Course 288 Petroleum Refining 2017

# **PROCESSES** CONVERSION AND BLENDING



The National Air Compliance Training Program

# **Petroleum Refining Process**

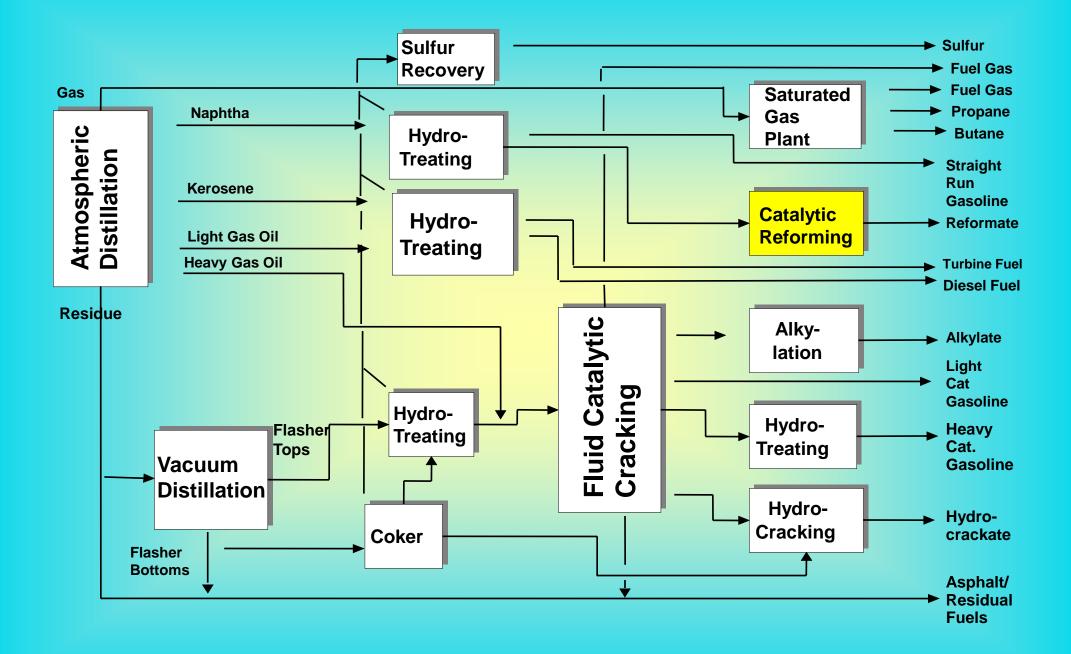
- Separation
- Treatment
- <u>Conversion</u>
- Blending

# **Types of Conversion Processes**

- Property Change
  - Catalytic Reforming
  - Isomerization
- Build-Up
  - Alkylation
- Break-Up
  - Fluid Catalytic Cracking
  - Hydrocracking
  - Coking
  - Visbreaking

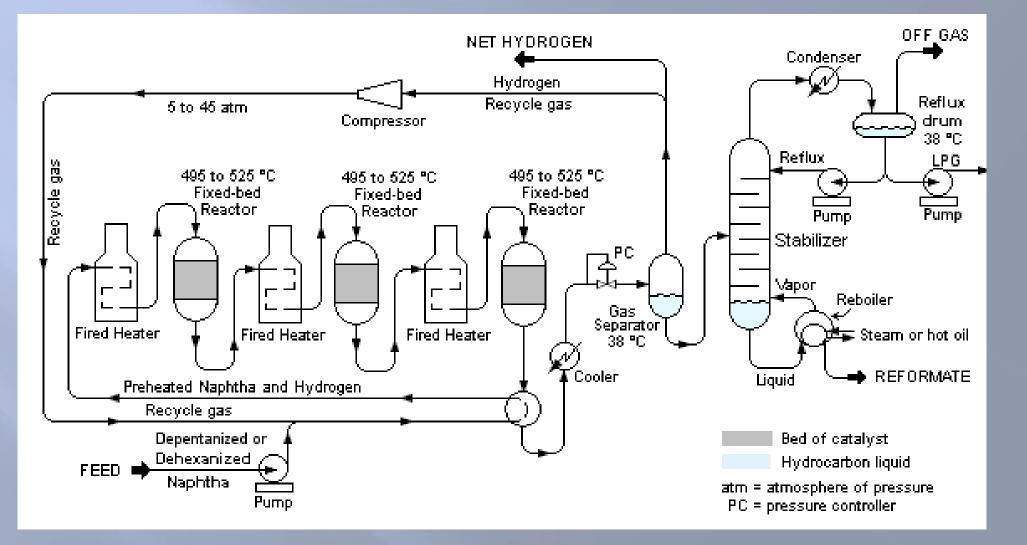
PURPOSE: Increase octane level of C3-C12 molecules by reshaping through use of a catalyst.





- REACTION: An endothermic process used to convert straight chained molecules to branched or cyclic/ring structures in a series of reactions. Takes place on a fixed bed of catalyst. Usually preceded by hydrotreating to protect the catalyst.
- PRODUCT: Reformate and H<sub>2</sub>

Page 51 of the Student Handbook



#### **Catalytic Reformer**



# Catalytic Reformer -Reactor Pre-Heat Furnaces



**REGULATIONS:** 

#### **INSPECTION POINTS:**

- Heaters
- Fugitive VOC's
- Cooling Towers
- Hazardous Waste

#### Isomerization

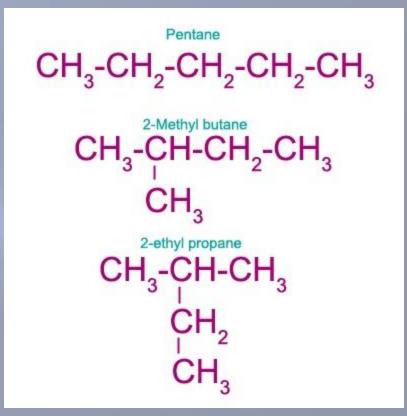
- PURPOSE: To increase the octane of a
  light hydrocarbon (sweet pen/hex) by
  rearranging molecules. Can also be used to
  provide feedstock to other processes such
  as conversion of n-butane to i-butane.
- REACTION: Molecular structures are changed over a solid or fixed bed of catalyst in the presence of hydrogen.

#### **Isomerization** Butane Isomerization Plant



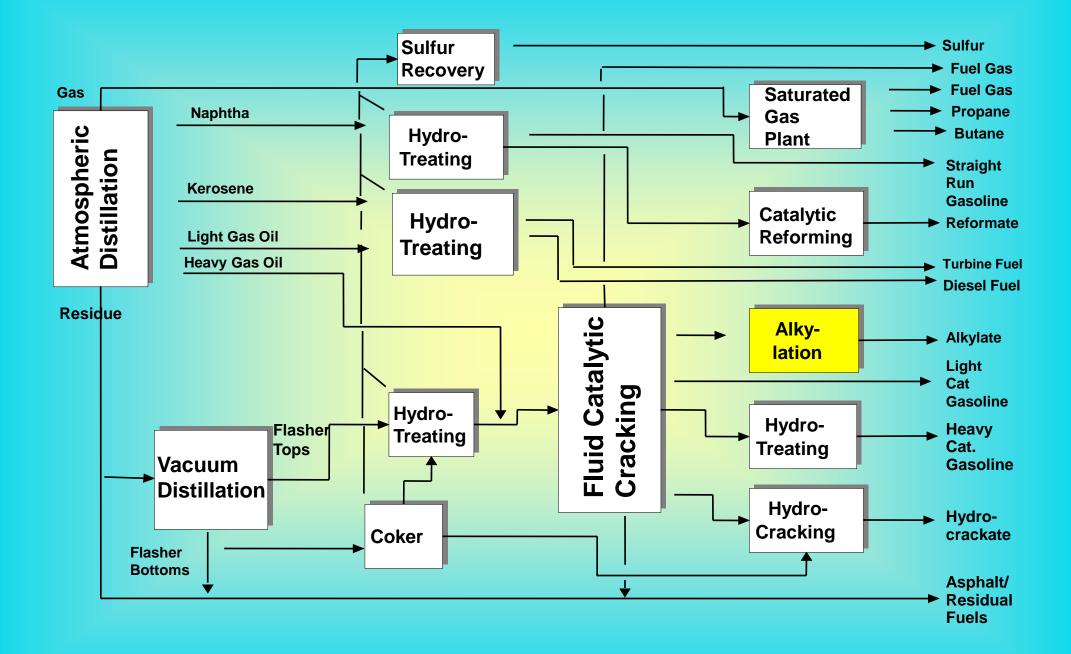
#### Property Change Isomerization

Rearrange atoms in low-octane normal parafins (30-35 octane) into higher octane isoparafins (65 octane) in the presence of hydrogen (H2) and hydrochloric acid (HCL).



# Alkylation

PURPOSE: Smaller molecules have higher vapor pressure therefore can not be blended into gasoline.
This process combines two small molecules into a larger high octane molecule.



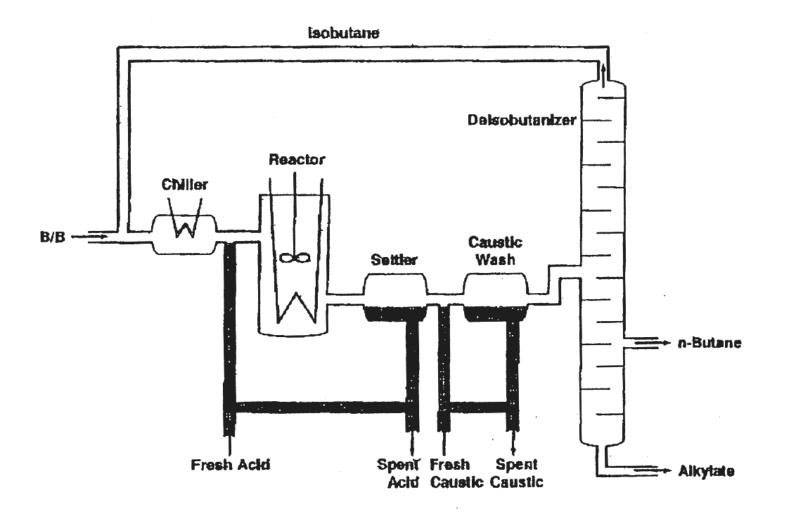
# Alkylation

REACTION: Isobutane (iso-C4) and isobutylene (unsaturated C4) are reacted to form iso-octane in the presence of a acid catalyst. The catalyst can be either hydroflouric acid or sulfuric acid.

#### Alkylation

- What does that mean?
- Shorter chain gasses: butene or propene and isobutane (predominately produced in the FCC) to make high octane gasoline

#### **Sulfuric Acid Alkylation**



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#### **Sulfuric Acid Alkylation Unit**

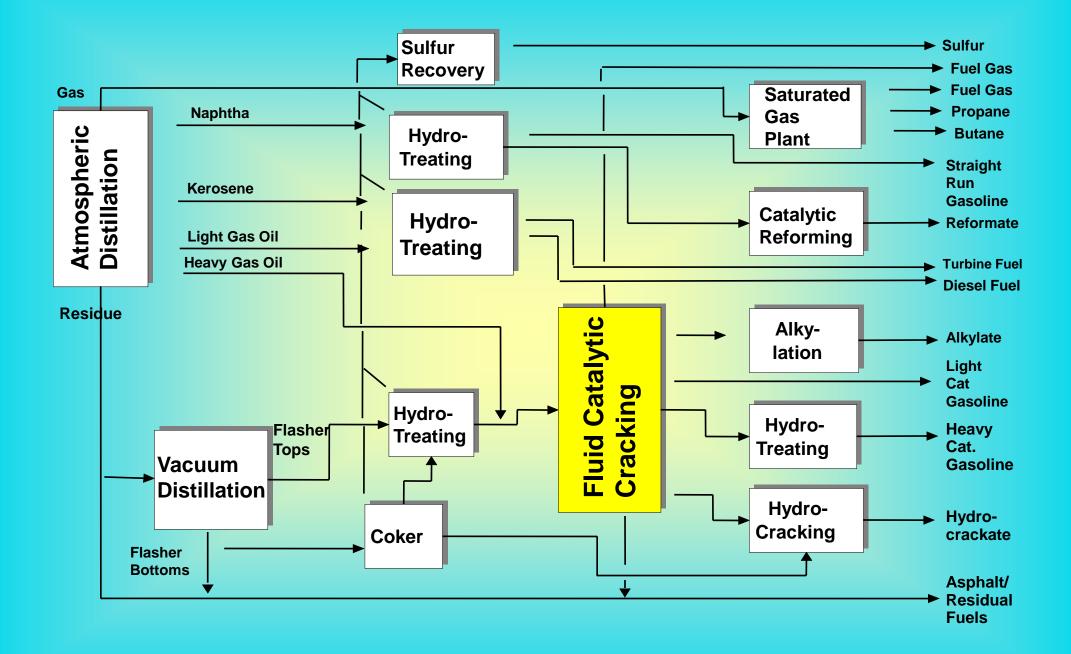


### **Catalytic Polymerization**

- Also combines smaller HCs to create a more valuable HC
- I.e., Combine three C3s to form one C9

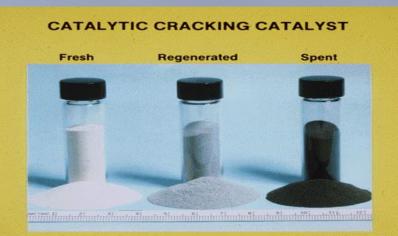


PURPOSE: To convert the heavier crude fractions to usable fractions that can be blended into gasoline, jet fuel or diesel.



REACTION: Heavy crude fractions (gas-oils) are cracked to form lighter molecules and carbon in the presence of a catalyst. The carbon is burned off the catalyst which is then recycled back into the process. The cracked products are condensed and distilled into blending fractions.

CATALYST: The catalyst used is a zeolite material (fine powder). This material is used through the process in a fluidized form.
Once the carbon is burned off the catalyst then any particulate matter needs to be removed from the flue gas stream.



**Crystalline Silica/Alumina Zeolite** 

#### **Gas-oil Feeds**

- Atmospheric gas-oil from atmospheric crude tower
- Vacuum gas-oil
- Coker gas-oil
- Unconverted oil (cycle oil) from FCC
- Light gas-oil
- Heavy gas-oil
- Hydrotreated gas-oil



# **Cracking Products**

- Ethane
- Propane, Proplyene
- Butane, Butene
- Gasoline
- Light Gas Oil



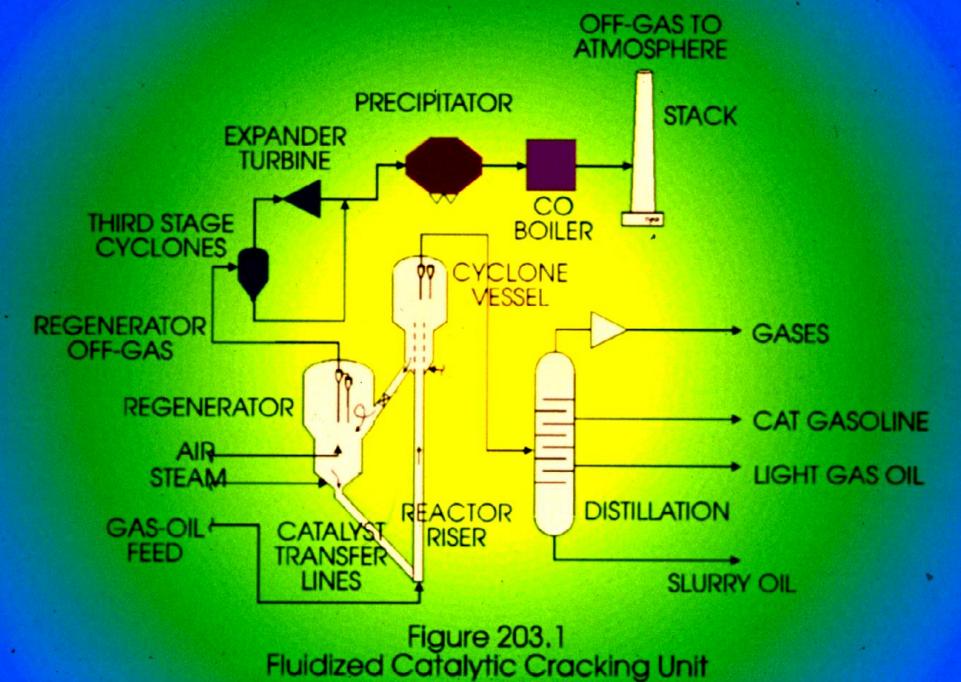
# Fluid Catalytic Cracker



# **Primary FCC Emissions**

- Particulates (Cat. Fines)
- CO
- SOx
- NOx





PARTICULATE CONTROL TECHNOLOGY: A
 cyclones with electrostatic precipitator or bag house
 are generally used to control the particulate matter.
 Wet scrubbers (also good for SOx) may also be used.

- NOx control technology: SCR
- SOx control technology: Feed desulfurization
- CO control technology: High temperature regeneration, or CO Boilers

#### **Catalytic Cracker With Scrubber**



# Electrostatic Precipitator on Fluid Catalytic Cracking Unit



### **Electrical Field Generation**

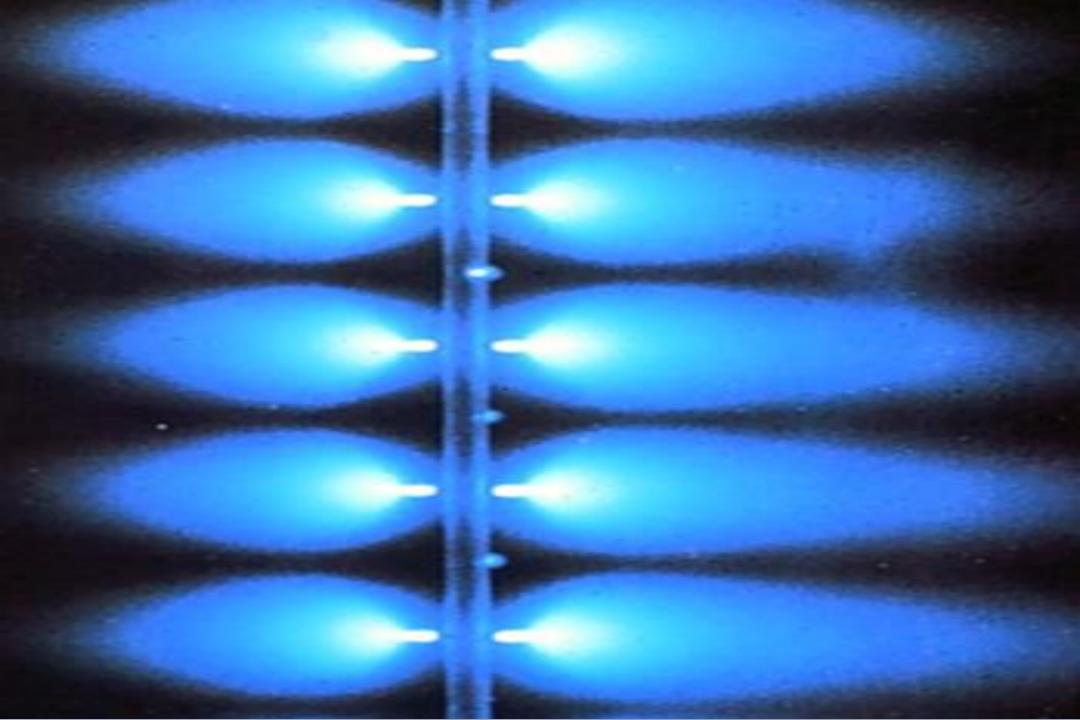
**Collection Electrode** 

Chaiging and Migration

**Electrical Field** 

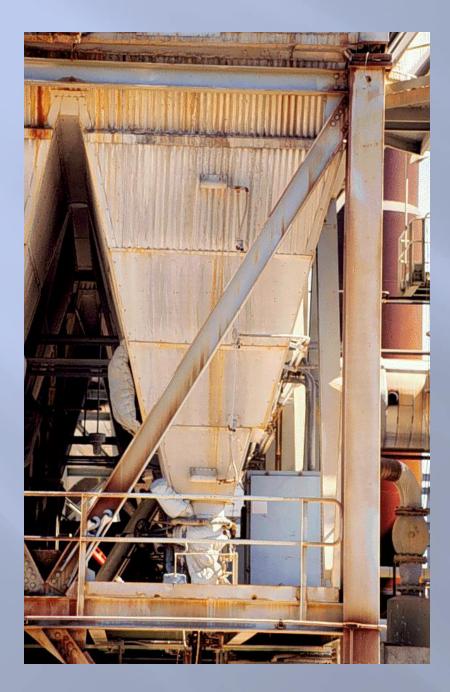
**Discharge Electrode** 

**Collection Electrode** 



#### **Electrostatic Precipitators**

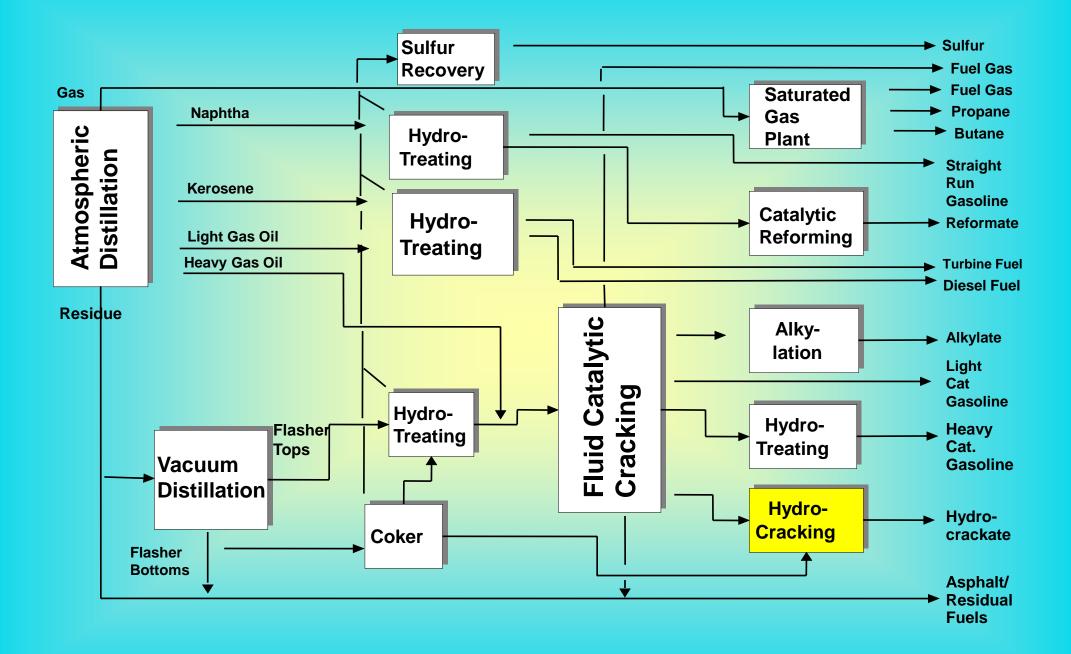




# Catalyst Fines Hopper

#### Hydrocracking

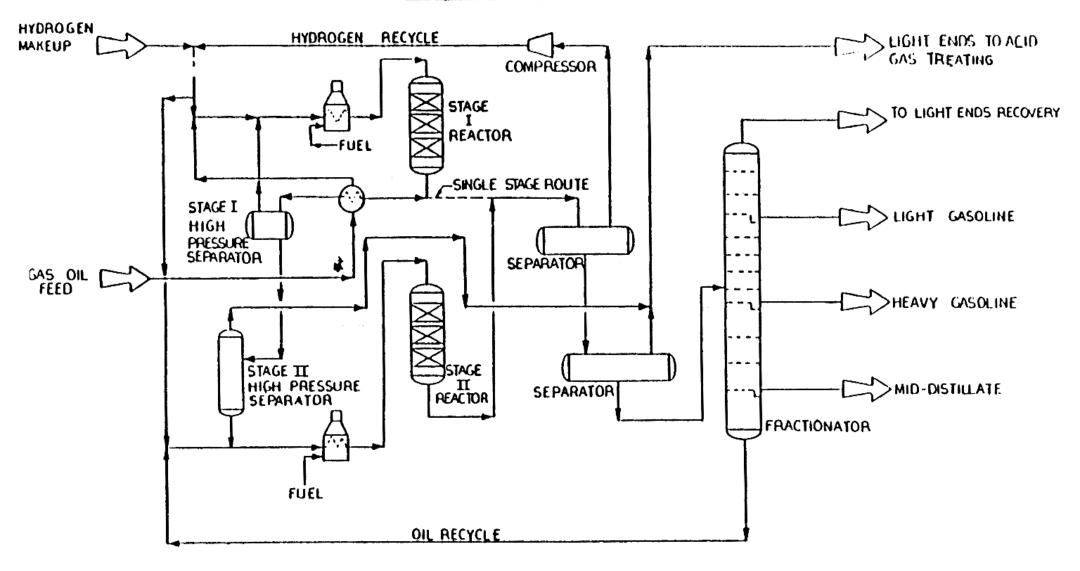
- PURPOSE: To convert heavier crude
  fractions into components that can be
  blended into gasoline and jet fuel.
- REACTION: Larger molecules are cracked
  into lighter components in the presence of a
  fixed bed catalyst, hydrogen, and high
  temperature and pressure.
- Useful for sour feedstocks



#### Hydrocracking

Go to page SHB-55 of the Student Handbook

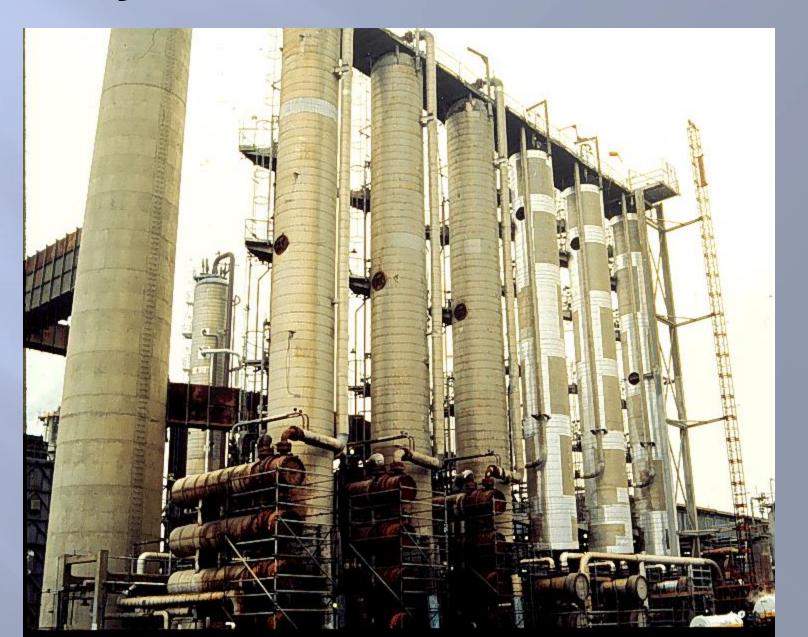
#### HYDROCRACKING UNIT



# Hydrocracker



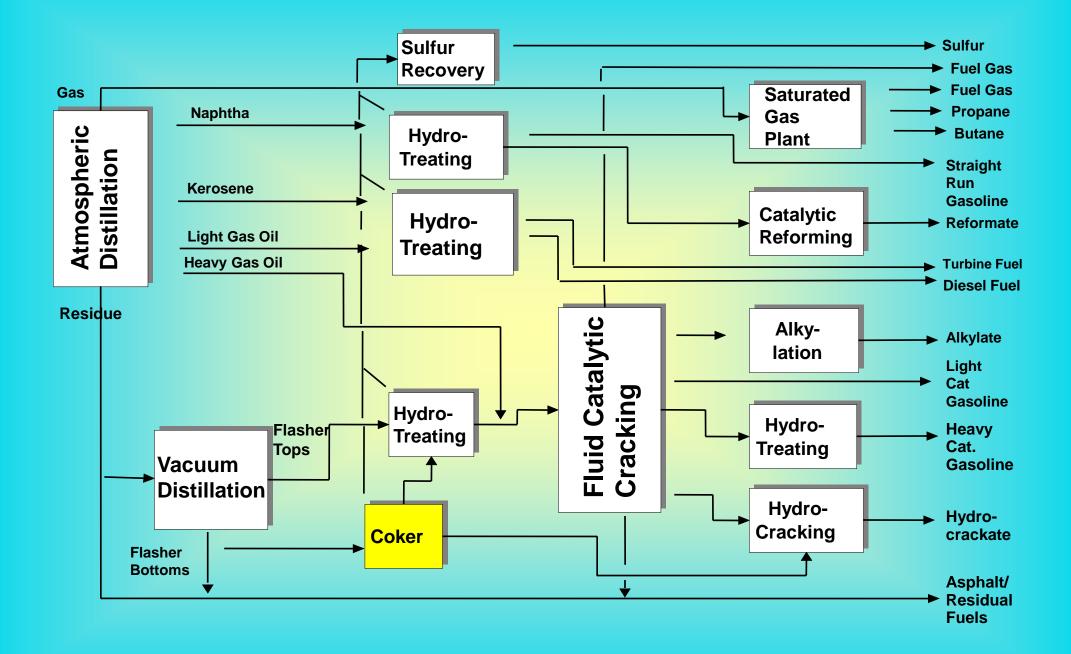
#### **Hydrocracker Reactors**



Coker

PURPOSE: To convert the heaviest crude fractions into lighter components for blending.

By products are coke or low BTU gas



### Coker

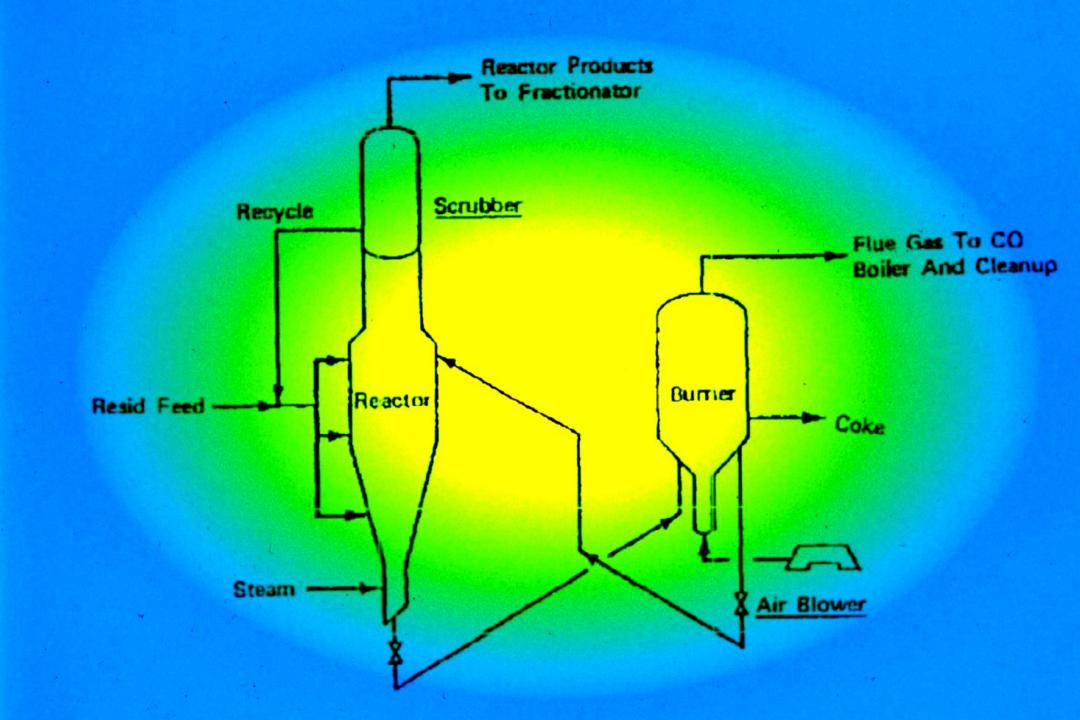
- REACTION: There are 3 types of coking processes utilized. Each process uses high temperature to crack the large molecules and form by-product coke.
  - Flexi-Coking
  - Fluidized Coking
  - Delayed Coking

# **Coking Process Comparison**

TYPE	FEED	PRODUCTS	USES FOR COKE
Flexi Coker		Low BTU Gas	No coke
Fluidized Coker		High quality coke	Weldin <u>g</u> anodes
Delayed Coker		Low quality coke	Steel

# **Fluidized Coking**





#### Fluid Coker

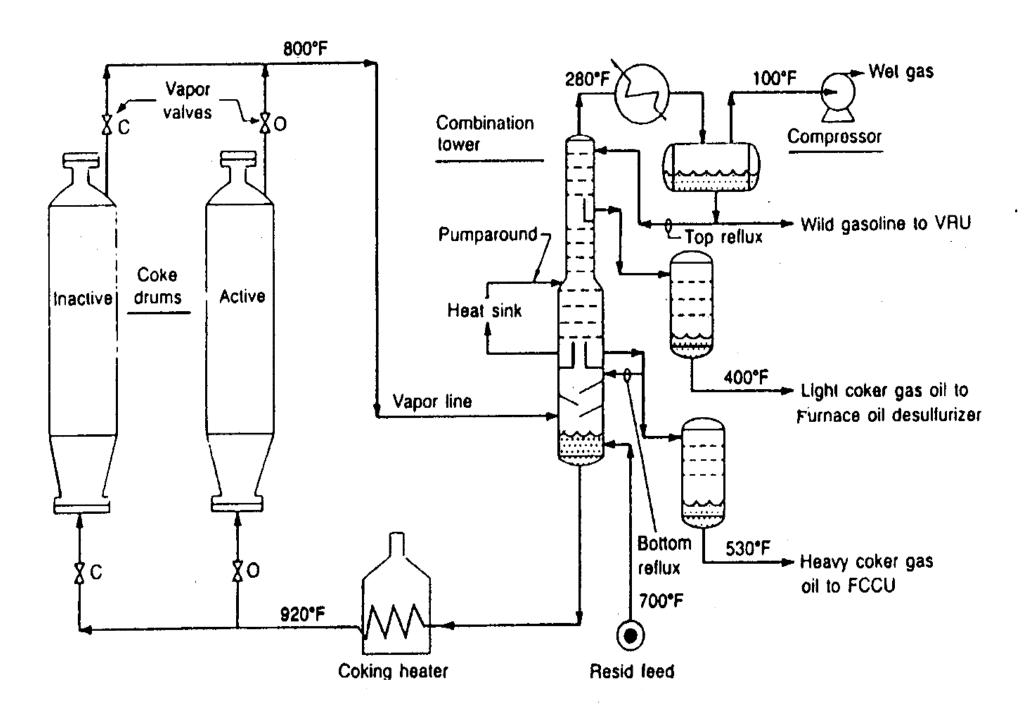


# **Delayed Coker**

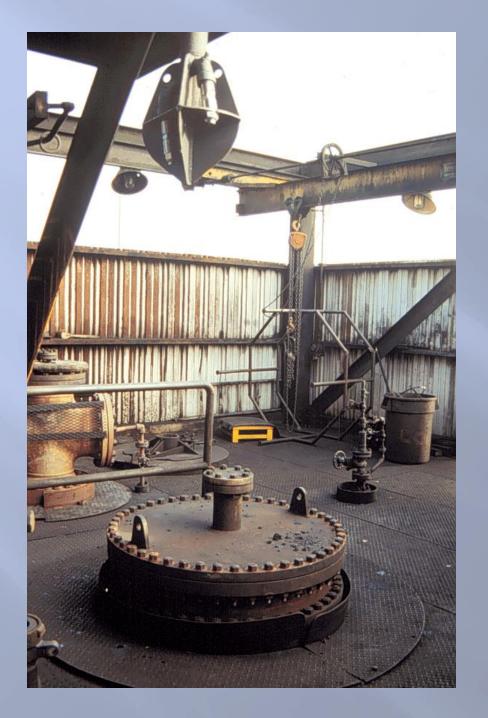


### **Delayed Coker**

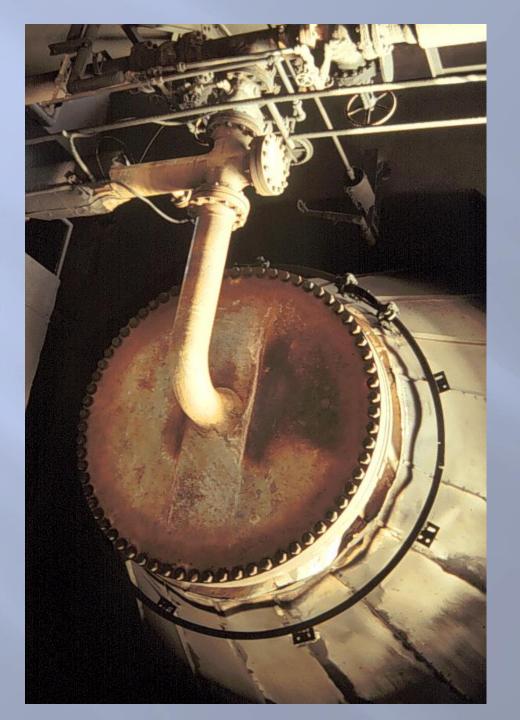




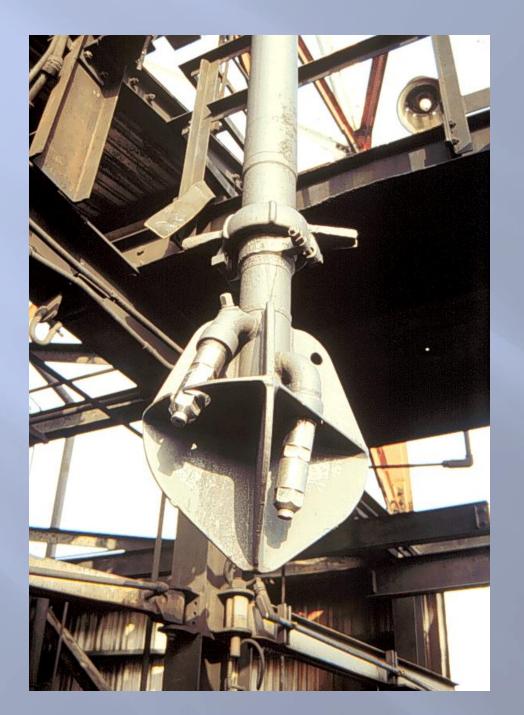




Top of Coke Drum -Hot



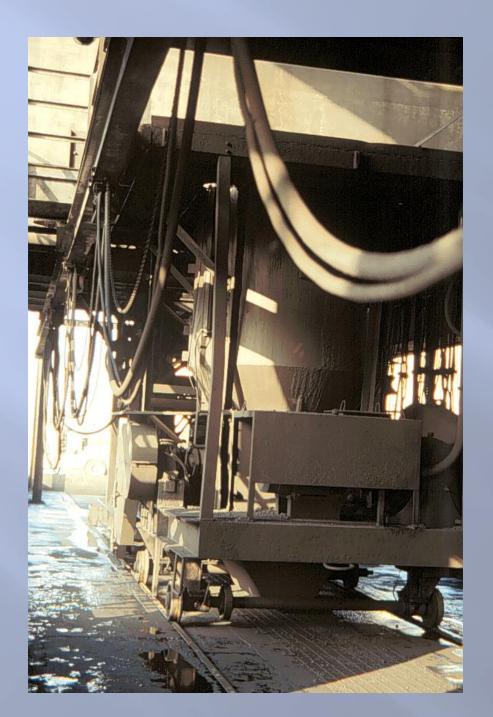
Bottom of Coke Drum



Hydro-Drill Assembly for Coke Removal

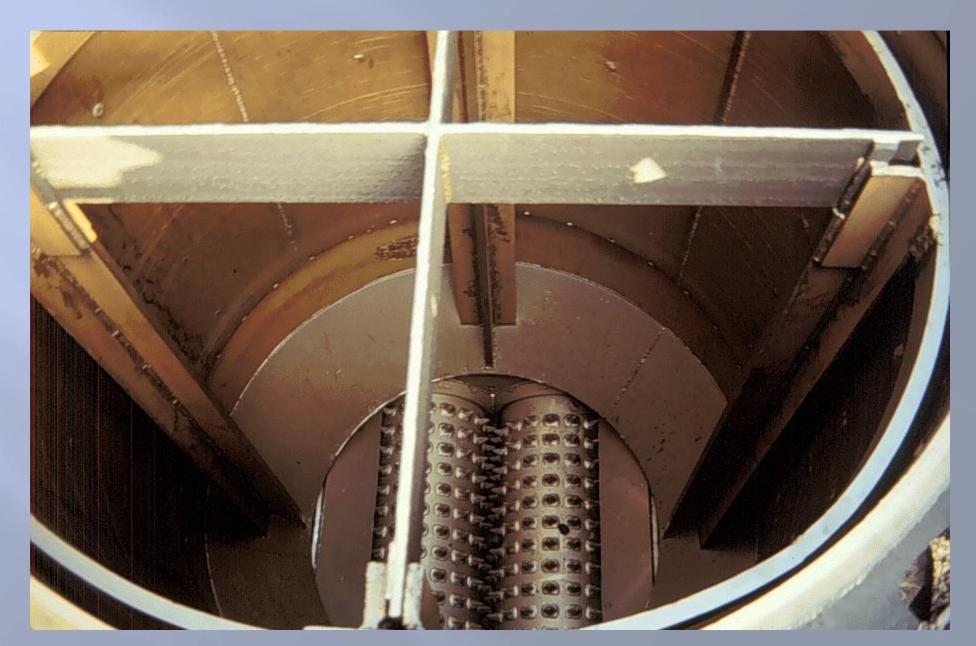






# Coke Treatment Facility

### **Coke Grinder**



#### **Coker Wastewater Treatment**



### **Coke Loading**

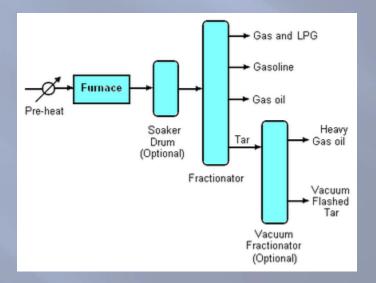


# Visbreaking

- PURPOSE: Heavier crude fractions are thermally treated to improve the viscosity and pour point of products.
- mild cracking/reforming of residium resulting in unsaturated products without coke laydown

# Visbreaking

REACTION: Through mild thermal cracking larger molecules are cracked to somewhat smaller molecules. Cracking takes place in a furnace at between 850 and 900 F.

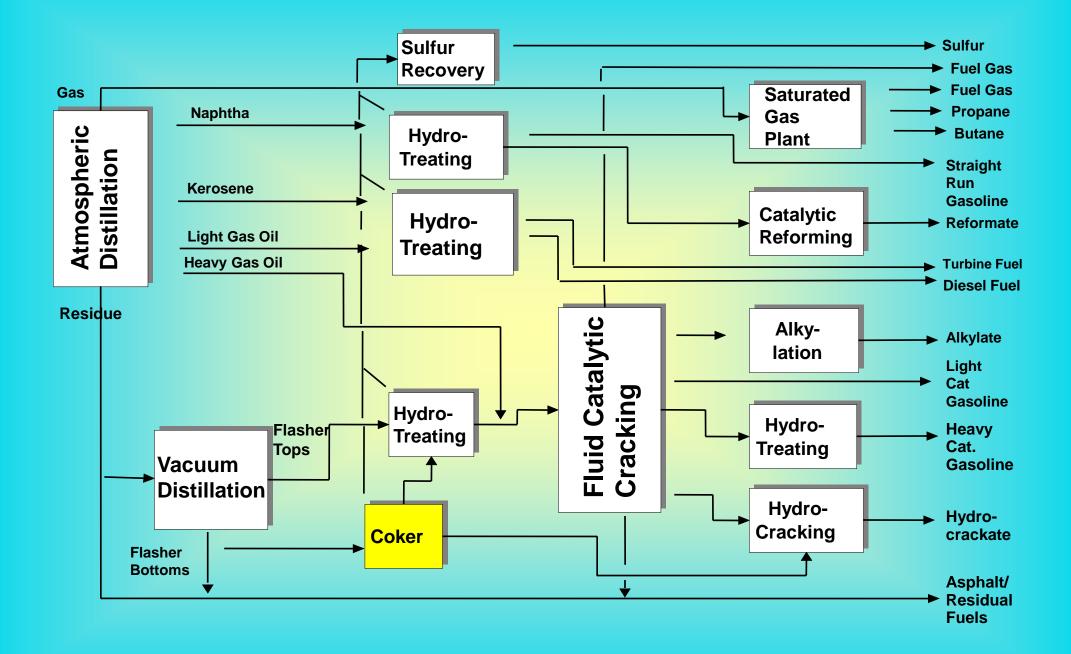


# **Quick Review 5**

- Conversion
  - Property change
    - Catalytic reforming
    - Isomerization
  - Build up
    - Alkylation
    - Polymerization
  - Break up
    - Fluid Catalytic Cracking
    - Hydrocracking
    - Coking
    - Visbreaking

# **Petroleum Refining Process**

- Separation
- Treatment
- Conversion
- Blending



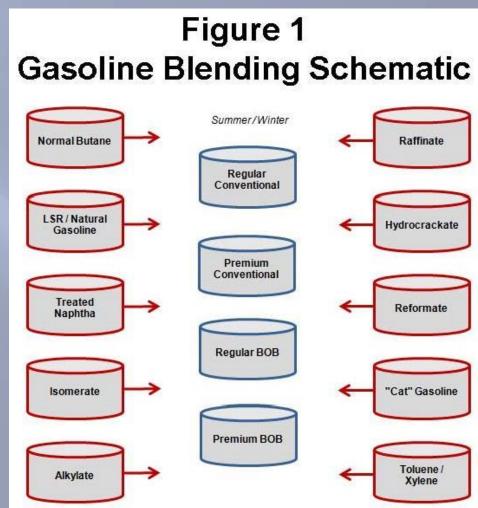
# Blending

 PURPOSE: Refinery products such as gasoline, jet fuel, diesel, and lubricating oils have specifications that must be met. The blending process utilizes all of the available components into end products which meet these specifications.



# Blending

**PROCESS: Blending** components are drawn from storage tanks into a blend manifold where they are metered into a final blend tank. The blend is mixed and sampled to determine if it meets the specifications prior to sale.



# **Gasoline Specifications**

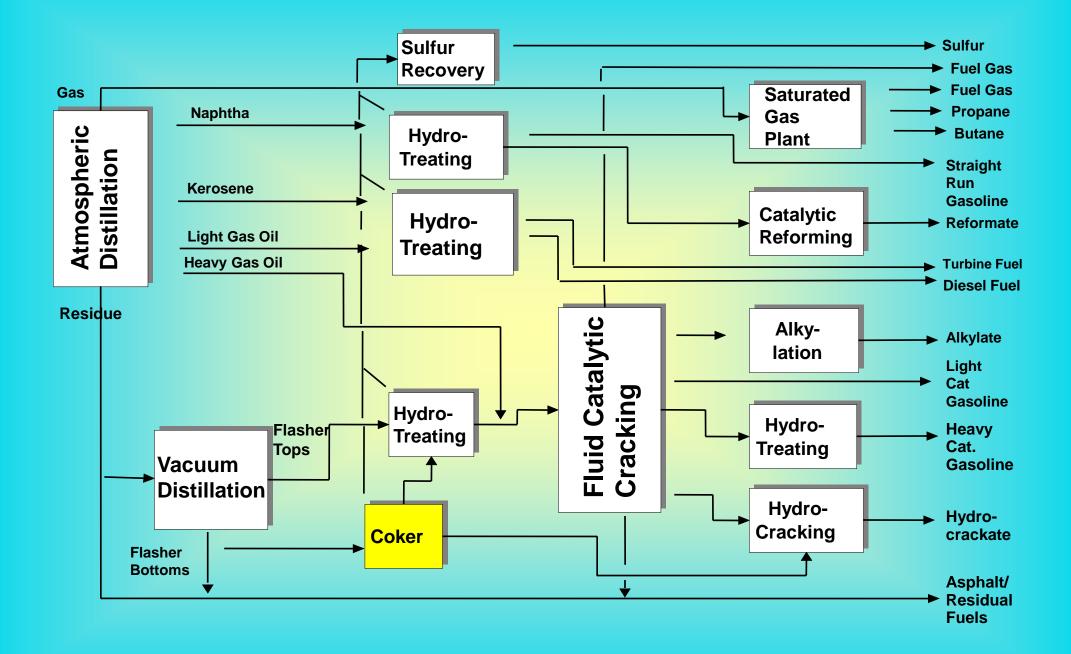
- Octane
- Reid Vapor Pressure
- Sulfur Content
- Benzene Content
- Olefin Content
- Oxygen Content
- Distillation
- Aromatic Content
- Lead Content
- Phosphorus
- Manganese Content
- Deposit Control Additive

# **Diesel Specifications**

- Cetane Number
- Sulfur Content
- Aromatic Hydrocarbon Content
- Polynuclear Aromatic HC Content
- Nitrogen Content

# **Quick Review 6**

- Separation
  - No molecular manipulation
- Treatment
  - Removes impurities
- Conversion
  - Molecular manipulation
    - Desired size
    - Desired properties
- Blending



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# **PROCESSES** THE REST OF THE PROCESS

The National Air Compliance Training Program

#### Wastewater Treatment

PURPOSE: To remove any traces of hydrocarbons from water used in the processes throughout the refinery prior to discharge.

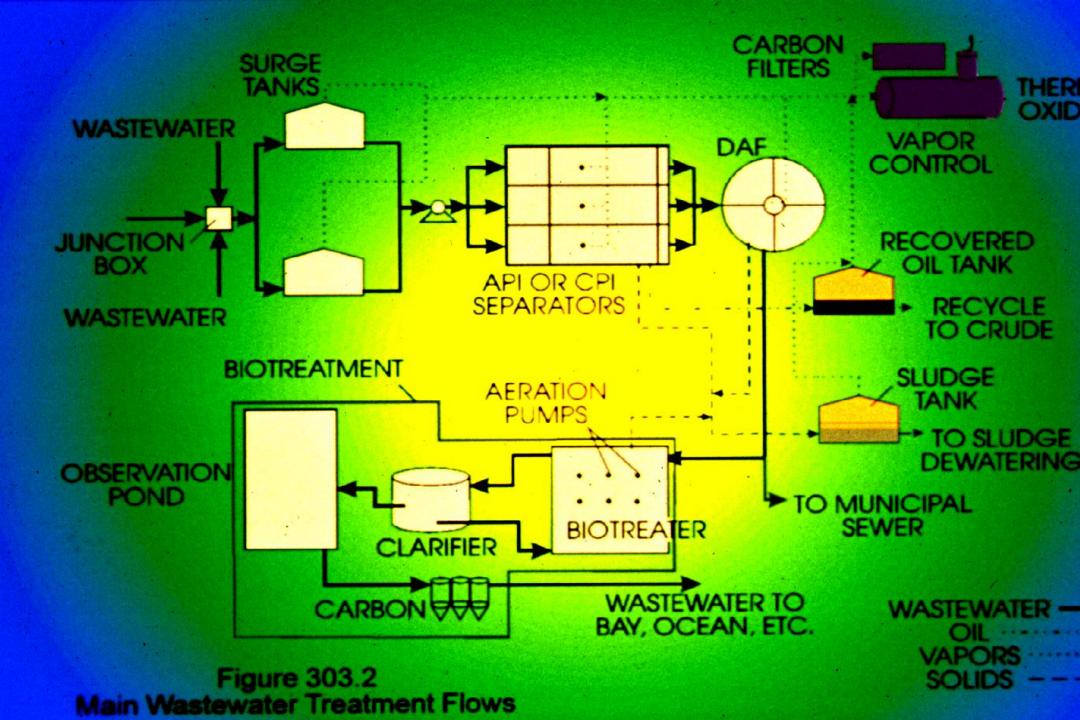


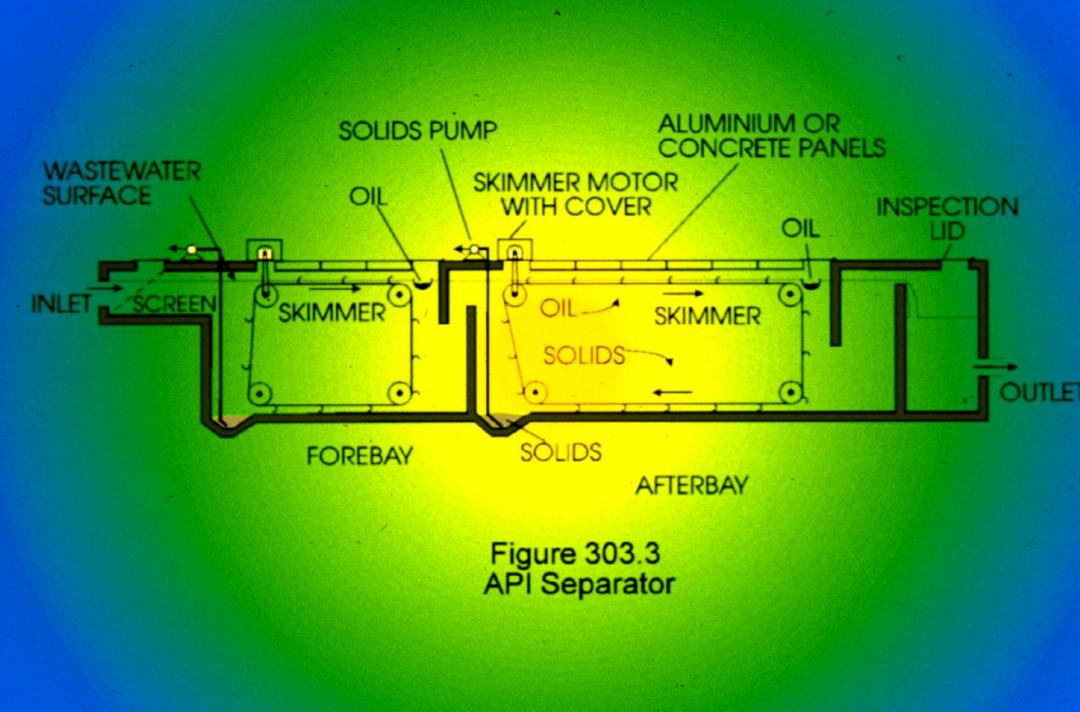
#### Wastewater Treatment

 PROCESS: A series of gravity separation and skimming steps followed by biological treatment to remove traces of hydrocarbons. Some refineries may use filtration or carbon adsorption depending on the permit requirements.

#### **Wastewater Sources**

- Crude Desalting Operations
- Process Water
- Steam Stripping Operations
- Equipment/Tank Washouts
- Storage Tank Roof Drains
- Unit Washdowns and Spills





### **API Separator Forebay**



# **API Separator**



# **API Separator**



# **API Separator with Open Cover**



## **Oil Skimmer Inside API**





Corrugated **Plate** Interceptor (CPI) Wastewater **Separator** 

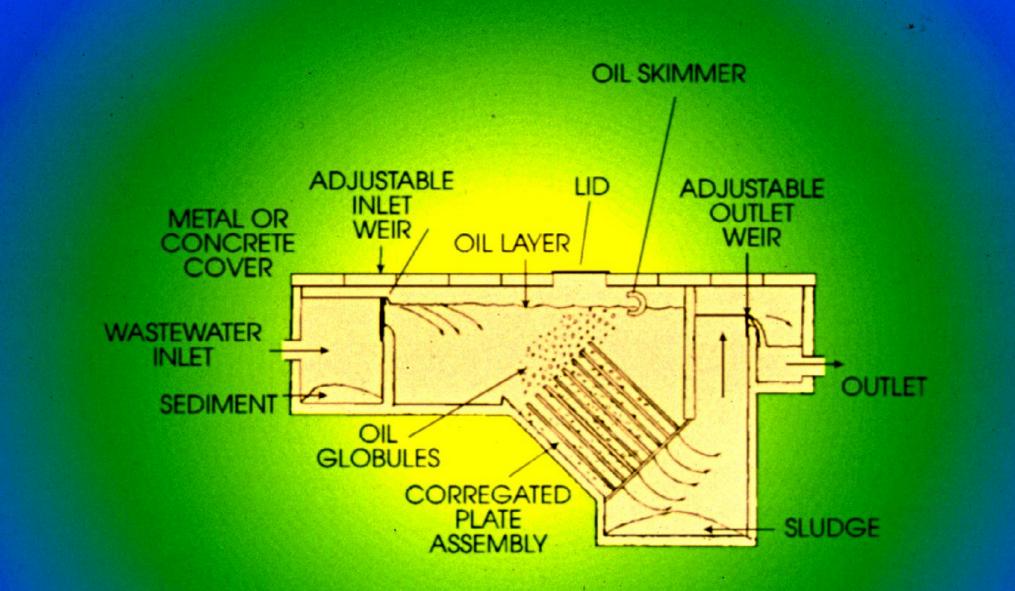
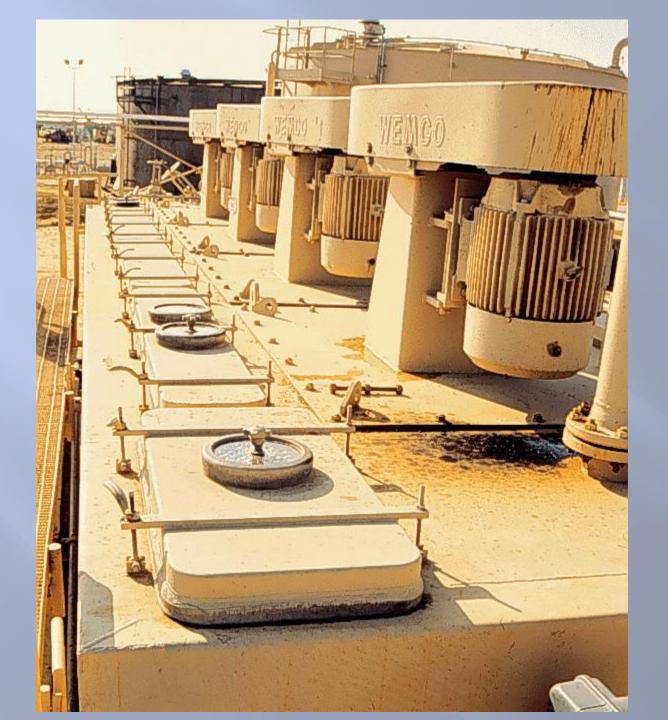
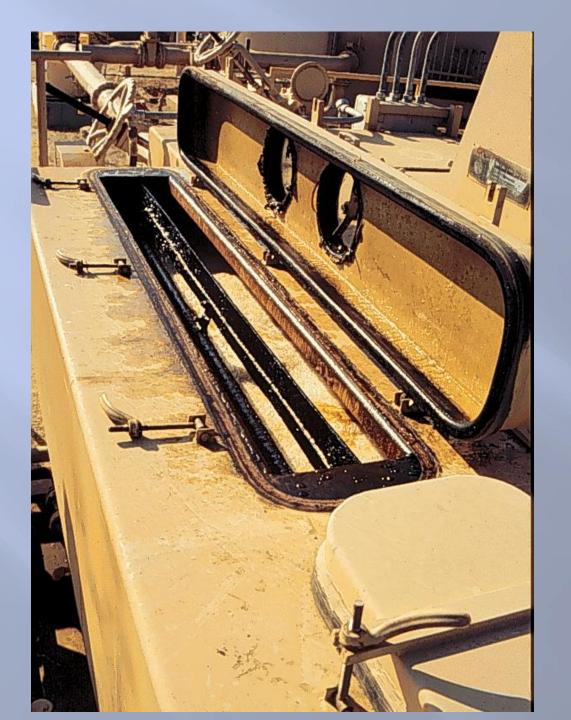


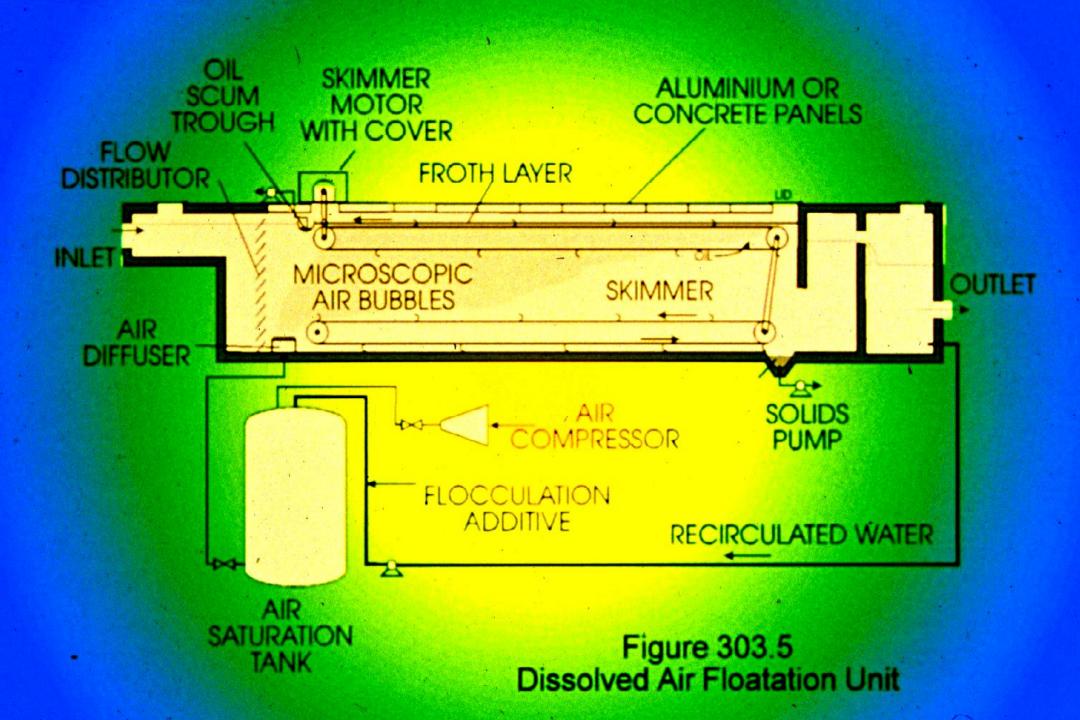
Figure 303.4 CPI Separator



# Induced Air Floatation (IAF)



Inside of a IAF



## Clarifier



# Biological Wastewater Treatment (Biotreaters)



### **Aerators on Treatment Ponds**



# **Bioreactor (Pond)**

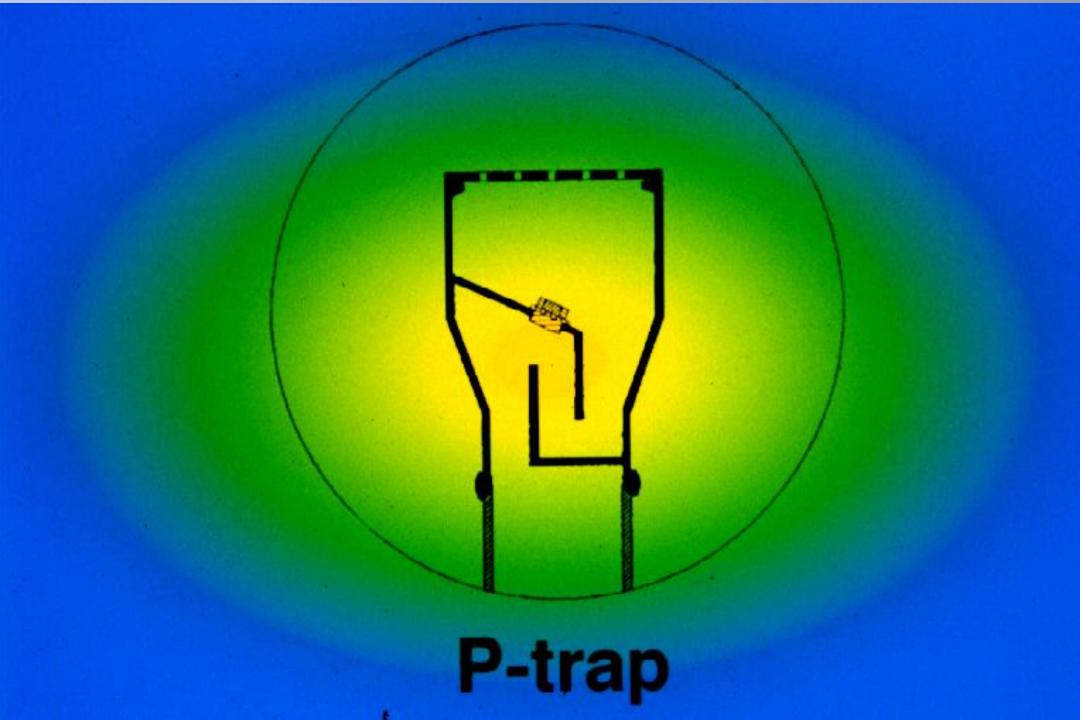


# Granular Activated Carbon (GAC) Treaters



#### **Process Sewer**

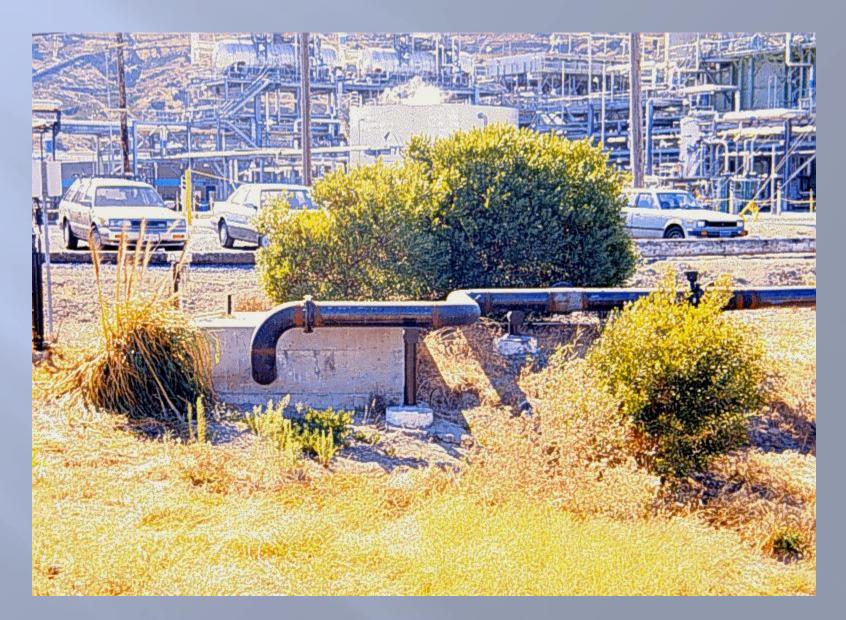




## **Process Drain with a P-Trap**



## **Junction Box for Wastewater**



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# REGULATIONS



The National Air Compliance Training Program

## **REFINERY REGULATIONS**

Refinery NSPS/NESHAP/MACT Standards (Also General SIP Requirements and NSR Consent Decree)					
NSPS - 40 CFR Part 60					
Subpart	Date	Affected Facility	Pollutant		
60.18(b)	Depends	Flares			
Cd (EG)	Pre 8/17/71	Sulfuric Acid Production Unit	H2SO4 mist		
D	8/17/71	Fossil Fueled Fired steam gen >250 mmbtu	PM, SO2, NOx		
Da	9/18/78	Elec Utility steam gen >250 mmbtu (may include certain combined cycle turbines)	PM, SO2, NOx, Hg		
Db	6/19/84	ICI steam gen >100 mmbtu	PM, SO2, NOx		
Dc	6/9/89	ICI steam gen 10 – 100 mmbtu	PM, SO2,		
Н	8/17/71	Sulfuric Acid Production Unit	H2SO4 mist		
1	'76 or '84	FCCU cat regen/fuel gas combustion devices/Claus Plant > 20 LTD	PM, CO (FCCU) SO2		
Ja	5/14/07	FCCU, Fluid Coking Unit, delayed coking units, fuel gas combustion devices (inc. flares and process heaters) and sulfur recovery plants	PM, NOx, SO2, CO (FCCU or FCU); <u>SO2</u> ( <u>SRU); SO2</u> , H2S, NOx (fuel gas comb. dev.)		
K	6/73 - 5/78	Storage Vessel for Pet. Liq. > 40K gal	VOC		
Ka	5/78 - 7/84	Storage Vessel for Pet. Liq. > 40K gal	VOC		
Kb	7/23/84	Organic Liquid Storage Vessel >= 75 m3	VOC		
GG	10/3/77	Stationary Gas Turbines	SO2, NOx		
UU	'80 or '81	Asphalt Processing and Asphalt Roof Mfg	PM		
VV	1/5/81	SOCMI Equipment Leaks	VOC		
XX	12/17/80	Bulk Gasoline Terminals	VOC		
GGGa	11/7/06	Equip Leaks of VOC in Petrol Refineries	VOC		
ш	10/21/83	VOC Emissions from SOCMI Air Oxidation Unit Processes	TOC		
NNN	12/30/83	VOC Emissions from SOCMI Distillation Operations	TOC		
QQQ	5/4/87	VOC Emissions from Petrol Refinery Wastewater Systems	VOC		
RRR	6/29/90	VOC Emissions from SOCMI Reactor Processes	TOC		
CCCC	11/30/99 m/r 6/1/01	Commercial and Industrial Solid Waste Incinerators	Dioxin/furans; 3 metals; PM; opacity; 3 acid gases; CO; ash		
DDDD (EG)	Pre 11/30/99	Commercial and Industrial Solid Waste Incinerators	Dioxin/furans; 3 metals; PM; opacity; 3 acid gases; CO; ash		

Subpart	Date	Affected Facility	Pollutant	
ш	Varies	Stationary Compression Ignition IC Engines	HC, NOx, CO, PM	
1111	Varies	Stationary Spark Ignition IC Engines	NOx, CO, VOC	
KKKK	2/18/05	Stationary Combustion Turbines	SOx, NOx	
NESHAPS – 40 CFR Part 61				
Subpart	Date	Affected Facility	Pollutant	
j	⇔'84	Equipment Leaks of Benzene	Benzene	
M	⇔'84	Asbestos	Asbestos	
v	⇔'84	Equipment Leaks	Benzene, Vinyl Chloride	
Y	<> '89	Benzene Storage Vessels	Benzene	
BB	⇔'90	Benzene Transfer Operations	Benzene	
FF	<> '90	Benzene Waste Operations	Benzene	
MACT - 40 CFR Part 63				
Subpart	Date	Affected Facility	Pollutant	
F	<b>'94</b>	SOCMI (HON)	HAPs	
G	<b>'94</b>	SOCMI Process Vents, Storage Vessels, Transfer Operations, Wastewater	HAPs	
Н	<b>'94</b>	Equipment Leaks	HAPs	
I	<b>'94</b>	Equipment Leaks (certain processes)	HAPs	
Q	<b>'94</b>	Industrial Cooling Towers	Chromium	
R	<b>'94</b>	Gasoline Distribution Facilities	HAPs	
Т	<b>'94</b>	Halogenated Solvent Cleaning	Halogenated Solvents	
CC	<b>'9</b> 5	Petroleum Refineries (MACT I)	HAPs	
EEE	<b>'99</b>	Hazardous Waste Combustors	HAPs	
υυυ	<b>'02</b>	Petroleum Refineries (MACT II – cat cracking, cat reforming, sulfur plant units)	HAPs	
EEEE	<b>'04</b>	Organic Liquids Distribution (non- gasoline)	HAPs	
FFFF	<b>'03</b>	Misc. Organic Chemical Mfg	HAPs	
YYYY	<b>'04</b>	Stationary Combustion Turbines	Formaldehyde	
ZZZZ	<b>'04</b>	Reciprocating Internal Combust. Engines	Formaldehyde	
DDDDD	<b>'04</b>	ICI Boilers and Process Heaters	HAPs	
GGGGG	<b>'03</b>	Site Remediation	HAPs	
LLLLL	<b>'03</b>	Asphalt Processing and Asphalt Roof Mfg	HAPs	
General SIP Rules				
NSR Consent Decree				
Construction and Operating Permit Requirements				

Thanks to the MWRPO for allowing reliance on a document developed for them by MACTEC.

## REGULATIONS

- Federal regulations are NSPS, **NESHAPS (MACT)**  State and local agencies may have additional regulations This course will touch on a few of
  - the applicable NSPS and NESHAPS applicable to refineries

# CONTINUOUS EMISSION MONITORING

CEM Requirements 40 CFR 60, Subpart J FCC Tailgas Units

#### **CEM Requirements**

40 CFR 60, Appendix B -Performance Specifications

 40 CFR 60, Appendix F - Quality Assurance Programs

#### **Purpose of CEMs -Regulators View**

- Determine emission compliance
- Identify periods of excess emissions
- Assess control equipment efficiency
- Monitor operating parameters
- Validate emission credits
- Public perception reports

## **Purpose of CEMs - Industry View**

- Comply with regulations
- Demonstrate compliance
- Monitor control equipment
- Monitor process parameters
- Validate emission credits
- Complaint protection
- Plant safety

# **MACT 1 AND MACT 2**

- 40 CFR 63 NESHAP, Subpart CC [MACT 1]
- 40 CFR 63 NESHAP, Subpart UUU [MACT 2]

Revised or brand-new standards for various refinery equipment (including storage tanks, flares, catalytic cracking units, & coking units), as well as the introduction of mandatory continuous fenceline monitoring for benzene.

# **MACT 1 AND MACT 2**

The EPA performed this rule revision in accordance with the mandatory technology and residual risk reviews that are required every 8 years by the Clean Air Act.

- Signed September 29, 2015.
- Compliance Dates
  - All new sources installed after February 1, 2016 -Upon start-up
  - All existing sources it depends, February 1, 2016 February 1, 2019
- Specific Information Be Found

EPA Website:

http://www3.epa.gov/airtoxics/petref.html

#### **WORK PRACTICE STANDARDS**

- Maintenance Vents
  - Maximum hydrocarbon limits prior to opening
- Pressure Relief Devices
  - Continuous monitoring & release reporting requirements
- Delayed Coking Units
  - Maximum pressure limits prior to opening
- Flares
  - Continuous monitoring & maximum flare tip velocity limits
- Fluid Catalytic Cracking Units (FCCU)
  - Minimum O<sub>2</sub> and cyclone face velocity operating limits during start-up, shut-down, & hot standby
- Sulfur Recovery Unit (SRU)
  - Minimum temperature and O<sub>2</sub> operating limits during startup & shut-down.

### PRESSURE RELIEF DEVICES (§63.648)

**Compliance Date: February 1, 2016** 

**Operating requirements:** Except during a pressure release, operate each pressure relief device in organic HAP gas or vapor service with an instrument reading of less than <u>500 ppm above background</u>

#### Returning to normal operation after a release episode:

The owner or operator must conduct instrument monitoring, <u>no later</u> <u>than 5 calendar days after the pressure relief device returns to</u> <u>organic HAP gas or vapor service following a pressure release</u> to verify that the pressure relief device is operating with an instrument reading of less than 500 ppm.

In addition to the above, if the pressure relief device consists only of a rupture disk, a <u>replacement disk must be installed no later than 5</u> <u>calendar days after the pressure release</u>. The owner or operator may not initiate start-up of the equipment served by the rupture disk until the rupture disc is replaced.

### **DELAYED COKING UNITS (§63.657)**

#### **Compliance Date: February 1, 2016**

Each owner or operator of a delayed coking unit shall depressurize each coke drum to a closed blowdown system until the following conditions are met:

For delayed coking units at an <u>existing affected source</u>, <u>meet either</u>: (A) An average vessel pressure of 2 psig determined on a rolling 60-event average; (B) An average vessel temperature of 220 degrees Fahrenheit determined on a rolling 60-event average.

For delayed coking units at a <u>new affected source</u>, <u>meet either:</u>

(A) A vessel pressure of 2.0 psig\* for each decoking event; or

(B) A vessel temperature of 218 degrees Fahrenheit for each decoking event.

#### \*Notice the additional significant digit for a new affected source, not trivial

Each operator of a delayed coking unit complying with these pressure limits <u>must</u> <u>install, operate, calibrate, and maintain a pressure monitoring system</u> to determine the coke drum vessel pressure.

#### FLARES (§63.670)

**Compliance Date: January 30, 2019** 

**Pilot flame presence:** The owner or operator shall operate each flare with a pilot flame present <u>at all times when regulated material is routed to the flare</u>.

**Pilot flame monitoring:** The owner or operator shall <u>continuously monitor the</u> <u>presence of the pilot flame</u> using a device (e.g. thermocouple, ultraviolet beam sensor, or infrared sensor) capable of detecting that the pilot flame is present.

Flare tip velocity: Actual flare tip velocity (Vtip) must be less than 60 feet per second OR less than 400 feet per second AND also less than the maximum allowed flare tip velocity (Vmax).

**Combustion zone operating limits:** For each flare, the owner or operator shall operate the flare to maintain the net heating value of flare combustion zone gas  $(NHV_{cz})$  at or above 270 Btu/scf.

Flare vent gas, steam assist and air assist flow rate monitoring: The owner or operator shall install, operate, calibrate, and maintain a monitoring system capable of continuously measuring, calculating, and recording the volumetric flow rate in the flare header or headers that feed the flare, as well as any supplemental natural gas used.

### FLARES (§63.670)

#### Compliance Date: January 30, 2019

**Visible emissions monitoring:** The owner or operator shall monitor visible emissions while regulated materials are vented to the flare. An <u>initial visible</u> <u>emissions demonstration must be conducted using an observation period of 2 hours</u> <u>using EPA Method 22</u>. The owner or operator must record and report any instances where visible emissions are observed for more than 5 minutes during any 2 consecutive hours. <u>Subsequent visible emissions monitoring must be performed</u> <u>either by daily visible emissions monitoring or continuous video surveillance</u>.

Flare vent gas composition monitoring: The owner or operator shall determine the concentration of individual components in the flare vent gas using a continuous monitoring or grab sampling system.

### FLARES (§63.670)

#### Compliance Date: January 30, 2019

If either criteria (i) or (ii) occurs, the refinery is now required to complete a <u>root cause</u> analysis and implement corrective action within 45 days of event.

#### **Violation Criteria:**

- (i) The vent gas flow rate exceeds the smokeless capacity of the flare and visible emissions are present from the flare for more than 5 minutes during any 2 consecutive hours during the release event.
- (ii) The vent gas flow rate exceeds the smokeless capacity of the flare and the 15minute block average flare tip velocity exceeds the maximum flare tip velocity.

The following are a violation of the emergency flaring work practice standards:

- (A) <u>Any flow event</u> for which a root cause analysis was required and the root cause was determined to be <u>operator error or poor maintenance</u>.
- (B) <u>Two visible emissions exceedance events meeting the criteria in (i)</u> that were not caused by a force majeure event from a <u>single flare in a 3 calendar year period for</u> <u>the same root cause for the same equipment.</u>
- (C) <u>Two flare tip velocity exceedance events meeting the criteria in (ii)</u> that were not caused by a force majeure event from a <u>single flare in a 3 calendar year period for</u> <u>the same root cause for the same equipment</u>.

#### FLUID CATALYTIC CRACKING UNITS (§63.1564 & §63.1565)

Compliance Date: February 1, 2016 → August 1, 2017?

New Standards for start-up, shut-down, and hot standby operation:

#### **PM Standard**

• Maintain the <u>inlet velocity to the primary internal cyclones</u> of the catalytic cracking unit catalyst regenerator <u>at or above 20 feet per second</u> as an alternative to the normally permitted PM limit.

#### **HAPS Standard**

 Maintain the <u>oxygen (O<sub>2</sub>) concentration in the exhaust gas</u> from your catalyst regenerator <u>at or above 1 volume percent (dry basis</u>) as an alternative to the normally permitted CO limit.

# FLUID CATALYTIC CRACKING UNITS (§63.1571)

Compliance Date: August 1, 2017

#### PM Performance Test Standard

 Conduct a <u>periodic performance test for PM or Ni</u> for each catalytic cracking unit at least <u>once every 5 years</u>\*. You must conduct the first periodic performance test <u>no later than August 1, 2017</u>.

\*Exempt from this requirement if monitoring with a PM CEMS

#### HAPS Performance Test Standard

 Conduct a <u>one-time performance test for HCN\*</u> for each catalytic cracking unit <u>no</u> later than August 1, 2017

\*If you conducted a performance test for HCN for a specific catalytic cracking unit <u>between March 31, 2011 and February 1, 2016</u>, you may submit a request to the Administrator to use the previously conducted performance test results. **Request must be submitted by March 30, 2016.** 

### SULFUR RECOVERY UNITS (§63.1568)

#### Compliance Date: February 1, 2016 → August 1, 2017?

New Standards for start-up & shut-down:

#### **HAPS Standard**

• Send any <u>start-up or shut-down purge gases to a flare</u> as an alternative to the normally permitted SOx limit.

#### OR

Send any <u>start-up or shut-down purge gases to a thermal oxidizer or incinerator</u> operated at a minimum hourly average temperature of 1,200 °F in the firebox and a minimum hourly average outlet O<sub>2</sub> concentration of 2 volume percent (dry basis) as an alternative to the normally permitted SOx limit.

#### **MISCELLANEOUS CHANGES**

#### MARINE VESSEL LOADING (Compliance Date: February 1, 2016)

 EPA has <u>deleted the exclusion for marine vessel loading operations at</u> <u>petroleum refineries in 40 CFR 63 NESHAP, Subpart Y</u>. Removing this exclusion will require <u>small marine vessel loading operations</u> (i.e., operations with HAP emissions less than major source thresholds) <u>and offshore marine</u> <u>vessel loading operations to use submerged filling</u> based on the cargo filling line requirements in 46 CFR §153.282.

#### **GROUP 1 STORAGE VESSELS (Compliance Date: May 1, 2016)**

 Group 1 storage vessels now include smaller tanks with lower vapor pressures. An <u>existing tank</u> is now classified as Group 1 if:

Capacity (gallons)	Maximum True Vapor Pressure (psia)	Annual Average Weight HAP Content
$20,000 \le X < 40,000$	1.9	4%
40,000 ≤ X	0.75	4%

If any existing tanks are now redefined as a Group 1 storage vessel, new control requirements specified in 40 CFR §63.646 will need to implemented.

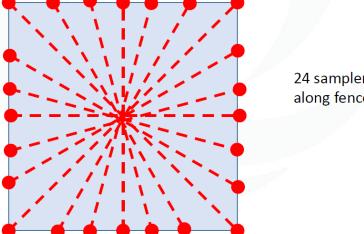
Fenceline Monitoring Requirements are now required as part of 40 CFR 63, NESHAPS CC. The requirements are part of the package the refinery MACT 1 and MACT 2 updates signed September 29, 2015. Refineries have three years to deploy monitoring. Significant emission reductions are expected to result from action plans in response to fenceline monitoring. https://www.epa.gov/stationary-sources-airpollution/petroleum-refinery-sector-risk-andtechnology-review-and-new-source

- The Benzene Fenceline Monitoring Rule was added as part of 40 CFR 63 NESHAP, Subpart CC [Refinery MACT 1] as a completely new subsection (§63.658) along with 2 new EPA Test Methods 325 A/B.
- This rule is being implemented to account for and monitor fugitive emissions from sources such as leaking equipment and wastewater treatment. The air concentration limit set forth by this continuous monitoring is 9 µg/m<sup>3</sup>.
- Any exceedance of this limit will trigger corrective action to be completed by the facility.

- **Program Requirements:** 
  - 12 month's worth of data must be obtained by February 1,
    2019 (i.e. must start sampling by February 1, 2018)
  - EPA 325A and 325B sorbent tube sampling (passive sampler (PS)), 14-day sampling period (next sampling event starts immediately after the end of previous event)
    - <750 acres 30 degrees (12 samplers)
    - >750, < 1500 acres 20 degrees (18 samplers)</li>
    - >1500 15 degrees (24 samplers)
    - \*Additional samplers may be required if there is an emission source that is close to the fence
  - Siting of monitors determined by proscribed methods, based on facility layout, at a minimum of 2,000 feet apart

- Refineries required to deploy passive timeintegrated benzene samplers – 14 day sample period
- Up to 24 monitoring locations distributed around the perimeter (fenceline) of the refinery
- Action level of 9 micrograms per cubit meter  $(\mu g/m3)$

Note that extra samplers are required near known sources of VOC emissions



24 samplers placed along fenceline

- Refineries required to deploy passive time-integrated benzene samplers 14 day sample period
- Up to 24 monitoring locations distributed around the perimeter (fenceline) of the refinery
- Reduction in frequency allowed if 2 years of samples are below 0.9  $\mu$ g/m3



Photo courtesy of Enthalpy Analytical, Inc.

- Corrective actions may include
  - Leak inspection using Method 21 and repair
  - Leak inspection using optical gas imaging and repair
  - Visual inspection to determine the cause of the high benzene emissions and repair
  - More frequent Methods 325A and 325B sampling or active sampling
  - Other measures



Photo courtesy of Camsco, Inc.

# Fugitive Leak Regulations Wastewater Systems

- Federal NSPS
  - 40 CFR 60 Subpart QQQ

https://www.epa.gov/stationary-sources-airpollution/volatile-organic-compounds-vocemissions-petroleum-refinery

### **Fugitive Leaks**

- NSPS Leak Detection and Repair (LDAR)
- 40 CFR Part 60, Subpart GGGa
- Equipment Leaks of VOC in Petroleum Refineries
  PURPOSE: To ensure that the numerous pump seals, compressor seals, valves, flanges and other components are not leaking beyond a threshold value.

https://www.epa.gov/stationary-sources-airpollution/equipment-leaks-volatile-organiccompounds-voc-petroleum-refineries

### Requirements

- Leakage Limits
- Self Inspection
- Leak Repair
- Component Identification
- Recordkeeping and Reporting

Handbook Pages 301-19 thru 301-33

### LDAR Equipment/Methods

- Organic Vapor Analyzer (OVA)
- Threshold Limit Value Meter (TLV)
- Flame Ionization Detector (FID)
- Photoionization Detector (PID)
- Infrared Detector (FLIR)
- Look and Listen
  - Soap Solution
  - Visual Distortion
  - Odor

### **Inspection Points**

- Screen as many components as possible
- Review records
- Verify repair tag dates

Handbook Page 301-39

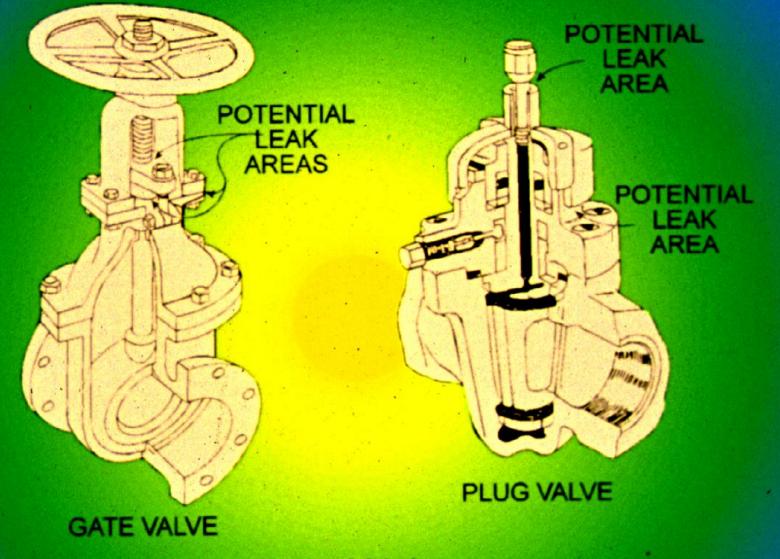
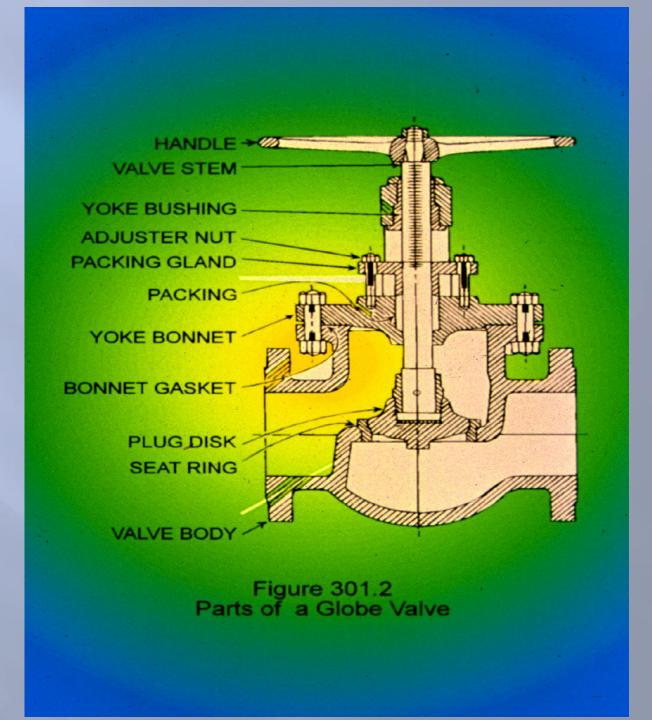


Figure 301.4 Potential Leak Areas for Gate Valves and Plug Valves



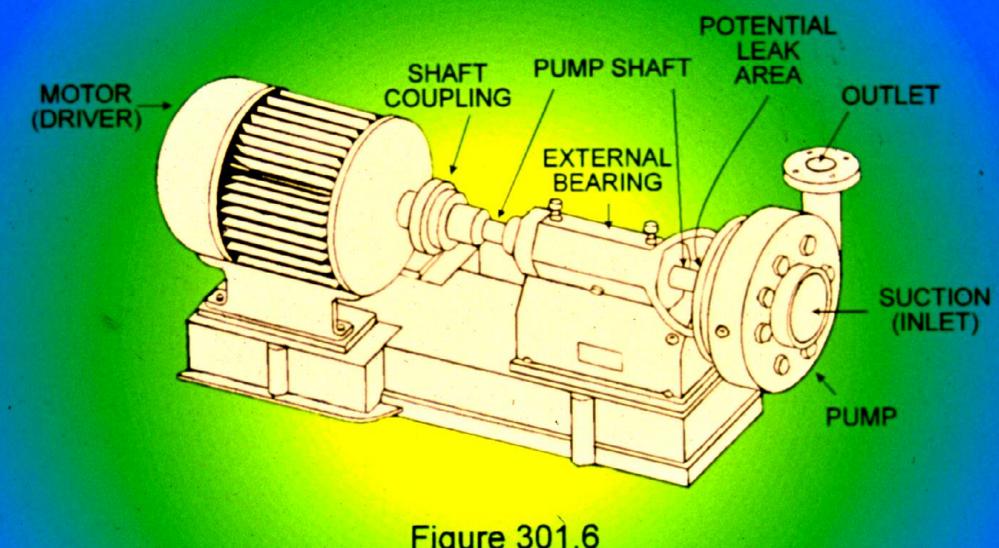
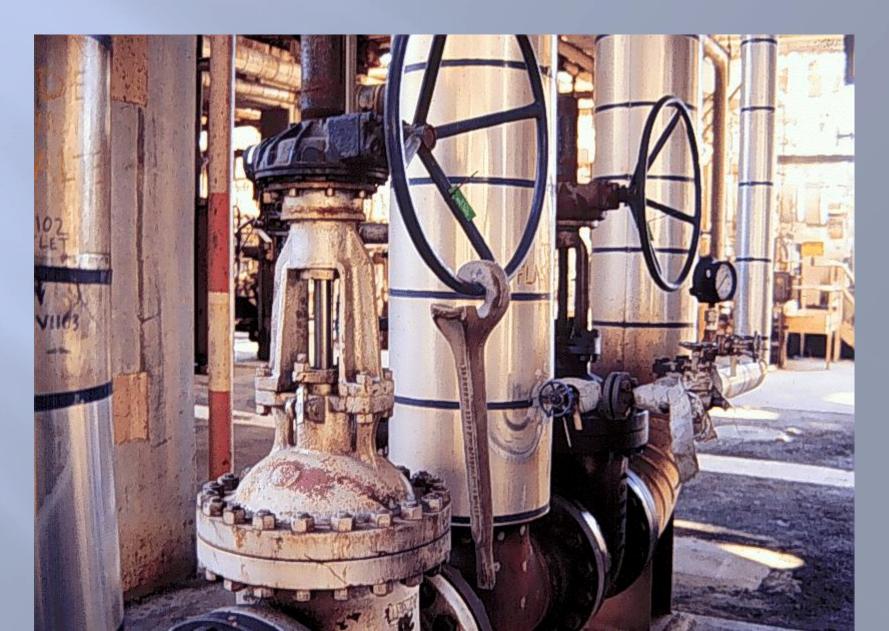


Figure 301.6 Centrifugal Pump



# Control Valve





### **Centrifugal Pump**



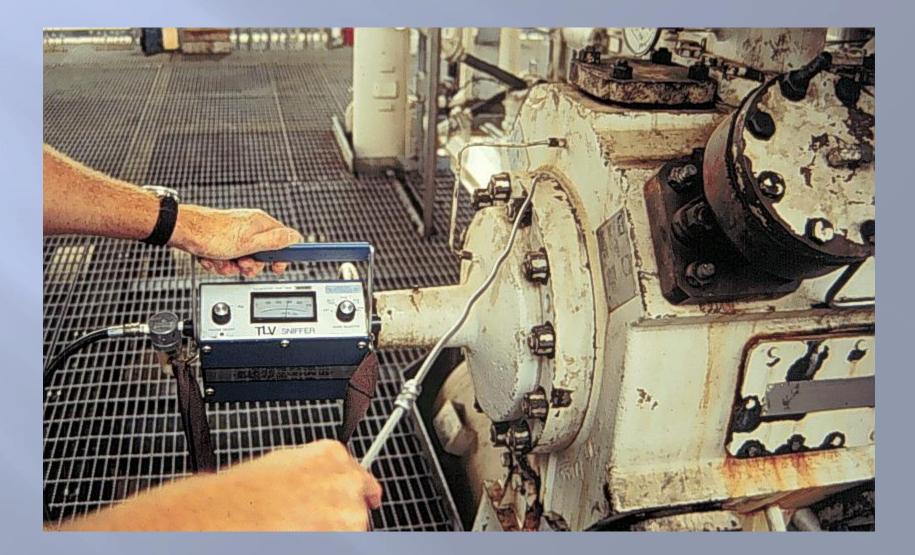
### **Reciprocating Compressor**





Fugitive VOC Testing with OVA

### Fugitive VOC Testing with TLV



# Fugitive VOC Testing - Leaky Valve



# Fugitive VOC Testing - Valve Tag



# Fugitive VOC Testing - Valve Tag

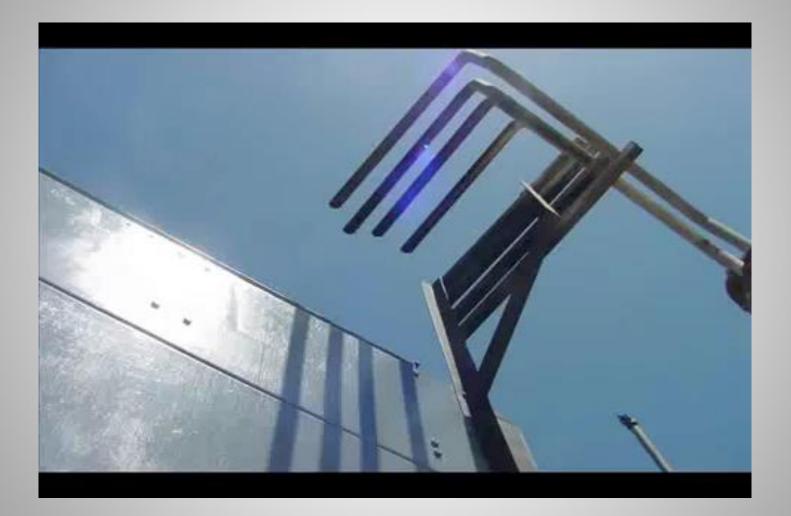


# Fugitive Leak Screening - FLIR

FLIR stands for forward looking infrared camera – it reads the thermal infrared signature of a plume. Although not a Method 21 device it can be used to screen for leaks quickly.



#### FLIR Video of Vent from Centrifugal Pump



# **INSPECTION SAFETY**



### **Refinery Hazards**

- Hydrogen Sulfide (H2S)
- Heat
- Hydrofluoric Acid
- Heights
- Asbestos
- Explosions
- Fires
- Noise
- Sulfur Dioxide



### FIELD VISIT

