

INTEGRATED WASTE MANAGEMENT FOR A SMART CITY
FOCUSED ON MSW, C&D AND E-WASTE MANAGEMENT

Welcome to Week-8

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During this week (Week-8)

- Thermal Treatment Details
- Landfill Basics, Liner Requirement

Stoichiometric Combustion

- the principle elements of solid waste are C, H, O, N and S
- under ideal combustion conditions, the following gases are formed:
 - carbon dioxide (CO₂),
 - water (H₂O and the water content of flue gas),
 - nitrogen (N₂), and
 - small amounts of sulphur dioxide (SO₂)
- oxygen is required for combustion of the:
 - carbon
 - hydrogen (minus the fraction that enters as H₂O with the SW)
 - sulphur

Stoichiometric Combustion

- the basic reactions for the stoichiometric combustion of the C, H and S in the organic fraction of waste are as follows:
 - for carbon:
$$\frac{C}{12} + \frac{O_2}{32} + \text{heat} \rightarrow CO_2$$
 - for hydrogen:
$$\frac{2H_2}{4} + \frac{O_2}{32} + \text{heat} \rightarrow 2H_2O$$
 - for sulphur:
$$\frac{S}{32.1} + \frac{O_2}{32} + \text{heat} \rightarrow SO_2$$

Stoichiometric Combustion

- if we assume that dry air contains 23.15% oxygen by weight, then the amount of air required for the oxidation of 1 kg of carbon would be 11.52 kg:

$$M_{\text{air, C}} = (1 \text{ kg C}) \left(\frac{32 \text{ g/mol}}{12 \text{ g/mol}} \right) \times \left(\frac{1}{0.2315} \right) = 11.52 \text{ kg air}$$
- and, the amount of air for hydrogen:

$$M_{\text{air, H}} = (1 \text{ kg H}) \left(\frac{32 \text{ g/mol}}{4 \text{ g/mol}} \right) \times \left(\frac{1}{0.2315} \right) = 34.56 \text{ kg air}$$
- and, the amount of air for sulphur:

$$M_{\text{air, S}} = (1 \text{ kg S}) \left(\frac{32 \text{ g/mol}}{32.1 \text{ g/mol}} \right) \times \left(\frac{1}{0.2315} \right) = 4.31 \text{ kg air}$$

Stoichiometric Combustion

- it should be noted that the amount of hydrogen must first be adjusted by subtracting 1/8th of the percentage of oxygen from the total percentage of hydrogen initially present in the waste
- this subtraction accounts for the oxygen in the waste combining with hydrogen to form water
 - remember, H₂O = (2 g/mol H) + (16 g/mol O)
- then the total O₂ can then be converted to total air
- these calculations define the exact (*stoichiometric*) amount of oxygen (or air) needed for complete combustion
- combustion with oxygen in excess of the stoichiometric amount is called *excess-air combustion*, and this is done to ensure that complete combustion occurs
- excess air is usually 120 – 200% of the design requirement

Excess Air

- the objective of combustion is the complete destruction of the organic waste to form harmless gases
 - the combustion process needs excess air as a result of the non-homogeneous mixture of waste
 - but, too much oxygen reduces the combustion temperature
- the 3 Ts of combustion are:
 - temperature** – high enough to ignite the constituents
 - less than 790°C and odorous compounds are released
 - greater than 980°C, and we get a reduction in dioxins, furans, and VOCs
 - time** – enough for complete combustion of the waste
 - in the combustion chamber
 - in the flue gas (secondary chamber)
 - turbulence** – mixing of the waste material with oxygen
 - provides for good mixing to promote more complete combustion
 - both in the primary chamber (MSW) and the secondary chamber (gases)



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Excess Air

- theoretical oxygen requirements can be determined by balancing chemical reactions, or
- using an easy-to-use formula for O_2 needed to combust a fuel or waste

$$O_2 \text{ (m}^3\text{/kg} \cdot \text{fuel)} = 24.6 \cdot \{ (C/12) + (H/4) + (S/32) - (O/32) \}$$

- C, H, S and O are the decimal fractions of each of the element in a kg of fuel on a dry weight basis
- for example, for C_aH_b :

$$\% C = \left(\frac{a(12)}{a(12) + b(1)} \right)$$
- this formula takes into account the water formed



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Mechanisms of Combustion

- the combustion process is accomplished in 4 phases
- Phase 1** – the drying phase
 - moisture is driven off at up to 150°C
- Phase 2** – volatilization of vapours and gases
 - combustible volatile organics diffuse out when their flash points occurs (150°C – 700°C)
- Phase 3** – burn-down of solids
 - when heated further, the remaining fixed carbon volatile material (partially oxidize cellulose, lignin, ...) are oxidized (700°C – 1100°C)
- Phase 4** – final burn-down of char
 - remaining char is burned down to bottom ash
 - this material is the end product of the reaction
 - after a short period of cooling on the grate this ash is dumped into a dry hopper (the ash-receiving system)



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Types of Incinerators

- there are generally 2 types of MSW incinerators:
- fixed-hearth incinerators – have a stationary kiln
 - first type of waste combustion system designed/used
 - normally used for smaller quantities of waste (medical, pharmaceutical, research and development, ...)
 - waste is charged into the incinerator and combusted as it passes through the combustion chamber
 - ash residue comes out the last hearth



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Types of Incinerators

- rotary kiln incinerators
 - consists of 2 chambers
 - waste is charged into the primary combustion chamber, which is a refractory-lined rotary kiln that is sloped slightly downhill
 - the kiln is slowly turned (at approximately 1 RPM)
 - material is burned while it tumbles down the kiln
 - the secondary chamber is used for gas phase combustion



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The Combustion Process

- combustion starts with the unloading of the MSW collection trucks into a storage pit
 - storage capacity is usually equal to the volume of waste for 2 days
- an overhead crane is used to batch load waste into the feed (charging) chute
- the chute directs wastes to the furnace
- the crane can be used to select wastes of different types to blend the C/N ratio
- solid waste lands on the grates where they enter the mass-fired furnace
- air may be introduced from the bottom (below the grates) or from above the grates by a forced draft fan
 - this controls the burning rate and temperature



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The Combustion Process

- gases and particulate matter are driven off the combusted organic material, and rise into the combustion chamber where they are burned in excess of 800°C
- heat recovery occurs using water filled tubes in the walls of the combustion chamber
- air pollution control equipment may include:
 - injected ammonia for NO_x control
 - dry scrubber for SO_2 and acid gas control
 - bag house (fabric filter) for particulate matter
- ashes and unburned material from the grates fall into a residue hamper
 - residue ash + fly ash (bag house) are mixed together for ash treatment



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Typical Combustor Design

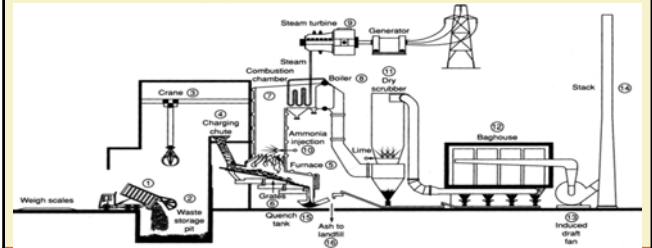
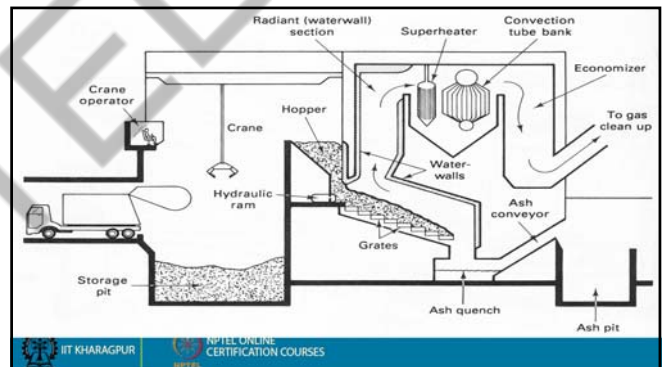


Figure 9.31
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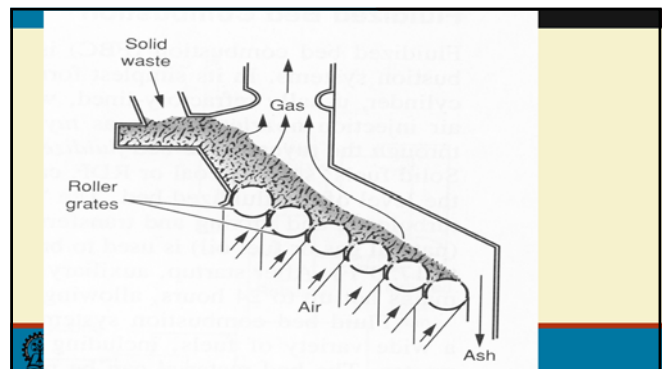
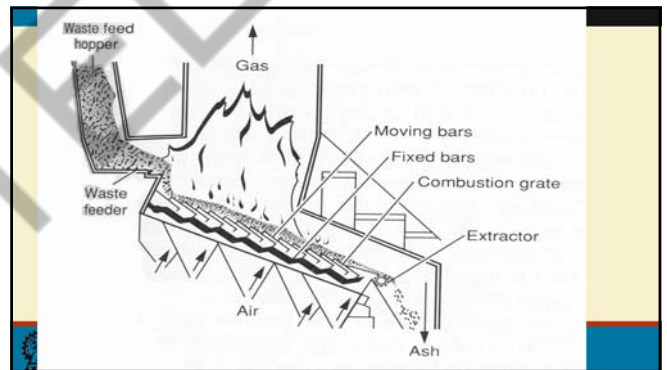
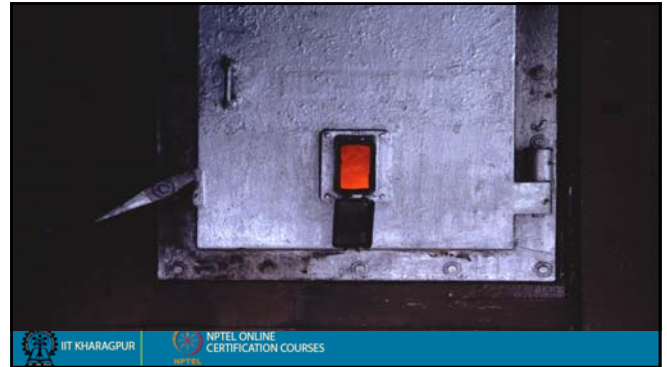


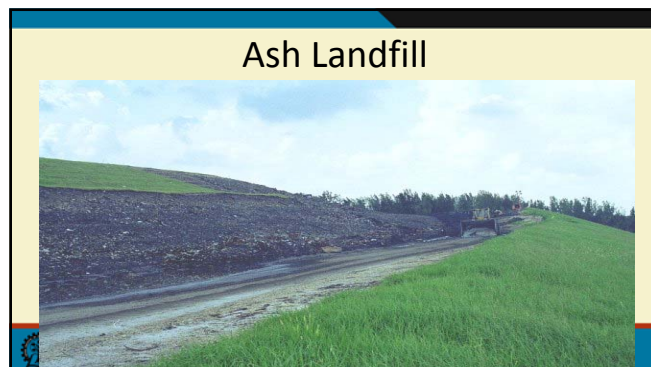
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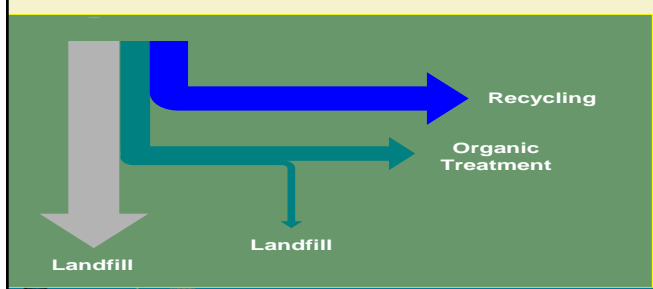
Terminology

- **Thermal treatment (or incineration):** a range of processes where temperature is used to reduce the volume of waste and to render it harmless.
- **Waste to Energy (WTE):** as above, with the recovery of heat energy to produce steam and/or generate electricity.
- **Conventional WTE:** mass burn, fluidized bed, modular, rotary kiln, (refuse derived fuel)
- **Advanced WTE:** gasification, pyrolysis, plasma

The role of thermal treatment

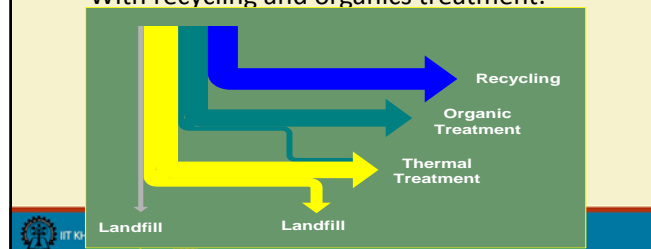
- Waste volume reduction, preservation of landfill space
 - Does NOT replace the need for a landfill
- Energy recovery from the solid waste stream
- Destruction of contaminants
- Reducing waste transportation requirements
- Dealing with waste here and now

The role of thermal treatment (2)



The role of thermal treatment (3)

- With recycling and organics treatment:



The role of thermal treatment (4)

- Last treatment of waste before land disposal
- Applied after recycling, organics management
- If recycling goal is 60%, then WTE can treat balance of waste
- Recovers remaining energy
- Converts energy into heat
- Electricity can be sold to the grid
- Offsets fossil fuel use for power generation



The role of thermal treatment (5)

- One tonne of waste can deliver 400 to 700 kWh of electricity to the grid
- One tonne of waste has the same energy as one barrel of oil, or a quarter tonne of coal
- 24 tonnes of waste can provide all the electricity for a Canadian home for a year
- One home of 4 persons: $(1.5 \text{ kg/person-day}) * 4 * 365 = 2.2 \text{ tonnes}$



How thermal treatment works

- Technologies offer different ways of releasing the energy in the waste
 - Conventional combustion/WTE
 - Advanced thermal treatment (Gasification/pyrolysis, plasma systems)
- WTE systems are essentially power plants using waste as fuel instead of coal, natural gas or uranium

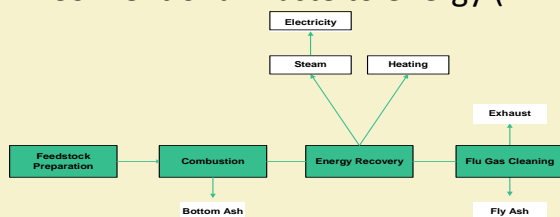


Conventional combustion technologies

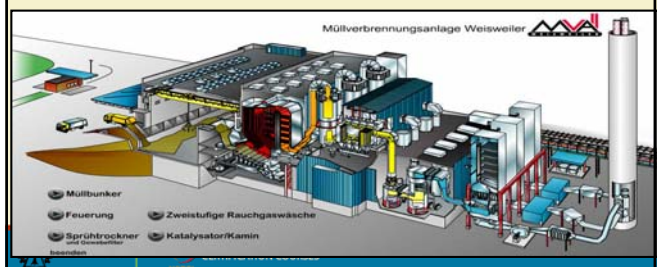
- Mass burn – most common (Burnaby, and upcoming York-Durham and Peel Region plan)
- Fluidized bed – mid sized and specialty applications (wood, coal)
- Modular – smaller systems
- Rotary kiln – hazardous and medical waste - rarely used for MSW
- Refuse Derived Fuel (RDF)– Dongara plant in GTA area



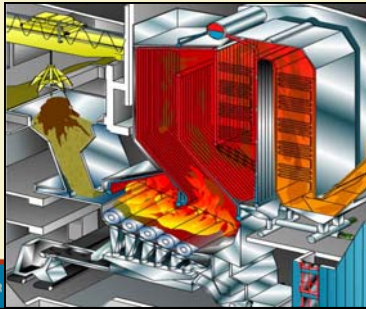
Conventional waste to energy (WTE)



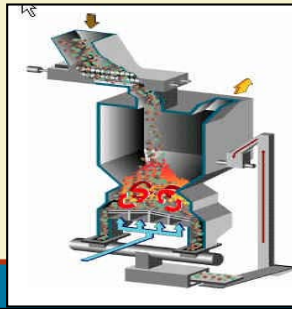
Mass burn: Facility overview



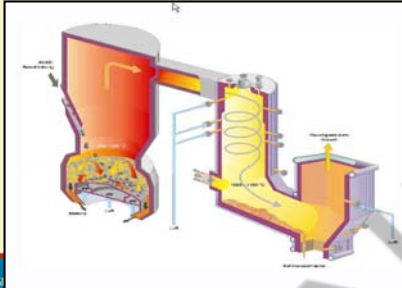
Mass burn: Furnace section



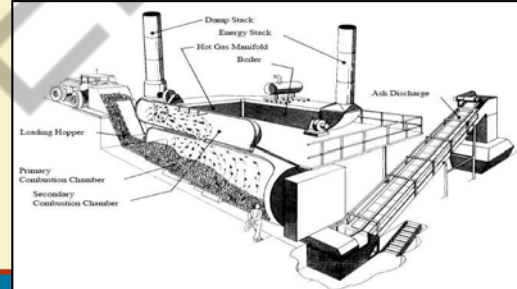
Fluidized bed furnace



Fluidized bed with ash melting



Modular controlled air combustion



Refuse derived fuel (RDF)

- Solid waste made into homogenous fuel
 - Can be sold and used off site, replacing other fuels such as coal or gas
 - Used by:
 - Cement kilns
 - Industry power boilers
 - Dedicated WTE plants

Ideal Incinerator

- rocking kiln incinerator
 - new variation of the rotary kiln
 - rotates only $\frac{1}{4}$ of a revolution, then reverses
 - thus, waste is subjected to less turbulence, but as a result releases less particulate matter into the secondary combustion chamber
 - no need to shred waste – less tumbling of larger items through the unit
- can handle a variety of waste (including hazardous waste)
 - solids, liquids, containers, ...
- slowly rotating refractory-lined cylinder that is slightly inclined and rocking:
 - insulation provided by fireclay, dense aluminum, silica, silicon carbide
- length to diameter ratio of 2:1 to 10:1
- rotation at 0.3 to 3 m/min
 - continually mixes waste, and entrains combustion air

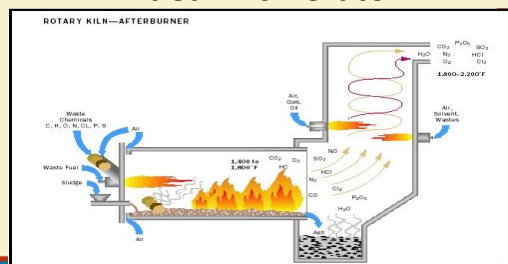
Ideal Incinerator

- combustion temperatures ranging from 800°C to 1100°C
- requires large amounts of excess air due to leakage
 - remember: excess air = 120 to 200%
- retention time varies in combustion chambers
 - 0.1 to 2 s for gases and liquids
 - minutes to hours for solids
- minimizing products of incomplete combustion (PICs) requires further oxidation of gases
 - afterburner (secondary combustion chamber)
 - residence time of up to 3 seconds
 - temperatures of 1100°C to 1300°C
- continuous ash generation
 - non-putrescible – does not decay
 - sterile – free of living organisms
 - inert – does not react in any way



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Ideal Incinerator



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www.pollutionissues.com/Ho-Li/Incineration.html

Disadvantages

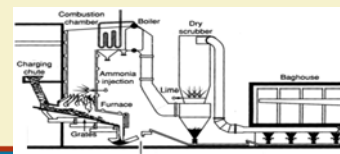
- high capital cost
- skilled operators required
- some materials are non-combustible
- some materials require supplemental fuel
- air contaminant potential
 - dioxin, mercury (90% reduction since 1995), particulate matter, ...
- volume of gas from incineration is 10 times as great as other thermochemical conversion processes
 - greater cost for gas clean-up/pollution control
- public disapproval
 - risk imposed rather than voluntary
 - incineration will decrease property value (perceived)
 - distrust of government/industry – ability to regulate



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Ash

- ash is the residue left over from the combustion of MSW
- bottom ash
 - recovered from combustion chamber
- heat recovery ash
 - collected in the heat recovery system (boiler, economizer, superheater)
- fly ash
 - particulate matter removed prior to sorbents
- air pollution control residue
 - combined with fly ash
- combined ash
 - most US facilities combine all ashes together



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Ash Treatment / Re-use

- there are a variety of methods of treating incinerator ash
- generally it is processed to standardize the material and remove contaminants
 - ferrous metal removal (magnetic separation)
 - non-ferrous metal (aluminum, copper) removal
 - screening and crushing
- so that it can be used as an aggregate
 - construction fill
 - road construction (asphalt, pavement concrete)
 - landfill daily cover
 - cement block production
 - treatment of acid mine drainage



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Flue Gas

- flue gas is the gas that exits the incinerator stack (i.e. the flue) to the atmosphere, and may include:
 - particulate matter
 - acid gases – principally H_2S and CO_2 (which produces H_2CO_3)
 - NO_x – $\text{NO} + \text{NO}_2$ – react to form smog, acid rain
 - CO – a weak greenhouse gas
 - organic hazardous air pollutants – e.g. PCDD, PCDF, ...
 - metal hazardous air pollutants – e.g. mercury, ...
- each of these components must be considered in the design of air pollution control facilities



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Particulates

- solid particles – usually defined as $PM_{2.5} < 2.5\mu m$
- condensable PM – vapours that condense to form particulates
- causes
 - too low of a combustion temperature (incomplete combustion)
 - insufficient oxygen or overabundant EA (too high temperature)
 - insufficient mixing or residence time
 - too much turbulence, entrainment of particulates in the air stream
- control
 - cyclones – can remove large particles, but not effective for removal of small particulates
 - electrostatic precipitator – efficient removal using an electrostatic charge
 - fabric filters (bag houses)



Acid Gases

- come from Cl, S, N, F in the refuse
 - plastics
 - textiles
 - rubber
 - yard waste
 - paper
- uncontrolled incineration generates 18 – 20% HCl with $pH = 2.0$
- acid gas scrubbers (SO_2 , HCl, HF) usually ahead of electrostatic precipitator or bag house
 - wet scrubber – like a venturi scrubber
 - spray dryer
 - dry scrubber injectors – injects a dry sorbent like lime



Air Pollution Control Process

- electrostatic precipitator
 - removes particles from a flowing gas using an induced electrostatic charge
- bag houses
 - uses engineered fabric filter long bags/tubes to remove particulate matter
- acid gas scrubbers
 - wet scrubber
 - dry scrubber
 - chemicals added in slurry to neutralize acids
- activated carbon
- selective non-catalytic reduction
- catalytic converters
 - recently started using them in Germany for converting NO_x and PCDD/F
 - using titanium dioxide (TiO_2) as a base

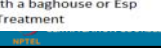


Comparison of Air Pollution Control Systems

| Parameter / Component | Spray Dry Absorber | Venturi | Packed Bed | Dry Electrostatic Precipitator |
|-----------------------|--------------------|---------------|--------------|--------------------------------|
| Particulate Removal | Poor to Fair | Good | Poor | Excellent |
| Heavy Metal Removal | Excellent * | Good | Poor | Good |
| Acid Gas Removal | Good to Exc. | Good | Excellent | Poor |
| Residue | Fly Ash | Scrub Liquor | Scrub Liquor | Fly Ash |
| Auxiliary Equipment | Bag House | Demister | Demister | Ash Handling |
| Needed | Handling | Liquid S&T ** | Liquid S&T | |
| Turndown | 3:1 | 2:1 | 5:1 | 5:1 |
| Plume Suppression | Easy | Difficult | Difficult | Easy |
| Pressure Drop | Low | High | Moderate | Low |
| Capital Cost | Moderate | Low | Low | High |

* when used with a baghouse or Esp

** Storage and Treatment

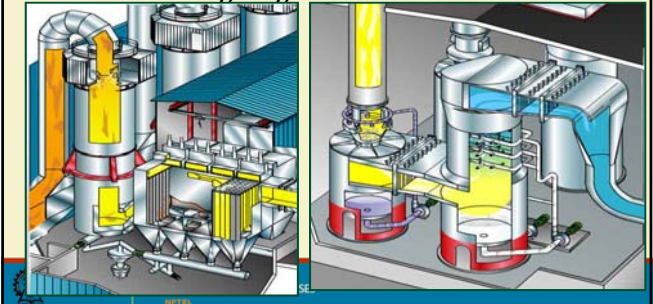


Major Components at WTE Plants (2)

- Air pollution control
 - Mature technology.
 - Systems available to meet most stringent air emission standards
 - Custom matched to combustion technology
 - WTE most highly regulated form of waste management
 - Emission standards more stringent than for most coal fired power plants or industrial boilers



Semi-dry, dry, and wet scrubbers



Major Components at WTE Plants (3)

- Solid Residues:
 - Conventional combustion
 - Metals recovered and recycled
 - Bottom ash and fly ash,
 - 25% by weight and 10% by volume of treated waste
 - Bottom ash suitable for road base, landfill cover or disposal
 - Fly ash usually needs to be stabilized before disposal
 - Advanced Combustion
 - Slag with varying amounts of fixed carbon, up to 30% by weight
 - Slag may be reduced by reprocessing
 - Plasma systems have almost no residue



Air Emissions

- WTE most highly regulated form of waste management
- Most countries have very strict standards
- EU and Ontario A7 guidelines considered to be the most stringent in the world
- Technologies have been developed and are applied to meet these standards
- In Europe, emissions from WTE are so low, that they are often considered irrelevant compared to industrial and transportation sources

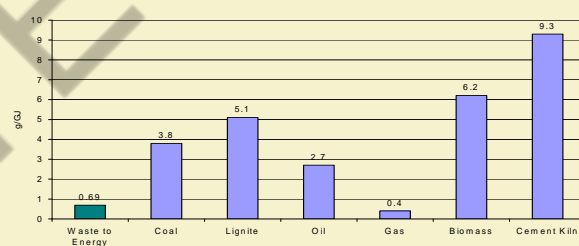


Comparison of Relevant Air Emissions from Selected Combustion Technologies

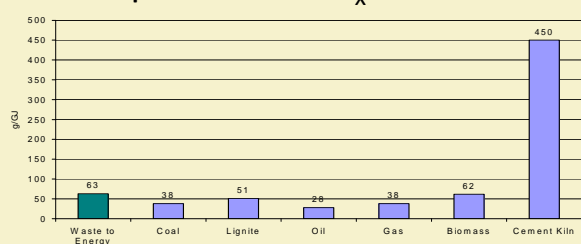
- Paper presented by Helmut Rechberger and Gerald Schoeller, Technical University of Vienna, 2006 CEWEP Congress
- Extensive emissions comparisons based on energy production (mg/GJ)
- WTE figures from 50 existing WTE facilities in Europe
- Cement kiln data from Association of German Cement Kilns
- Other data from literature



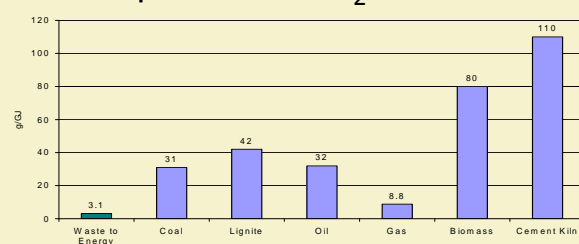
Comparison of Dust/Particulate Emissions



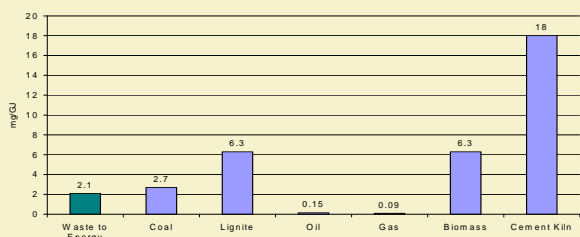
Comparison of NO_x Emissions



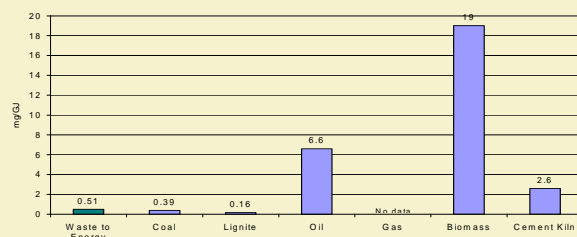
Comparison of SO₂ Emissions



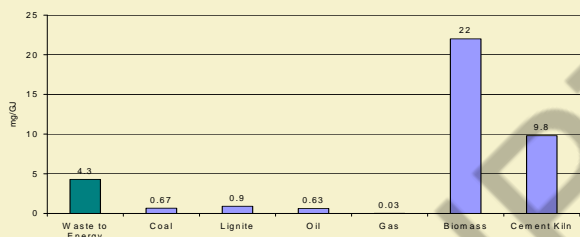
Comparison of Mercury Emissions



Comparison of Cadmium Emissions



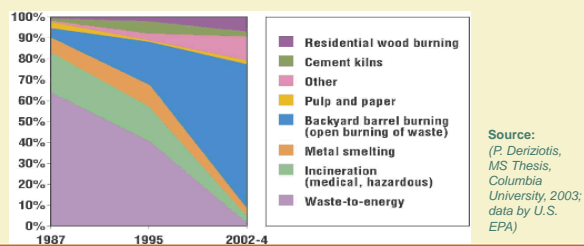
Comparison of PCDD/F Emissions



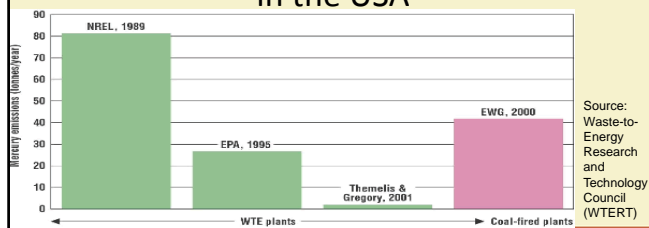
Notes to emissions slides

- Values shown in previous slides are for existing facilities, some of which are older
- Newer facilities are made to meet more stringent emission targets
- Metro Vancouver's Burnaby WTE facility often has no detectable dioxins
- New technologies exist to remove mercury from flue gas

Dioxin Emissions in the USA



Reduction of Mercury from WTE in the USA



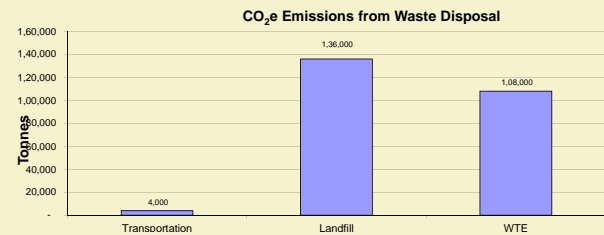
Carbon Dioxide (CO₂)

- WTE emits CO₂ like any other combustion process
- 40 to 60% is biogenic and is therefore part of the active carbon cycle
 - Unlike CO₂ from fossil fuels, this does not count as contributing towards climate change
- Electricity from WTE reduces the need to generate power from other sources (fossil fuels, nuclear)
- Generally, WTE results in less CO₂ equivalents than landfilling
- One European study calculated that in the EU:
 - WTE emits 0.348kg CO₂ eq. / kg of waste
 - Landfills emit 0.69 kg CO₂ eq. / kg of waste

Flact Management Consultants.
Waste to Energy and the revision
of the Waste Framework Directive.
Opportunities to reduce climate
change by using energy from
waste. FF/KW/2006.023-final.
Delft, January 2007



CO₂ of Transportation and WTE



Costs of WTE

- High initial capital costs
- Operating costs generally offset by energy sales (for larger facilities)
- Tipping fees must generally cover capital repayment
- Once paid for, WTE can be revenue generator
- Facility life 20 to 50 years

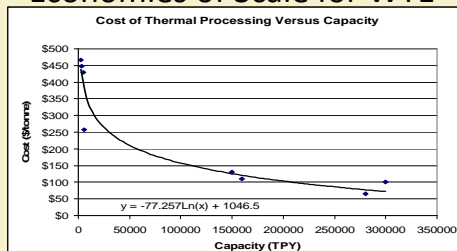


Revenues from WTE

- Tipping fees
- Electricity sales
- Steam sales (cogeneration, if available)
- District heat (if feasible)
- Recycled metals from ash or upfront processing
- CO₂ credits (future)



Economies of Scale for WTE



Political/social acceptance of WTE as diversion

- Europe
 - In practice used as diversion
 - Looking for official recognition to capitalize on tax credits
- USA
 - In some states considered renewable fuel
 - In other states not recognized as diversion
- Japan
 - Over 90% of solid waste combusted, mostly for energy
- Canada
 - Alberta recognizes WTE as diversion, Ontario does not, BC is undecided



Issues: Opposition and hurdles

- Negative public perception
- Lack of public awareness of technological progress and high regulated standards
- Large initial investment needed
- Higher operating costs than most local landfills
- Need for long term waste supply contracts



Issues: Opposition and hurdles (2)

- Full cost accounting and long term benefits rarely considered
- Waste has not yet been defined as renewable energy in Canada
- GHG credits are difficult to define and do not flow into the economics calculations



WTE in the USA

- 65 mass burn plants
 - 20 million tonnes per year capacity total
- 9 modular and 10 RDF plants
 - About 5 million tonnes per year capacity
- 15 RDF plants
 - 6 million tonnes per year
- 13% of USA waste managed by WTE



Comparison of WTE with selected Renewable Energy Sources in USA

| Energy Source | % of Renewable energy |
|-----------------|-----------------------|
| – Geothermal | – 28% |
| – WTE | – 28% |
| – Landfill gas | – 14% |
| – Wood/biomass | – 17% |
| – Solar thermal | – 2% |
| – Wind | – 11% |

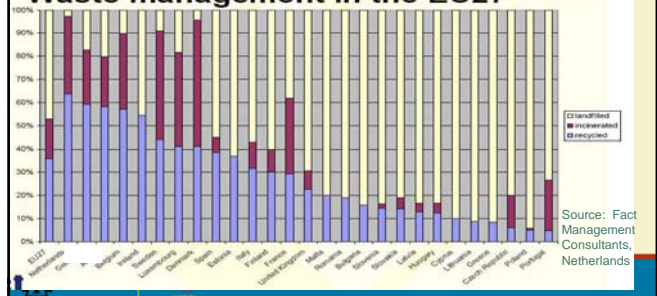


WTE in Europe

- More than 370 WTE plants with total annual capacity over 53 million tonnes
- Average EU recycling rate 36% - long term goal 60%
- EU WTE rate 17%
- Landfilling in EU 48%
- Landfill Directive progressively prohibits landfilling of organic materials
- High cost of energy = good revenue from heat and electricity
- Carbon credits enhance economics of WTE and help meet national reduction goals



WTE, Recycling and Landfilling in Europe Waste management in the EU27



Isle of Man, UK

- 200 tonnes per day



Lille, France



Karlsruhe, Germany

- Gasification Plant (shut down, but similar



Paris, France

- 350 tonnes per day



Vienna Austria

- Designed by famous artist Hundertwasser



The Japan Experience

- Very strict land disposal guidelines
 - No raw waste
 - No ash without stabilization
- Over 90 % of solid waste combusted, mostly with energy recovery
- 2300 combustion facilities in Japan
- 23 WTE facilities in Tokyo
- High standards for social integration and environmental performance
- Double typical north American/European costs



Japanese WTE and Sludge burning



Miscellaneous Issues

- diversity of waste
 - requires that better incineration controls are used to ensure that complete combustion occurs
- delivering constant waste to the combustion chamber
- maintaining sufficient temperature
 - can not start and stop operation
- replacement of refractory material is expensive
- competing against 3Rs for input material
 - common sense would suggest it's a problem
 - the same material that is useful from recycling is also combustible
 - practical experience suggests it's not
 - we do not capture all the recyclable material
 - combination of recycling and incineration is a more complete and integrated waste management approach

Effectiveness of Incinerators

- destruction efficiency (DE)
 - takes place in the combustion chamber
- removal efficiency (RE)
 - takes place in the APCE
- DRE is a combination of both
- required efficiencies:
 - most organic compounds at 99.99%
 - PCBs at 99.9999%
 - required to minimize formation of dioxins
 - in addition, HCl
 - particulate matter
- example of PCB DRE
 - inlet = 637 kg/h
 - outlet = 0.012 kg/h

$$DRE = \left(\frac{637 - 0.012}{637} \right) \times 100 = 99.9981 \rightarrow \text{FAILS}$$

Energy from Waste

- energy from waste (EFW) facilities use high temperature to extract energy from the trash
 - generate electricity or heat/steam from the waste combustion
 - and employ sophisticated emissions control systems
- incinerators only reduce the volume of the trash
- typical mass-burn 1,500 tonne per day (TPD) EFW facility produces enough clean, renewable electricity for 45,000 homes
 - which saves more than 450,000 barrels of oil each year
- offers net benefits in terms of greenhouse gases
 - emits 2/3 less carbon dioxide (CO₂) than coal fired power plants
 - emits 2/3 less carbon dioxide (CO₂) than oil and natural gas power plants
- life-cycle analysis of a 1,500 TPD EFW facility
 - annually reduces about 270,000 tons of CO₂ equivalent emissions
 - Europe ~ 400 plants and 60,000,00 tonnes per year (500 TPD)

Co-generation

- takes advantage of lost heat for another process
 - thermal power plants reject 50 – 65% heat to the environment
- local generation station near a facility that needs heat
 - transport the heat via low temperature hot water
 - use steam if distance less than 4 – 5 kilometers
- various fuels can be used to generate electricity
 - natural gas, coal, wood residue, MSW, biofuels
- captured heat used for secondary processes
 - greenhouse operation, paper producing facilities, petrochemical industry, bio energy, fuel cells, WWTP, Universities
- an example – Stuttgart WWTP
 - dries it's sludge and incinerates it with MSW – produces heat
 - also heat from anaerobic digestion
 - used to produce electricity
 - methane gas from anaerobic digestion piped into the natural gas system

Co-incineration

- waste can also be burned at facilities that are not dedicated to waste incineration
- the main candidates are:
 - steam and electricity producers
 - blast furnaces
 - lime and cement kilns
- few technical barriers exist as long as the waste is pre-treated to suit the primary (existing) process
- the main limitation is the composition of the waste and its possible contamination that might impact the industrial process


Pyrolysis, Combustion, Gasification

- Pyrolysis, gasification and combustion have been considered as separate thermochemical conversion processes for general classification. **However, in a gasification process, both pyrolysis and combustion processes take place.**
- Pyrolysis** is the destructive distillation of carbon-based materials with heat in the absence of


oxygen into gaseous products, char, condensable

Advanced thermal technologies

- Gasification and pyrolysis
 - Converts solids into synthetic gas
 - Gas is cleaned before combustion or other uses
 - Complex technology
- Plasma
 - Ultra high temperature process, total organics destruction
 - Makes synthetic gas
 - Creates vitrified slag
 - **Lowest residuals**



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Advanced thermal technologies. gasification/pyrolysis

```


graph LR
    A[Feedstock Preparation] --> B[Gasification or Pyrolysis]
    B --> C[Syngas Cleaning]
    B --> D[Char / Ash]
    C --> E[Residue / Ash]
    C --> F[Energy Recovery]
    F --> G[Steam]
    F --> H[Gas Turbine or Recip. Engine]
    F --> I[Exhaust]
    G --> J[Electricity]
    H --> J
  
```

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Pros and cons of advanced thermal technologies

| <u>Pros</u> | <u>Cons</u> |
|---|---|
| <ul style="list-style-type: none">• Few air emissions during syngas generation• Lower CO₂ generated when syngas formed• Ash can be vitrified with some processes• Recovery of energy from waste• Better environmental perception | <ul style="list-style-type: none">• Syngas must be cleaned, leaving residues• CO₂ formed when syngas burned• Vittrification has high energy requirement/cost• Often lower energy recovery efficiency than conventional combustion systems• No real environmental advantages over combustion if syngas is used for heat/power |



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Thermochemical Conversion Processes

The diagram illustrates the thermochemical conversion process, starting with Feedstock entering a yellow box labeled 'Conversion Technology'. This box is divided into 'Pyrolysis' and 'Gasification'. The output of this stage is a blue box labeled 'Primary Products', which contains 'char', 'tars and oils', and 'gas'. Arrows from 'char' and 'tars and oils' point to a pink box labeled 'Product Recovery'. Arrows from 'gas' and 'tars and oils' point to a pink box labeled 'Energy Recovery'. Both 'Product Recovery' and 'Energy Recovery' have arrows pointing to a green box labeled 'Secondary Products', which contains 'chemicals', 'gasoline', 'methanol', and 'ammonia'. Below the primary products, there is a section for 'Combustion' (labeled 'Balgiorne at 200°C') which includes 'heat' and 'power'. This section is connected to a 'Boiler'.

Conversion Technology

Primary Products

Secondary Products

Feedstock

Pyrolysis

Gasification

char

tars and oils

gas

Product Recovery

Energy Recovery

chemicals

gasoline

methanol

ammonia

Combustion

Boiler

heat

power

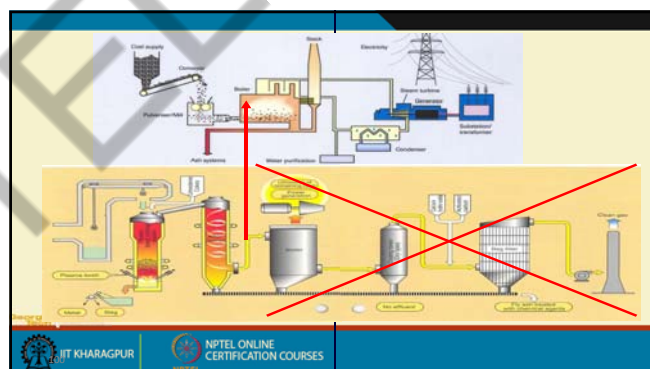
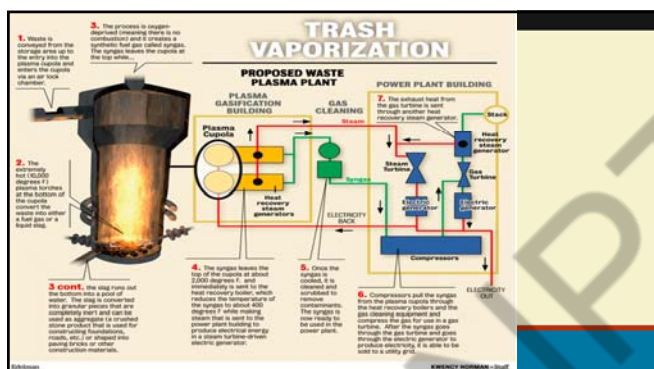
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Plasma Arc

- plasma is a state of matter created by heating a gas in which a certain portion of the particles are ionized
- plasma can be created using almost any kind of gas
 - nitrogen, oxygen, air
- heated to extreme temperature
 - 2000 – 6000°C
- ionized gas conducts electricity
- this energy is transferred to the waste material
 - excites molecular bonds such that the materials break into elemental atoms
 - all waste constituents are completely melted into a solid mass called dross
- minerals can be recovered from the dross
- all known contaminants effectively treated
 - municipal, hazardous, hospital

Plasma Arc

- the process generates PCG – plasma converted gas
 - hydrogen fuel as energy
- lower volume of gas requiring treatment
- power consumption
- double the cost of
- but, a smaller environmental footprint



Example Problem #1

- determine the stoichiometric amount of air required for the combustion of an organic solid waste
 - the waste is defined as C_5H_{12}
 - excess air requirements set at 150% per tonne
- first, write the stoichiometric equation:

$$C_5H_{12} + 8O_2 \rightarrow 5CO_2 + 6H_2O$$

$$\frac{72}{256} \rightarrow \frac{3555.6}{3555.6}$$
 - so \rightarrow 256 grams of O_2 oxidizes 72 grams of C_5H_{12}
- therefore, oxygen = $\left(\frac{256}{72}\right) \cdot \left(1000 \frac{\text{kg waste}}{\text{tonne}}\right) = 3555.6 \text{ kg } O_2/\text{tonne}$
- assume that oxygen is 23.15% of air by weight

Example Problem #1

- therefore, $\text{air} = \left(\frac{3555.6 \text{ kg } O_2/\text{tonne}}{0.2315 \text{ kg } O_2/\text{kg air}}\right) = 15,359 \text{ kg air/tonne}$
- the density of air is $\left(\frac{15,359 \text{ kg air/tonne}}{1.2928 \text{ kg air/m}^3}\right) = 11,880 \text{ m}^3 \text{ air/tonne}$
- therefore, 1 tonne of waste requires 11,880 m³ of air for complete combustion

$$\text{m}^3/\text{kg} \cdot \text{fuel} = 24.6 \left[\left(\frac{C}{12}\right) + \left(\frac{H}{4}\right) + \left(\frac{S}{32}\right) - \left(\frac{O}{32}\right) \right]$$
- or we can use the simplified equation:

Example Problem #1

- element percentages: $\% C = \left(\frac{5(12)}{5(12) + 12(1)} \right) = \frac{60}{72} = 0.833$
- $\% H = \left(\frac{12}{5(12) + 12(1)} \right) = \frac{12}{72} = 0.167$
- oxygen requirements: $O_2 \text{ (m}^3/\text{kg} \cdot \text{fuel)} = 24.6 \left[\left(\frac{0.833}{12} \right) + \left(\frac{0.167}{4} \right) \right]$
 $= 2.733 \text{ m}^3/\text{kg fuel}$
- air requirements: $\text{air} = \left(\frac{2.733}{0.2315} \right) = 11.808 \text{ m}^3/\text{kg} = 11,808 \text{ m}^3/\text{tonne}$
- excess air @ 150%: $\text{air} = (11,808 \times 1.50) = 17,710 \text{ m}^3/\text{tonne}$



Example Problem #2

- determine the amount of air required for the combustion of an organic solid waste
 - an organic waste is defined as $C_{760}H_{1980}O_{875}N_{13}S$
 - excess flowrate/hour if 500 tonnes/day waste (dry basis)
 - excess air requirements set at 175% per tonne
 - for efficiency, incinerator operation is $\frac{24}{7}$ day
- waste processing: $= \left(500 \frac{\text{tonne}}{\text{day}} \times \frac{1}{24 \text{ hour}} \times \frac{1000 \text{ kg}}{\text{tonne}} \right) = 20,833 \text{ kg/hr}$
- molar mass = $(760 \times 12.0) + (1980 \times 1.0) + (875 \times 16.0) + (13 \times 14.0) + (32 \times 1.0)$
 $= 25314.1 \text{ g/mol}$



Example Problem #2

- element percentages: $\% C = \left(\frac{760(12)}{25314} \right) = \frac{9120}{25314} = 0.360$
- $\% H = \left(\frac{1980(1)}{25314} \right) = \frac{1980}{25314} = 0.078$
- $\% O = \left(\frac{875(16)}{25314} \right) = \frac{14000}{25314} = 0.553$
- $\% N = \left(\frac{13(14)}{25314} \right) = \frac{182}{25314} = 0.007$
- $\% S = \left(\frac{1(32.1)}{25314} \right) = \frac{32.1}{25314} = 0.001$ ← * very small contribution



Example Problem #2

- using the simplified equation: $O_2 \text{ (m}^3/\text{kg} \cdot \text{fuel)} = 24.6 \left[\left(\frac{C}{12} \right) + \left(\frac{H}{4} \right) + \left(\frac{S}{32} \right) - \left(\frac{O}{32} \right) \right]$
 $= 24.6 \left[\left(\frac{0.360}{12} \right) + \left(\frac{0.078}{4} \right) + \left(\frac{0.001}{32} \right) - \left(\frac{0.553}{32} \right) \right]$
 $= 24.6 \left[(0.0300) + (0.0196) + (0.00004) - (0.0173) \right]$
 $= 24.6 (0.0323) = 0.7953 \text{ m}^3/\text{kg fuel}$
- oxygen = $(20,833 \text{ kg fuel/hr} \times 0.7953 \text{ m}^3/\text{kg fuel}) = 16,566 \text{ m}^3/\text{hr}$
- amount of O_2 per hour:



Example Problem #2

- amount of air per hour: $\text{air} = \left(\frac{16,566 \text{ m}^3/\text{kg}}{0.2315} \right) = 71,570 \text{ m}^3/\text{hr}$
- based on excess air = 175% $\text{air} = 1.75 \cdot (71,570 \text{ m}^3/\text{hr}) = 125,248 \text{ m}^3/\text{hr}$
- power requirements: $Q = \frac{125,248 \text{ m}^3/\text{hr}}{3600 \text{ sec/hr}} = 34.8 \text{ m}^3/\text{s}$
- $P = \frac{Q \cdot \gamma \cdot H}{\eta}$
 $\gamma = \text{specific weight} = 11.8 \text{ N/m}^3$
 $H = \text{head drop} = 10 \text{ m}$
 $\eta = \text{efficiency} = 0.70$
 $P = \frac{(34.8 \times 11.8 \times 10)}{0.70} = 4,130 \text{ N} \cdot \text{m/s} = 4.13 \text{ kW}$



Example Problem #2

- using a centrifugal fan, with an efficiency of 70%:
 $P = \frac{5.7 \text{ kW}}{0.70} = 7.6 \text{ Hp}$
- we need to blow 125,000 m³/hour of air into the combustion chamber, using a 7.6 Hp blower



Example Problem #3

Let's Look at Wood from C&D Debris

- Wood:

%C = 41.20, %H = 5.03, %O = 34.55, %N = 0.24, %Cl = 0.09
%S = 0.07, %Moisture = 16, %Ash = 2.82

$$VS = \frac{M_{VS}}{M_{Dry}} \quad VS = \frac{100 - 16 - 2.82}{100 - 16}$$

VS = 96.6%



Let's Find the Heat Value

- Use the Dulong Equation

$$BTU / lb = 145C + 610\left(H - \frac{O}{8}\right) + 40S + 10N$$

$$BTU / lb = 145(41.2) + 610\left(5.03 - \frac{34.55}{8}\right) + 40(0.24) + 10(0.07)$$

BTU per pound = 6,418

Heat Value in KJ/kg = 14,890



Let's Compare to Value in Table

| ULTIMATE ANALYSIS OF MUNICIPAL SOLID WASTE COMPONENTS (percent by weight) | | | | | | | | | |
|--|-------|------|-------|------|------|------|----------|-------|--------------|
| MATERIAL | C | H | O | N | Cl | S | Moisture | Ash | HHV (Btu/lb) |
| Mixed Waste | 27.5 | 3.7 | 20.6 | 0.45 | 0.5 | 0.83 | 23.2 | 23.4 | 4,830 |
| Corrugated | 36.79 | 5.08 | 35.41 | 0.11 | 0.12 | 0.23 | 20 | 2.26 | 6,322 |
| Newsprint | 36.62 | 4.66 | 31.76 | 0.11 | 0.11 | 0.19 | 25 | 1.55 | 6,233 |
| Magazines | 32.93 | 4.64 | 32.85 | 0.11 | 0.13 | 0.21 | 16 | 13.13 | 5,466 |
| Other Paper | 32.41 | 4.51 | 29.91 | 0.31 | 0.61 | 0.19 | 23 | 9.06 | 5,481 |
| Plastics | 56.43 | 7.79 | 8.05 | 0.85 | 3.00 | 0.29 | 15 | 8.59 | 11,586 |
| Rubber/Leather | 43.88 | 6.37 | 11.62 | 1.34 | 4.97 | 1.17 | 10 | 22.49 | 8,433 |
| Wood | 41.20 | 5.03 | 34.55 | 0.24 | 0.09 | 0.07 | 16 | 2.82 | 6,933 |
| Textiles | 37.23 | 5.02 | 27.11 | 3.11 | 0.27 | 0.28 | 25 | 1.98 | 6,595 |
| Yard Waste | 23.29 | 2.93 | 17.54 | 0.89 | 0.13 | 0.15 | 45 | 10.07 | 4,065 |
| Food Waste | 17.93 | 2.55 | 12.85 | 1.13 | 0.38 | 0.06 | 60 | 5.10 | 3,265 |

Source: D.A. Tillman, "The Combustion of Solid Fuels & Wastes," Academic Press, San Diego, 1991

BTU per pound = 6,418



BTU Value

- Values are close (note that Dulong Equation is only an estimate and other methods of estimation are out there).
- But what about water? Water as not factored in.
- Remember, energy is consumed in the evaporation of water. 1,040 BTU/lb



Heat Consumed by Evaporation

$$\text{Heat Consumed by Evaporation} = \left(\frac{0.16 \text{ lb water}}{1 \text{ lb waste}} \right) \left(\frac{-1,040 \text{ BTU}}{\text{lb water}} \right)$$

$$\text{Heat Consumed by Evaporation} = -166 \frac{\text{BTU}}{\text{lb waste}}$$

- Actual Heat Value = 6,252 BTU/lb = 14,505 KJ/kg



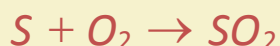
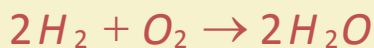
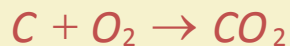
C&D Debris Wood is not Pure Wood

- Other components: Metal, Dirt
- If we assume X% contamination, we could reduce our heat value by that amount.
- But for this example, let's work with pure wood



Process to Calculate Air Requirement

- Use basic equations:



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Assumptions for Solution

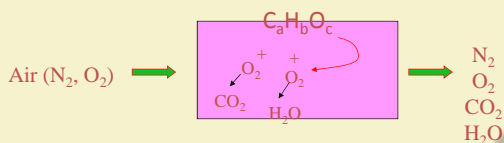
- Only consider oxidation of carbon and hydrogen for air demand.
- Let's go ahead and work everything from a total weight standpoint.
- We will operate at 50% excess air.
- Remember, oxygen in the waste contributes some oxygen toward combustion.



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Contribution of O₂



Wood:

%C = 41.20, %H = 5.03, %O = 34.55, %N = 0.24, %Cl = 0.09
 %S = 0.07, %Moisture = 16, %Ash = 2.82, VS = 96.6%



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Start With Carbon

$$41.2 \left(\frac{\text{kg C}}{100 \text{ kg waste}} \right) \left(\frac{1 \text{ kg-mole C}}{12 \text{ kg C}} \right) \left(\frac{1 \text{ kg-mole O}_2}{1 \text{ kg-mole C}} \right) \left(\frac{22.4 \text{ cum O}_2}{1 \text{ kg-mole O}_2} \right) \left(\frac{100 \text{ cum air}}{21 \text{ cum O}_2} \right)$$

→ $\left(\frac{366 \text{ cum air}}{100 \text{ kg waste}} \right)$

- Note: this is the air demand from Carbon only!!!



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Next, Hydrogen

$$5.03 \left(\frac{\text{kg H}}{100 \text{ kg waste}} \right) \left(\frac{1 \text{ kg-mole H}}{1 \text{ kg H}} \right) \left(\frac{1 \text{ kg-mole H}_2}{2 \text{ kg-mole H}} \right) \left(\frac{1 \text{ kg-mole O}_2}{2 \text{ kg-mole H}_2} \right) \left(\frac{22.4 \text{ cum O}_2}{1 \text{ kg-mole O}_2} \right) \left(\frac{100 \text{ cum air}}{21 \text{ cum O}_2} \right)$$

→ $\left(\frac{134 \text{ cum air}}{100 \text{ kg waste}} \right)$

- Note: this is the air demand from Hydrogen only!!!



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Now, O₂ Contribution

$$34.55 \left(\frac{\text{kg O}}{100 \text{ kg waste}} \right) \left(\frac{1 \text{ kg-mole O}_2}{32 \text{ kg O}} \right) \left(\frac{22.4 \text{ cum O}_2}{1 \text{ kg-mole O}_2} \right) \left(\frac{100 \text{ cum air}}{21 \text{ cum O}_2} \right)$$

→ $\left(\frac{115 \text{ cum air}}{100 \text{ kg waste}} \right)$

- Note: This is the air contribution from oxygen only!!!



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Net Air Demand

$$\left(\frac{366 \text{ cum air}}{100 \text{ kg waste}} \right) + \left(\frac{134 \text{ cum air}}{100 \text{ kg waste}} \right) - \left(\frac{115 \text{ cum air}}{100 \text{ kg waste}} \right)$$

$$= \left(\frac{385 \text{ cum air}}{100 \text{ kg waste}} \right)$$



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Factor in Excess Air

- We want to use 50% excess air

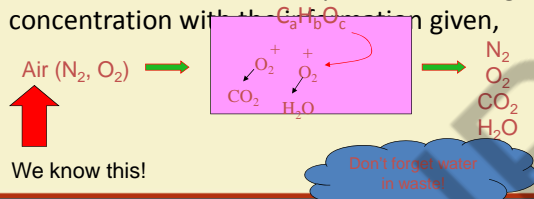
$$\left(\frac{385 \text{ cum air}}{100 \text{ kg waste}} \right) * 1.5 = 578 \frac{\text{cum air}}{100 \text{ kg waste}}$$



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Exit Gas Concentration

- You should be able to easily solve the exit gas concentration with the $C_a H_b O_c$ given,



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Future of thermal treatment

- Rising energy costs will make WTE attractive for power generation/heat utilization
- Increasing costs and long-term environmental concerns with landfills will support WTE
- Energy recovery increasingly recognized as logical and integral part of WM process
- Waste increasingly recognized as renewable energy with GHG benefits
- European legislation supports WTE as opposed to landfilling



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Future challenges of thermal treatment

- Education required to achieve a balanced public perception and acceptance
- Increasing thermal efficiencies
- Finding markets for heat
- Reducing operating costs and increasing revenues from sale of energy
- Regionalization required to achieve economies of scale
- Regulatory and policy support needed
- Acceptance of WTE as renewable energy



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What is a Landfill?

- Concept fostered in early 20th century
- An area of land that has solid waste deposited on it in such a quantity to noticeably change the surface elevation.



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Why to use a landfill?



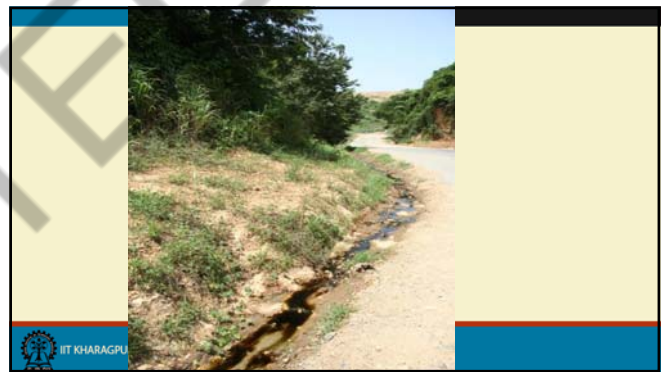
Potential Landfill Problems

- Landfill can present problems with respect to:
 - Spread of disease
 - Odors
 - Fires

} Controlled by sanitary
landfill techniques

- Contamination of groundwater
- Gas emissions

} Controlled by
modern landfill
design



Sanitary Landfill

- Landfills may be:
 - Excavated and filled
 - Fill existing depressions
 - Built up from the ground
 - A combination of above
- Operate landfills in a controlled safe fashion
 - Use cover soil
 - Excavate cells
 - Compact the waste
 - Control access

Modern Landfills are Engineered Structures

- Designed to Contain Leachate and Minimize Release of Pollutants from the Landfill

Leachate

- Leachate is the liquid (or wastewater) that forms when water (rainfall, groundwater) travels through solid waste
- Leachate can migrate into underlying groundwater, resulting in contamination
- Leachate can contain many different chemicals, depending on what is in the solid waste



Landfill Gas

- Landfill gas consists primarily of methane and carbon dioxide
- Results from the anaerobic decomposition of biodegradable solid wastes



Typical Regulatory Requirements

- Location restrictions
 - Airports
 - Wetlands
 - Fault lines
 - Unstable areas
 - Endangered species



Typical Regulatory Requirements

- Liners -- Low permeability barrier layers
 - Compacted soil (clay)
 - Geomembranes (plastic)
 - Composites of both
- Liner keep leachate from migrating out of the landfill
- Leachate must be collected and removed



Single Liner System

- One liner consisting of compacted soil **or** geomembrane



Composite Liner

- A single liner consisting of compacted soil and geomembrane in intimate contact



Double Liner

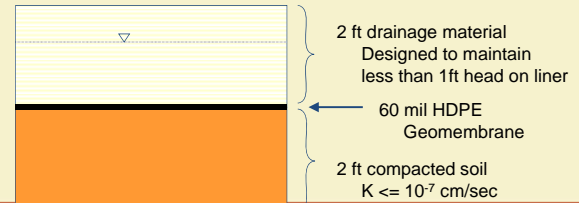
- A liner system with low permeability barrier layers with a leak detection system layer in between. The upper and lower components are either compacted soil, geomembrane, or composite.



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Typical Subtitle D Liner

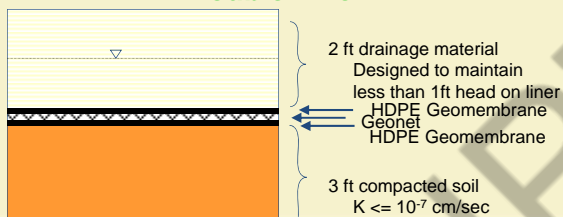
Single Composite Liner



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Typical Subtitle C Liner

Double Liner



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Types of Geomembrane Materials

- HDPE
- PVC
- VLDPE
- PP



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What Controls Head on the Liner?

- Liner Slope
- Pipe Spacing
- LCS Hydraulic Conductivity
- Impingement Rate



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What is a Geosynthetic Clay Liner?


- A manufactured product that contains a soil component (dry bentonite) contained in a fabric or affixed to a geomembrane.



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INTEGRATED WASTE MANAGEMENT FOR A SMART CITY
FOCUSSED ON MSW, C&D AND E-WASTE MANAGEMENT

End of Week-8

BRAJESH KUMAR DUBEY
DEPARTMENT OF CIVIL ENGINEERING



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