocks or various plastic shapes) over which the waste is trickled.
- An active biological growth forms on the media, and the organisms obtain their food from the waste stream dripping over the bed.
- Air is either forced through the media.
- Wastewater is sprayed and runs over a plastic media and organisms clinging to the media remove organic matter from the wastewater.

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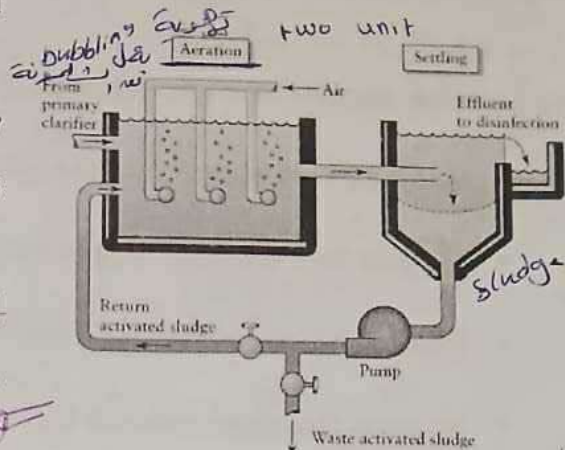
Activated sludge system

➤ Air is bubbled into this tank (called the aeration tank)

➤ The microorganisms use the energy and carbon by decomposing this material to CO_2 and H_2O .

➤ The microorganisms are separated from the liquid in a settling tank, called a secondary or final clarifier

➤ The separated microorganisms exist on the bottom of the final clarifier without additional food and become hungry waiting for more dissolved organic matter. These microorganisms are said to be activated.



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هو sludge الذي رجناه في secondary clarifier الى aeration tank من جديد، يوضع في BOD tank الذي يعمل على ازالة BOD

Final Clarifier

- The activated sludge process is a continuous operation
- one of the end products of this process is excess microorganisms. If the microorganisms are not removed, their concentration eventually increases to the point where the system is clogged with solids.



low flow rate

biomass في

لازم sludge الذي في Bottom اقبه
من ما يتكسر ويحل layers تقلل من
volume flow rate لأنه يده resist time

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لازم اميل sludge الذي في Bottom
كل وقت يمس من زفير volume كافي لئلا يتصل هذا volume clarifier

Basic Ingredients & Conditions

BOD
microorganisms

الكائنات
الحيوية

لازم يكون عدد الكائنات الحية عاليًا لكي يكون النظام فعالًا

الحياة على الكائنات الحية في النظام

بubbling
من فقاعات

- High density of microorganisms (keep organisms in system)

- Good contact between organisms and wastes (provide mixing)

bubbling
probability
احتمالية

- Provide high levels of oxygen (aeration)

تتلاقى BOD بكميات كبيرة
على الأغذية

- Favorable temperature, pH, nutrients (design and operation)

- No toxic chemicals present (control industrial inputs)

تدخل كيميائيات ضارة

أكثر تآكلية للبكتيريا فتتلف
وتتلف أكثر على الأغذية

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Biological Growth Type

نوع النمو البيولوجي

- Suspended Growth - Activated Sludge

Secondary Biomass
Clarifier

- Bacteria are kept in suspension by mixing

- Sludge age is controlled by reactor volume or biomass recycle following sedimentation

- Attached Growth - Biofilter (Trickling Filters, Rotating Biological Contactors)

- Bacteria are attached on supporting media or entrapped between supporting media

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Dispersed growth vs. Fixed Growth

Dispersed Growth - suspended organisms

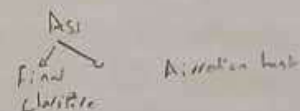
- ✓ Activated sludge
- ✓ Oxidation ditches/ponds
- ✓ Aerated lagoons, stabilization ponds

Fixed Growth - attached organisms

- ✓ Trickling filters
- ✓ Rotating Biological Contactors (RBCs)

Design of Activated Sludge Systems

- Wastewater is aerated in a tank
- Bacteria are encouraged to grow by providing
 - ✓ Oxygen
 - ✓ Food (BOD)
 - ✓ Nutrients
 - ✓ Correct temperature
 - ✓ Time



- As bacteria consume BOD, they grow and multiply
- Treated wastewater flows into secondary clarifier
- Bacterial cells settle, removed from clarifier as sludge
- Part of sludge is recycled back to activated sludge tank, to maintain bacteria population
- Remainder of sludge is wasted

The microbial growth curve

Biochemical reaction



Nutrients such as carbon, nitrogen, phosphorus, sulfur, and the elements required for the synthesis of proteins, nucleic acids, and other structural parts of the cells

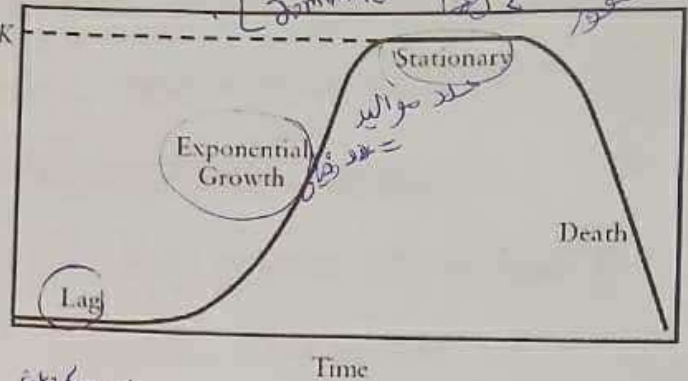
The number of microorganisms is proportional to the growth rate:

$$\frac{dX}{dt} = \mu X$$

$$X = X_0 e^{\mu t}$$

where X = number of microorganisms
 μ = instantaneous, specific growth rate

Number of Organisms (X)



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تقسم عملية تكاثر البكتيريا بعد ظروف الزرع إلى أربع مراحل (تأخر، نمو متسارع، ثبات، وموت) وتسمى كل مرحلة لتتبع فلتس (جدولتين) وتسمى كذا

Bacteria reproduce

fission (each cell divides into two new cells), the increase in cell number follows in geometrical progression

1 → 2 → 4 → 8 → 16

1 → 2 → 4 → 8 → 16

The log growth phase

substrate becomes

toxic byproducts

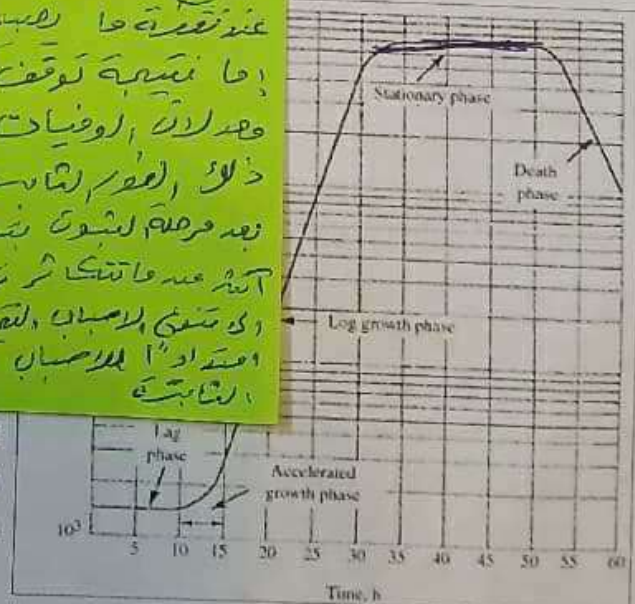
some point the population

constant either as a result

of fission or a balance in death and reproduction rates. This is depicted by the stationary phase on the growth curve.

Following the stationary phase, the bacteria begin to die faster than they reproduce. This death phase is due to a variety of causes that are basically an extension of those that lead to the stationary phase.

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The Monod Equation

$$\mu = \frac{\mu_{max} S}{K_s + S} = \frac{1}{\theta_c}$$

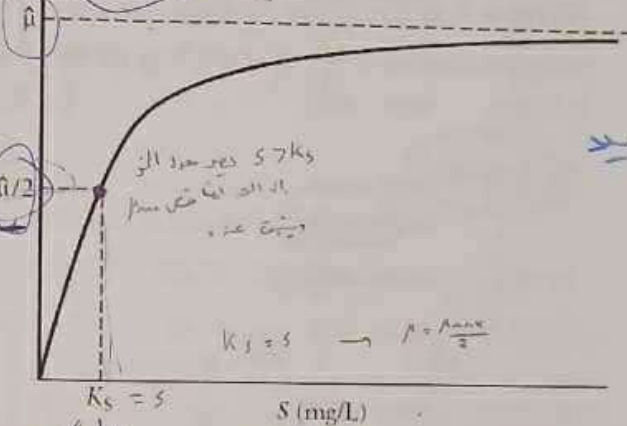
where μ = maximum specific growth rate (at nutrient saturation)

S = substrate or nutrient concentration

K_s = saturation, or half-velocity, constant

Two constants are used to describe the growth rate:

- μ_{max} (mg/L) is the maximum growth rate constant (the rate at which the substrate concentration is not limiting)
- K_s is the half-saturation constant (mg/L) (i.e., concentration of S when $\mu = \mu_{max}/2$)



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Biomass production

$$\frac{dX}{dt} = \text{growth rate} - \text{death rate} = \mu X - k_d X$$

Where k_d represents the endogenous decay rate (d^{-1}) (i.e., microorganism death rate).

Substituting the growth rate constant:

$$\frac{dX}{dt} = \left(\frac{\mu_{max} S}{K_s + S} \right) X - k_d X$$

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- استخدم الميكروبيولوجيا
- Substrate utilization

$$-\frac{dS}{dt} = \frac{1}{Y} \frac{dX}{dt}$$

Where Y is the yield factor →

(mg of biomass produced / mg of food consumed)

- Y range:

Aerobic: 0.4 - 0.8 mg/mg

Anaerobic: 0.08 - 0.2

$$\frac{dS}{dt} = \frac{1}{Y} \frac{dX_{growth}}{dt} = \frac{1}{Y} \left(\frac{\mu_m SX}{K_s + S} \right)$$

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- نسبة الغذاء إلى الميكروبيولوجيا
- Food to microorganism ratio (F/M)

تتمثل النسبة اليومية للغذاء (المادة العضوية) المتاحة للتغذية المزدودة للكائنات الحية في وحدة المساحة

- Represents the daily mass of food supplied to the microbial biomass, X, in the mixed liquor suspended solids, (MLSS)

- Units are Kg BOD₅/Kg MLSS/day

$$\frac{F}{M} = \frac{BOD_5 (Kg/m^3) \times Influent \text{ flow} (m^3/d)}{Reactor Solids (Kg/m^3) \times Reactor Volume (m^3)}$$

$$F/M = \frac{Q S_0}{V X} = \frac{S_0}{\bar{t} X} \quad \text{or } Kg/m^3$$

Where hydraulic retention time,

$$\theta_c = \frac{V X}{Q X} = \frac{V}{Q}$$

average time the liquid remains in the reactor. → $\bar{t} = \frac{V}{Q}$

➤ Typical F/M ratio is between 0.2 - 0.5 kg BOD₅/Kg MLSS/day

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Design Parameters for ASS

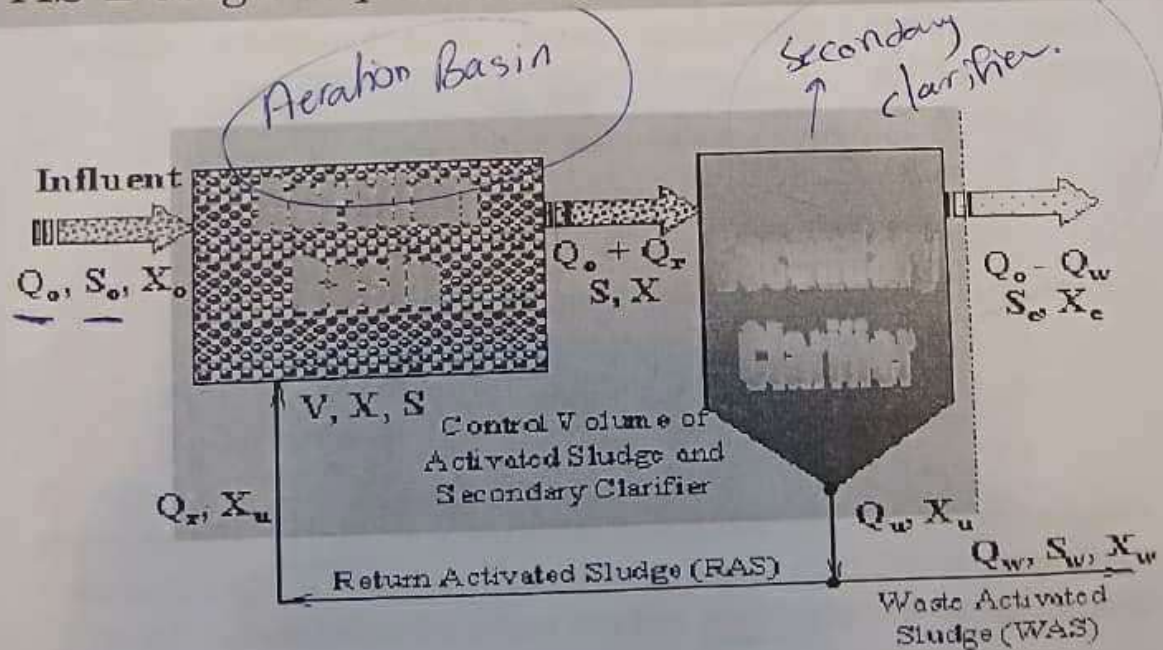
Design parameters for activated sludge treatment processes

Type of process	Mean cell residence time (days)	F/M (kg BOD ₅ /kg MLSS)	Loading (kg BOD ₅ /m ³ -d)	Hydraulic retention time (hr)	MLSS (mg/L)	Recycle ratio
Conventional	5 - 15	0.2 - 0.4	0.3 - 0.6	4 - 8	1500 - 3000	0.25 - 1.0
Step aeration	5 - 15	0.2 - 0.4	0.6 - 1.0	3 - 5	2000 - 3500	0.25 - 0.75
Completely mixed	5 - 30	0.1 - 0.6	0.8 - 2.0	3 - 6	2500 - 4000	0.25 - 1.5
Contact stabilization	5 - 15	0.2 - 0.6	1.0 - 1.2	0.5 (contact) 3 - 6 (stabil)	1000 - 3000 4000 - 10000	0.50 - 1.5
High-rate	5 - 10	0.4 - 1.5	1.6 - 16	2 - 4	4000 - 10000	1.0 - 5.0
Extended aeration	20 - 30	0.05 - 0.15	0.16 - 0.4	18 - 36	3000 - 6000	0.75 - 1.5
Pure oxygen	8 - 20	0.25 - 1.0	1.6 - 3.2	1 - 3	3000 - 8000	0.25 - 0.5

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AS Design Equations



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Mass balance of biomass production (CSTR)

Influent biomass + biomass production = effluent biomass + sludge wasted

$$Q_o X_o + V \frac{dX}{dt} = (Q_o - Q_w) X_e + Q_w X_w$$

Substitute biomass production equation

$$Q_o X_o + V \left(\frac{\mu_m S}{K_s + S} X - k_d X \right) = (Q_o - Q_w) X_e + Q_w X_w$$

Assume that influent and effluent biomass concentrations are negligible and solve

$$\frac{\mu_m S}{K_s + S} = \frac{Q_w X_w}{VX} + k_d$$

Mass balance of food substrate

Influent substrate + substrate consumed = effluent substrate + sludge wasted substrate

$$Q_o S_o + V \frac{dS}{dt} = (Q_o - Q_w) S_e + Q_w S_w$$

Substitute substrate removal equation

$$Q_o S_o + \frac{V}{Y} \left(\frac{\mu_m XS}{K_s + S} \right) = (Q_o - Q_w) S_e + Q_w S_w$$

Assume that no biochemical action takes place in clarifier. Therefore the substrate concentration in the aeration basin is equal to the substrate concentrations in the effluent and the waste activated sludge. Solve:

$$\frac{\mu_m S}{K_s + S} = \frac{Q_o Y}{VX} (S_o - S)$$

Overall equations

- Combine the mass balance equations for food and biomass:

$$\frac{Q_w V_w}{VX} + k_d = \frac{Q_o Y}{VX} (S_o - S)$$

The cell residence time is:

$$\theta_c = \frac{VX}{Q_w X_w}$$

and the hydraulic retention time is, $\theta = V/Q_o$

Substitute and rearrange:

$$X = \frac{\theta_c (Y) (S_o - S)}{\theta (1 + k_d \theta_c)}$$

Compute the F/M ratio

$$\frac{F}{M} = \frac{S_o}{X}$$

$$\frac{F}{M} = \frac{S_o}{(V/Q_o)X} = \frac{Q_o S_o}{VX}$$

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Values of growth constants for domestic wastewater

Parameter	Basis	Value ^a	
		Range	Typical
K_s	mg/L BOD ₅	25-100	60
k_d	d ⁻¹	0-0.30	0.10
μ_m	d ⁻¹	1-8	3
Y	mg VSS/mg BOD ₅	0.4-0.8	0.6

^aValues are for 20°C

Sources: Metcalf & Eddy, 2003; and Shahriar et al., 2006.

S in the effluent is exactly the same as S in the reactor is assumed to be perfectly mixed.

✓ Ref. Book: Vesilind

✓ Example: 11.1 ✓

Example: 11.2 ✓

Example: 11.3 ✓

Example: 11.4 ✓

Example: 11.5 ✓

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Phosphorus Removal

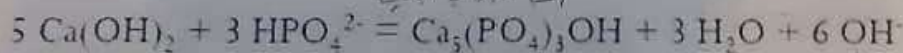
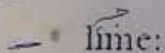
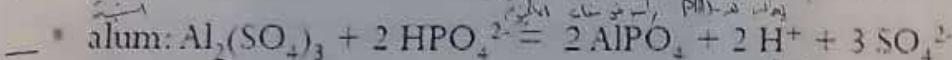
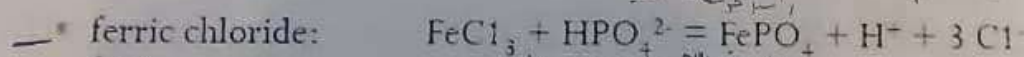
- Phosphorus is typically found as mono-hydrogen phosphate (HPO_4^{2-}) in wastewater.

نفسات الحادى راسه لاسنا

- The removal of phosphorus to prevent or reduce eutrophication is typically accomplished by chemical precipitation using one of three compounds.

تتم ازالة الفوسفور لمنع ازدياد تغلغل الطحالب
عادة عند صرف المياه الجارية بكميات كبيرة
باستخدام احد المركبات الثلاثة

- The precipitation reactions for each are shown below:



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- You should note that ferric chloride and alum reduce the pH while lime increases it.

The effective range of pH for alum and ferric chloride is between 5.5 and 7.0.

- If there is not enough naturally occurring alkalinity to buffer the system to this range, then lime must be added to counteract the formation of H^+ .

The precipitation of phosphorus requires:

- a reaction basin and
- a settling tank to remove the precipitate.

- When ferric chloride and alum are used, the chemicals may be added directly to the aeration tank in the activated sludge system.

- Thus, the aeration tank serves as a reaction basin
- The precipitate is then removed in the secondary clarifier with biological solids.

This is not possible with lime since the high pH required to form the precipitate is detrimental to the activated sludge organisms !!

- In some wastewater treatment plants, the $FeCl_3$ (or alum) is added before the wastewater enters the primary sedimentation tank.

- This improves the efficiency of the primary tank

- But: this may deprive the biological processes of needed nutrients.

Nitrogen Control

- Nitrogen in any soluble form (NH_3 , NH_4^+ , NO_2^- , and NO_3^- , but not N_2 gas) is a nutrient and may need to be removed from wastewater to help control algae growth in the receiving body.

- In addition, nitrogen in the form of ammonia:
 - exerts an oxygen demand and
 - can be toxic to fish.
- Removal of nitrogen can be accomplished either biologically or chemically:
 - The biological process is called nitrification / denitrification.
 - The chemical process is called ammonia stripping.

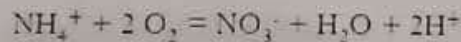
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Nitrification/de-nitrification (Bio.):

The natural nitrification process can be forced to occur in the activated sludge system by maintaining a cell detention time of 15 days or more.

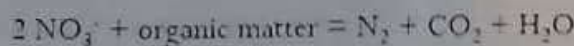
The nitrification step is expressed in chemical terms as follows:



- Bacteria must be present to cause the reaction to occur. This step satisfies the oxygen demand of the ammonium ion.

If the nitrogen level is not of concern for the receiving body, then wastewater can be discharged after settling.

If nitrogen is of concern, the nitrification step must be followed by anoxic denitrification by bacteria:



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Sludge Treatment & Disposal

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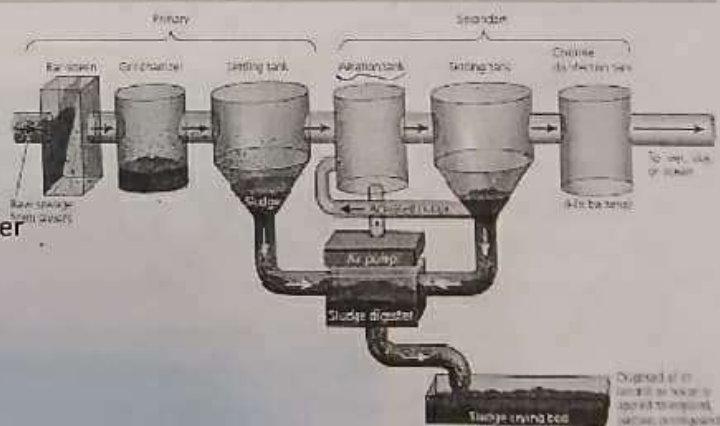
Univ. of Jordan/ Chem. Eng. Dept.

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Sludge Sources

Sources of sludge

- Primary sedimentation tank
- Aeration basin or secondary clarifier
- Screening and grinder
- Filter backwash water



Sludge must be treated because:

- they are aesthetically displeasing,
- they are potentially harmful,
- and they contain too much water.

Sludge Types

لا يوجد نوعين متماثلين من مياه الصرف الصحي في جميع الجوانب

- No two wastewater sludges are alike in all respects.
- Sludge characteristics change with time.
- There is no "average sludge."

Primary sludge

- 3 to 8% solids
- About 70% organic material

Secondary sludge

- Consists of wasted microorganisms and inert materials
- About 90% organic material
- Trickling filter sludge: 2-5% solids

Tertiary sludge

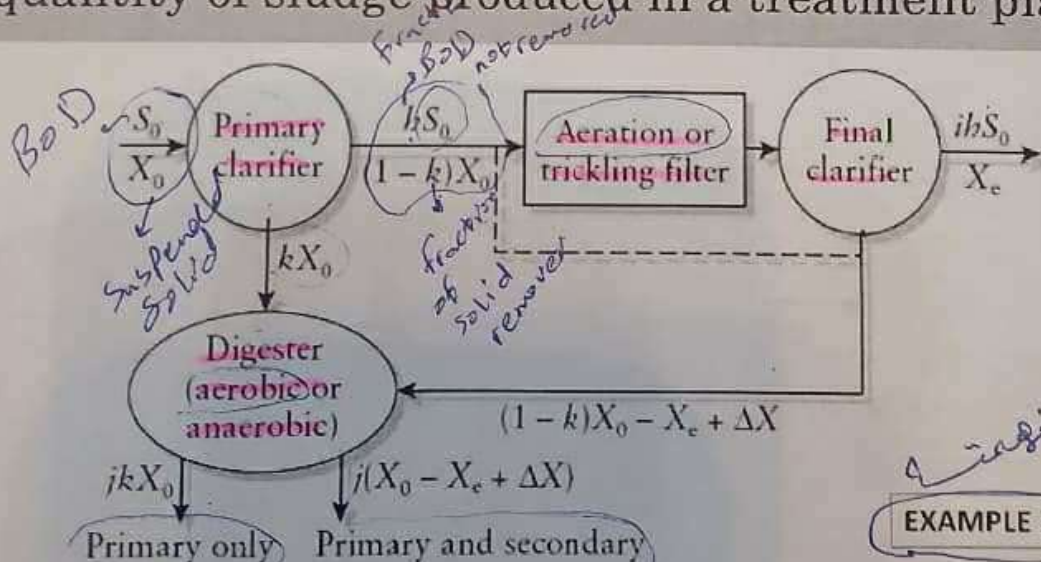
- If secondary clarifier is used to remove phosphate, this sludge will also contain chemical precipitates (more difficult to treat)
- De-nitrification sludges - similar to WAS sludge

Wast activated sludge -

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The quantity of sludge produced in a treatment plant



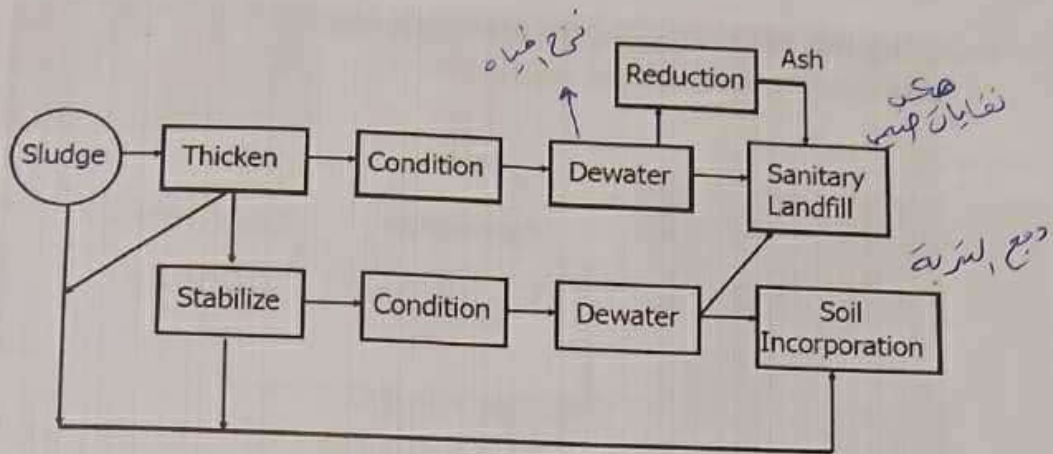
EXAMPLE 11.9

- S_0 = influent BOD, lb/d (kg/h)
- X_0 = influent suspended solids, lb/d (kg/h)
- h = fraction of BOD not removed in the primary clarifier
- i = fraction of BOD not removed in activated sludge system
- X_e = plant effluent suspended solids, lb/d (kg/h)
- k = fraction of influent solids removed in the primary clarifier
- ΔX = net solids produced by biological action, lb/d (kg/h)
- Y = yield, or the mass of biological solids produced in the aeration tank per mass of BOD destroyed, or

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Sludge Treatment

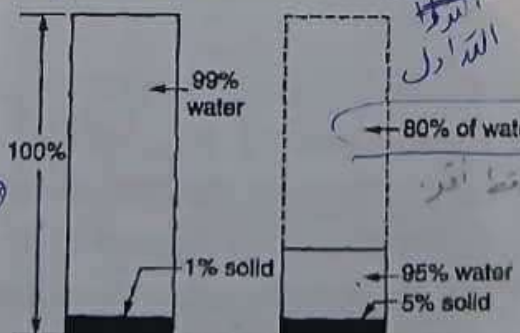


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Sludge Thickening

- Sludge thickening is a process in which the concentrations of solids is increased and the total sludge volume is decreased, but the sludge still behaves like a liquid instead of a solid.
- The advantages of sludge thickening are substantial. When sludge with 1% solids is thickened to 5%, the result is an 80% reduction in volume. Reducing from 1% solids to 20% solids, which might be achieved by sludge dewatering, would result in a 95% reduction in volume. The volume reduction translates into considerable savings in treatment, handling, and disposal costs.

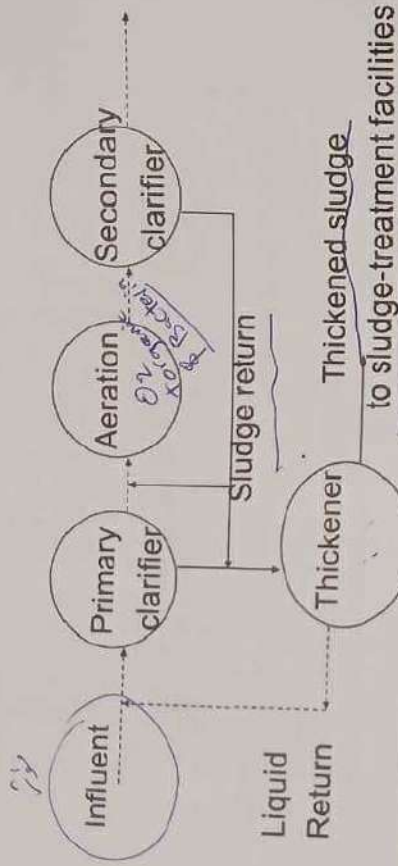


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Sludge Thickening

- Reducing the water in primary and secondary sludges.



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Sludge Thickening Methods

Methods

- Gravity thickening : using an additional clarifier to remove more water

- Dissolved air flotation (DAF)

- Used to concentrate secondary sludges

- Sludge is pressurized and injected with air; releasing into a settling tank (when the pressure is released, the extra air comes out of solution; attaching to the sludge particles as microscopic bubbles); the sludge is floated to the surface in a concentrated form; the underflow returns to the head of the plant.

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Gravity Thickener

Gravity thickening

- Best with primary sludge
- Increases solids content from 1-3% to 10%

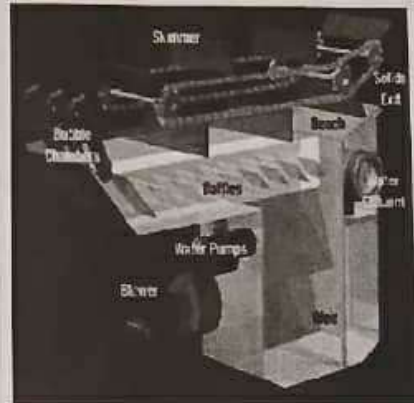
→ there is more solid particles rather than secondary

Flotation

DAF

- Especially effective on activated sludge
- Increases solids content from 0.5 -1% to 3-6%

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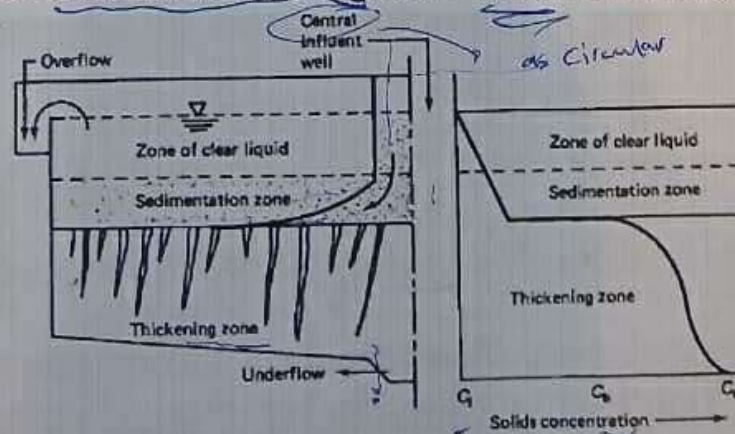


Flotation

Gravity Thickening

- Accomplished in circular sedimentation basins
- Degree of thickening: 2~5 times the incoming solids conc.
- Max. achievable solids concentration: < 10%
- Chemical and waste activated sludges are difficult to thicken under gravity.

→ under flow
Thickened solid
→ overflow
water nearly clear



G_1 - Influent solids concentration
 G_2 - Lowest concentration at which sludge thickening begins
 G_3 - Underflow concentration

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Gravity Thickener Equipment

➤ Generally circular concrete tanks with bottom sloping toward the center.

Equipment

- Rotating bottom scraper arm
- Vertical pickets
- Rotating scum collection mechanism with scum baffle plates
- Overflow weir

Other configurations

- Circular steel tank: Generally cheaper because of simplicity of construction, equipment installation, and operation and maintenance
- Rectangular concrete and steel tanks

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Dissolved Air Flotation (DAF)

Primarily used to thicken the solids in chemical and WAS

Waste Activated Sludge

Separation of solids is achieved by introducing fine air bubbles created under pressure of several atmosphere into the liquid, attaching to solids to cause flotation of solids

Degree of thickening: 2~8 times the incoming solids concentration

Max. solids concentration: 4~5%

DAF Variations

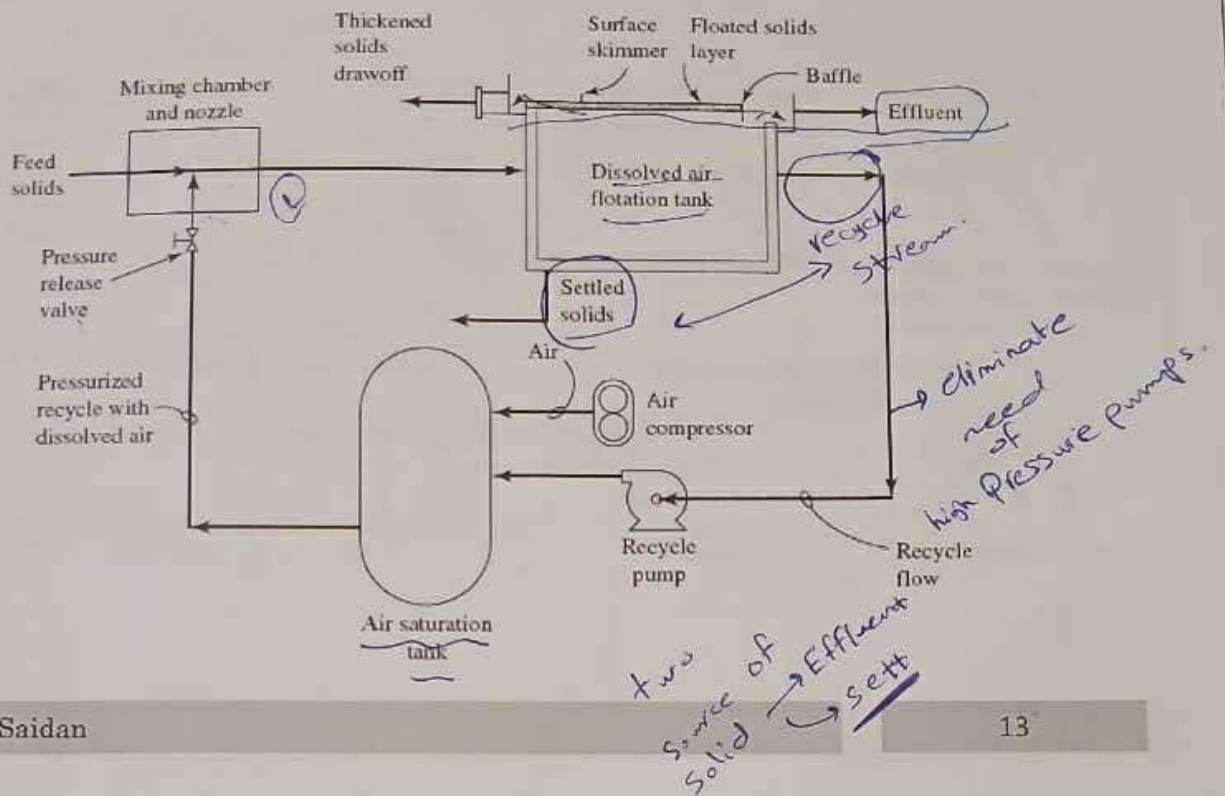
Pressurize total or only a small portion of the incoming sludge

Pressurize the recycled flow from the flotation thickener - preferred because it eliminates the need for high-pressure sludge pumps.

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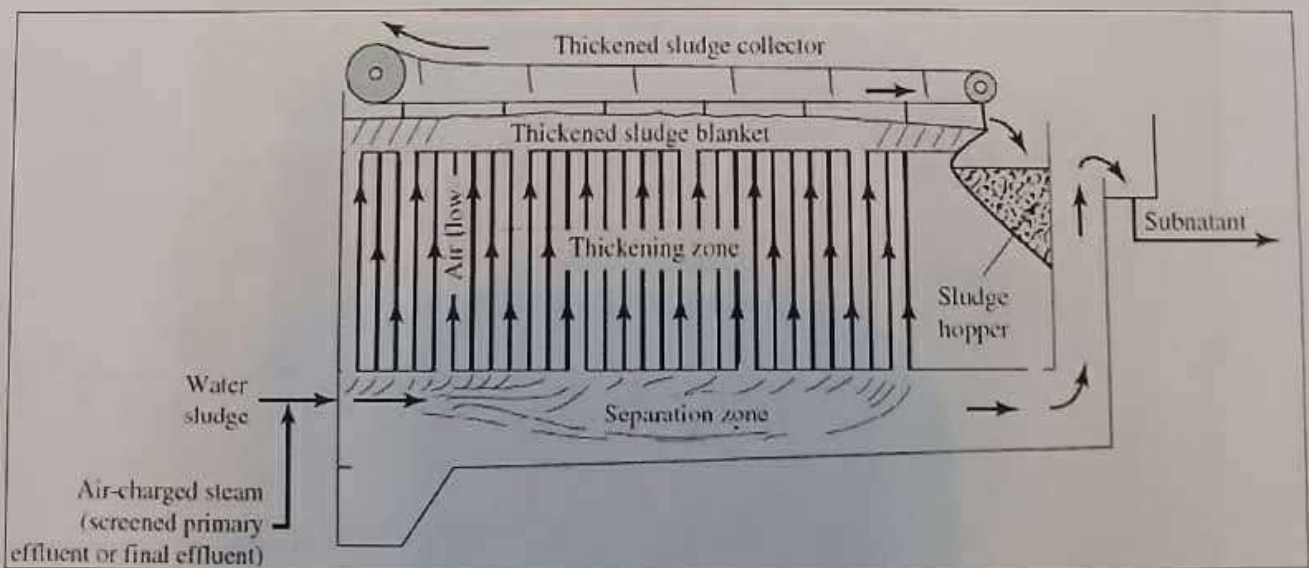
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Schematic of DAF thickener system



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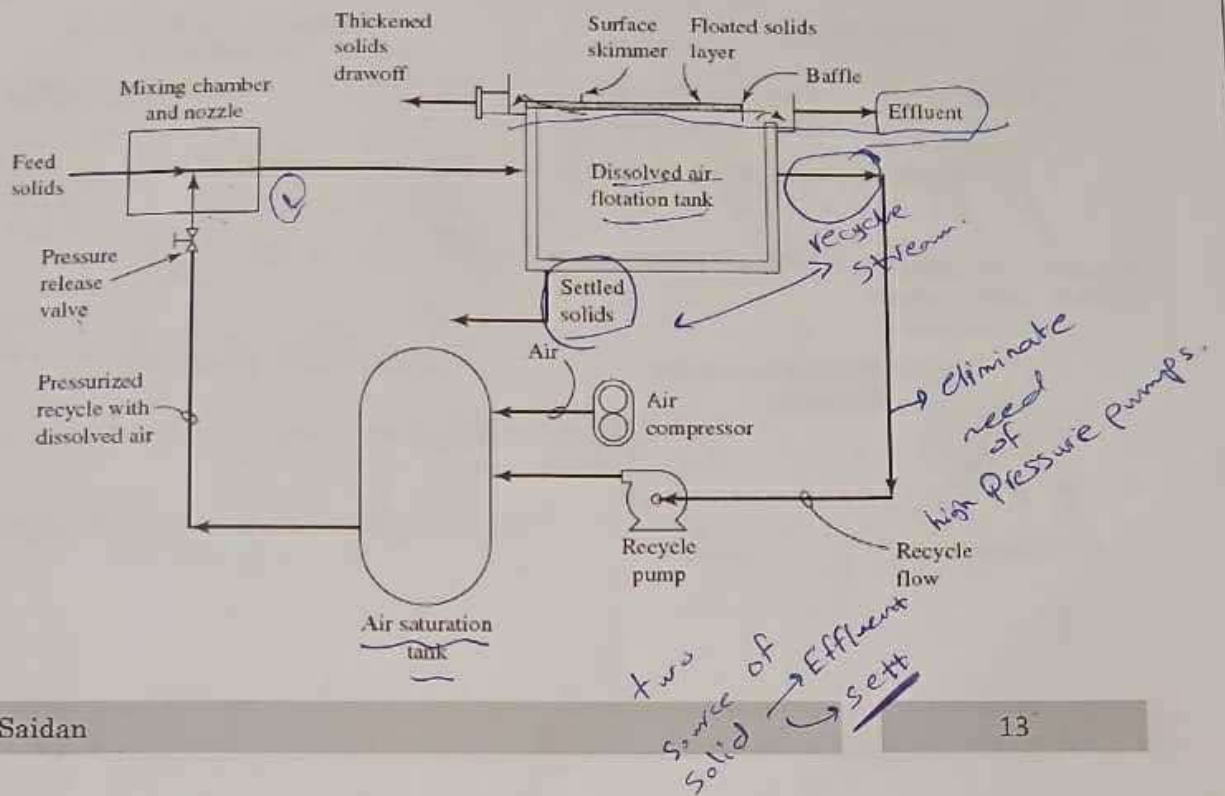
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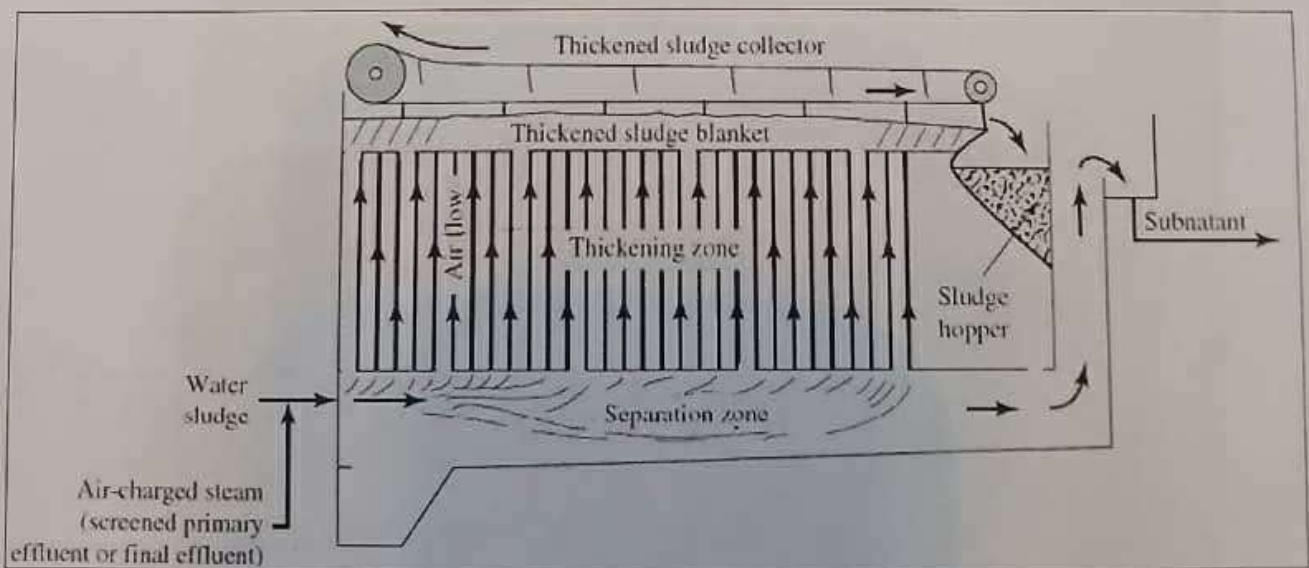
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Schematic of DAF thickener system



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Sludge Stabilization Means

Biological Treatment

1. Aerobic Digestion
2. Anaerobic digestion
3. Lagoons
4. Composting

Chemical Treatment

1. Wet combustion
2. Lime stabilization
3. Chlorination
4. Heat stabilization
5. Irradiation

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Time
Aerobic
Anaerobic

Lime stabilization is achieved by adding lime to the sludge, which raises the pH to about 11 or above. This significantly reduces the odor and helps in the destruction of pathogens. The major disadvantage of lime stabilization is that it is temporary. With time (days), the pH drops and the sludge once again becomes putrescent.

Aerobic digestion is merely a logical extension of the activated sludge system. Waste activated sludge is placed in dedicated aeration tanks for a very long time, and the concentrated solids are allowed to progress well into the endogenous respiration phase, in which food is obtained only by the destruction of other viable organisms.

This results in a net reduction in total and volatile solids. Aerobically digested sludges are, however, more difficult to dewater than anaerobic sludges.

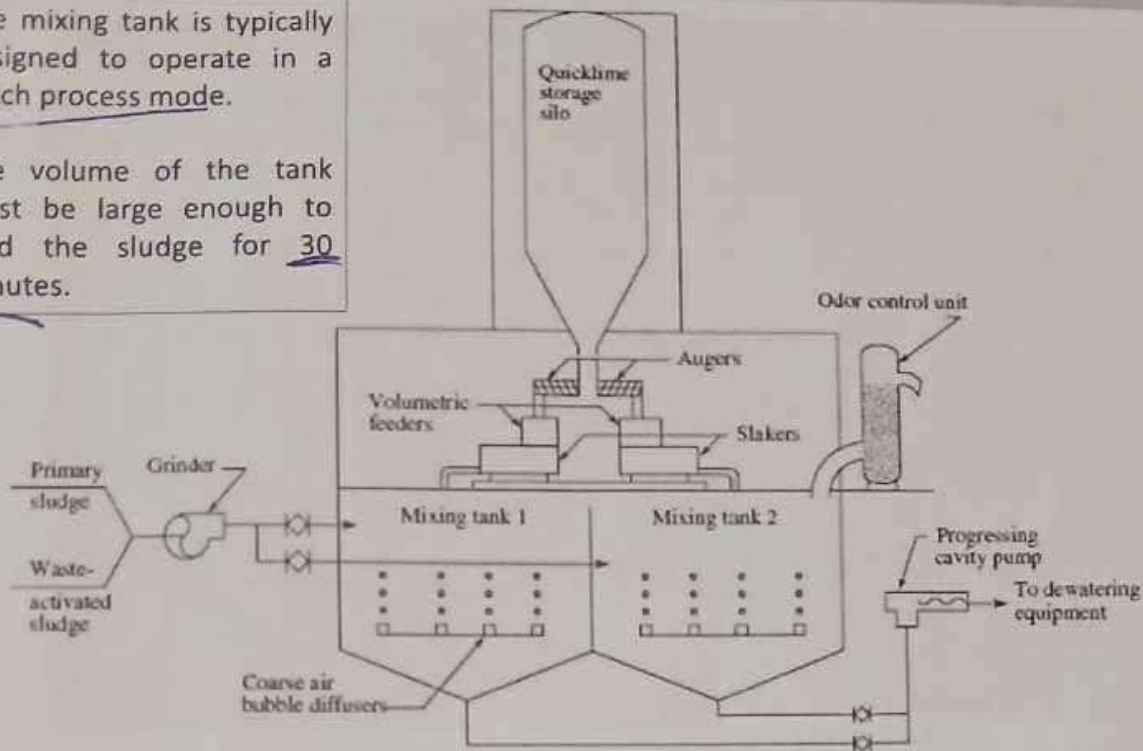
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Conceptual design for a lime stabilization facility

The mixing tank is typically designed to operate in a batch process mode.

The volume of the tank must be large enough to hold the sludge for 30 minutes.



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Second + Final.

المرحلة الثانية Anaerobic Digestion

ثلاث مراحل صغيرة .

The anaerobic treatment of biological sludges involves three distinct stages:

- In the first stage, complex waste components, including fats, proteins, and polysaccharides, are hydrolyzed to their component subunits.
 (هضم أولي) تحلل المواد العضوية المعقدة
- The second stage is commonly referred to as acid fermentation. In this stage, organic material is simply converted to organic acids, alcohols, and new bacterial cells.
 (تخمير حمضي) التحلل الهوائي البسيط
- In the third stage, the end products of the second stage are converted to gases (mainly methane, CH_4 , and carbon dioxide, CO_2) by several different species of strictly anaerobic bacteria.
 (تخمير اللاهوائي) التخمير اللاهوائي البسيط

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Theory of Anaerobic Digestion

1. Acid forming bacteria break down complex organic substances to simpler compounds, notably fatty acids.

Carbohydrates $\longrightarrow$ fatty acids

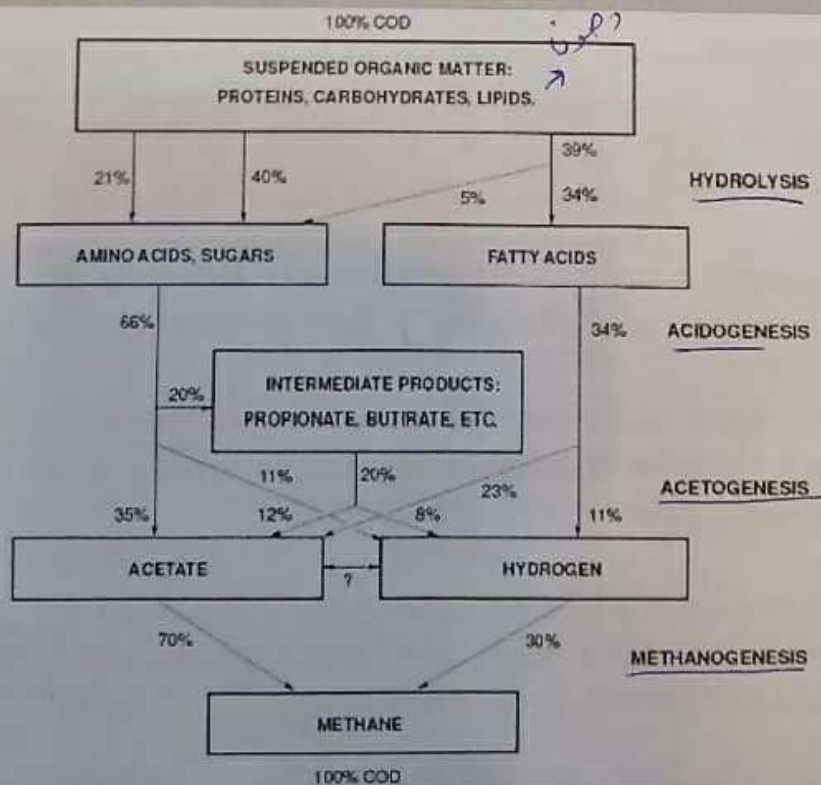
Proteins $\longrightarrow$ Amino acids $\longrightarrow$ NH_3 + fatty acids

2. Products of the first stage are further broken down, by methane forming bacteria, to methane and carbon dioxide.

NH_3 + fatty acids $\longrightarrow$ CH_4 + CO_2

- Methane forming bacteria work in pH range 6 to 8 (better in 7.2 - 7.4).
- Lime is added to adjust pH.

Decomposition of excess activated sludge



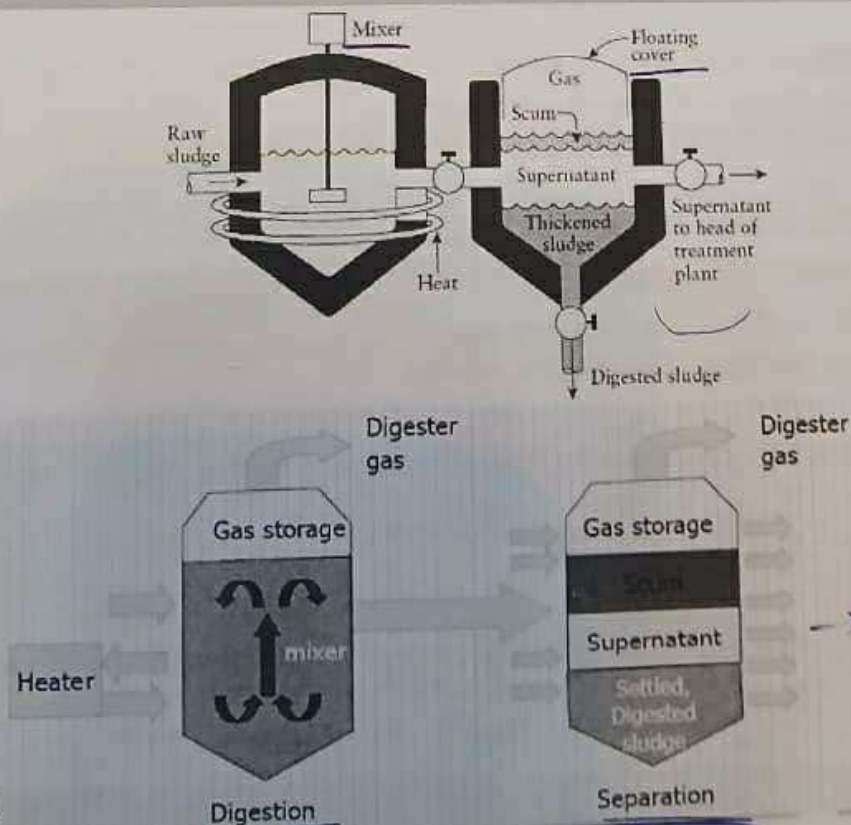
Anaerobic Digestion

- Anaerobic digestion of sludge decrease the volatile organics by 40-50% and reduce the numbers of pathogenic organisms in sludges.
 المواد العضوية المتطايرة
 تقليل عدد الكائنات المرضية
- Accomplished by holding the sludge in closed tanks for periods of 10 to 90 days.
 دفع زلال
 مدة احتجاز طويلة
- Old process : unmixed, unheated, long detention time (30-90 days)
- Recent process : complete mixing, heating (35-45°C), detention time 10-20 days
 مختلطة
 الحرارة

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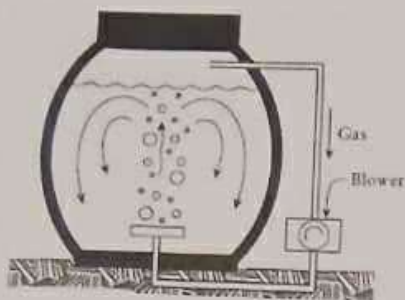
Two-stage anaerobic digestion



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Egg-shaped anaerobic digesters



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Second Time or later

Egg-shaped anaerobic digesters

Advantages

- High degree of waste stabilization at high organic loading rates
- Very little sludge production (< 5% of biodegradable organic matter being converted to cell material) (10% of aerobic sludge production)
- Easy dewatering of the excess sludge
- Low nutrient requirement (10% of aerobic process requirement)
- No aeration equipment
- Methane production – very low energy input (if the methane gas is used to heat the digester)

Egg-shaped anaerobic digesters

Disadvantages

- ^{ناتج بكتيري منخفض} Low bacterial yield
- ^{نقصان سهولة في مرآة حوضه بيل} prolonged periods of biomass build-up,
- ^{فترة بدء تشغيل طويلة} requiring longer start-up period (8 to 12 weeks).
- ^{حساسة للتغيرات البيئية} Temperature, pH, toxic sensitive
- ^{تكلفة استثمار مرتفعة} High capital costs
- ^{تحتاج وحدة لاحتاج اذ متفكر دمرة} Complex operation requiring skilled operators

Digester Heating

رفع درجة حرارة الحماة الداخلية ^{أي} ~~الداخلية~~ حرارة العمز ولتحويض الفقد

- ^{المرآة من طود الحماة والأرضية وسقف الممرات ولتحويض الفاقة التي قد يحدث في الماكينة} To raise the incoming sludge to digestion tank temperatures, to compensate for the heat losses through the walls, floor, and roof of the digester, and to make up the losses that might occur in the piping between the source of the heat and the tank.
- ^{المبادلة حرارية داخلية وخارجية} Internal or external heat exchangers
- ^{متطلبات الحرارة} Heat requirements

$$q = UA \Delta T$$

where

q = heat loss, BTU/h (W);

U = overall coefficient of heat transfer, BTU/ft²·hr·°F (W/m²·°C);

A = cross-sectional area through which the heat loss is occurring, ft² (m²); and

ΔT = temperature drop across the surface in question, °F (°C).

Sludge Conditioning

لزيادة كفاءة التصفية

Chemical Conditioning: Chemical additives may be used to improve sludge dewaterability by acting as coagulants. Chemicals commonly used for this are ferric chloride (FeCl_3), lime (CaO), and organic polymers

إضافة مواد كيميائية

Physical Conditioning: Physical conditioning is primarily by heat. Heat conditioning involves heating at High temperatures ($175-230^\circ\text{C}$) and High pressures (10 to 20 atmospheres).

تكييف بالحرارة

بشكل دائم

Dewaterability is improved dramatically and pathogens are destroyed as well.

بشكل كبير
بالإضافة إلى القضاء على مسببات الأمراض

Sludge Dewatering and Drying

لزيادة كفاءة التخلص من الحمأة بعد المعالجة

الافتتاحية
المرتبطة
بمستوى
أدوية الأرض

Dewatering reduces the moisture content of the sludge so that it can more easily be disposed of by landfill, incineration, heat drying, composting or other means.

المسحوق

Unlike sludge thickening, where the treated sludge continues to behave as a liquid, dewatered sludge will behave like a solid after treatment.

Dewatering is seldom used as an intermediate process unless the sludge is to be incinerated. Most wastewater plants use dewatering as a final method of volume reduction before ultimate disposal.

The objective is a moisture content of 60 to 80 percent depending on the disposal method.

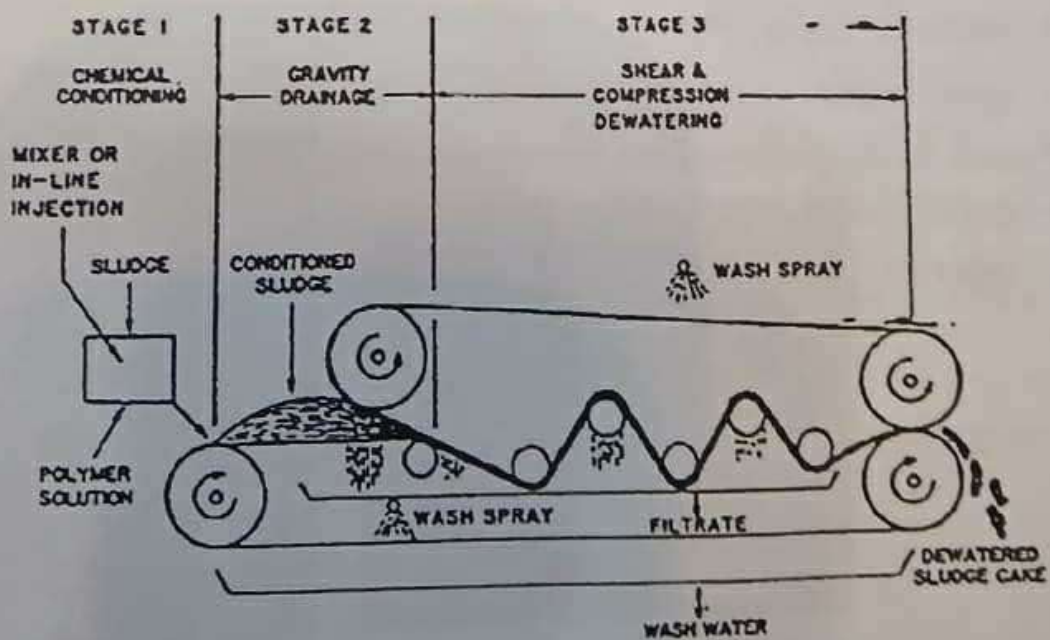
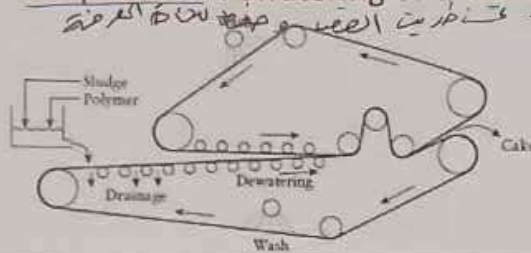
By applying the sludge to sand drying beds

By using mechanical dewatering equipment

Belt Press Filtration

- Belt filter presses employ single or double moving belts to continuously dewater sludges through one or more stages of dewatering.
نظام حزام الترشيح بالخرام خزاناً واحد أو أكثر حيث يمر من خلاله الماء
- All belt press filtration processes include three basic operational stages:

- ➔ chemical conditioning of the feed sludge;
معالجة كيميائية للمخلفات المدخلة
- ➔ gravity drainage to a non-fluid consistency;
تصفية الجاذبية للمخلفات لتصل إلى حالة غير سائلة
- ➔ shear and compression dewatering of the drained sludge.
الضغط والقص للمخلفات لتصفيته



Sludge drying beds

- Sludge drying beds rely on drainage and evaporation to effect moisture reduction.

These beds are open; and as such, are very susceptible to climatic conditions such as precipitation, sunshine, air temperature, relative humidity, and wind velocity. For example, sludge drying in 6 weeks in the summer would take at least 12 weeks to dry in the winter.

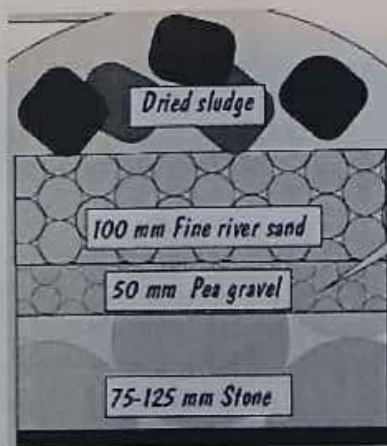
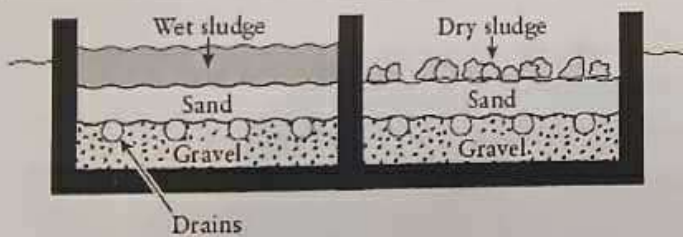
- Sludge bed drying efficiency can be improved significantly by covering the bed with glass or plastic and by providing artificial heat.

Heat could be supplied using waste biogas as a fuel or waste heat from the base power plant.

يمكن توفير الحرارة بالطاقة من غاز حيوي ناتج من نفايات كمواردات خلال الصيف.

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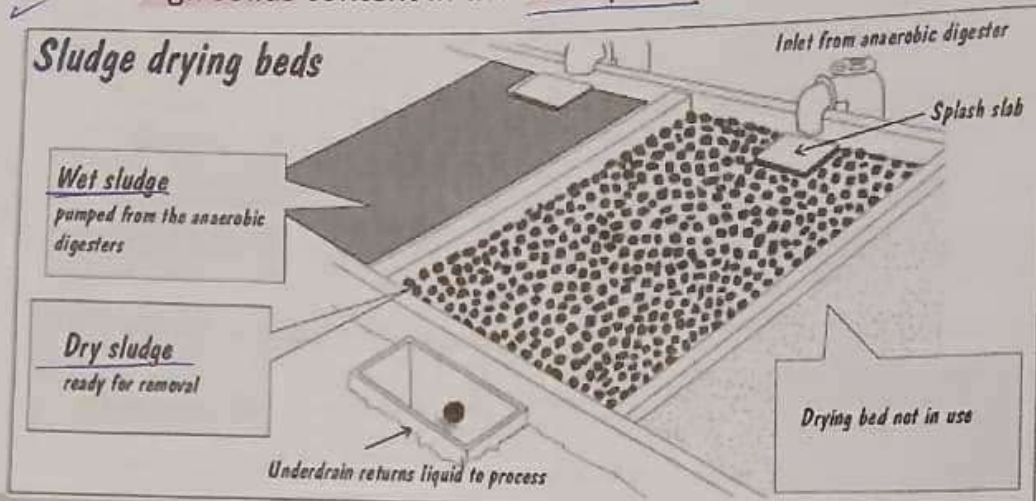
© Courtesy of P. Auerne Mrs. Lind

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The principle advantages of drying beds

- ✓ 1. Low cost,
- ✓ 2. Infrequent attention required, and
- ✓ 3. High solids content in the dried product



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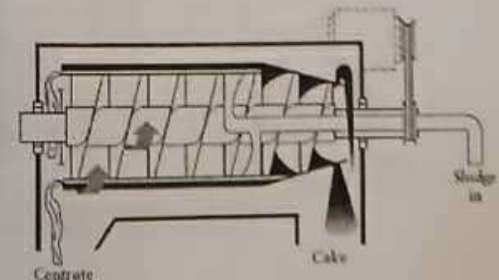
Centrifugal dewatering

في هذه العملية يتم فصل السوائل عن المواد الصلبة باستخدام قوة الطرد المركزي.

■ **Centrifugal dewatering of sludge** is a process which uses the force developed by fast rotation of a cylindrical drum or bowl to separate the sludge solids from the liquid.

■ **In the basic process, when sludge slurry is introduced to the centrifuge, it is forced against the bowl's interior walls, forming a pool of liquid.**

- **Density differences cause the sludge solids and the liquid to separate into two distinct layers. The sludge solids "cake" and the liquid "centrate" are then separately discharged from the unit.**



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Air Pollutants & Human Health Effects

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Univ. of Jordan/ Chem. Eng. Dept.

1

Stausmelter (Cook). 370 1 me mi
370 1 me mi



Natural طبيعي
Anthropogenic صنع الإنسان

Air Pollution Definitions

- Air Pollution Definitions**
- Transfer of harmful and/or of Natural/Synthetic materials into the atmosphere as a direct/indirect consequences of human activity (OECD).
مواد ضارة طبيعية أو اصطناعية
الآثار المباشرة وغير المباشرة
البيئة
مركبات عضوية متطايرة
 - The introduction of chemicals, particulate matter, or biological materials that cause harm or discomfort to humans or other living organisms, or damages the natural environment into the atmosphere.
عوامل بيئية
البيئة الطبيعية

- The overwhelming scientific consensus is that the earth's atmosphere is warming rapidly mostly because of human activities, and that this will lead to significant climate change during this century.
- M. saidan

Because the Human activity for example.

- increasing in Burning Fossil fuels, use oil, coal, drive cars that release huge amount of carbon dioxide (CO_2)
- Deforestation: Trees absorb and store CO_2 when forests are cut down, this CO_2 back into atmosphere.
- Global Temperature Rise.

Air Pollutants - Rising sea level due to melting ice and thermal expansion of water.

chemical reaction can be triggered from light (give it energy) → Thermal expansion of water. Smoke + fog → industrial smog, mostly the result of burning coal, and photochemical smog, caused by motor vehicle, industrial, and power plant emissions. combustion plant → source of pollution.

Neutral gas → An air pollutant can be a gas or a particulate.
or Aerosol →

Sources: liquid يكون سائل
نالحو زينة صف لما نرشد خط

- Natural sources
 ✓ Dust blown by wind
 ✓ Pollutants from wildfires and volcanoes
 ✓ Volatile organics released by plants

- Human sources: mostly in industrialized and/or urban areas
- Stationary sources
- Mobile sources

Air Pollutants Classification

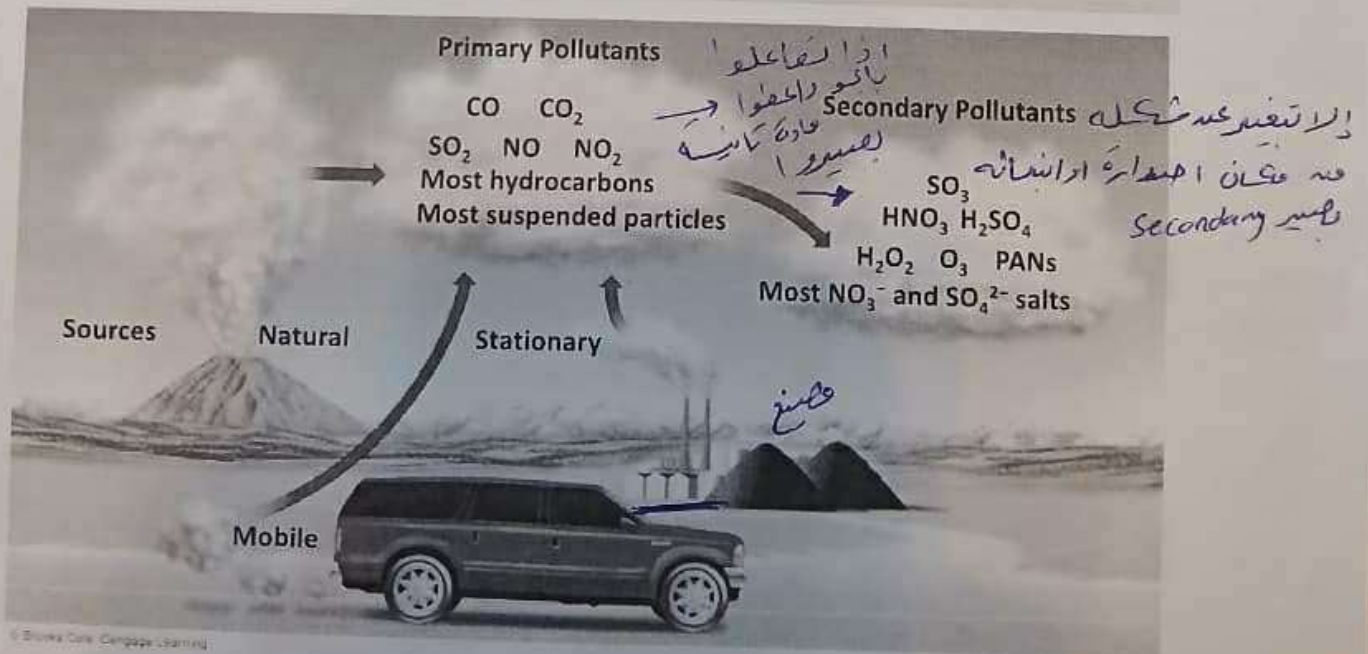
Classified as:

Fixed

1. **Primary pollutants:** *stable* foreign matter injected into the atmosphere by human activities. *اي gases, materials pollutant يقع على الجو ويظل مستقر*
2. **Secondary pollutants:** resulting from chemical transformations, typically with primary pollutants and (often) sunlight. *نتيجة عنه لتفاعلات كيميائية عادة بالملوثات الأولية وغالباً ضوء الشمس*

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5



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PM_{2.5} → combustion particles, organic compounds, metals.
 PM₁₀ → Dust, pollen, mold.
 الغبار، الحبوب، الفطريات

Major Air Pollutants

particulate matter → Air pollutants solid state.
 suspended → Suspended particulate matter (SPM):

Size of SPM → Consists of a variety of solid particles and liquid droplets small and light enough to remain suspended in the air.
 حجم الجسيمات، إذا كانت صغيرة وخفيفة بما يكفي لتبقى عالقة في الهواء.

PM₁₀ → 10 μm
 The most harmful forms of SPM are fine particles (PM-10, with an average diameter < 10 micrometers) and ultrafine particles (PM-2.5). Smaller than PM₁₀

Environmental protection Agency

According to the EPA, SPM is responsible for about 60,000 premature deaths a year in the U.S.
 وفقاً للوكالة البيئية

Smoke → Sources → Heating, vaporization, condensation of solid
 main composition → Fine solid particles (metal oxides) that have cooled from a gas.

particle size → much smaller
 example → welding, often in visible

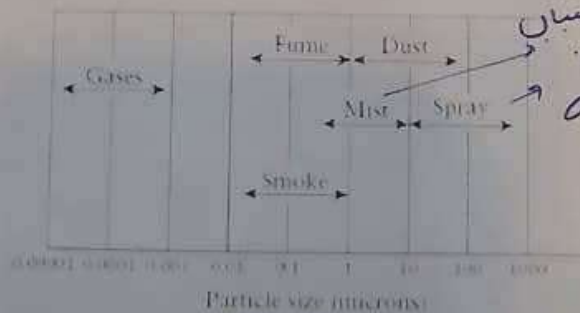
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Smoke → Source → incomplete combustion. (burning)
 main composition → Solid and liquid particles. (soot, tar) ultrafine solid.
 particle size → Generally larger.
 visibility → easily visible, cloudy
 Example → Campfire, car exhaust, burning cigarette.

Major Air Pollutants

Size

Particulate matter or pollutants can be further classified as dusts, fumes, mists, smoke, or spray.
 الغبار، الدخان، الضباب، الدخان، الرذاذ



Terminology	Abbreviation	Diameter Range (μm)
Ultrafine	—	< 0.1
Fine	PM _{2.5}	< 2.5
Coarse	PM ₁₀	2.5 - 10

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Particulate Matters

- Dust is defined as solid particles.

- A fume is also a solid particle, frequently a metallic oxide, formed by the condensation of vapors by sublimation, distillation, calcination, or chemical reaction processes. The particles in fumes are quite small, with diameters from 0.03 to 0.3 μ .

- A mist is an entrained liquid particle formed by the condensation of a vapor and perhaps by chemical reaction. Mists typically range from 0.5 to 3.0 μ in diameter.

- Smoke is made up of entrained solid particles formed as a result of incomplete combustion of carbonaceous materials. Smoke particles have diameters from 0.05 to approximately 1 μ .

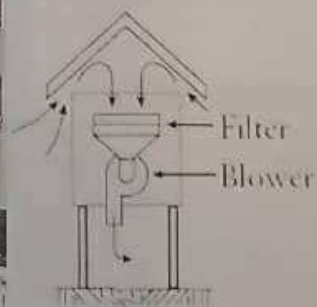
- a spray is a liquid particle formed by the atomization of a parent liquid. Sprays settle under gravity

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Measurement of Particulates

- The measurement of PM10 has been historically done by using bottom the high-volume sampler (or hi-vol).

- The high-volume sampler operates much like a vacuum cleaner, forcing up to 86,000 ft³ of air through a filter in 24 hr. The analysis is gravimetric; the filter is weighed before and after, and the difference is the particulates collected.



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Gaseous Pollutants

- Gaseous pollutants include substances that are gases at normal temperature and pressure as well as vapors of substances that are liquid or solid at normal temperature and pressure.

Name	Formula	Properties of Importance	Significance as Air Pollutant
Sulfur dioxide	SO ₂	Colorless gas, intense choking odor, highly soluble in water - forming sulfurous acid, H ₂ SO ₃ .	Damage to property, health, and vegetation.
Sulfur trioxide	SO ₃	Soluble in water - forming sulfuric acid, H ₂ SO ₄ .	Highly corrosive.
Hydrogen sulfide	H ₂ S	Rotten egg odor at low concentrations, odorless at high concentrations.	Highly poisonous.
Nitrous oxide	N ₂ O	Colorless gas, used as carrier gas in aerosol bottles.	Relatively inert; not produced in combustion.
Nitric oxide	NO	Colorless gas.	Produced during high-temperature, high-pressure combustion; oxidizes to NO ₂ .
Nitrogen dioxide	NO ₂	Brown to orange gas.	Major component in the formation of photochemical smog.
Carbon monoxide	CO	Colorless and odorless.	Product of incomplete combustion; poisonous.
Carbon dioxide	CO ₂	Colorless and odorless.	Formed during complete combustion; greenhouse gas.
Ozone	O ₃	Highly reactive.	Damage to vegetation and property; produced mainly during the formation of photochemical smog.
Hydrocarbons	C _x H _y or HC	Many.	Emitted from automobiles and industries; formed in the atmosphere.
Methane	CH ₄	Combustible, odorless.	Greenhouse gas.
Chlorofluorocarbons	CFCs	Inert, excellent thermal properties.	Deplete ozone in upper atmosphere.

Ozone Depleting Substances

Air Pollution Effects

Effects on Human Health

- Respiratory problems
- Allergies
- Risk for cancer

Effects on the environment

- Acid rain (Regional)
- Ozone depletion (Stratospheric)
- Greenhouse Effect (Global warming)

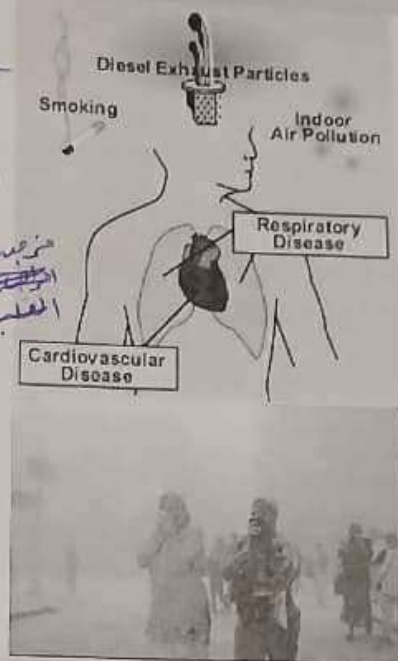
Human Health Effects

التعرض
Exposure to air pollution is associated with numerous effects on human health, including pulmonary, cardiac, vascular, and neurological impairments. عائية اعاقات

الموئل
The health effects vary greatly from person to person. High-risk groups such as the elderly, infants, pregnant women, and sufferers from chronic heart and lung diseases are more susceptible to air pollution. المرضية

Children are at greater risk because they are generally more exposed to outdoor environment and their lungs are still developing stage. الرئة

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المعززة
welfare → Ecosystem → انسان وسية

Pollutant	Description	Sources	Health Effects	Welfare Effects
Carbon Monoxide (CO)	Colorless, odorless gas غاز عديم الرائحة واللون	Motor vehicle exhaust, indoor sources include kerosene or wood burning stoves.	Headaches, reduced mental alertness, heart attack, cardiovascular diseases, impaired fetal development, death.	Contribute to the formation of smog.
Sulfur Dioxide (SO ₂)	Colorless gas that dissolves in water vapor to form acid, and interact with other gases and particles in the air.	Coal-fired power plants, petroleum refineries, manufacture of sulfuric acid and smelting of ores containing sulfur.	Eye irritation, wheezing, chest tightness, shortness of breath, lung damage.	Contribute to the formation of acid rain, visibility impairment, plant and water damage, aesthetic damage.
Nitrogen Dioxide (NO ₂)	Reddish brown, highly reactive gas.	Motor vehicles, electric utilities, and other industrial, commercial, and residential sources that burn fuels.	Susceptibility to respiratory infections; irritation of the lung and respiratory symptoms (e.g., cough, chest pain, difficulty breathing).	Contribute to the formation of smog, acid rain, water quality deterioration, global warming, and visibility impairment.
Ozone (O ₃)	Gaseous pollutant when it is formed in the troposphere.	Vehicle exhaust and certain other fumes. Formed from other air pollutants in the presence of sunlight.	Eye and throat irritation, coughing, respiratory tract problems, asthma, lung damage.	Plant and ecosystem damage.
Lead (Pb)	Metallic element	Metal refineries, lead smelters, battery manufacturers, iron and steel producers.	Anemia, high blood pressure, brain and kidney damage, neurological disorders, cancer, lowered IQ.	Affects animals and plants, affects aquatic ecosystems.
Particulate Matter (PM)	Very small particles of soot, dust, or other matter, including tiny droplets of liquids.	Diesel engines, power plants, industries, windblown dust, wood stoves.	Eye irritation, asthma, bronchitis, lung damage, cancer, heavy metal poisoning, cardiovascular effects.	Visibility impairment, atmospheric deposition, aesthetic damage.

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والتي تسمى حمض الكبريتيك
صنفاً

معدن أوروبا ← حمض الكبريتيك

Acid Rain

sulfuric acid
هو المسطر
لهون

One way in which SO_2 is removed from the atmosphere is the formation of acid rain

- contains high levels of sulfuric or nitric acids

- contaminate drinking water and vegetation

- damage aquatic life

- erode buildings

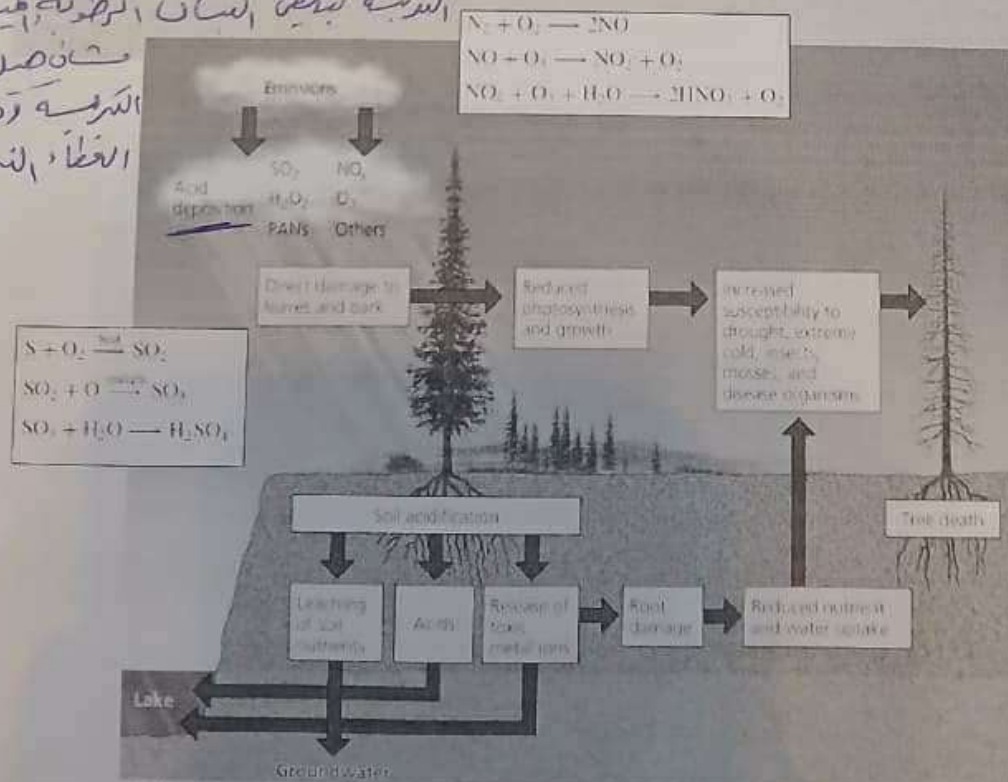
- Alters the chemical equilibrium of some soils



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يصير عن فاصية السموم كثيرة بدال ما
التي تسمى بـ حمض الكبريتيك
التي تسمى بـ حمض الكبريتيك
التي تسمى بـ حمض الكبريتيك
التي تسمى بـ حمض الكبريتيك

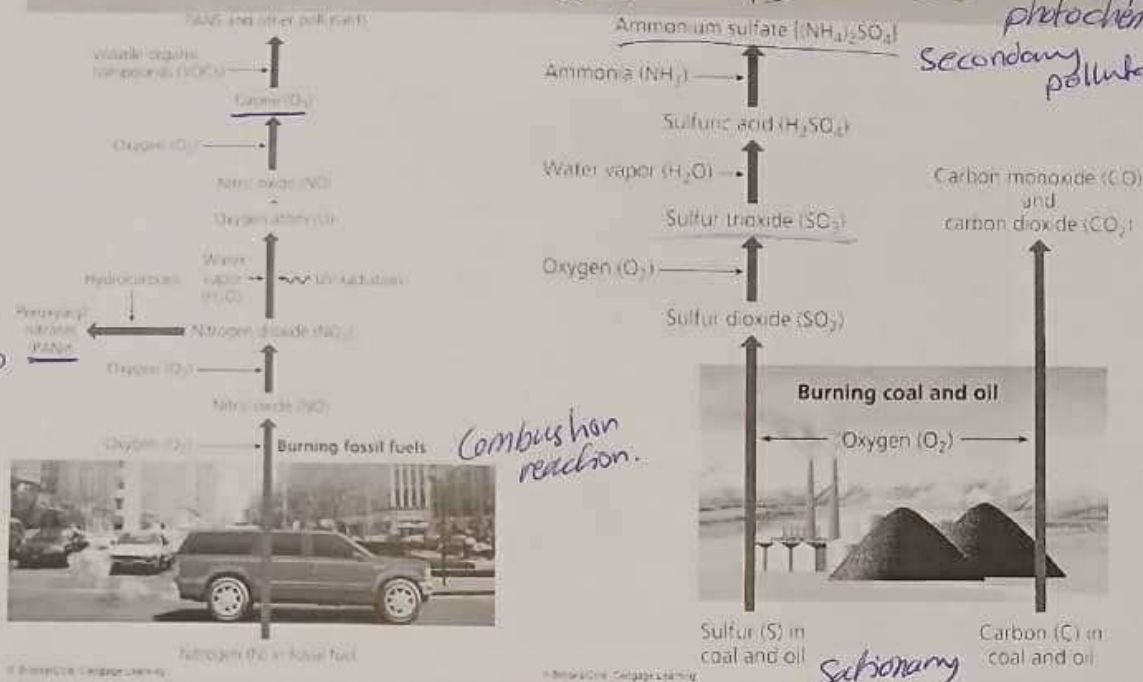


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smog $\Rightarrow$ smoke + fog $\Rightarrow$ الدخان والضباب
 اللون فيه على سواد
 smoke + fog

Photochemical Smog $\Rightarrow$ تفاعل كيميائي ضوئي
 secondary pollutant. ملوث ثانوي



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Secondary Source • 17

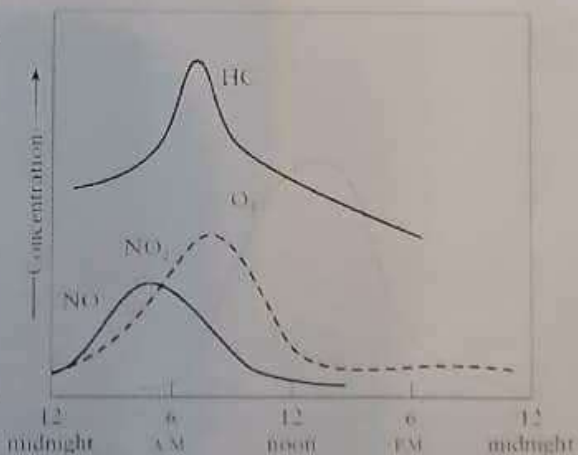
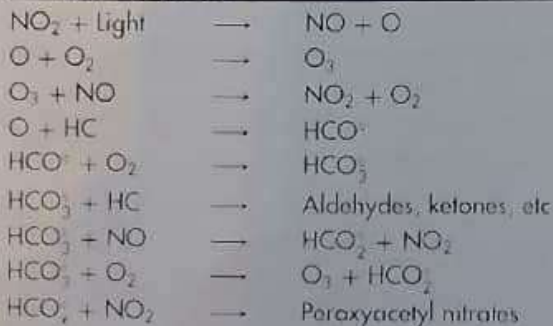
Simplified Reaction Scheme

Hydrocarbons + Nox + Sunlight $\rightarrow$ photochemical smog (oxidants)

المركبات الهيدروكربونية، أكاسيد النيتروجين

primary oxidants produced: ozone (O₃), formaldehyde, peroxyacetyl nitrate (PAN)

intermediate reaction.



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الضباب الكيميائي الضوئي

Photochemical Smog in Santiago, Chile



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Ozone Depletion

0.5 - 1.5 cm

مقياس للعين

- Ozone (O_3) is an eye irritant at usual urban levels, but urban O_3 should not be confused with stratospheric O_3 , 7 to 10 mile above the earth's surface.
- The latter acts as an ultraviolet radiation shield, and its alteration can increase the risk of skin cancer as well as change the ecology in unpredictable ways.

كذلك عند الارتفاع فوق الغلاف الجوي

12 km or 7 to 10 miles

Ultraviolet

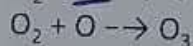
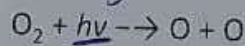
تغير البيئة

هناك الغلاف

Cyclic

Ozone in the upper atmosphere is created when oxygen reacts with light energy

(hv):

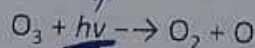


يعتمد على وجود الأشعة

التي توجد على ارتفاعات Stratosphere

Light energy also destroys ozone:

ultraviolet

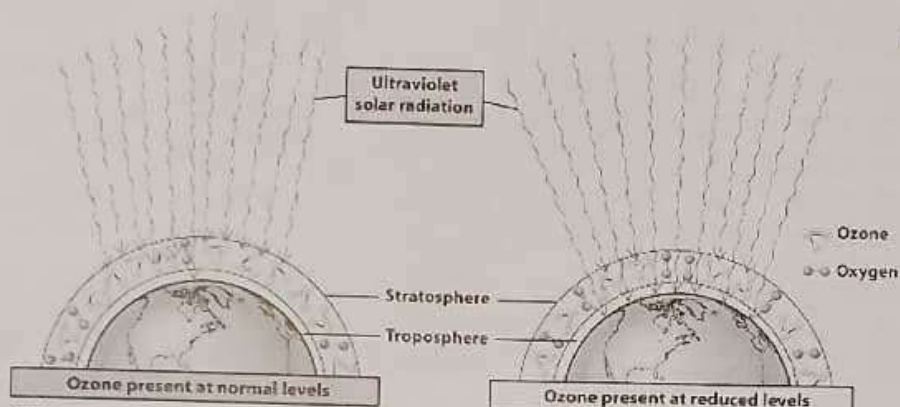


This is the mechanism by which ozone prevents ultraviolet radiation from reaching earth's surface.

UV

✓ Ozone Protects earth from UV radiation Part of the electromagnetic spectrum with wavelengths just shorter than visible light

جزء من الطيف الكهرومغناطيسي بأطوال موجية أقصر قليل من الضوء المرئي



(a) Stratospheric ozone absorbs about 99% of incoming solar ultraviolet (UV) radiation, effectively shielding the surface.

(b) When stratospheric ozone is present at reduced levels, more high-energy UV radiation penetrates the atmosphere to the surface, where its presence harms organisms.

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كل ما اتوجه (انقره فترت مع الدائري كسيرة
لاكون اشتر) يكون الى unit لحاسة صفيه O₃ intensity

Dobson units (DU) were developed to measure ozone concentration:

1 DU = 0.01 mm of O₃ at 1 atm and 0°C

* Certain Value = صا

قطر الغزن لوصف

القطب الجنوبي

طبقة الاوزون

نهاد المكان

At midlatitudes the ozone concentration is typically about 350 DU, at the equator it is 250 DU, and in the Antarctic region it is only 100 DU.

خط الاستواء لأن كبر السكاليه
لا يكون كذا اعلى تراكيز للاوزون في هاتين layer



total ozone (Dobson units)

110 220 330 440 550

© Stockton Science Center

على الجوانب
كسيرة O₃ اعلى

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Ozone Depleting Substances (ODS)

تعمل على تفتيت طبقة الأوزون
 $O_3 \rightarrow O_2$

Montreal protocol

بروتوكول مونتريال

How O_3 is Depleted in the Stratosphere?

Widespread use of certain chemicals has reduced ozone levels in the stratosphere, which allows for more harmful ultraviolet radiation to reach the earth's surface

Halons, CFCs

Halone 101

102

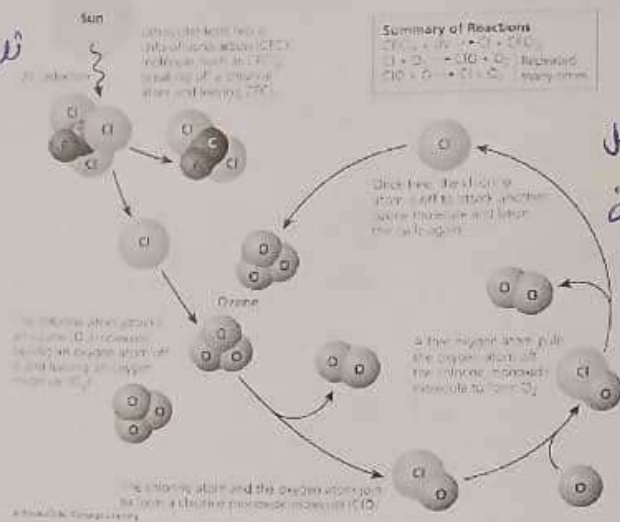
دوليت

تستخدم في

The chlorofluorocarbons (CFCs) chemicals that found wide use in aerosols and refrigeration systems and are responsible for climate change as well as the depletion of the protective ozone layer in the stratosphere.

Two of the most important CFCs are trichlorofluoromethane, $CFCl_3$, and dichlorodifluoromethane, CF_2Cl_2 , both of which are inert and not water soluble and, therefore, do not wash out of the atmosphere.

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The ozone hole & risks

There indeed was a huge monster hole right in the middle of the South Pole.



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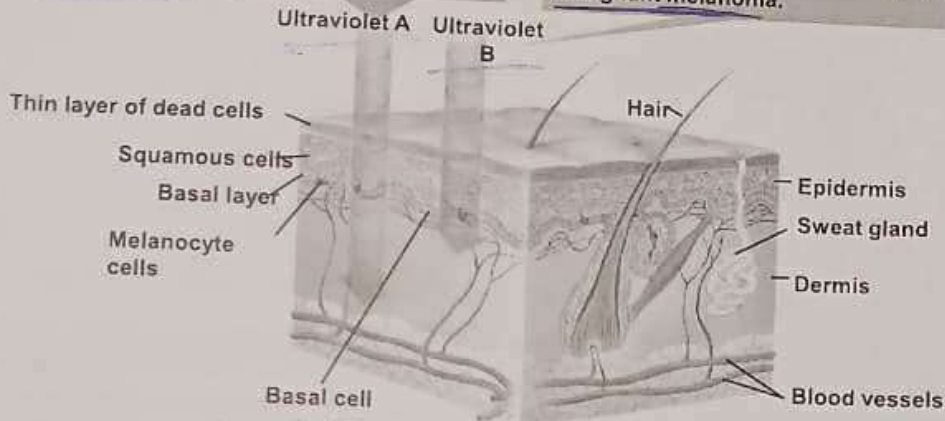
24

شعيرته بجلد

شعيرته بجلد و شحاله

This long-wavelength (low-energy) form of UV radiation causes aging of the skin, tanning, and sometimes sunburn. It penetrates deeply and may contribute to skin cancer.

This shorter-wavelength (high-energy) form of UV radiation causes sunburn, premature aging, and wrinkling. It is largely responsible for basal and squamous cell carcinomas and plays a role in malignant melanoma.



Squamous Cell Carcinoma



Basal Cell Carcinoma



Melanoma



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التغير المناخي
هو عبارة عن ارتفاع
درجة حرارة الارض
اذ الطبقة الخارجية
من الغلاف الجوي
تكون تيارات الهواء
در طبقاته

المسيرة العالمية

Climate Change

The overwhelming scientific consensus is that the earth's atmosphere is warming rapidly, mostly because of human activities, and that this will lead to significant climate change during this century.

Sept. 1979



Sept. 2007



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ارتفاع الحرارة في القمم الجليدية Temperatures in the Past ?

كل سنة
تاريخ وكي تاريخ
عينة
في CO2
هنا عينة يكون
قائمة
CO2 في الهواء
تجاري للفلسف
للأرض
تركيز CO2 على
الطبقات التي فوقه
يكون أكثر من
الطبقات التي
تحت



Scientists analyze tiny air bubbles trapped in ice cores learn about past:

- troposphere composition.
- temperature trends.
- greenhouse gas concentrations.
- solar, snowfall, and forest fire activity.

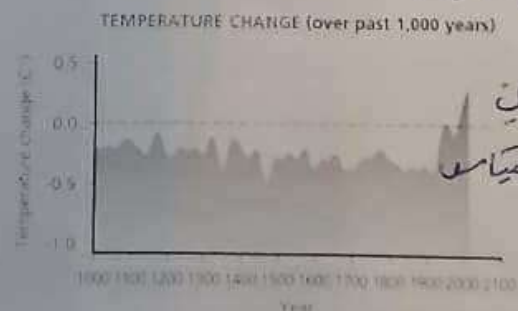
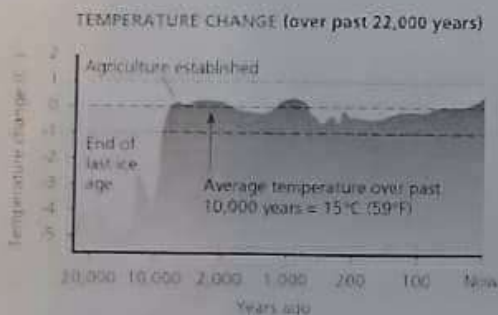
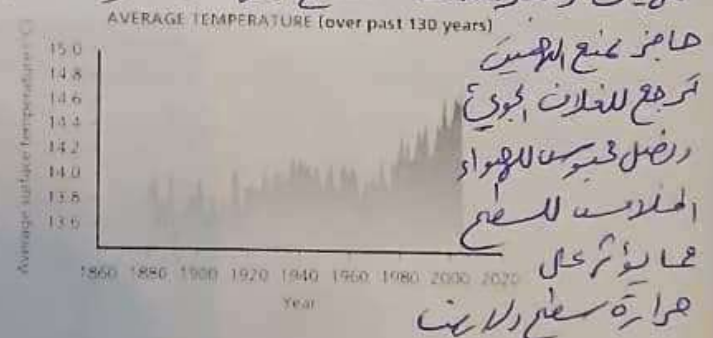
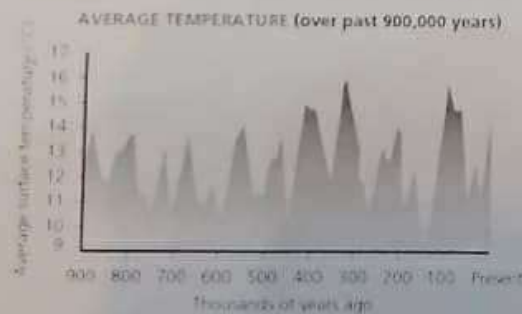
CO2 الموجود في الجو
equilibrium value
ما بينه وبين liquid لا CO2 دائما
Concentration بالاطراد معتدلة

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عدم وجود الغيوم أو هطول الذي يجعل انحرافات Estimated Changes للحسين وتغيرها في سطح الأرض وخصائصها قبل التلوث

ΔT
هنا مسؤولة
عن تغير
المناخ
increase
تلك الانحراف
السوي
الذي تحت



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Carbon Dioxide at highest levels

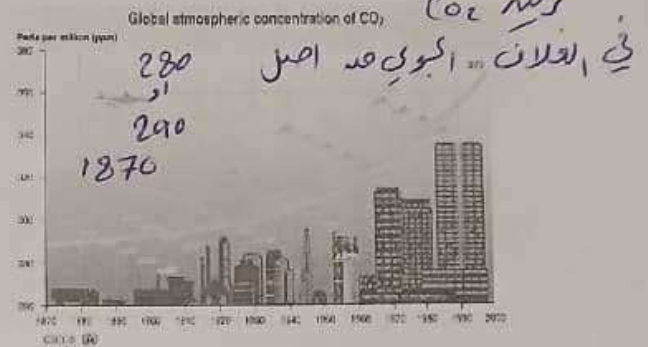


In 2005, an ice core showed that CO_2 levels in the troposphere are the highest they have been in 650,000 years.

2020 → 405

جود منه مليون

كربنة CO_2



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Climate Change Consequences

فقط زيادة

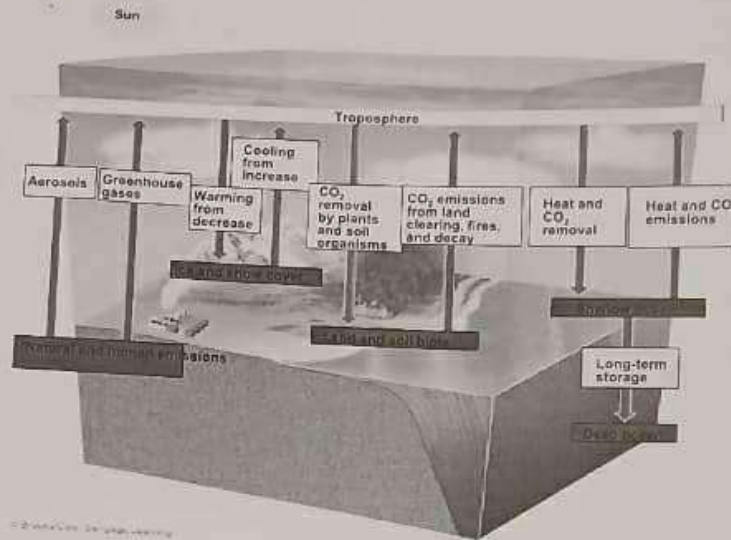
- 90–99% likely that lower atmosphere is warming
- 1906–2005: Ave. temp increased about 0.74°C
- 1970–2005: Annual greenhouse emissions up 70%
- Past 50 years: Arctic temp rising almost twice as fast as the rest of the earth
- Melting of glaciers and floating sea ice
- Prolonged droughts: increasing
- Last 100 years: sea levels rose 10–20 cm
- Warmer temperatures in Alaska, Russia, and the Arctic are melting permafrost releasing more CO_2 and CH_4 into the troposphere.

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Greenhouse Gases Effect

جدار بلاستيكي

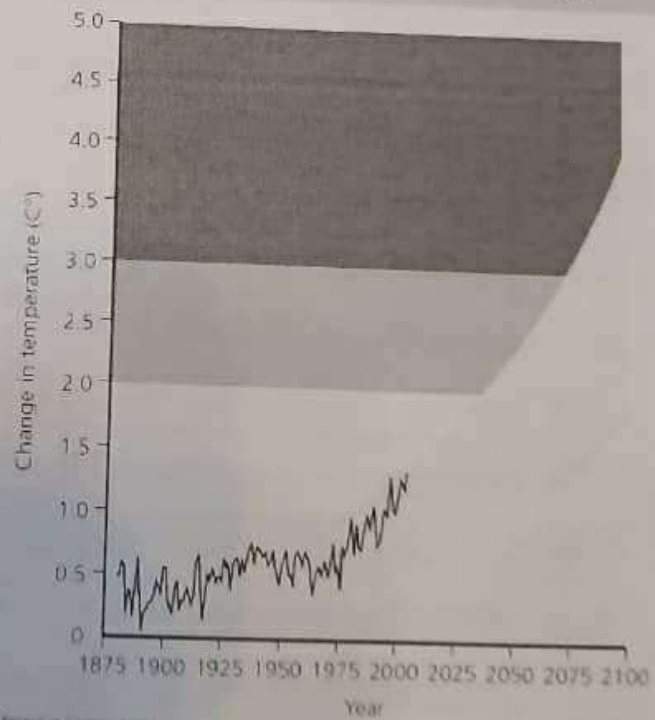


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Measured Temperature and Projected Changes

- The projected rapid change in the atmosphere's temperature during this century is very likely to increase drought and flooding, shift areas where food can be grown, raise sea levels, result in intense heat waves, and cause the premature extinction of many species.



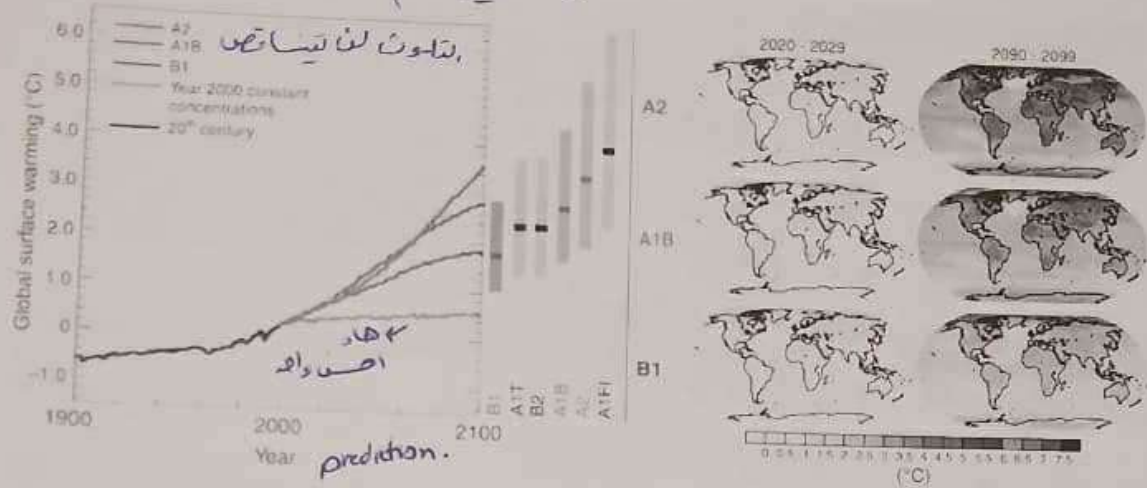
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Temperature projections

الوقت حاليًا تثبت بلونات كائنات
عليه في عام 2000



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Changing Ocean Currents

ظاهرة طين
تتميز بتأثير
حالة صلبة بالمقارنة
وتنوع دفنها وبيروت



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Treatment *علاج*
or purification *أو تنقية*

Air Pollution:

Flow *تدفق*
gas *غاز*

Control of Stationary Sources

pollutant *ملوث*
gases *غازات*, particulate *جسيمات*
matter *مادة*
emission *انبعاث*

Dr. Motasem Saidan

physical height *ارتفاع فيزيائي*

M.Saidan@gmail.com

the vertical measurement
of a person or object from
its base to its highest point

Univ. of Jordan/ Chem. Eng. Dept.

1

Control of Stationary Sources

Control of Stationary Sources:

- Particulate Matter
- Gaseous Emissions

Particulate Matter

يكون عناء كبير في عمله process
والتالي يكون efficiency اذ عناء في raw materials او reactant التي
تدخل فيه

- Most of the air pollution produced by stationary sources results from the efficiency ← incomplete combustion of fuel or industrial processing → raw materials

- The types of inorganic and organic air pollutants stationary sources emit are dependent on the specific process operations

Examples: Fossil fuel fired boilers emit ash, sulfur dioxide, nitrogen oxides and mercury, and or vanadium if contained in the fuel.

صانع المعادن Metallurgical plants can emit a variety of metal dusts, including iron oxides and sometimes fluorides and chlorides. Industrial plants manufacturing inorganic chemicals will emit various waste gases depending on their product. Odorous organic waste gases can also be emitted from organochemical and petrochemical plants.

تسبب في tank, اذا كان مفتوح او في duct, اذا كان عن فيه leak
والغازات اخطر, فواء, اللع, هيرن, اللع, فافيت مقاومة
وسيد 3 liquid ثم solid

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Vegetative Fugitive Sources.
مصادر التهرب

Industrial pollutant sources

Industrial pollutant sources such as these and others can be categorized into several groups based on their specific process operation:

Process Operations:

Process operations with incomplete chemical reactions, which include combustion due to unconverted reactants, or a reaction having a final yield that is less than expected theoretical conversion.

Atmospheric Releases:

Atmospheric releases of process's secondary components or impurities of raw materials.

Auxiliary Losses:

The auxiliary losses of compounds such as volatile organic solvents from fugitive sources or inorganics such as carbon disulfide and hydrogen sulfide in rayon production and fluorine compounds as in the production of aluminum.

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4 فيكون مائع في pollutant ناتبة في اجترابه gaseous
او لاشد في material التي تطلق drying او calcination وعادة في صانع
الاصح

لأننا نحتاج فقط على موضوعه خلال جمع أو ducts تلك وبلغ one point وهي بتركيب على unit
 حتى أغل Treatment أو على كل مصدر - ثم يقل emissions بتركيب unit حتى أغل capturing
 في pollutant minimization حتى يكون قسمة مع standard أو قبل eliminate بقايا
 at all بشكل كامل.

Waste Emissions:

Thermal نوع انه يطلق في انجوة أو unit
 Emissions of malodorous substances or oxidation compounds in the exhaust from oxidation, heating or drying processes.

Releases from these categories can originate from a variety of emission points and may not be centrally collected before entering the atmosphere.

Air release emission points, from industrial process operations, can be categorized as stack, duct, vent, fugitive and area

Fugitive emission points are release points, which are unconfined in a stack or duct before reaching the atmosphere.

Therefore, whether an emission source is considered a point or fugitive source is dependent on whether the release is confined or unconfined in a stack or duct prior to atmospheric release.

M. Saidan Booths → solvent كالأفج كمنصة 5

Vacuum System → driving Force بفعل Flow هاد، Flare
 point الذي تان يتم اجتواءم راج يطلق بالهواء، أعلقه

Blow molding → على لمبيد نيزك هو القالب، البلاستيك ماسورة في فتحة تان تأخذ شكل
 القالب بعدين بنسبة ويطبع وفت شكل عليه هاد

spray drying عبارة عن أمطاره طويلاً

20m عند فوهة في Nozzle يطبعن slurry

و drops from solid particles وفت تحت طالع صواد صافين

فتان يعلق لمبيد أي جوا، drops مقبلة

تجفيف ولازم هاد الهواء يطلق في هاد

dust ولازم أغل Treatment تان أصعب

cycles وجمع استعملت مرة ثانية

heating fluid في غل عن

هو هاد و إذا دخل مع يطلق منه جهة ثانية

ويحرقه ويزد منه يطلق منه ملازم انقله

فتان ما يكون الجو

melting وفتان

تطبع

الغالب مقبلة

لنفسه

تطبع

لنفسه

Generation Air Emission Points and Categories

Fugitive Sources	
Valves	صاير، التوربين
Pump seals	أجله
Flanges/connectors	في هاد
Compressors	مضغمة
Open ended lines	خطوط مفتوحة
Pressure relief devices	أجهزة تخفيف الضغط
Equipment cleaning/maintenance	تنظيف وصيانة المعدات

Handling, Storage, Loading	
Storage tank breathing losses	خسائر تنفس خزان التخزين
Loading/unloading	تحميل/تفريغ
Line venting	تفريغ الخطوط
Packaging and container loading	تغليف وتحميل الحاويات

Exhaust
المخارج
الغازات

تغيير عمليات الإنتاج
تطبيق تقنيات تحكم فعالة
حلول عملية

■ Pollution control can be achieved through common sense solutions such as the installation of effective control technology, changes in production processes, and the implementation of pollution prevention techniques.

معايير ISO
تطبيق تقنيات منع التلوث

■ Compliance with emission standards and the successful attainment of air quality standards depends in large part on the application of appropriate stationary source control measures.

تطبيق تدابير فاعلة للتحكم في مصادر التلوث الثابتة

Select proper control units on Air sources emissions.

particulate matter

PM Control Procedures

dispersion
انتشار

Control procedures for stationary sources of pollution include:

- the use of tall smokestacks, ^{ارتفاع 100m minimum}
- changes in plant operations, and
- installation of effective control devices

إجراءات التي يتبعها في مصادر التلوث الثابتة ؟؟
تطبيق تدابير فاعلة للتحكم في مصادر التلوث الثابتة

➤ The control strategy required for an industrial environmental impact is a four step process:

- (1) elimination of the problem source or operation,
 - (2) modification of the source operation,
 - (3) relocation of the source, and
 - (4) selection and application of the appropriate control technology.
- إزالة مصدر المشكلة أو العملية
تعديل عمليات المصدر
نقل المصدر
اختيار وتطبيق التكنولوجيا المناسبة للتحكم

Don't mix the waste → liquid waste or solid waste

لا تخلط النفايات السائلة مع النفايات الصلبة

stacks على ارتفاعات أعلى، وإذا كان المناخ صديقاً واد $\leftarrow$ Condition $\leftarrow$ stable $\leftarrow$ يكون سهل تفرق
 لأنه بسبب توهين الهواء، لا تكون التراكيز منخفضة، فمارع تأثير على الناس
 dispersion. Disperses it

Exhaust Stacks

end of pipe $\leftarrow$ نهاية $\leftarrow$ pipe $\leftarrow$ أنبوب $\leftarrow$ cleaner production $\leftarrow$ إنتاج أنظف

Exhaust stacks do not reduce emissions from a stationary source; rather they reduce the local effects of the pollution by elevating the exhaust stream to a point where it can be more effectively dispersed.

Solution $\leftarrow$ حل $\leftarrow$ ما من تفرق، موثوق به، لا تمان. ١٠ سنوات قدام

High exhaust stacks were an inexpensive solution in the absence of expensive control technology. For years elevated stacks were used with the nearby communities in mind. A belief was widely held that elevated stacks reduced the likelihood that pollutants would have any effect on neighboring populations.

Utility and smelter operations have traditionally used tall stacks (200m to 400m) in order to reduce the amount of ground-level concentrations of sulfur dioxide (SO₂). However, this did not always eliminate the problem, but instead simply transferred it to another location.

This approach was used for many years, until concern arose over the regional and transboundary spread of harmful toxics. For example, there is great concern over the spread of acid rain from one region to another.

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Persistence
Carcinogenic

Pollutants

Plant Operations

Compliance with emission standard may require the use of control technology, but many industrial operations have reduced emissions by changing operational methods.

Some of these changes include pre-treating process materials, fuel or material substitution, and changes in the manufacturing process. As an example of how pre-treating raw materials can be an inexpensive solution to pollution control, industry has discovered that significant reductions in particulate matter and sulfur emissions can be achieved by a technique called coal washing. Pre-treating raw materials in this manner not only reduces the amount of fly ash released from coal, but it also reduces the amount of inorganic sulfur released as well.

Another way to comply with emission standards is to substitute cleaner fuels during the refining process. Natural gas and low-sulfur fuel oil are just two examples of fuels that emit less pollution during combustion. However, cleaner fuels can be more expensive and can increase national reliance on foreign fuel sources.

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not effective

الصيداء، لدورته تقال صد اشماليه، ومنع حوادث الاربعين صد ضلالت انتانت افعال في
المعدان قبل حدوثه.

- Reduction in emissions from stationary sources can also be accomplished through increased attention to **plant maintenance**. 45/20

الحمد لله

فقطه لیلوت نسو
سور صبیانه افعدا

Plants that release significant quantities of pollutants into the environment frequently do so as the result of improperly maintained equipment. This is especially true of combustion equipment.

أصوات

ISO requirement

- فصل اول در بیان
مبانی کلی
ادبیات

Adequately scheduled maintenance must be performed to reduce both the exhaust and amount of fugitive emissions released from vats, valves, and transmission lines. **Periodic maintenance** also reduces the likelihood of spill-related accidents by discovering faulty equipment before problems occur.

اقتضاه

Adsorption = capturing of pollutant by solid material
 Adsorbent = solid material, Adsorbate = pollutant
 Fixed bed = solid media, Fluidized bed = solid particle in motion

هو عبارة عن عزل التلوث عن Capturing في Solid part أو gases Absorption في liquid part
يكون على شكل فوم أو Exhaust gas خارج من bottom يلتصق بجزء من pollutant أو كرات dust
أو pollutant particle size

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Control Technology

- A final way to reduce emissions from stationary sources is through the use of advanced, add-on control technology. استخدام تقنيات التحكم المتقدمة

اصفا دم تدعير

- Control devices can destroy or recover gaseous compounds or particulate matter for proper disposal or re-use. → إعادة استغلال

النقاء

- The pollution control operations used to destroy or capture gases include combustion, adsorption, absorption, and condensation. → ^{solid material} gas → liquid

solid material

→ gas → liquid

Heat exchanger

هاتقان مراد

- Control devices that implement these processes include thermal incinerators, catalytic incinerators, flares, boilers, process heaters, carbon absorbers, spray towers, and surface condensers.

activate carbons

مکان اصلی
در

The most important process parameters for selecting air pollution control equipment are the exhaust gas characteristics obtained from emissions tests and process or site characteristics obtained from a field survey.

فلسفہ

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Control Devices for Particulate Emissions

➤ Technologies used to control particulate matter focus on removing particles from the effluent gas stream.
تكنولوجيا إزالة الجسيمات من تيار الغازات الناتجة

هناك طرق مختلفة
لإزالة الجسيمات

Many factors (such as particle size and chemical characteristics) determine the appropriate particulate control device for a process.

➤ Devices most commonly used devices to control particulate emissions include:

- Gravity settlers (often referred to as settling chambers)
- Mechanical collectors (cyclones)
- Electrostatic precipitators (ESPs)
- Scrubbers (Absorption)
- Fabric filters
- Hybrid systems

على استخدام كل هذين
سلسلة
من أجل كفاءة أعلى

➤ In many cases, a combination of multiple devices yields the best collection efficiency. For example, a settling chamber can be used to remove large particles from the exhaust stream before it enters an electrostatic precipitator where smaller particles are removed.

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EQUIPMENT

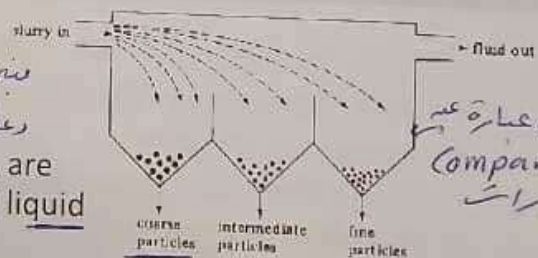
Gravity settlers (often referred to as settling chambers)

تستخدم الجاذبية لفصل الجسيمات عن الغازات
لا تحتاج سرعة عالية

physical system
(mechanical)
pumps, heating, cooling, no need any electric power.

Gravity settlers, or gravity settling chambers, are used industrially for the removal of solid and liquid waste materials from gaseous streams.

- Advantages accounting for their use are simple construction, low initial cost and maintenance, low pressure losses, and simple disposal of waste materials.



Mechanical collectors (cyclones)

Flow gas have high temperature
هناك تيارات غازية عالية الحرارة

hydrocyclones → liquid cyclones

- Centrifugal separators, commonly referred to as cyclones, are widely used in industry for the removal of solid and liquid particles (or particulates) from gas streams.

gravity → particles



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- Fabric filters (bag houses)** *cloth compartment* *High pressure*
- Filtration process may be conducted in many different types of fabric filters. Differences may be related to:
 - Type of fabric
 - Cleaning mechanism
 - Equipment
 - Mode of operation
- The Temperature should be moderate.*

- Gases to be cleaned can be either "pushed" or "pulled" through the bag house.
- In the pressure system (push through) the gases may enter through the cleanout, hopper in the bottom or through the top of the bags.
- In the suction type (pull through) the dirty gases are usually forced through the inside of the bag and exit through the outside.

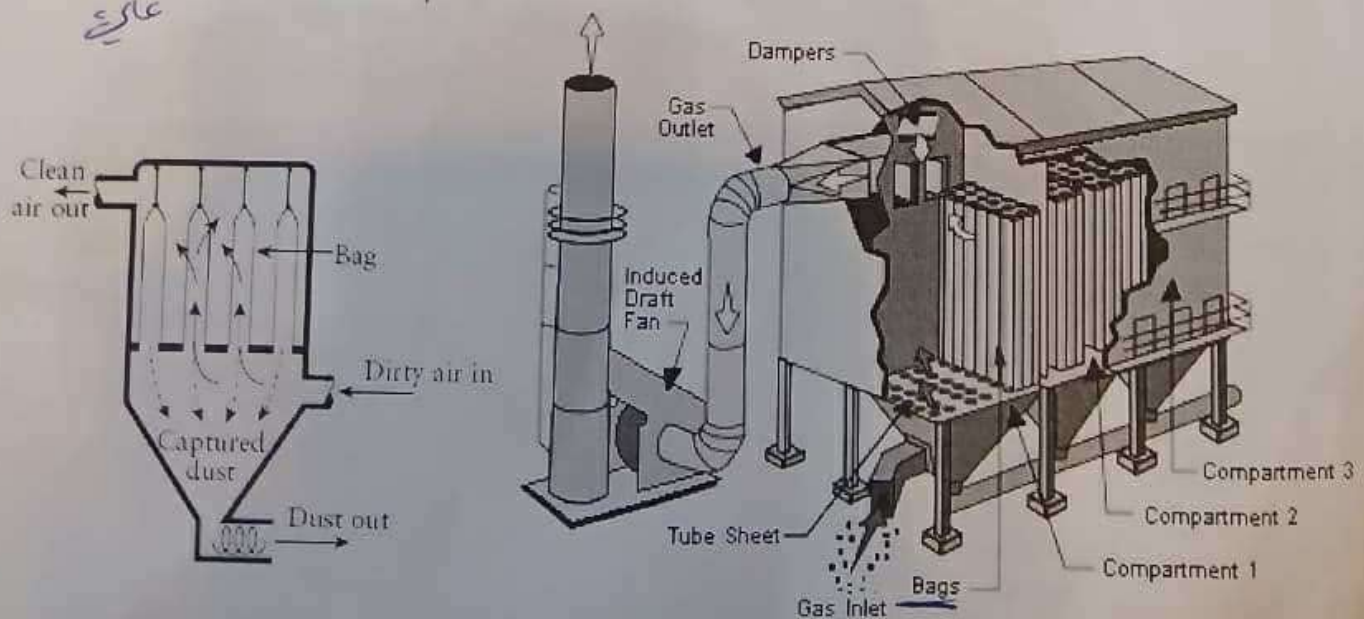
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Baghouse Filter *Vacuum*

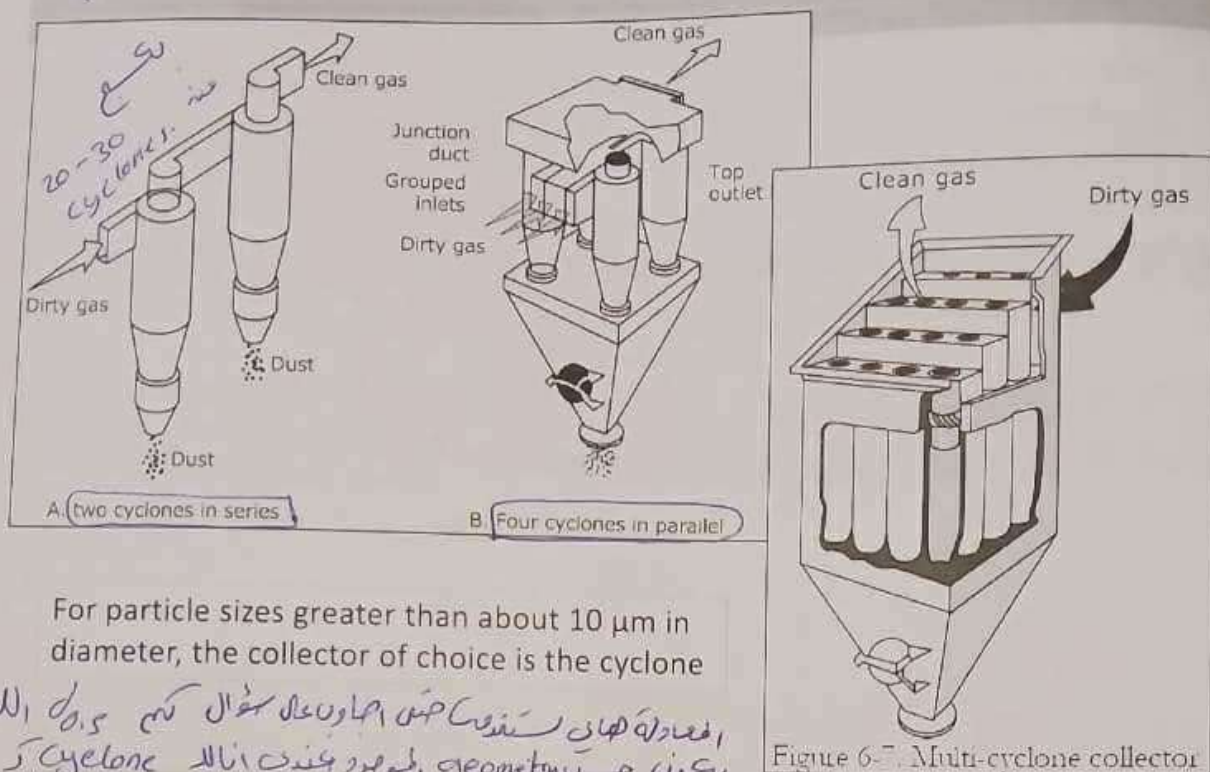
- similar to conventional home vacuum cleaner
- Efficiency:**
 - >99.5% for $<1 \mu\text{m}$ diameter
 - >99.8% for $>5 \mu\text{m}$ diameter

Figure 13. Reverse Air Fabric Filter



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For particle sizes greater than about 10 μm in diameter, the collector of choice is the cyclone

الهندسة هي فن اختيار الحل الأمثل للمشكلة الهندسية
تكون من geometry الموهبة عند انال cyclone design
M.saidan efficiency

rectangular, مثلث و متصل من دون زوايا حادة friction عالي

The particle size collected with 50% efficiency, termed the cut diameter: *radius of inlet*

$$d_{0.5} = \left[\frac{9\mu B^2 H}{\rho_p Q_g \theta} \right]^{1/2}$$

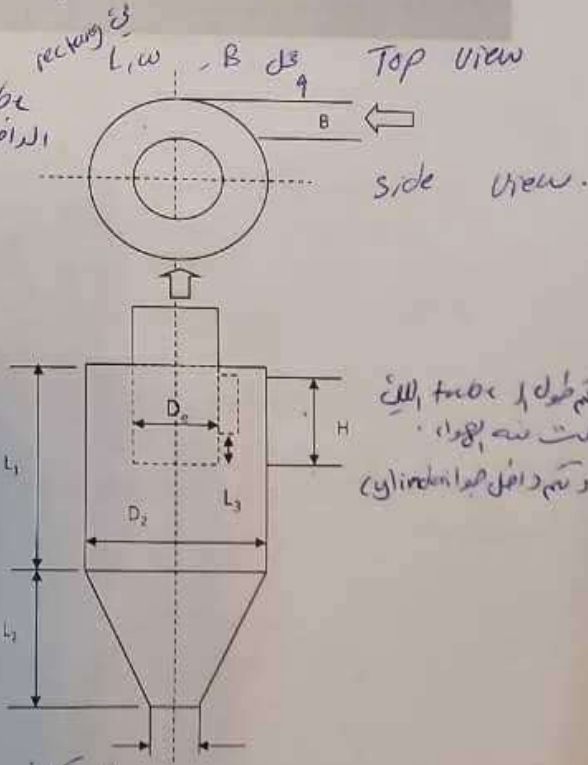
height of cylindrical shape $\theta = \frac{\pi}{H} (2L_1 + L_2)$ height of ~~cylindrical~~ conical shape

$d_{0.5}$ = cut diameter at 50% removal
 μ = dynamic viscosity of gas, Pa-s
 B = width, m
 H = height, m
 ρ_p = particle density, kg/m³
 Q_g = gas flow rate, m³/s
 θ = effective number of turns

B = width, m

ρ_p = particle density, kg/m³

o = effective number of turns

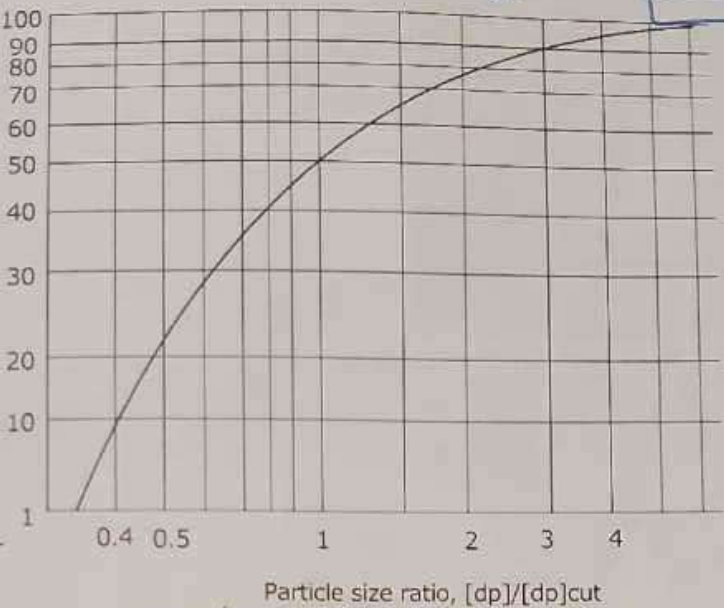


M.saidan → number of rotations of cylindrical part of particle.

Efficiency, η (%) of removed
 90% efficiency
 $\frac{dp}{dp_{0.5}}$
 $3 = \frac{dp}{dp_{0.5}}$
 $3 = \frac{10 \text{ microns}}{dp_{0.5}}$
 $dp_{0.5} = 3.33 \text{ microns}$
 train and error
 H, L_1, L_2
 Data Base
 standard.

Chart مع الملاحظات

n	$d_p/d_{p,0.5}$	d_p
75	10	
30	20	
70		



مع مقياس بالعمود
الحساب

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diameter
 flow gas.
 removal.
 $d_{0.5}$
 التي
 في
 عت
 اياه

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Example

Given:
 $D_2 = 0.5 \text{ m}$
 $Q_g = 4 \text{ m}^3/\text{s}$
 $T = 25^\circ\text{C}$
 $\rho_p = 800 \text{ kg/m}^3$

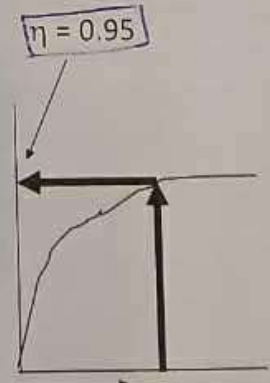
For standard Cyclone:
 $B = 0.25 D_2 = 0.13 \text{ m}$
 $H = 0.5 D_2 = 0.25 \text{ m}$
 $L_1 = L_2 = 2 D_2 = 1 \text{ m}$

$$\theta = \frac{\pi}{0.25} (2(1) + 1) = 37.7$$

Q = What is the removal efficiency
 for particles with ave
 diameter of $10 \mu\text{m}$?

$$d_{0.5} = \left[\frac{9(18.5 \times 10^{-6})(0.13)^2(0.25)}{(800)(4)(37.7)} \right]^{0.5} = 2.41 \times 10^{-6}$$

$$= 2.41(\mu\text{m})$$



@ $d = 10 \mu\text{m}$
 $\frac{d}{d_{0.5}} = \frac{10}{2.41} = 4.15$
 removal

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Electrostatic Precipitator (ESP)

- high efficiency, dry collector of particulates
- high electrical direct current potential (30-75 kV)

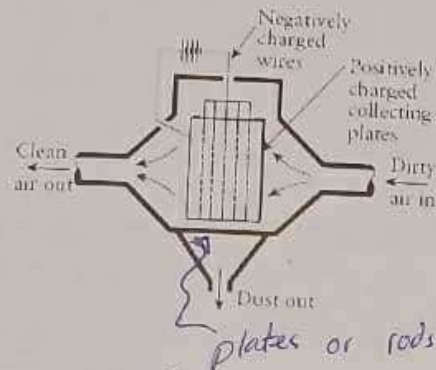
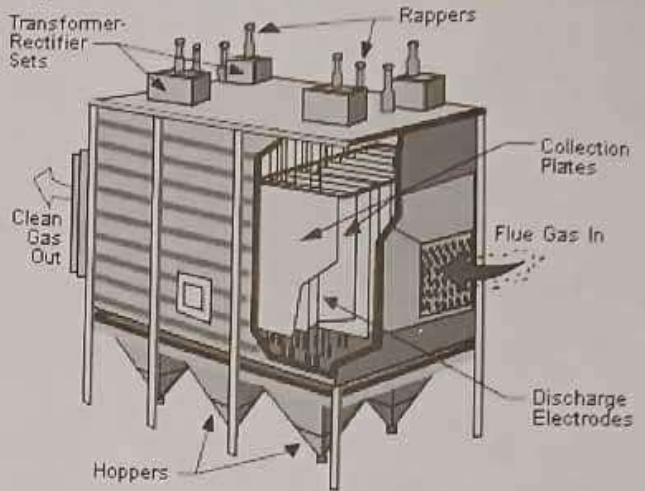


Figure 9. Conventional Electrostatic Precipitator



M.saidan يا نقل vibration من تحت الى اعلى
او تدفع على سطح من تحت الى اعلى

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Electrostatic precipitators (ESPs)

- They are satisfactory devices for removing small particles from moving gas streams at high collection efficiencies. They have been used almost universally in power plants for removing fly ash from the gases prior to discharge.

Two major types of high-voltage ESP configurations currently used are tubular and plate. Tubular precipitators consist of cylindrical collection tubes with discharge electrodes located on the axis of the cylinder. Vast majority of ESPs installed are of the plate type.

- Collected particles are usually removed by rapping.



افضل الهواء من فوق الى اسفل
الغالبية العظمى من ESP هي من نوع plate
بالقوة

Scrubbers (venturi scrubbers)

liquid drop capturing → dust or particulate matter
القطرات السائلة بالهواء

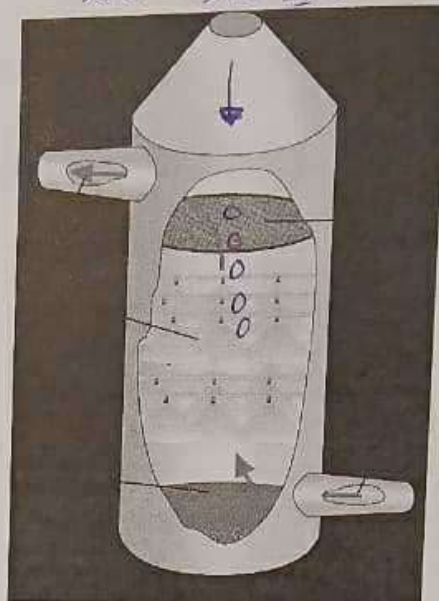
هون مع غبار شامان عار

Wet scrubbing involves the technique of bringing a contaminated gas stream into intimate contact with a liquid.

Wet scrubbers include all the various types of gas absorption equipment.

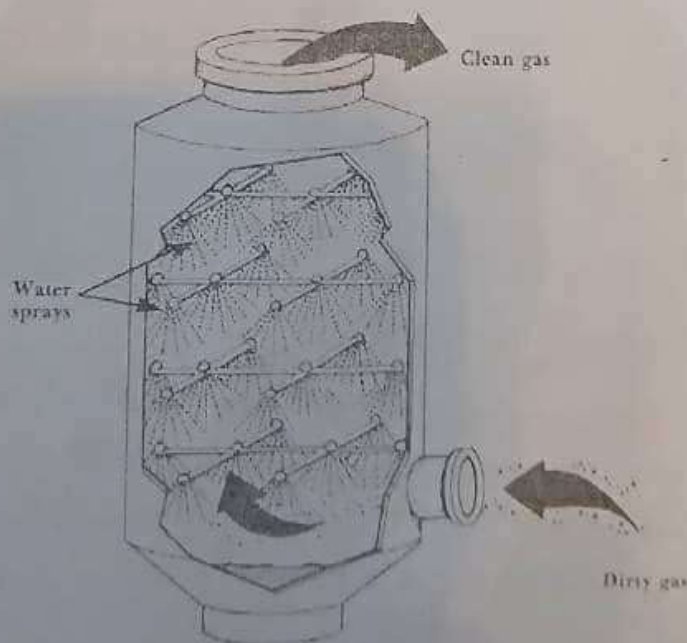
The term "scrubber" will be restricted to those systems which utilize a liquid, usually water, to achieve or assist in the removal of particulate matter from a carrier gas stream.

هذا الجهاز كفاءة عالية في إزالة الجسيمات من الغاز الناقل، فعملية إزالة الجسيمات من الغاز الناقل تتم باستخدام الماء، عادةً، كسائل تنظيف. هذه العملية تسمى scrubbing.

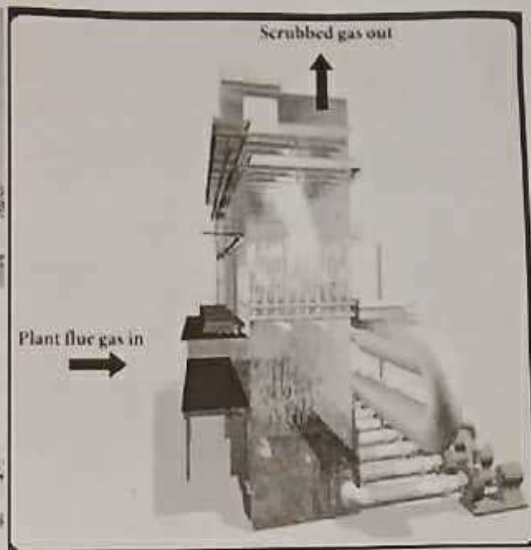


الهواء
من قبل
من
Bottom
ورفع
من
Top

مقت داخل الجسيمات
عملية



Scrubbers Absorption system



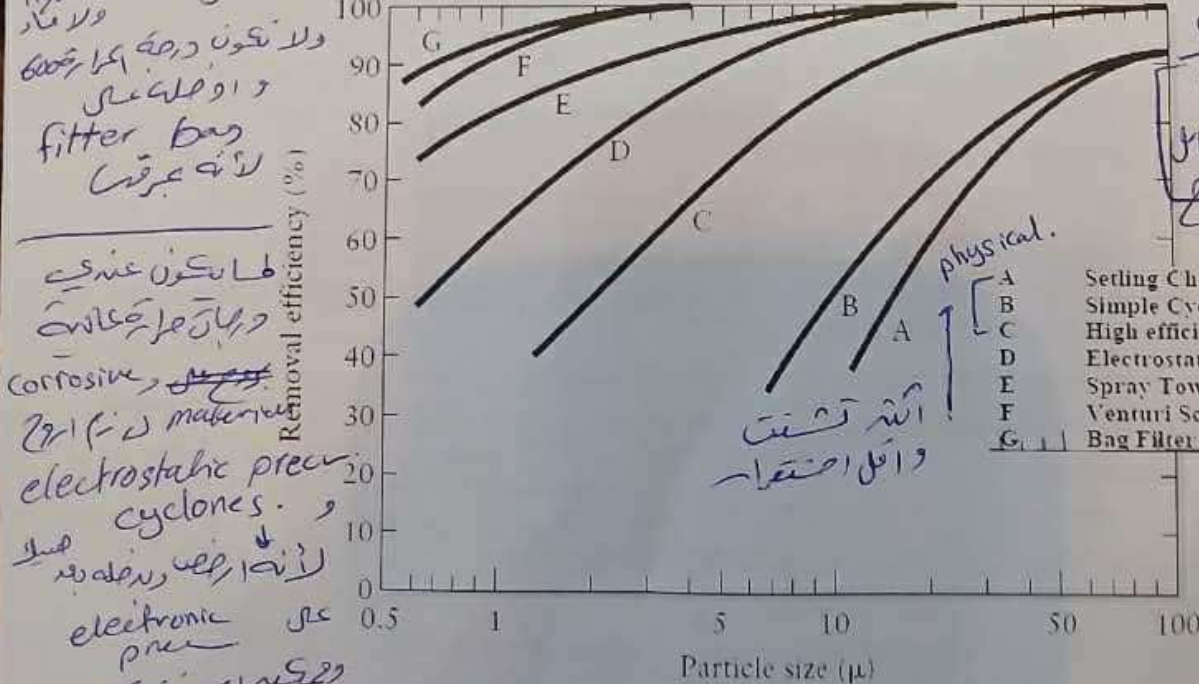
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إذا كان Flow gas الحرارة moderate يعني أقل من 250 °C، فإن flow path بإمكانه أن يجمع على (التيار) ليتم تنظيفه moderate. إذا كان Flow gas الحرارة moderate.

Comparison of Air Pollution Control Devices

وإذا كان Flow gas الحرارة moderate يعني أقل من 250 °C، فإن flow path بإمكانه أن يجمع على (التيار) ليتم تنظيفه moderate. إذا كان Flow gas الحرارة moderate.



Qualitative or Orientation.

لا يمكن معرفة نوعية المواد التي تدخل في العملية، ولكن يمكن معرفة نوعية المواد التي تخرج.

لأنها عملية كيميائية.

physical + electrical

physical

electrical

chemical

particle size

في هذا المخطط

نرى أن كفاءة إزالة الجسيمات

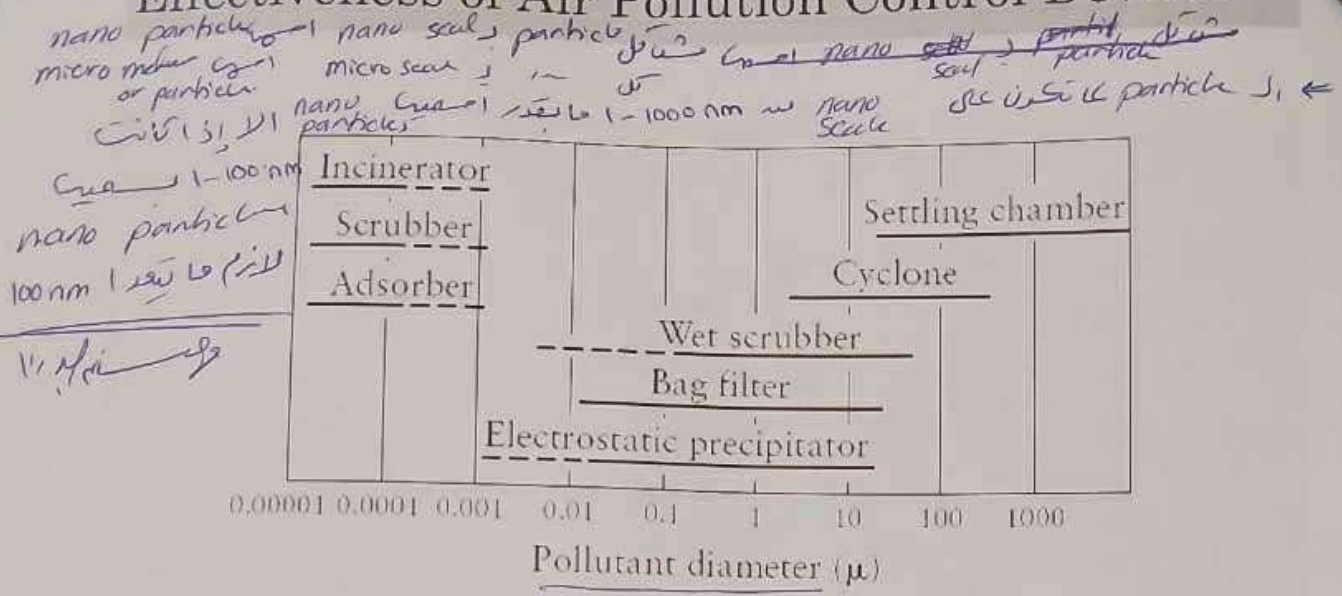
تزداد مع زيادة حجم الجسيمات.

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nano scale for nano particle.
 nano scale $\Rightarrow 1 \times 10^9$ scale unit of length.
 nano particle $\Rightarrow$ particle in nano At nano scale

Effectiveness of Air Pollution Control Devices



Gaseous Emissions

The preferred method for controlling gaseous pollutants is with add-on control devices used to destroy or recover the pollutant.

Control Equipment for Gaseous Emissions:

- Thermal Incinerators
- Catalytic Incinerators
- Flares
- Boilers and Process Heaters
- Carbon Adsorbers
- Absorbers
- Condensers

Combustion as purpose
 Heat Content
 fuel

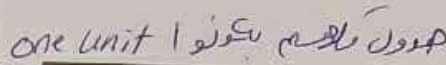
Thermally
 Adsorber
 solid matter

Absorber
 liquid material
 active matter

Convert from gases to liquid.

Boiler. نفوس نظام.

- Complete Combustion.



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فلا حول ولا قوة الا بالله

- 1000c no rule
complete composition

- للتزكية بعد لقاء القارئ العام بعد ثمانية دهر

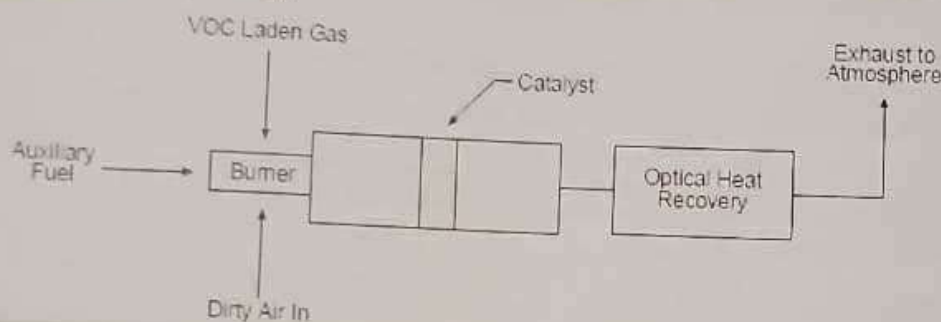
- هذا تان عن halogenated
organics
الافرع (عاصر) (الزئبق)
M. Saidan
يكون فيه 1000-10000 غير صلب
وتتبع للمو وتكون للقتل
persistent

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Catalytic Incinerators

Catalytic incinerators operate very similar to thermal incinerators. The primary difference (is that the gas, after passing through the flame area, passes through a catalyst bed to destroy essentially any oxidizable compound in an air stream.

The catalyst has the effect of increasing the oxidation reaction rate, enabling conversion at lower reaction temperatures than in thermal incinerator units. Therefore, catalysts also allow for smaller incinerator size. Catalysts typically used for VOC incineration include platinum and palladium. Other formulations include metal oxides, which are used for gas streams containing chlorinated compounds (EPA, 1998).



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- The waste stream must be preheated to a temperature sufficiently high (usually from 300 to 900°F) to initiate the oxidation reactions.
- Catalyst Temperatures Required for Oxidizing 80% of Inlet VOC to CO₂, °F for Two Catalysts

Compound	Temperature, °F	
	CO ₃ O ₄	Pt - Honeycomb
acrolein	382	294
n-butanol	413	440
n-propylamine	460	489
toluene	476	373
n-butyric acid	517	451
1, 1, 1-trichloroethane	661	661
dimethyl sulfide	-	512

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use for low recyclable value difficult to combust.

Flares

(VOCs) ^{مواد التي لا يمكن استخدامها}
 (غير قابلة لإعادة الاستخدام) ^{في الإنتاج}

plume

- Flaring is a VOC combustion control process in which the VOCs are piped to a remote, usually elevated, location and burned in an open flame in the open air using a specially designed burner tip, auxiliary fuel, and steam or air to promote mixing for nearly complete ($>98\%$) VOC destruction.

- Flares are typically used as a last resort to dispose of gases that are of little recyclable value or are not easily combustible.

Gases flared from refineries, petroleum production, and the chemical industry are composed largely of low molecular weight VOC and have high heating values.

As in all combustion processes, an adequate air supply and good mixing are required to complete combustion and minimize smoke. The various flare designs differ primarily by their ability to mix air with the combustibles.

بدون كل هذه المستويات على compartment أي اشتراك في عائل كـ liquid (moisture) أو solid. بدونه على unit ثانية (موضعية فاد) ويكون زي نظام الشفوية الدراجة نقل dubbling

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43 ^{مضان} حافظ على ^{مضان} flow rate و ^{مضان} pressure ثابت ^{مضان} VOC مع أي

والغاز هاد عادة يكون fuel.

اشترط فيه (نظام فلترة)

مضان، مجلة ما تقع تحت مضان هيك بعلم كـ شوي

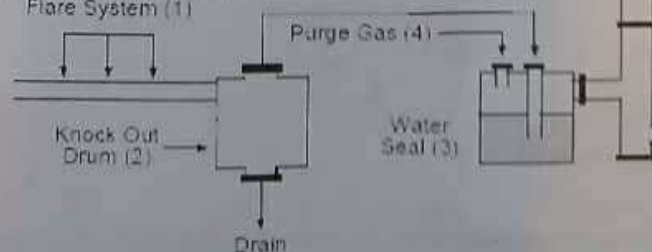
Steam-assisted Elevated Flare System

Version gas.



plume

Gas Collection Elevated Flare System (1)



بال Top يكون على steam دتفان صوا steam بيت high Temp and mixing efficiency of combustion high

الهدف من
 - filtration
 - To control pressure.
 - fuel

ادفعه لـ version غاز مضان (شوية) نقل شفوية VOCs مضان و (مضان) ادفعه version gas.

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شعلة غير مساعدة

- The non-assisted flare consists of a flare tip without any auxiliary provision for enhancing the mixing of air into its flame. Its use is limited to gas streams that have low heat content and low carbon/hydrogen ratio that burn readily without producing smoke. These streams require less air for complete combustion have lower combustion temperatures.

ليكون الشعلة غير مساعدة عند طوف شعلة بدون اي تجهيزات، انها صيغت لتعزز خلط الهواء في الشعلة
ليقتصر استعماله على تيارات الغاز منخفضة الحرارة ونسبة الكربون الى الهيدروجين ورائحة منخفضة
في الهواء دون امداد الاكسجين. وتطلب التيارات هواء أقل مما تتطلبه للارتعاش، وبالتالي
وتنقسم بدرجات حرارة اقل من تلك التي تتطلبها

الهيدروجين والوقود fuel للمرق

Boilers and Process Heaters

الوقود مع قضيب
الوقود
الوقود

- Boilers and process heaters are commonly used by production facilities to generate heat and power.

المساهمة في عمليات التصنيع

- Although their primary purpose to contribute to plant operations, they can also be used quite effectively as a pollution control device by recycling the pollutant for fuel.

عدم تدوير
الوقود كوقود

- However, the only pollutants that can be used for fuel are those that do not affect the performance of the burner unit. For example, an exhaust stream can be used as supplementary fuel, but only if its fuel value is sufficient to maintain the combustion process. All volatile organic compounds (VOCs) have different heating values. If the pollutant stream is large and the heating value is high, the exhaust can be a primary source of fuel for the plant.

يمكن استخدامه
تكون قيمته
الوقود بكمية كافية
للمساهمة في
العمليات

يمكن أن يكون الوقود في مصنع