

Summer School on Fire Safety Science

– Wildland/WUI Fire Behavior

Combustion

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Outline

- Introduction
- Pyrolysis and Combustion
- Combustion: the Thermodynamics Viewpoint
- Combustion Chemistry
- Flame Structure
- Flame Effects



Introduction to Combustion

- The purpose of this lecture is to:
 - Introduce the basic physical concepts used to describe combustion phenomena
 - Gas/liquid/solid-fueled combustion; flaming vs non-flaming combustion; premixed vs non-premixed/diffusion flame; momentum-driven vs buoyancy-driven flame; laminar vs turbulent flame
 - Coupling between pyrolysis and combustion: thermal feedback; mass loss rate (MLR) vs heat release rate (HRR); thermal degradation processes inside the solid biomass (drying, pyrolysis, char oxidation)



Introduction to Combustion

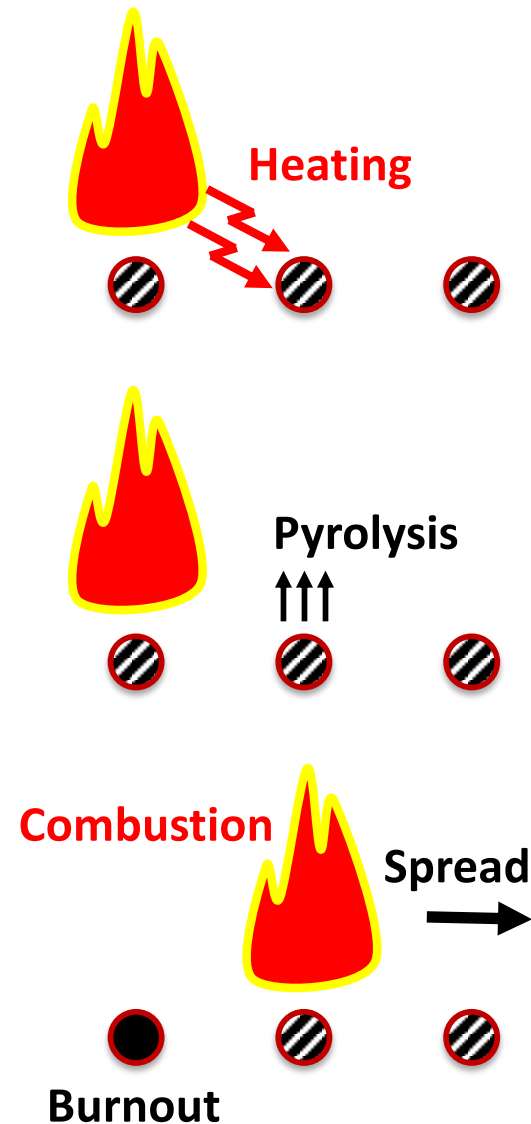
- The purpose of this lecture is to:
 - Introduce the basic physical concepts used to describe combustion phenomena
 - Global combustion equation model; over-ventilated vs under-ventilated fire conditions
 - Combustion chemistry; infinitely fast chemistry assumption
 - Structure of diffusion flames: mixture fraction variable; correlations for flame height; air entrainment; transport of combustion heat through convection, thermal radiation and firebrands; emissions of chemical species, including soot particles



Ignition and Combustion

■ Fire spread

- A repeated cycle of: (1) heating; (2) production of flammable vapors by the thermal degradation of vegetation biomass; and (3) combustion of these flammable vapors with ambient air
 - *Pyrolysis* = production of flammable vapors from thermally-degrading solid-phase vegetation biomass
 - *Ignition* = start of pyrolysis or start of combustion
 - *Combustion* = gas-phase oxidation of flammable vapors with heat release

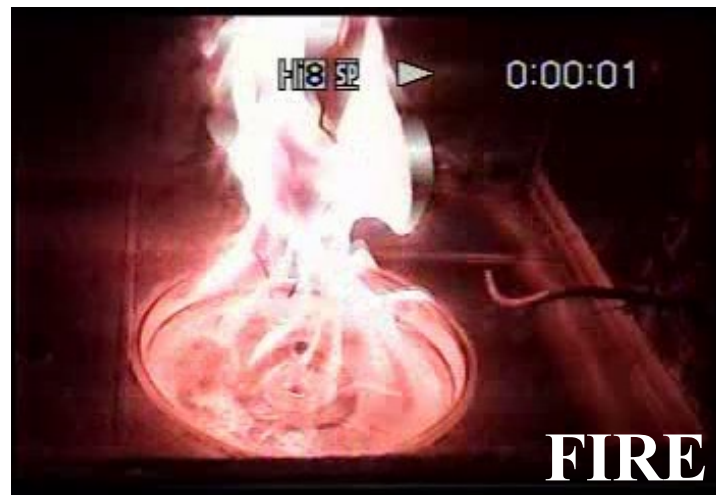
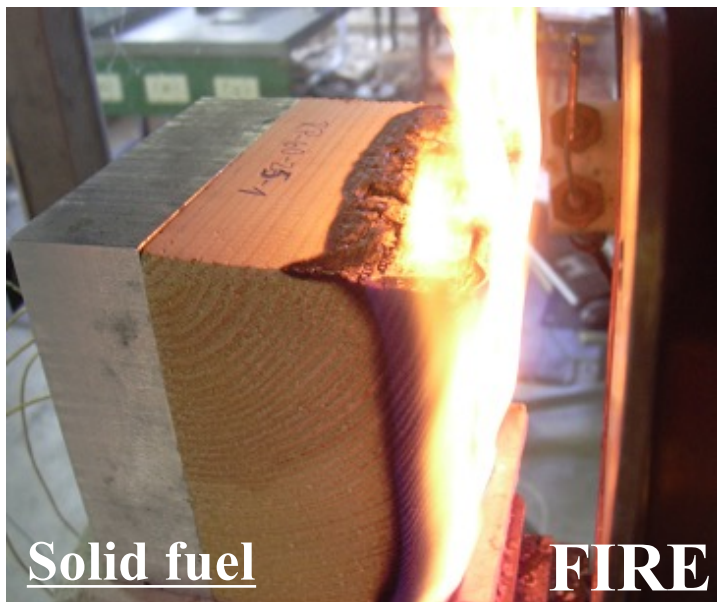




Introduction to Combustion

■ Basic features

- Source of oxygen (oxidizer): ambient air
- Source of fuel (hydrocarbons): gas/liquid/solid
 - From thereon assume solid fuel (vegetation biomass)





Introduction to Combustion

■ Basic features

- Non-flaming combustion mode (glowing/smoldering combustion): solid-gas phase reactions
- Flaming combustion mode: gas-phase reactions
 - From thereon assume flaming combustion



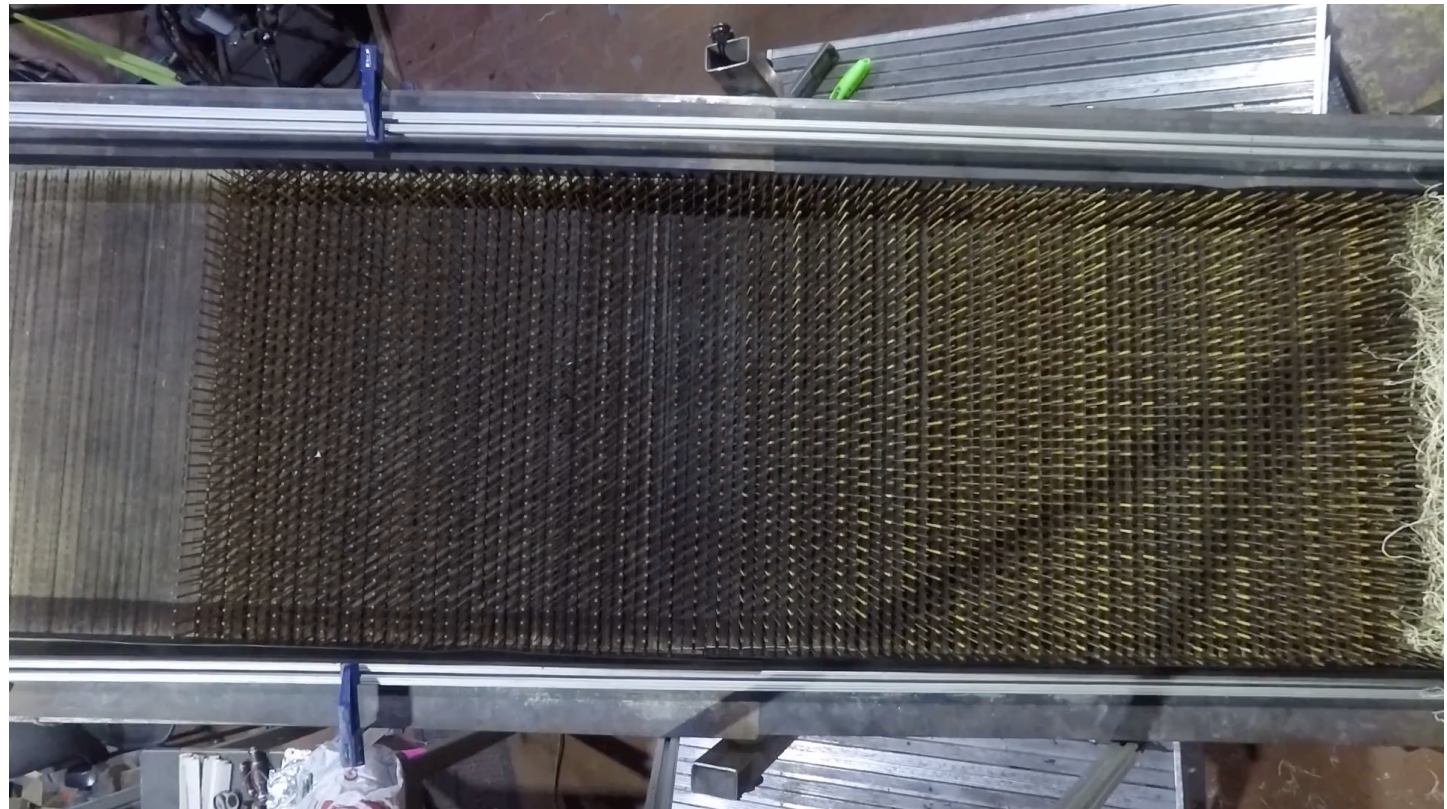


Introduction to Combustion

■ Basic features

● Flaming vs non-flaming combustion modes in flame spread

- Experiment performed at Missoula Fire Sciences Laboratory, USDA Forest Service
- Inclined surrogate fuel bed corresponding to an array of cardboard sticks
(Courtesy of M. Finney)

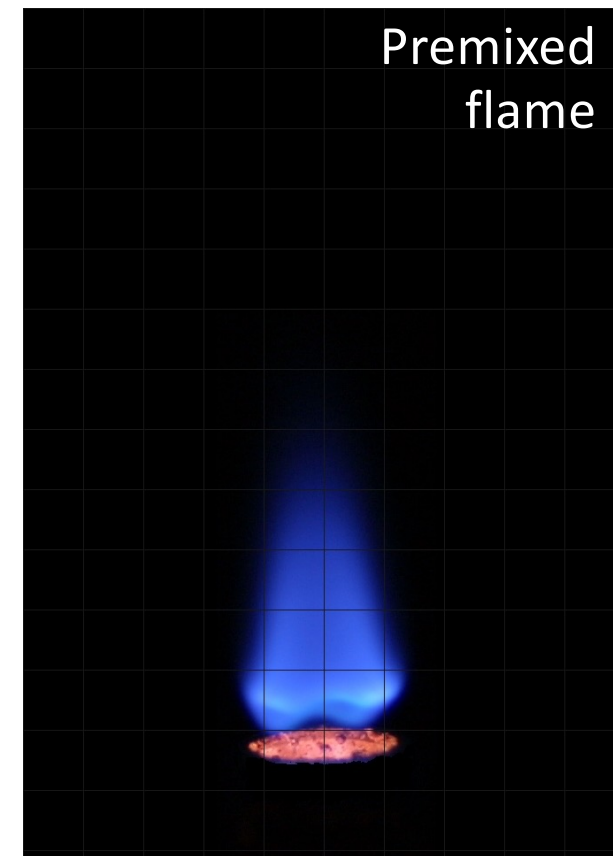
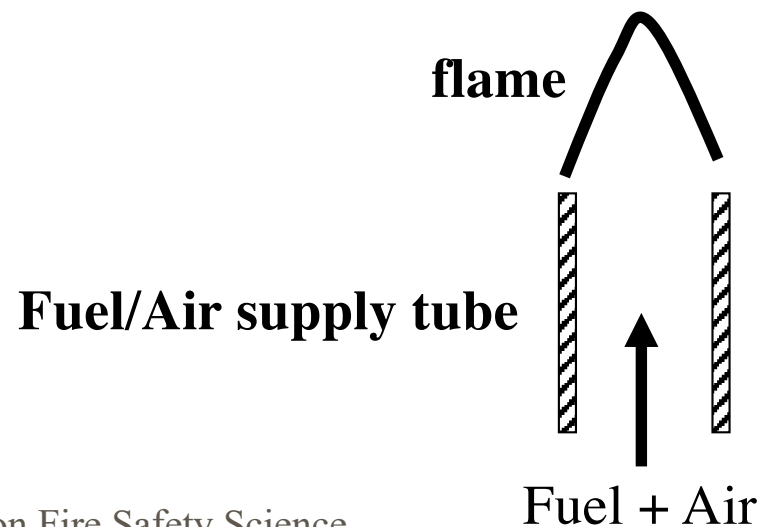




Introduction to Combustion

■ Basic features

- Premixed flame: fuel and air are mixed prior to entering the combustion zone

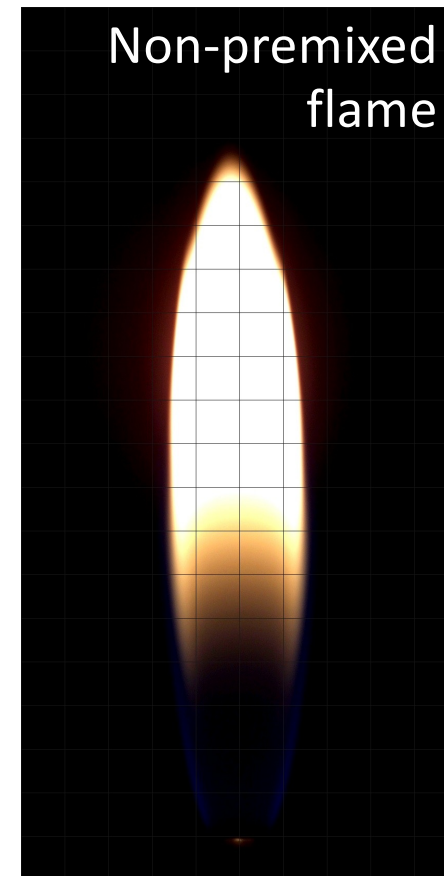
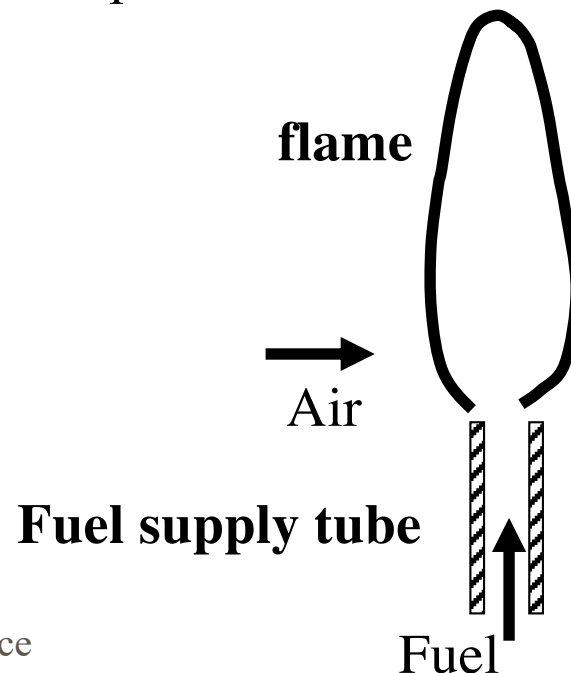




Introduction to Combustion

■ Basic features

- Non-premixed/diffusion flame: fuel and air remain separated prior to entering the combustion zone
 - From thereon assume non-premixed flame



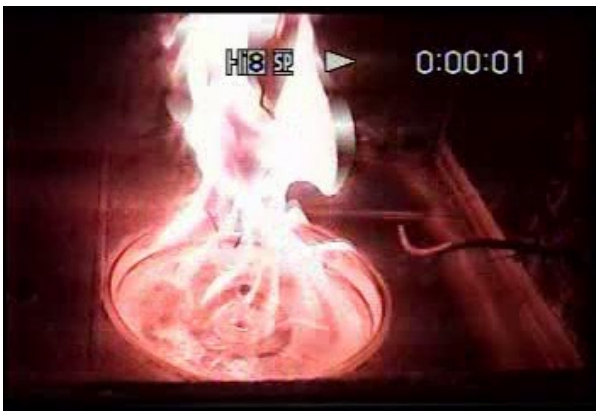


Introduction to Combustion

■ Basic features

- High-velocity fuel release: momentum-driven flame
- Low-velocity fuel release: buoyancy-driven flame
 - From thereon assume buoyancy-driven flame

$$Fr = \frac{u_{fuel}}{\sqrt{gL_{flame}}} \sim \frac{momentum}{buoyancy}$$



small fuel velocity/large diameter
low Froude number

large fuel velocity/small diameter
high Froude number





Introduction to Combustion

■ Basic features

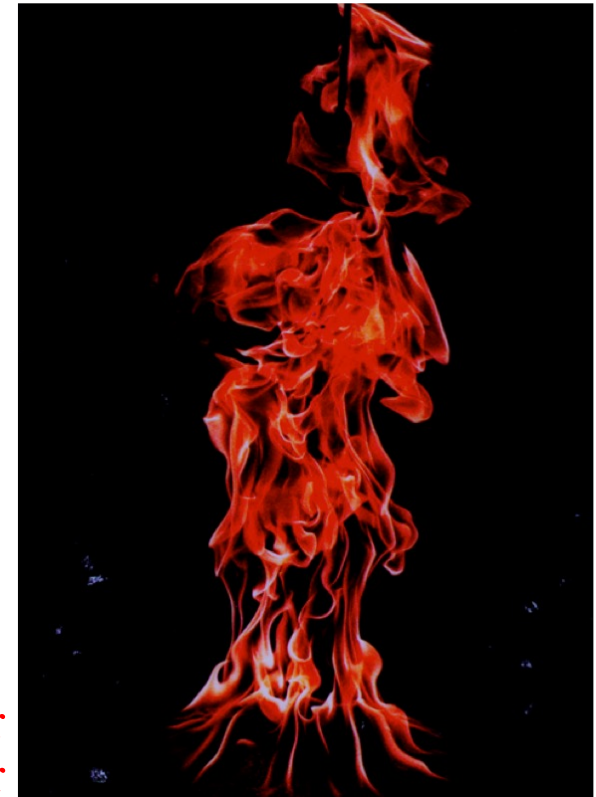
- Low-Reynolds-number flow: laminar flame
- High-Reynolds-number flow: turbulent flame
 - From thereon assume turbulent flame

$$Re = \frac{\sqrt{gL_{flame}} \times L_{flame}}{(\mu/\rho)} \sim \frac{convection}{diffusion}$$



small fuel velocity/small diameter
low Reynolds number

large fuel velocity/large diameter
large Reynolds number

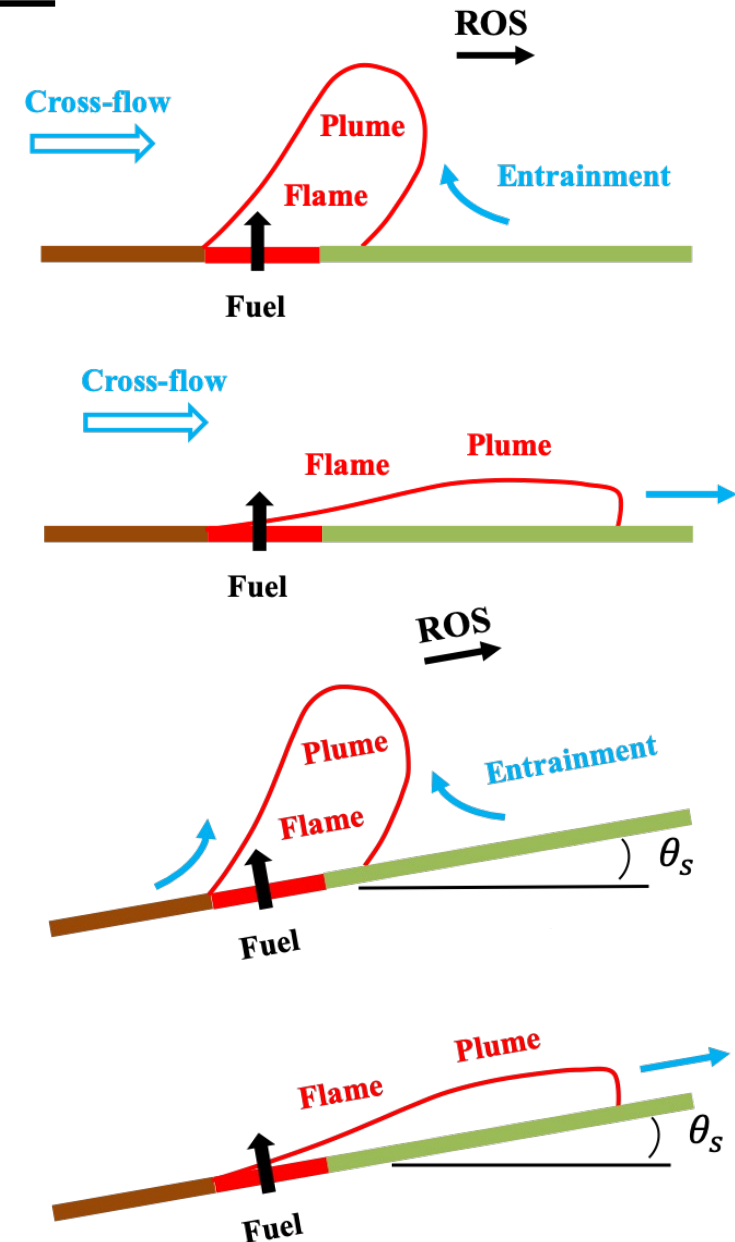




Introduction to Combustion

■ Basic features

- Flames are sensitive to their environment (wind, terrain topography)
 - Two limiting flame regimes: the *plume-dominated* regime in which the flame is mostly detached from the vegetation bed; and the *wind-driven* or *slope-driven* regime in which the flame is attached to the vegetation bed
 - From thereon assume flame under no wind and flat terrain conditions

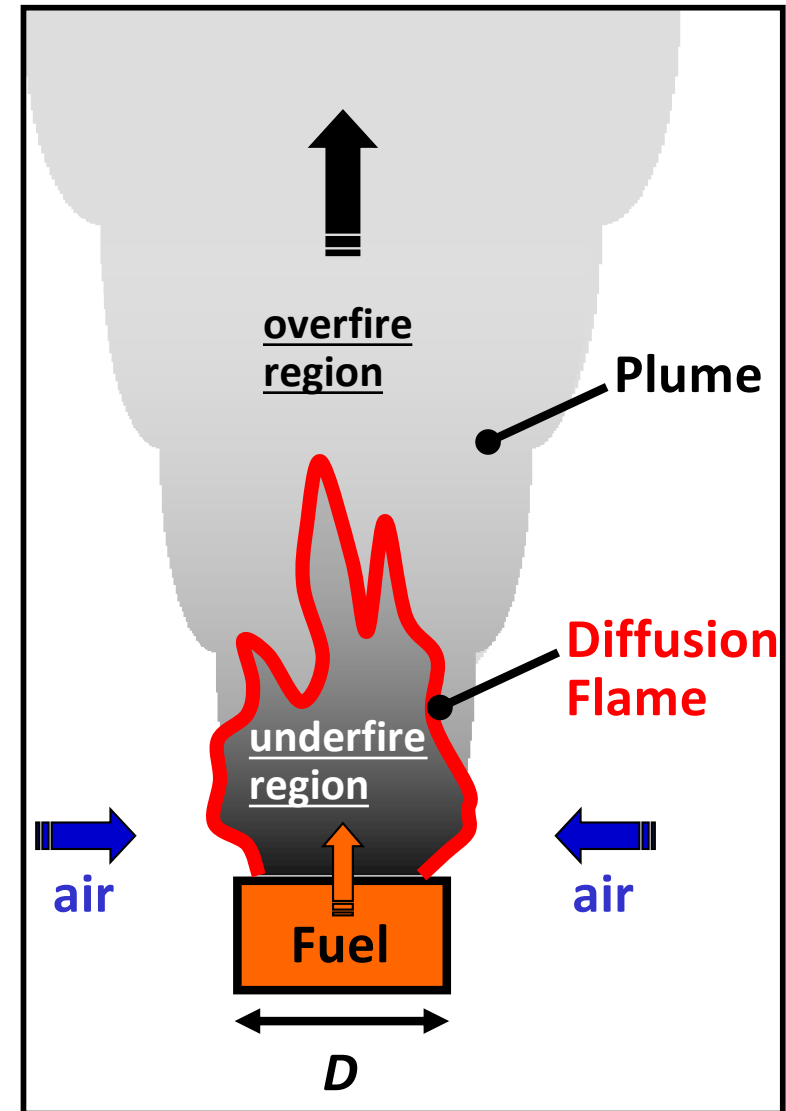




Flame Structure

■ Main features

- Fuel is flammable vapors produced by pyrolysis of vegetation biomass ($T_s \geq 500 - 600 \text{ K}$)
- Fuel reacts with ambient air; combustion occurs in flaming mode ($T_{flame} \sim 2000 \text{ K}$)
- Flame is non-premixed
- Fuel source velocity is small ($\sim 0.1-1 \text{ cm/s}$); buoyancy effects accelerate the flow up to several m/s
- Flow regime corresponds to moderate-to-high turbulence intensities





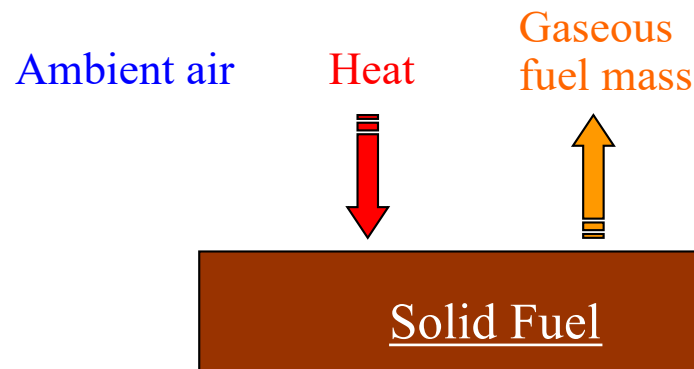
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- Combustion: the Thermodynamics Viewpoint
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Pyrolysis and Combustion

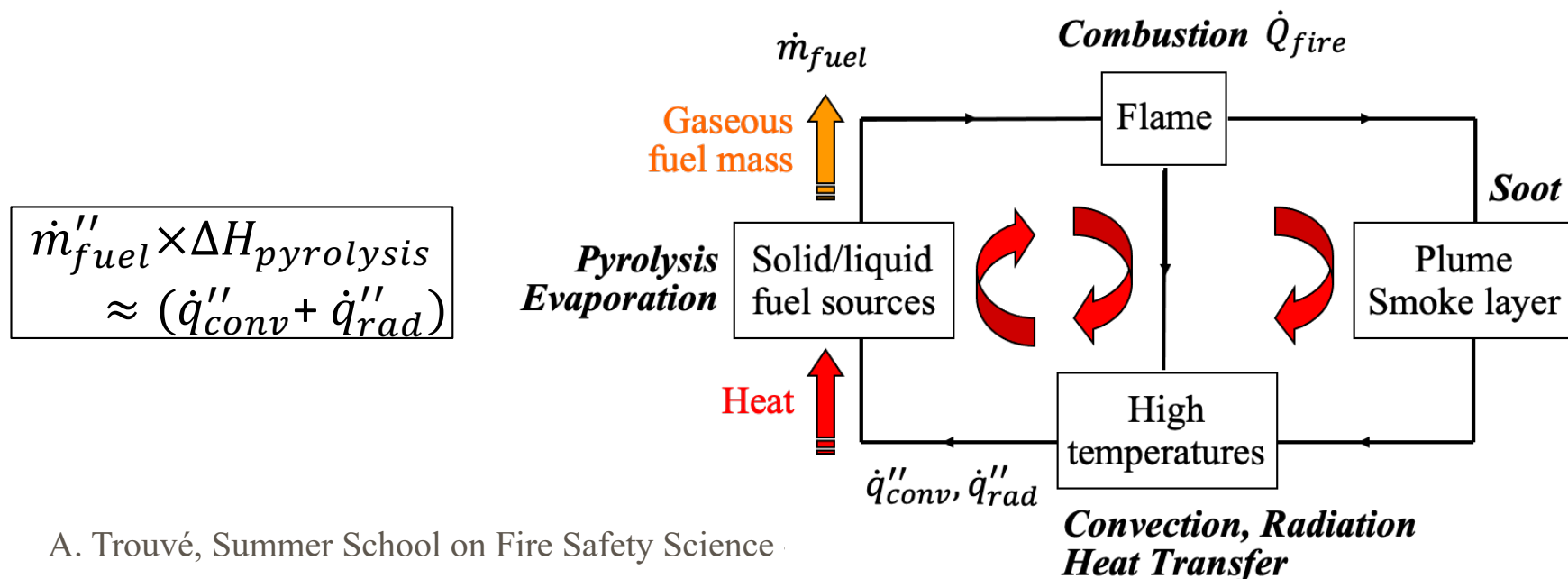
- Fire is an uncontrolled combustion process characterized by coupled solid-gas phase phenomena
 - Typical production of flammable vapors in a fire
 - Consider a flammable solid object/material that is a potential fuel source
 - At ambient temperature, the fuel is in solid form, the oxygen (from air) in gaseous form, and there is no combustion
 - At moderately elevated temperatures (~200-400 degrees Celsius), a complex thermal degradation process is initiated in the solid object/material, that corresponds to a phase change and produces fuel in gaseous form. This gasification process is called pyrolysis.





Pyrolysis and Combustion

- Fire is an uncontrolled combustion process characterized by a thermal feedback loop
 - Typical production of flammable vapors in a fire
 - Fuel gasification is an endothermic process and heat comes from the gas-to-solid heat transfer; the fuel gasification rate is controlled by the rate of gas-to-solid heat transfer (called the heat/thermal feedback)



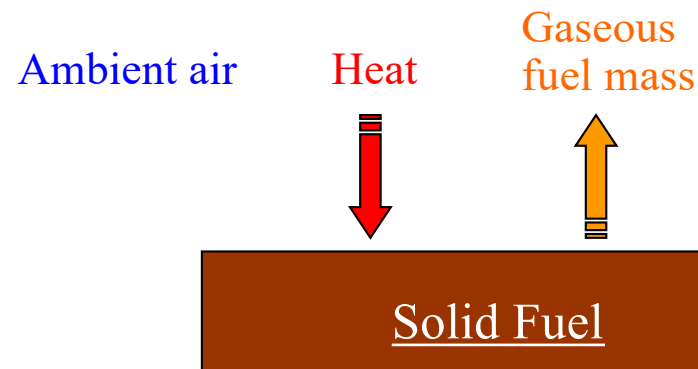


Pyrolysis and Combustion

- The fuel mass loss rate (MLR) [kg/s]
 - A property of the fuel source
 - Characterizes the intensity of the pyrolysis process

$$\dot{m}_{fuel}'' \approx \frac{(\dot{q}_{conv}'' + \dot{q}_{rad}'')}{\Delta H_{pyrolysis}}$$

$$\dot{m}_{fuel} = \iint_{A_{fuel}} \dot{m}_{fuel}'' dA_{fuel} \approx \frac{\iint_{A_{fuel}} (\dot{q}_{conv}'' + \dot{q}_{rad}'') dA_{fuel}}{\Delta H_{pyrolysis}}$$





Pyrolysis and Combustion

- The heat release rate (HRR) [W]
 - A property of the flame
 - Characterizes the intensity of the combustion process

$$\dot{Q}_{fire} = \dot{m}_{fuel} \times \Delta H_{combustion}$$





Pyrolysis

■ Thermally-driven decomposition of vegetation biomass

- Drying reaction

$(1 \text{ kg wet solid}) \rightarrow (\eta_{H_2O,Rd} \text{ kg water vapor}) + (\eta_{ds,Rd} \text{ kg dry solid})$

- Thermal/oxidative pyrolysis reactions (production of fuel)

$(1 \text{ kg dry solid}) \rightarrow (\eta_{f,Rp} \text{ kg fuel}) + (\eta_{c,Rp} \text{ kg char})$

$(1 \text{ kg dry solid}) + (\eta_{O_2,Rop} \text{ kg } O_2) \rightarrow (\eta_{f,Rop} \text{ kg fuel}) + (\eta_{c,Rop} \text{ kg char})$

- Char oxidation reaction

$(1 \text{ kg char}) + (\eta_{O_2,Rco} \text{ kg } O_2) \rightarrow (\eta_{p,Rco} \text{ kg products}) + (\eta_{a,Rco} \text{ kg ash})$





Pyrolysis

■ Thermally-driven decomposition of vegetation biomass

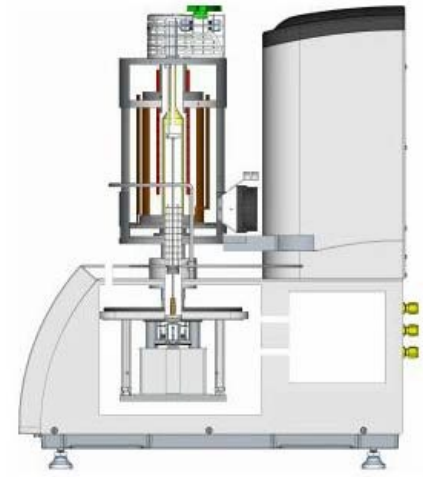
• Input data

Thermal properties of wet solid	$\rho_{s,ws}, k_{s,ws}, c_{s,ws}, \epsilon_{ws}$
Thermal properties of dry solid	$\rho_{s,ds}, k_{s,ds}, c_{s,ds}, \epsilon_{ds}$
Thermal properties of char	$\rho_{s,c}, k_{s,c}, c_{s,c}, \epsilon_c$
Thermal properties of ash	$\rho_{s,a}, k_{s,a}, c_{s,a}, \epsilon_a$
Porosity and permeability of wet solid	ψ_{ws}, K_{ws}
Porosity and permeability of dry solid	ψ_{ds}, K_{ds}
Porosity and permeability of char	ψ_c, K_c
Porosity and permeability of ash	ψ_a, K_a
Drying reaction	$A_{Rd}, E_{Rd}, n_{Rd}, \Delta H_{Rd}, \eta_{ds,Rd}$
Thermal pyrolysis reaction	$A_{Rp}, E_{Rp}, n_{Rp}, \Delta H_{Rp}, \eta_{c,Rp}$
Oxidative pyrolysis reaction	$A_{Rop}, E_{Rop}, n_{Rop}, n_{O_2,Rop}, \Delta H_{Rop}, \eta_{c,Rop}, \eta_{O_2,Rcp}$
Char oxidation reaction	$A_{Rco}, E_{Rco}, n_{Rco}, n_{O_2,Rco}, \Delta H_{Rco}, \eta_{a,Rco}, \eta_{O_2,Rco}$



Pyrolysis

- Basic study of phenomenology; evaluation of MLR and/or model input data
 - Micro-scale experiments (*e.g.* Thermogravimetric Analysis – TGA, Differential Scanning Calorimetry – DSC, Microscale Combustion Calorimetry – MCC)
 - Bench-scale experiments (*e.g.*, cone calorimeter, Fire Propagation Apparatus – FPA)
 - Fuel package experiments





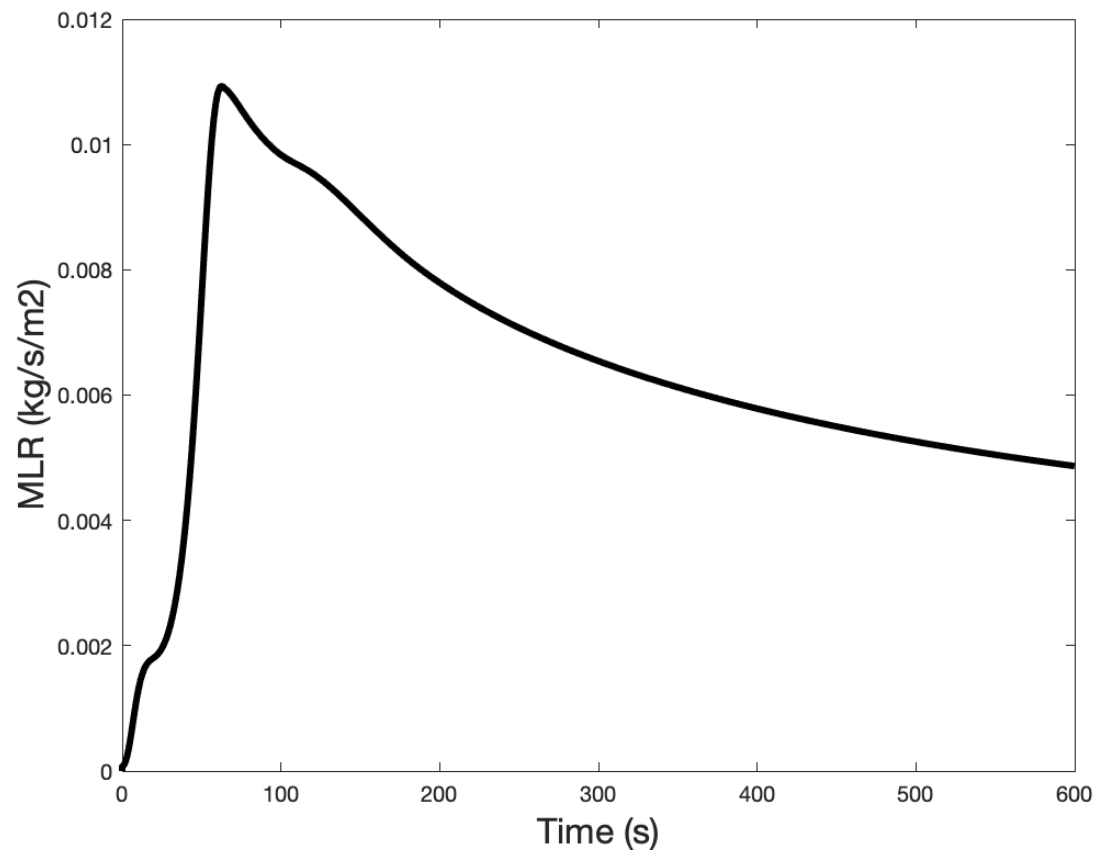
Pyrolysis

- Example of results from cone calorimeter test (model)
 - Example: white pine (Lautenberger & Fernandez-Pello, *Combust. Flame* 156:1503-1513 (2009))

$$\Delta = 3.8 \text{ cm}$$

$$G = 40 \text{ kW/m}^2$$

$$x_{O_2,g} = 0.21$$





Pyrolysis

■ Example of results from cone calorimeter test (model)

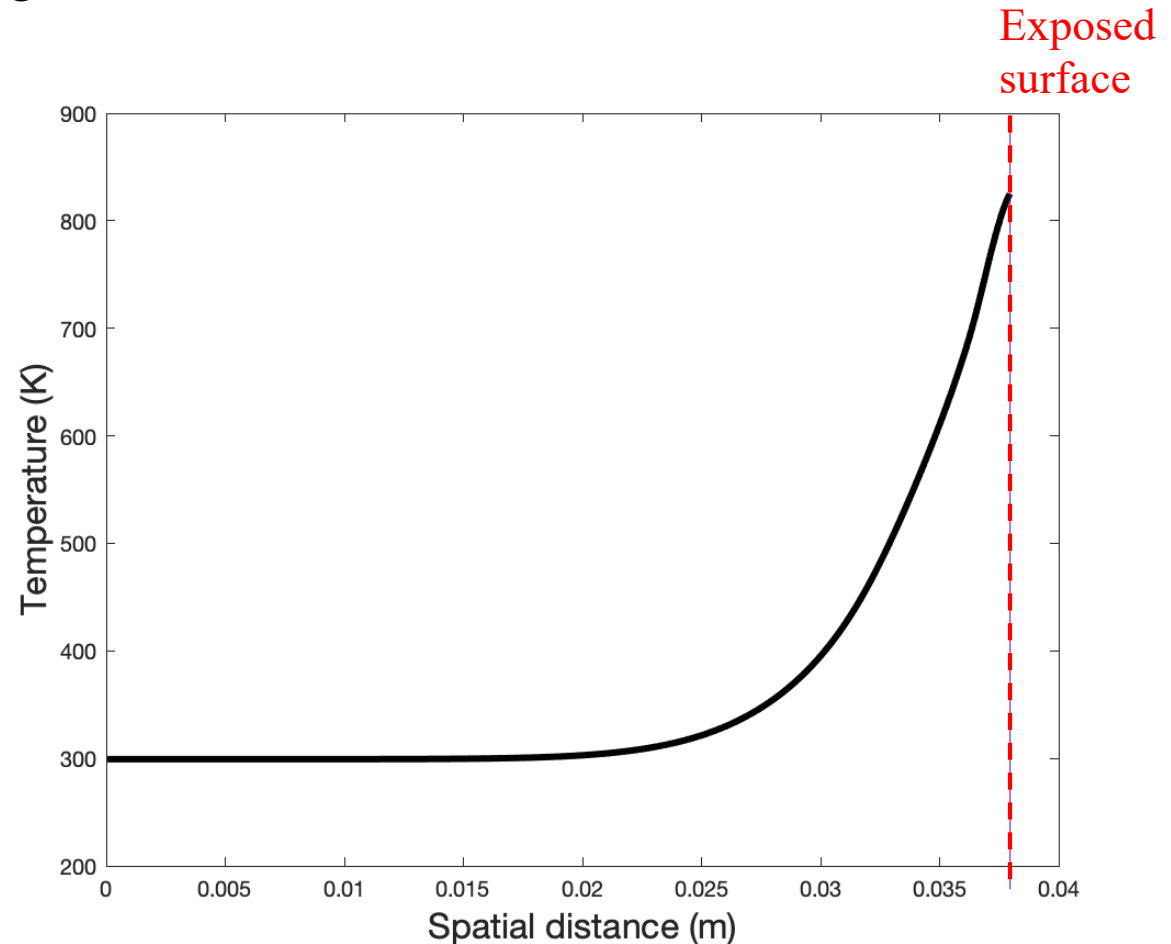
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$$x_{O_2,g} = 0.21$$

$$t = 100 \text{ s}$$





Pyrolysis

■ Example of results from cone calorimeter test (model)

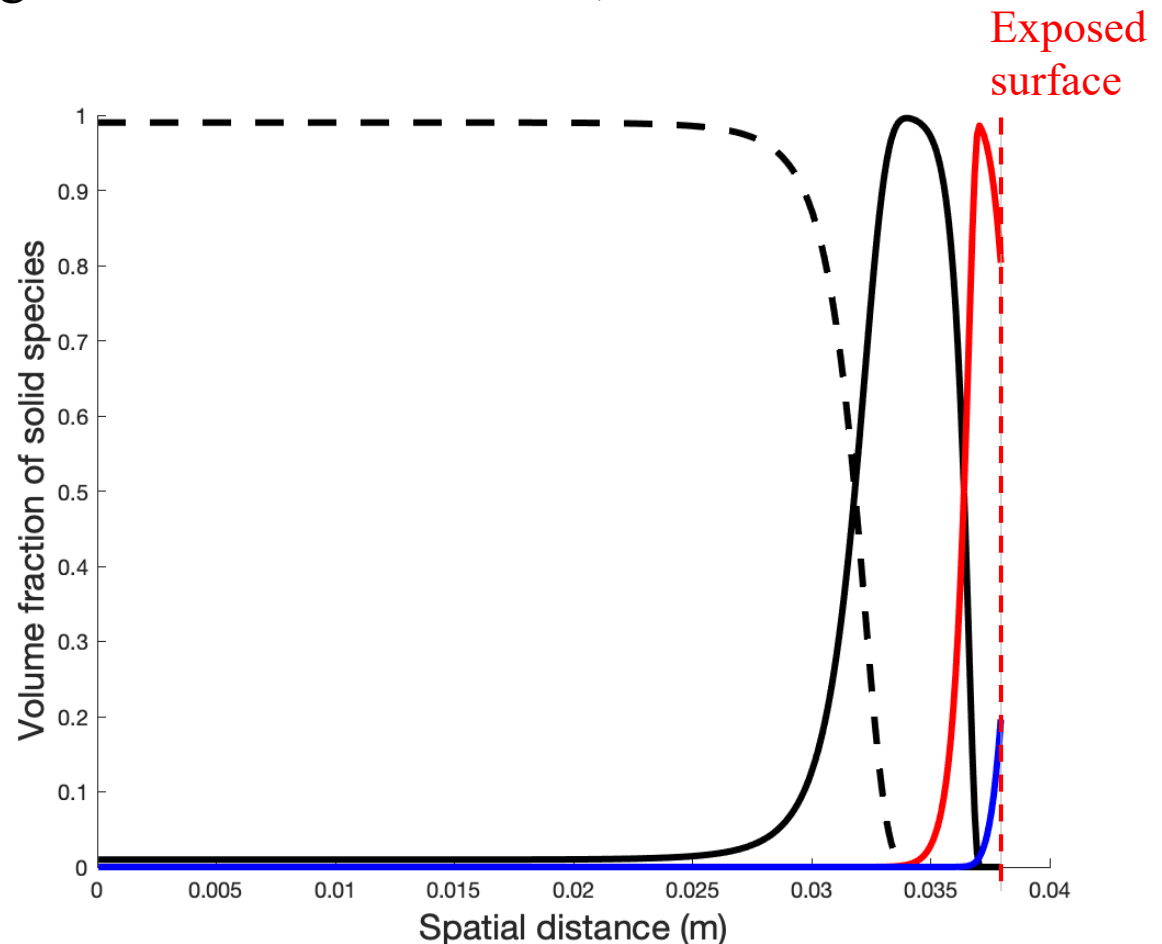
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Pyrolysis

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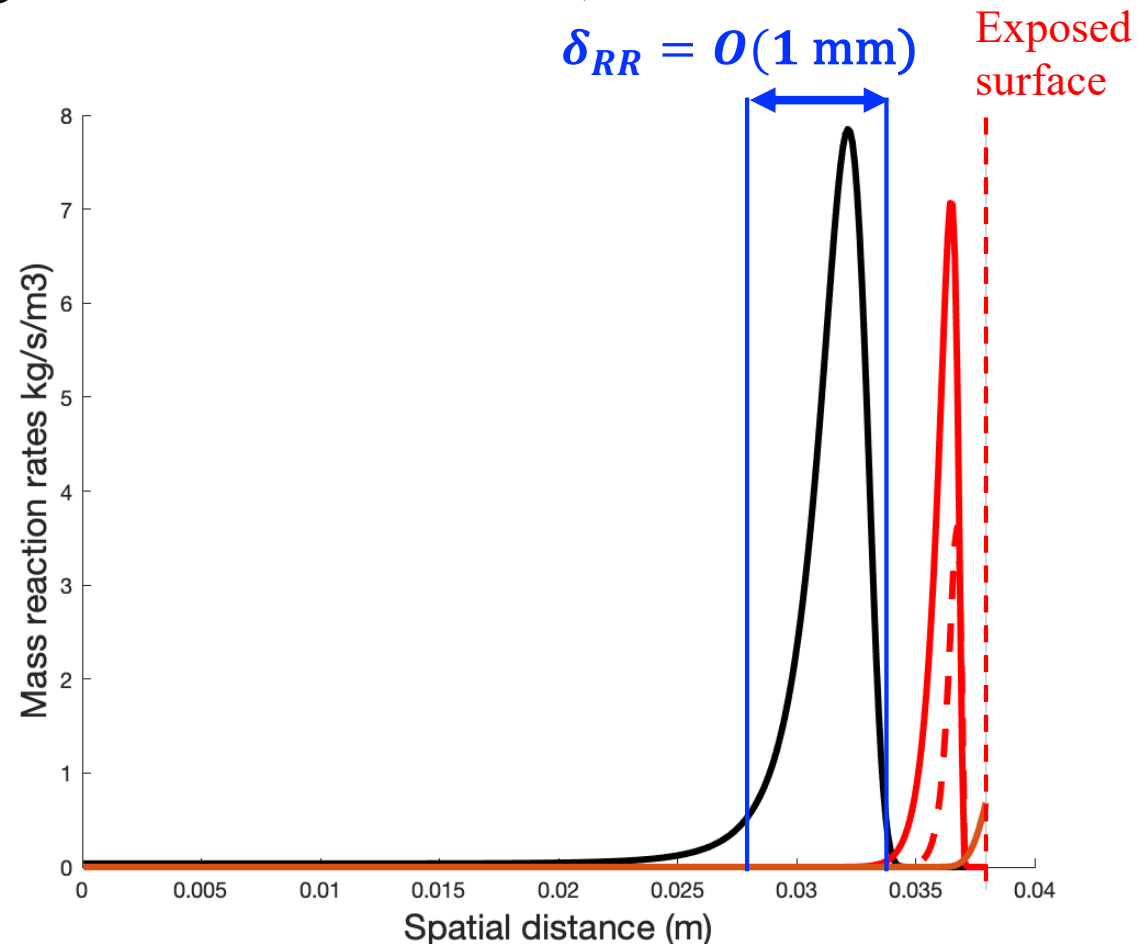
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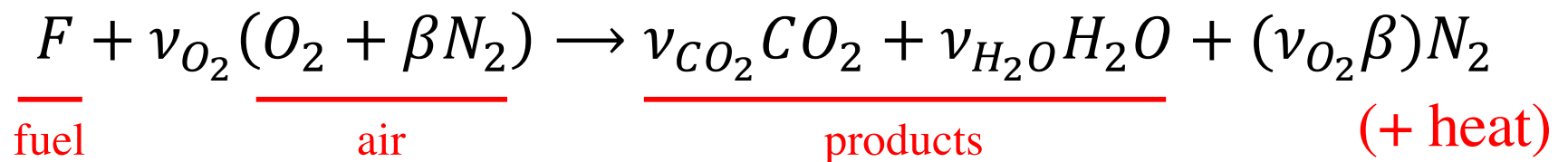
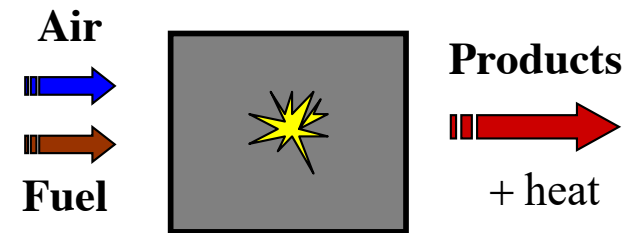
Combustion



■ Global combustion equation

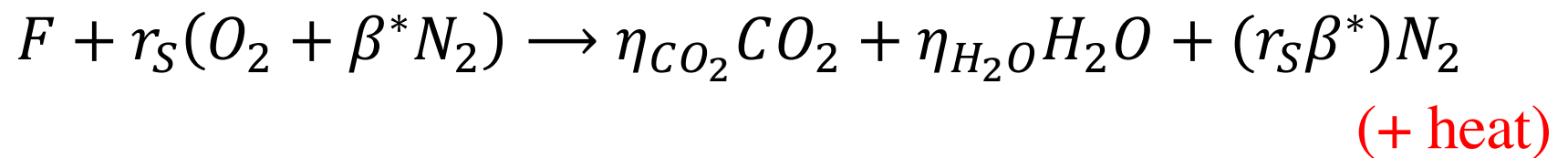
- Black box model

- written for 1 mole of fuel



where $\beta = (n_{N_2}/n_{O_2})_{air}$

- or written for 1 kg of fuel



where $\beta^* = ((n_{N_2}MW_{N_2})/(n_{O_2}MW_{O_2}))_{air}$

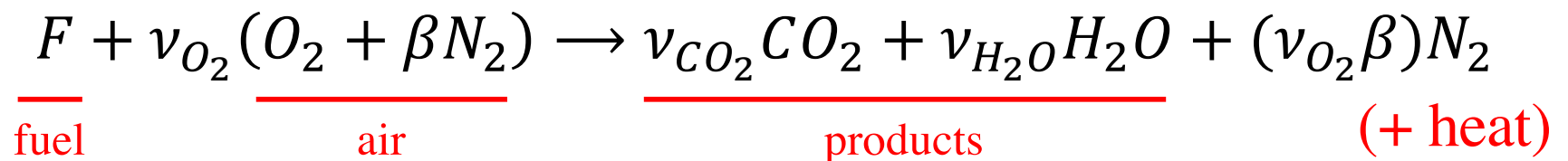
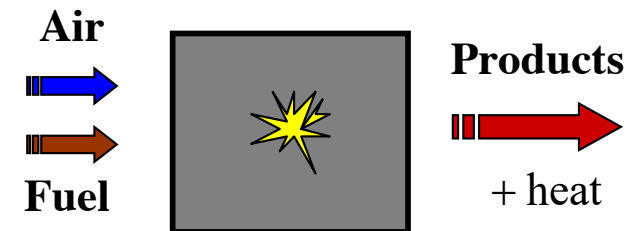


Combustion

■ Global combustion equation

- Assume $F = C_n H_m O_p$ (known)

➤ for 1 mole of fuel



➤ Conservation of C , H , O and N atoms

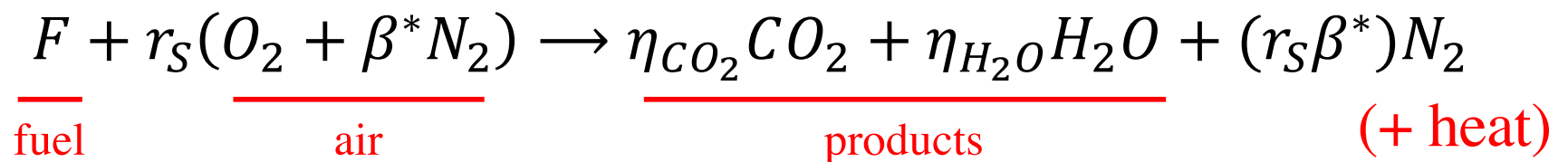
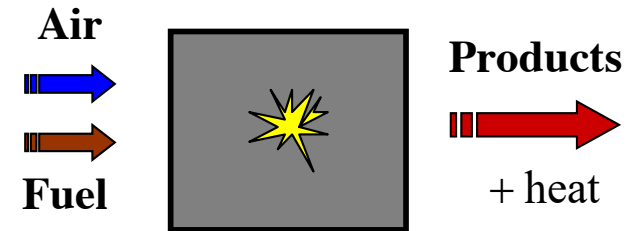
$$\nu_{CO_2} = n; \quad \nu_{H_2O} = \frac{m}{2}; \quad \nu_{O_2} = n + \frac{m}{4} - \frac{p}{2}; \quad \nu_{N_2} = \nu_{O_2} \times \beta$$



Combustion

■ Global combustion equation

- Assume $F = C_n H_m O_p$ (known)
 - for 1 kg of fuel



- Conservation of C , H , O and N atoms

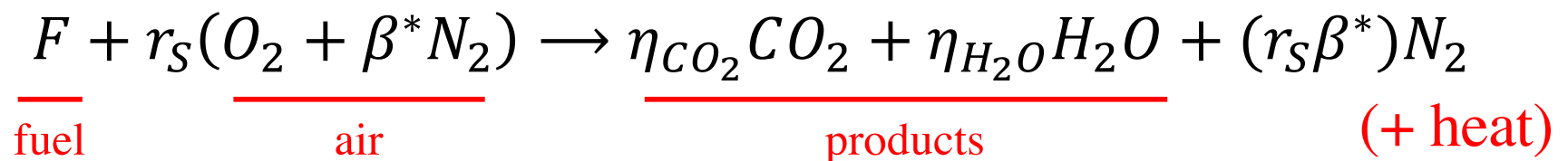
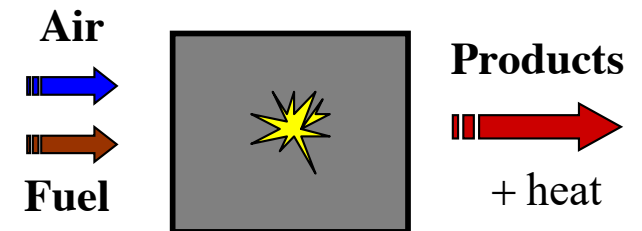
$$\eta_{CO_2} = \frac{\nu_{CO_2} MW_{CO_2}}{MW_F} ; \quad \eta_{H_2O} = \frac{\nu_{H_2O} MW_{H_2O}}{MW_F} ; \quad r_S = \frac{\nu_{O_2} MW_{O_2}}{MW_F}$$



Combustion

■ Global combustion equation

- Assume $F = C_n H_m O_p$ (known)
 - for 1 kg of fuel



- Heat of combustion (per unit mass of fuel)

$$\Delta H_F = r_S \times \Delta H_{O_2}$$

$$\Delta H_{O_2} \approx 13.1 \text{ MJ/kg}[O_2]$$



Combustion

■ Global combustion equation

- *Stoichiometric* conditions

- The proportions of fuel and oxygen are those required by the combustion equation
- All the fuel and oxygen mass are consumed by the combustion process
- Corresponds to a chemical optimum: maximum emissions of carbon dioxide/water vapor as well as maximum heat release (maximum flame temperature)



Combustion

■ Idealized combustion equation

- *Non-stoichiometric* conditions

$$\phi = \frac{(n_F^i/n_{O_2}^i)}{(1/\nu_{O_2})} = \nu_{O_2} \left(\frac{x_F^i}{x_{O_2}^i} \right)$$

$$\phi = \frac{(m_F^i/m_{O_2}^i)}{(MW_F/(\nu_{O_2} MW_{O_2}))} = r_S \left(\frac{Y_F^i}{Y_{O_2}^i} \right)$$

Equivalence ratio
(control volume analysis)



Combustion

■ Idealized combustion equation

- *Fuel lean* (fuel-limited, over-ventilated) conditions

$$\phi \leq 1$$

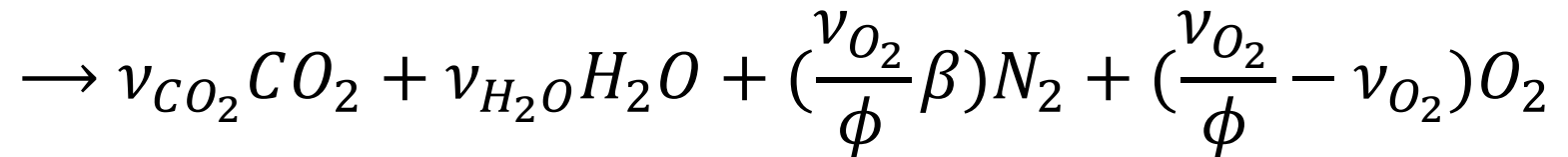
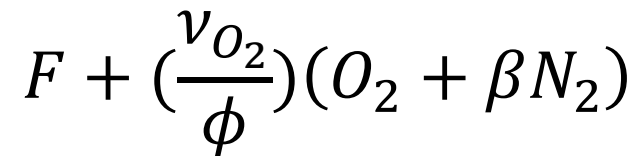
- There is more oxygen than required by the combustion chemistry
- All the fuel mass is consumed by the combustion process; there is some excess oxygen mass in the products



Combustion

■ Idealized combustion equation

- *Fuel lean* (fuel-limited, over-ventilated) conditions



excess oxygen



Combustion

■ Idealized combustion equation

- *Fuel rich* (oxygen-limited, under-ventilated) conditions

$$\phi \geq 1$$

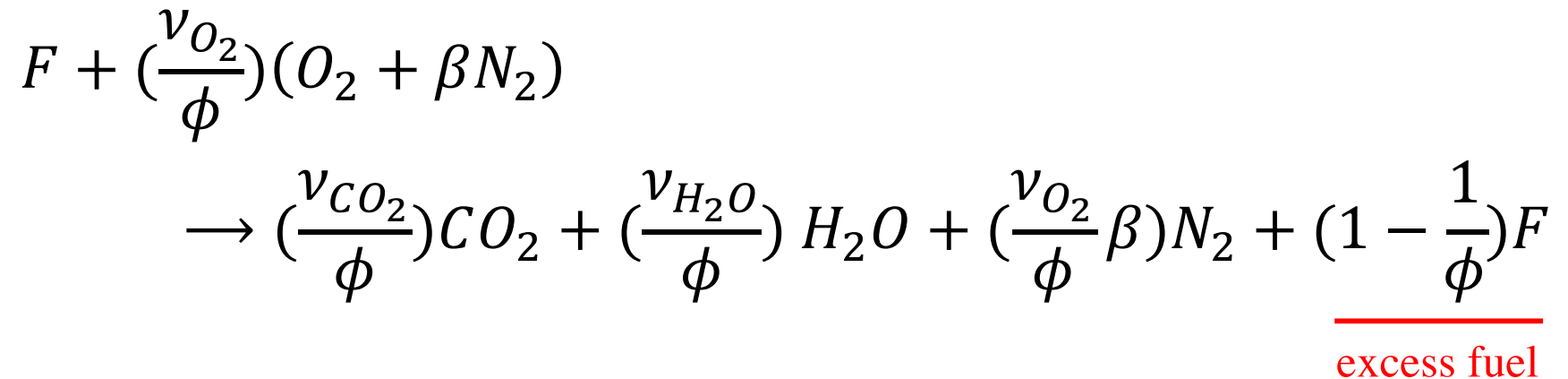
- There is less oxygen than required by the combustion chemistry
- All the oxygen mass is consumed by the combustion process; there is some excess fuel mass in the products



Combustion

■ Idealized combustion equation

- *Fuel rich* (oxygen-limited, under-ventilated) conditions





Combustion

■ Idealized combustion equation

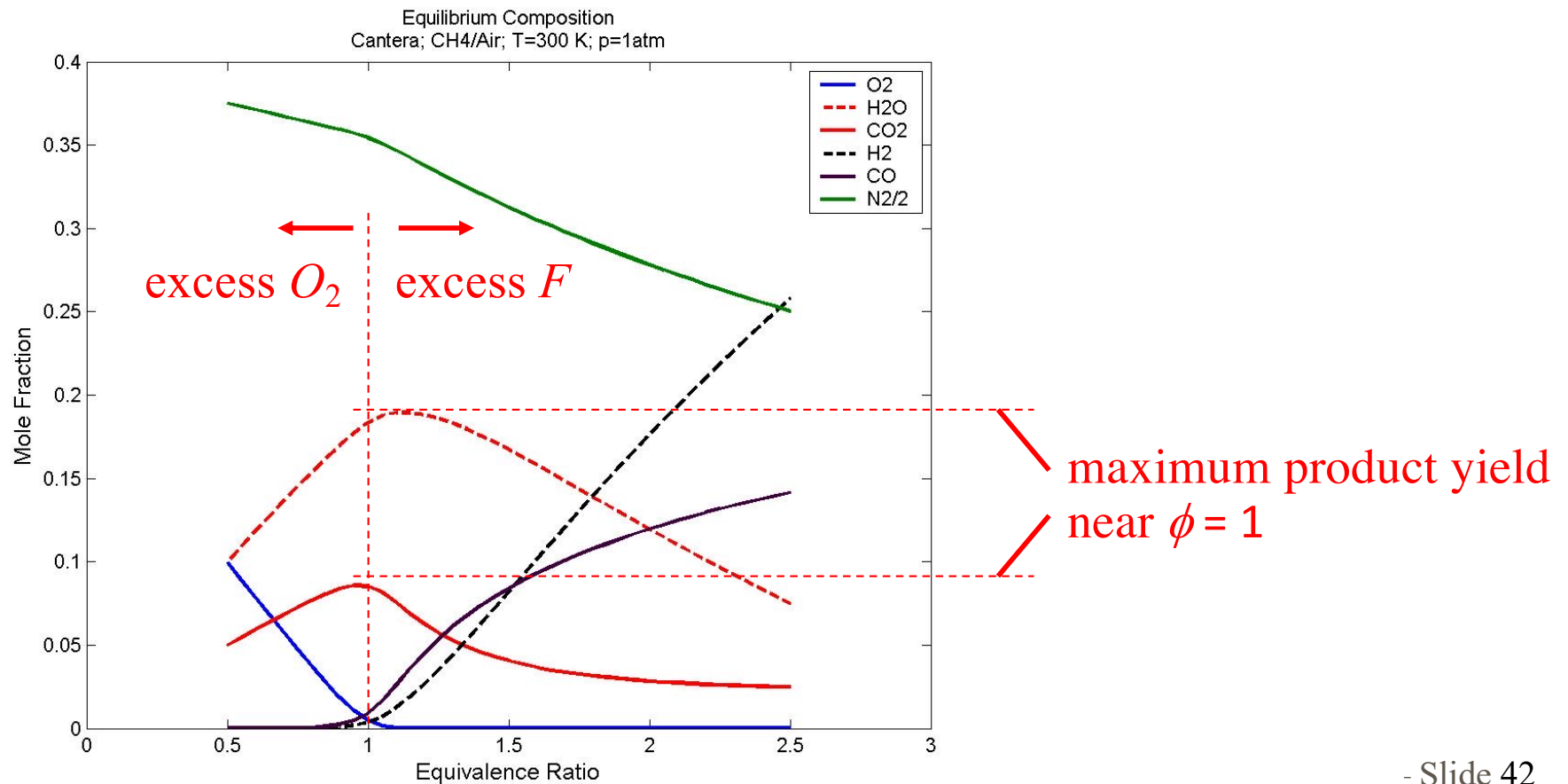
• Limitations

- Under high temperature conditions, CO_2 and H_2O experience chemical *dissociations* (*i.e.* breakdown of large molecules into smaller molecules and atoms); combustion products include CO_2 and H_2O , as assumed so far, but also additional minor species (CO , H_2 , OH , H , O , *etc*)
- Excess fuel mass present in fuel-rich combustion does not remain as virgin fuel, as assumed so far, but decomposes into CO , H_2 , and also other hydrocarbon species (uHC), and soot



Combustion

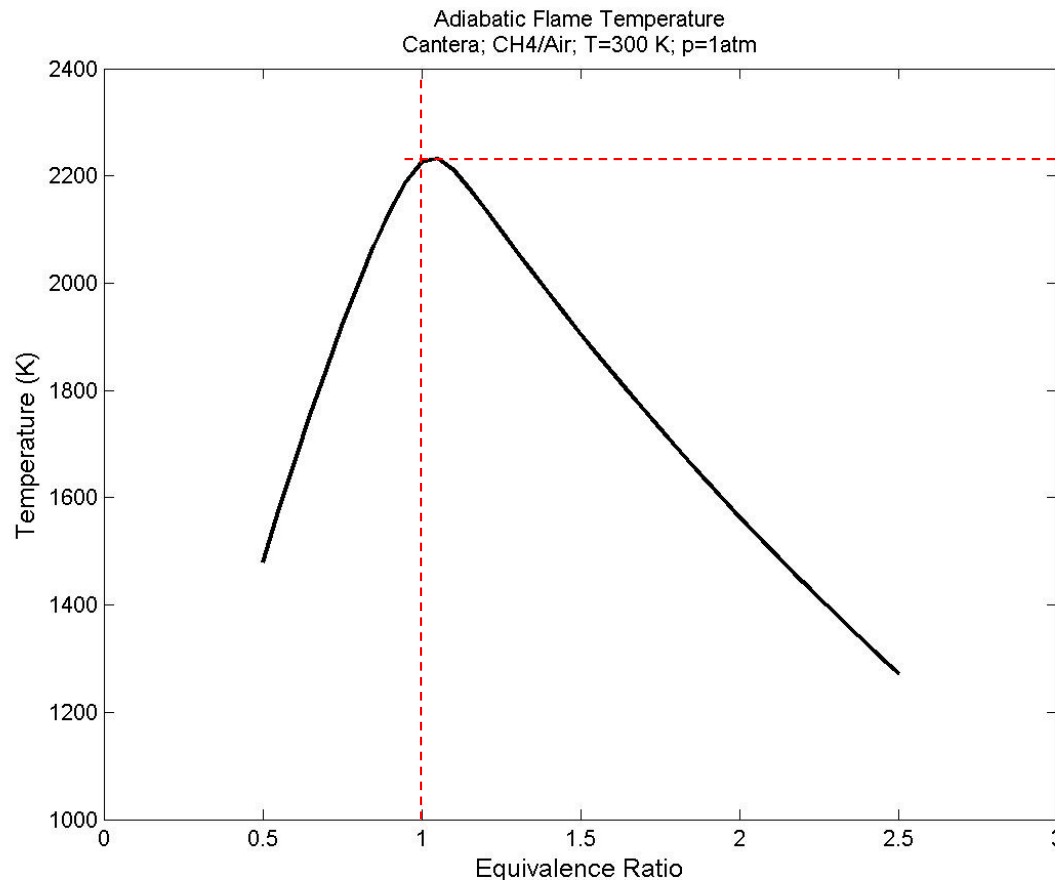
- Calculation of equilibrium mixture composition with state-of-the-art software (CHEMKIN, Cantera, *etc*)





Combustion

- Calculation of adiabatic flame temperature with state-of-the-art software (CHEMKIN, Cantera, *etc*): measure of heat released by combustion process



maximum temperature
near $\phi = 1$
maximum temperature
 $\sim 2,200\text{--}2,300$ K
for most practical fuels
(STPC)



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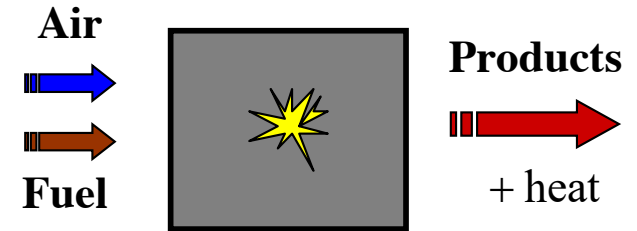
Combustion



■ Global combustion equation

- A black-box model

- Describes the inputs and outputs of the combustion transformation
- Also a statement on global mass (and energy) conservation
- But not a statement on the speed of combustion
 - The speed of combustion is described in studies of combustion chemical kinetics and/or flame studies





Chemistry

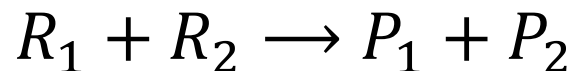
- Detailed chemical kinetic mechanism
 - A “first-principles” description of combustion chemistry (*elementary reactions*)
 - A series of statements about probable outcome of collisions between molecules at the quantum level
 - Based on the kinetic theory of gases (statistical thermodynamics)
 - Provides estimates of the reaction rates (RR) for each elementary reaction (*i.e.*, estimates of the speed of combustion)



Chemistry

■ Detailed chemical kinetic mechanism

- Reaction rate (RR) of an elementary bimolecular reaction



➤ Kinetic theory of gases

$$-\frac{dC_{R_1}}{dt} \sim \left\{ \begin{array}{l} \text{frequency of collisions} \\ \text{between } R_1 \text{ and } R_2 \end{array} \right\} \times \left\{ \begin{array}{l} \text{probability of a} \\ \text{successful collision} \end{array} \right\}$$

$$\sim C_{R_1} C_{R_2}$$

$$\sim \exp(-E_A/RT)$$

activation energy

$$-\frac{dC_{R_1}}{dt} \sim C_{R_1} C_{R_2} \times k(T)$$

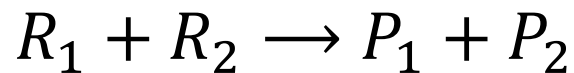
RR coefficient



Chemistry

■ Detailed chemical kinetic mechanism

- Reaction rate (RR) of an elementary bimolecular reaction



➤ *Arrhenius* model

$$k(T) = AT^b \exp\left(-\frac{E_A}{RT}\right)$$

temperature exponent

activation energy [J/mol]

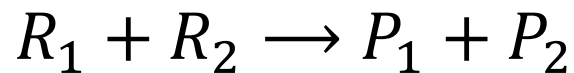
pre-exponential factor



Chemistry

■ Detailed chemical kinetic mechanism

- Reaction rate (RR) of an elementary bimolecular reaction



- *Arrhenius* model: RR varies exponentially with temperature

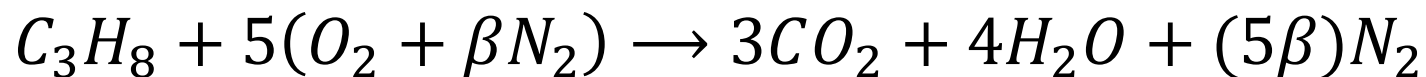
$$-\frac{dC_{R_1}}{dt} = C_{R_1} C_{R_2} \times AT^b \exp\left(-\frac{E_A}{RT}\right)$$

$$\text{where } C_k = \left(x_k \frac{p}{RT}\right) = \left(Y_k \frac{\rho}{MW_k}\right)$$



Chemistry

- Arrhenius model applied to global combustion equation (example: propane-air combustion)



- Reaction rate [mol/cm³/s] (C.K. Westbrook, F.L. Dryer, *Combustion Science and Technology* 1981)

$$-\frac{dC_{C_3H_8}}{dt} = (C_{C_3H_8})^{0.1} (C_{O_2})^{1.65} \times 8.6 \times 10^{11} \exp\left(-\frac{15,098}{T}\right)$$

- Heat release rate [W/m³]

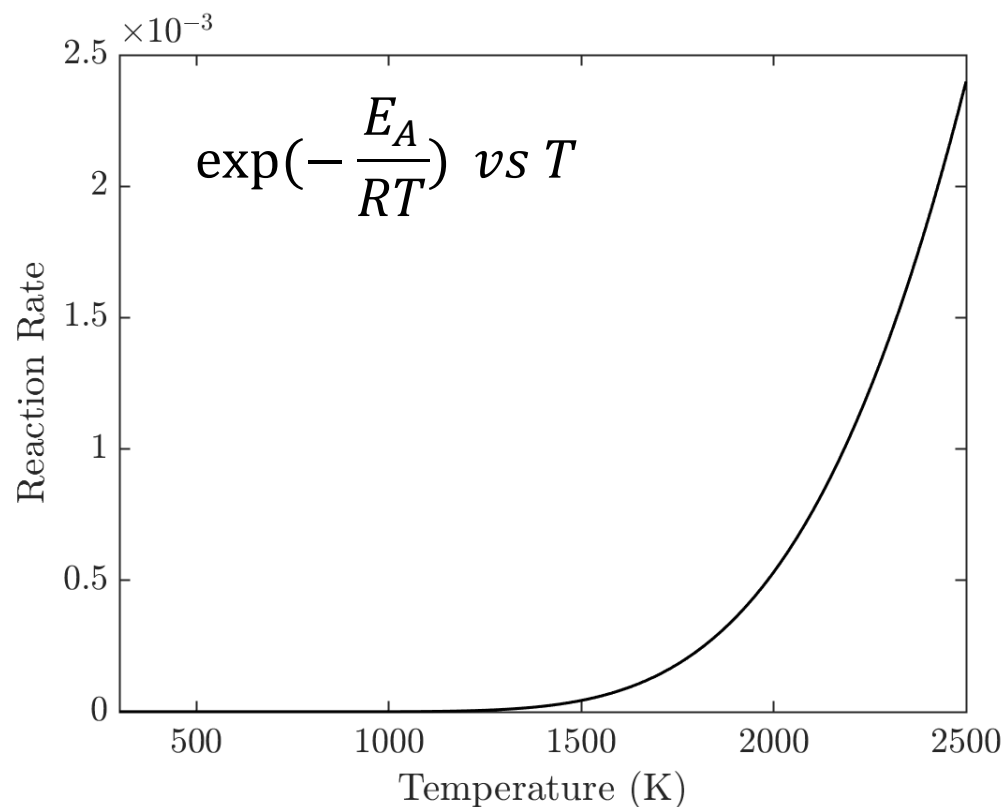
$$\dot{q}''''_{combustion} = \left(-\frac{dC_{C_3H_8}}{dt}\right) \times MW_{C_3H_8} \times 10^6 \times \Delta H_{C_3H_8}$$

Chemistry



■ Arrhenius model

- High temperatures correspond to (exponentially) fast chemistry

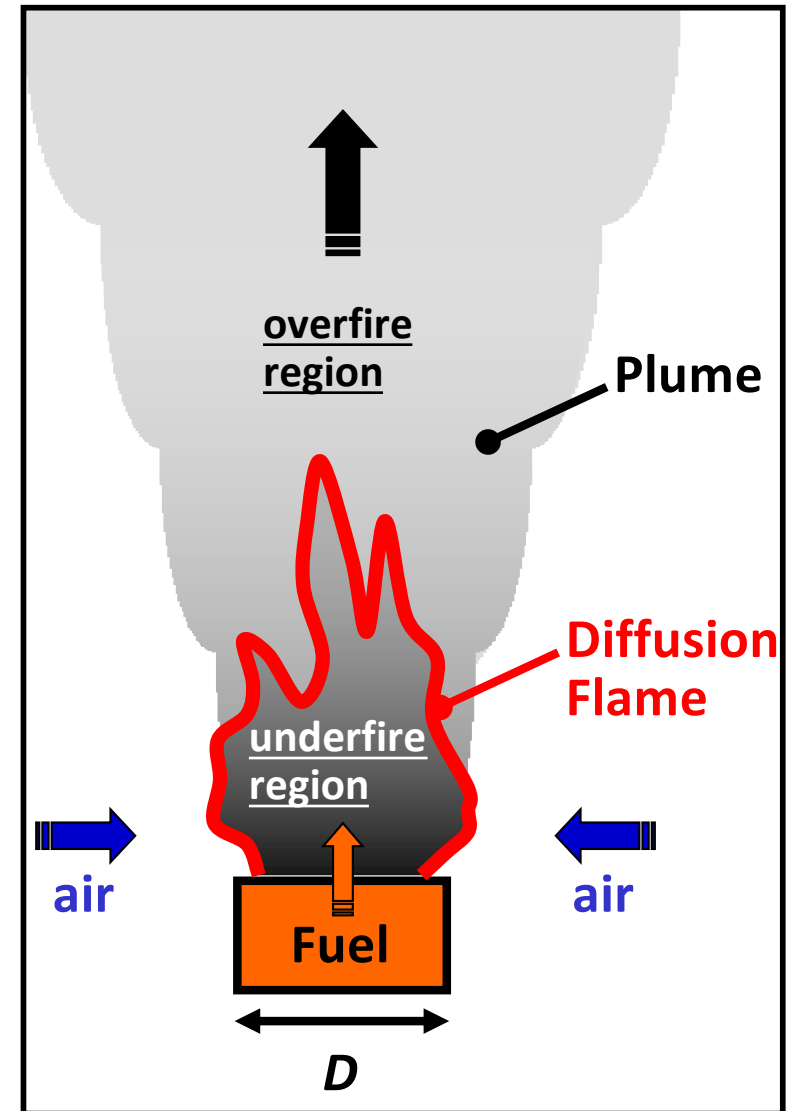




Flame Structure

■ Main features

- Flame topology corresponds to a thin surface (sheet)
- Burning rate limited by diffusion of fuel and air into the flame surface (combustion chemistry is *usually* very fast)
- The flame structure is characterized in terms of a mixing variable called the *mixture fraction*





Outline

- Introduction
- Pyrolysis and Combustion
- Combustion: the Thermodynamics Viewpoint
- Combustion Chemistry
- **Flame Structure**
- Flame Effects



Flame Structure

■ Mixture fraction

- Non-dimensional variable Z used to describe the mixture composition
- Field variable, $Z(x, y, z, t)$, defined as the local fraction of mass that originates from the fuel supply stream (convention: $Z = 1$ in pure fuel; $Z = 0$ in pure air)

$$Z = \frac{r_S Y_F - Y_{O_2} + Y_{O_2,air}}{r_S + Y_{O_2,air}}$$



Flame Structure

■ Mixture fraction

- Assuming combustion chemistry to be infinitely fast
 - The flame topology corresponds to a surface (*i.e.* a front or a sheet)
 - The flame is located where fuel and oxygen mass meet in stoichiometric proportions, at $Z = Z_{st}$

$$Z_{st} = \frac{Y_{O_2,air}}{r_s + Y_{O_2,air}}$$

- Fuel and oxygen mass do not coexist ($Y_F \times Y_{O_2} = 0$)

$Y_F = Y_{O_2} = 0 (Z = Z_{st})$
Flame surface

$Y_F = 0 (Z < Z_{st})$
Air side

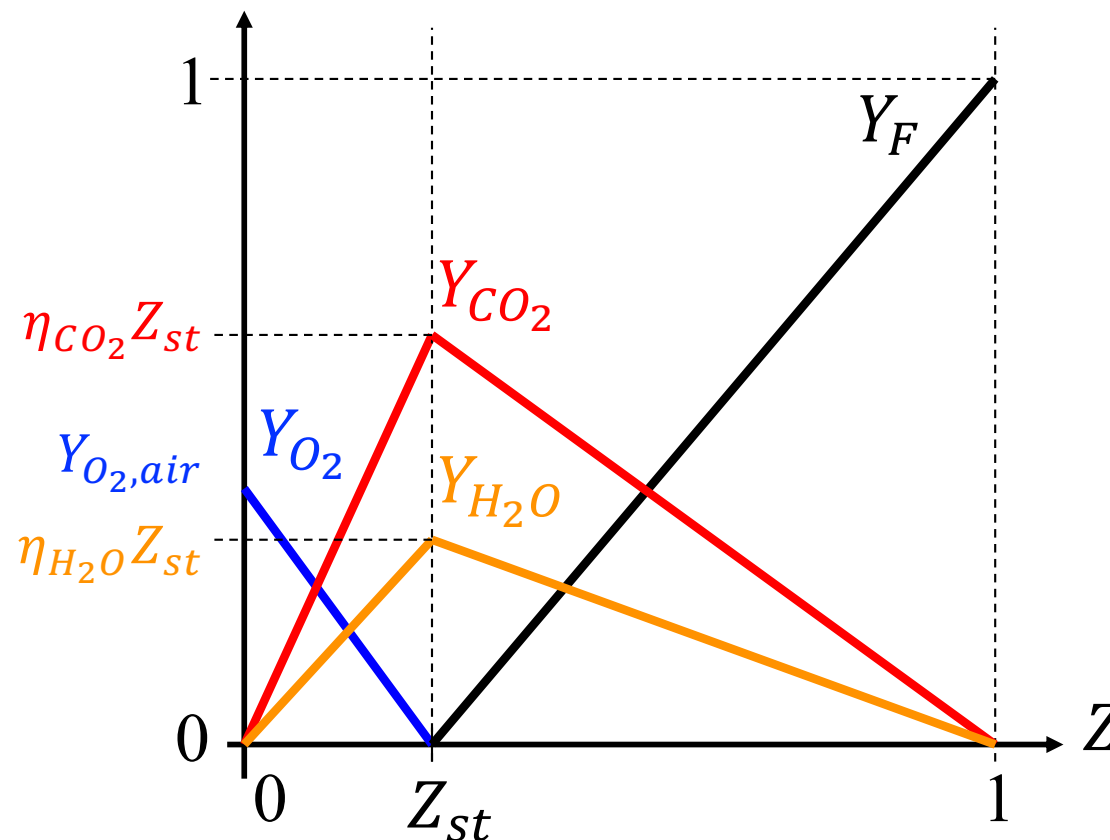
$Y_{O_2} = 0 (Z > Z_{st})$
Fuel side



Flame Structure

■ Mixture fraction

- Description of mixture composition in terms of the mixture fraction (Burke-Schumann model)

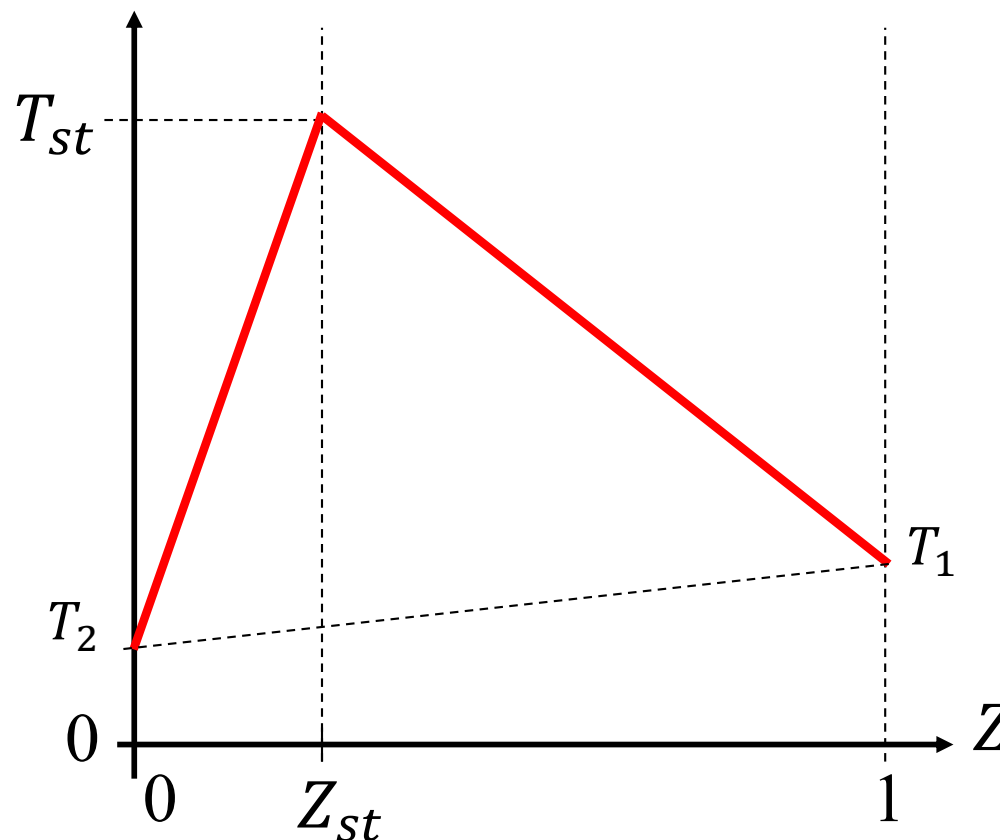




Flame Structure

■ Mixture fraction

- Description of temperature in terms of the mixture fraction (Burke-Schumann model with assumption of adiabatic combustion)

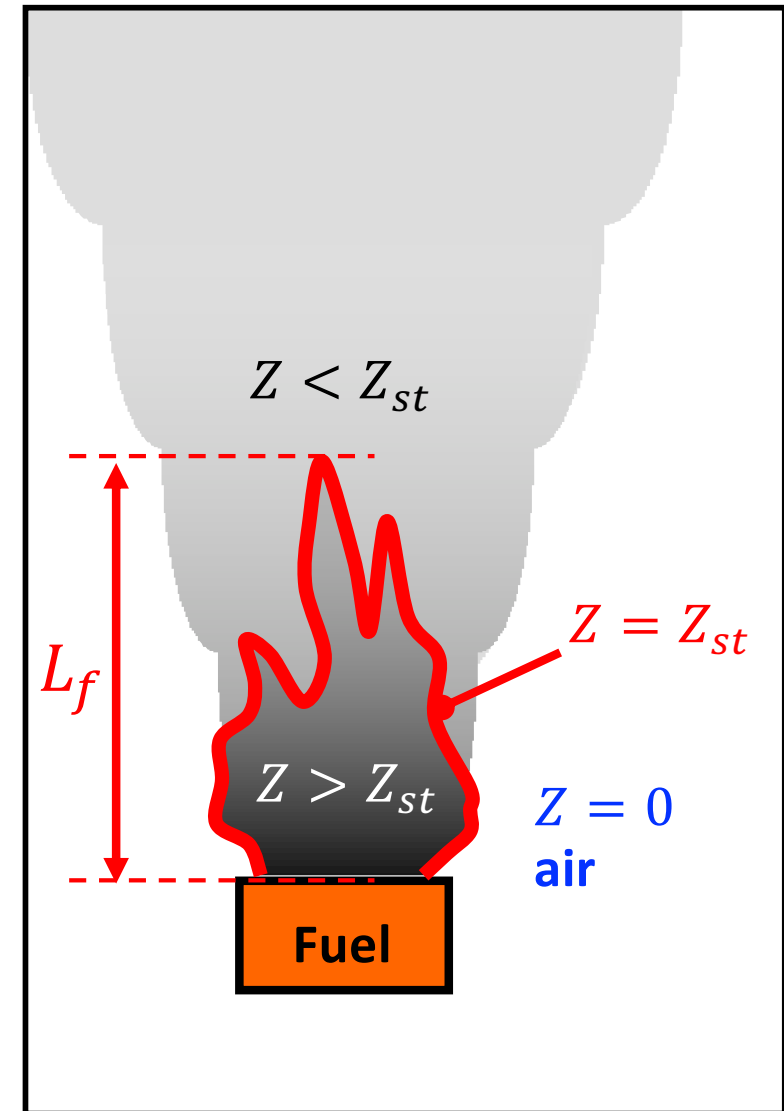




Flame Structure

■ Mixture fraction

- Field variable, $Z(x, y, z, t)$, solution of an unsteady convection-diffusion equation
- Characteristic length scale of the flame geometry: the mean flame height L_f
- Characteristic time scale of the flame: the mean flame residence time $\sim (L_f / \sqrt{gL_f}) = \sqrt{L_f/g}$





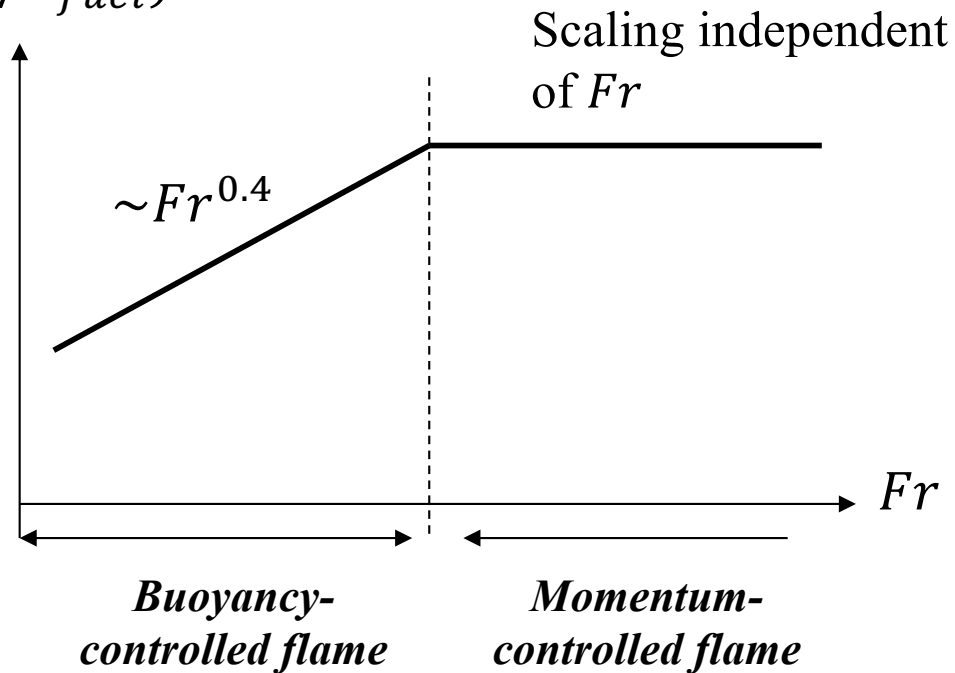
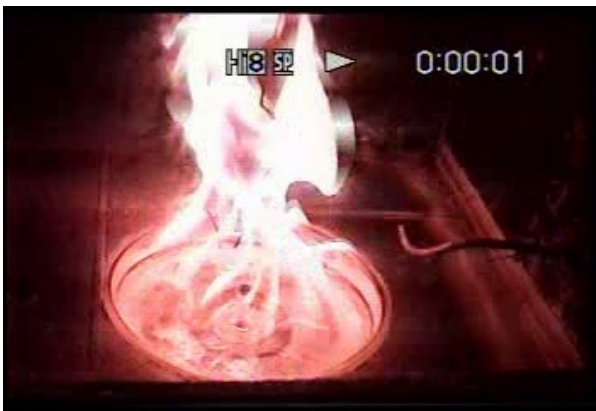
Flame Structure

- Mean flame height (pool fire configuration)

Non-dimensional
flame length (L_f/D_{fuel})

$$Fr = \frac{u_{fuel}}{\sqrt{gD_{fuel}}} \sim \frac{\text{momentum}}{\text{buoyancy}}$$

low Froude number



high Froude number





Flame Structure

- Mean flame height (pool fire configuration)
 - For fire applications (buoyancy-controlled flames, small values of Fr), flame length correlations based on the following Q -star parameter

$$\dot{Q}^* = \frac{\dot{Q}_{fire}}{(\rho c_p T)_{air} \sqrt{g D_{fuel}} D_{fuel}^2}$$



Flame Structure

■ Mean flame height (pool fire config.)

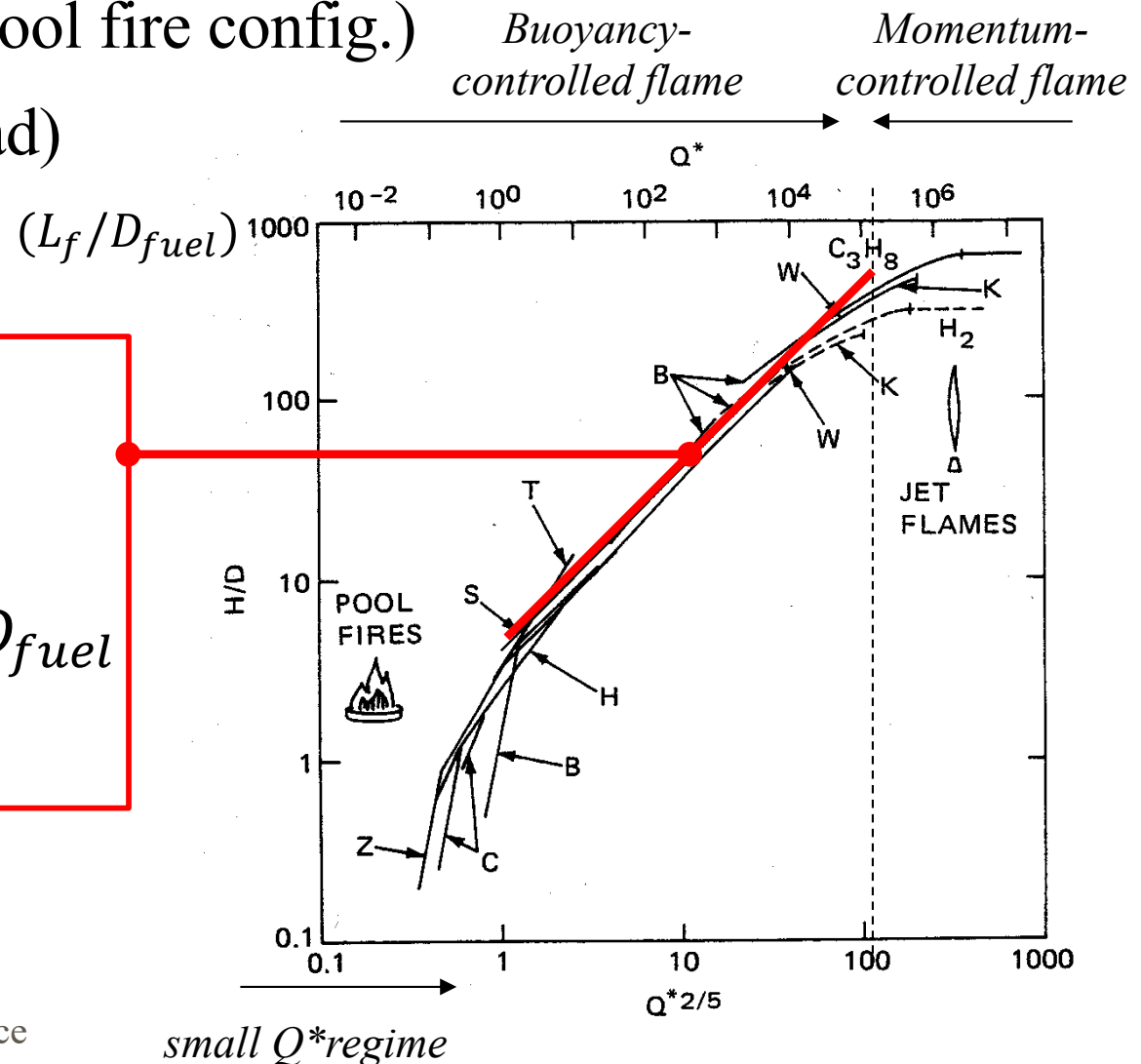
- Correlation (Heskestad)

$$(1 \leq \dot{Q}^* < 10^5)$$

$$\left(\frac{L_f}{D_{fuel}}\right) = 3.7 \times \dot{Q}^{*2/5} - 1.02$$

$$L_f = 0.235 \times \dot{Q}_{fire}^{2/5} - 1.02 \times D_{fuel}$$

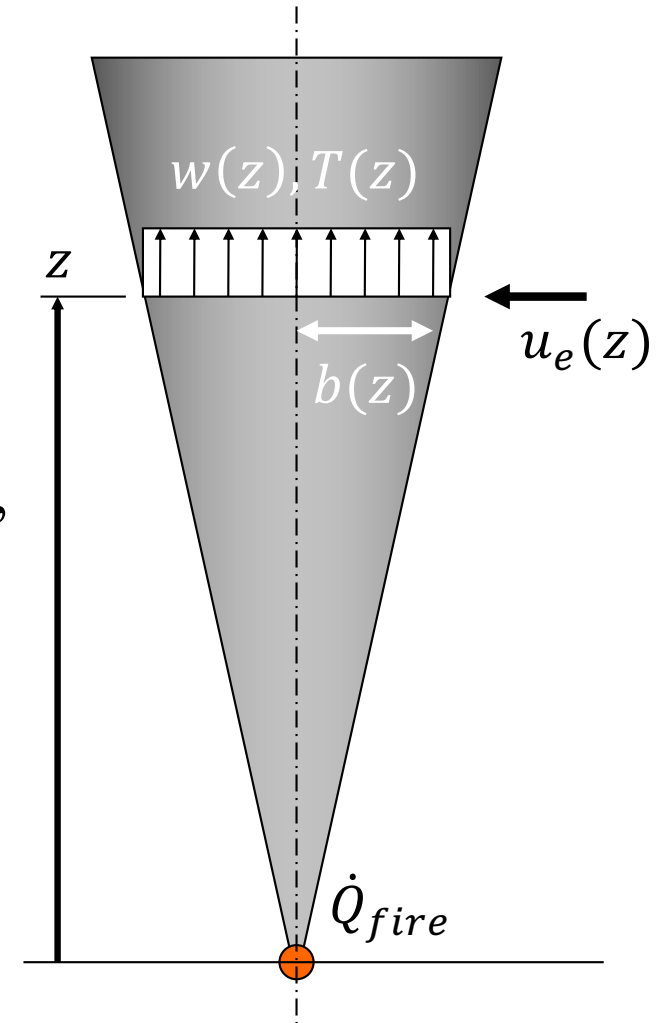
(\dot{Q}_{fire} in kW)





Plume Structure

- Ideal buoyant plume (pool fire config.)
 - Axisymmetric geometry
 - Point source release of heat (far field plume theory)
 - “Fire induced wind”: $u_e(z) = \alpha \times w(z)$, empirical entrainment factor α ($\alpha \sim 0.1-0.2$)





Plume Structure

- Ideal buoyant plume (pool fire config.)

$$b(z) = C_b \times z$$

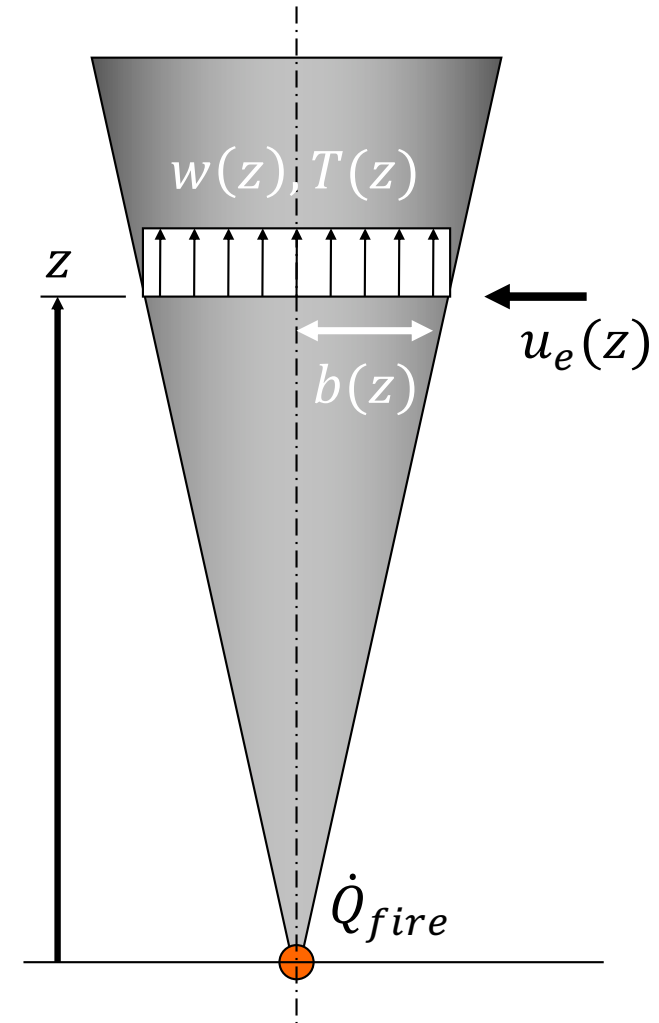
$$w(z) = C_w \times \dot{Q}_{fire}^{1/3} \times z^{-1/3}$$

$$\frac{T(z) - T_{air}}{T_{air}} = C_T \times \dot{Q}_{fire}^{2/3} \times z^{-5/3}$$

$$\dot{m}_z(z) = \rho(\pi b^2)w = C_{\dot{m}} \times \dot{Q}_{fire}^{1/3} \times z^{5/3}$$

$$C_b = \frac{6\alpha}{5}; \quad C_w = \left(\frac{25}{48\pi\alpha^2} \frac{g}{(\rho c_p T)_{air}} \right)^{1/3}$$

$$C_T = \frac{1}{\pi C_b^2 C_w (\rho c_p T)_{air}}; \quad C_{\dot{m}} = \frac{1}{C_T (c_p T)_{air}}$$





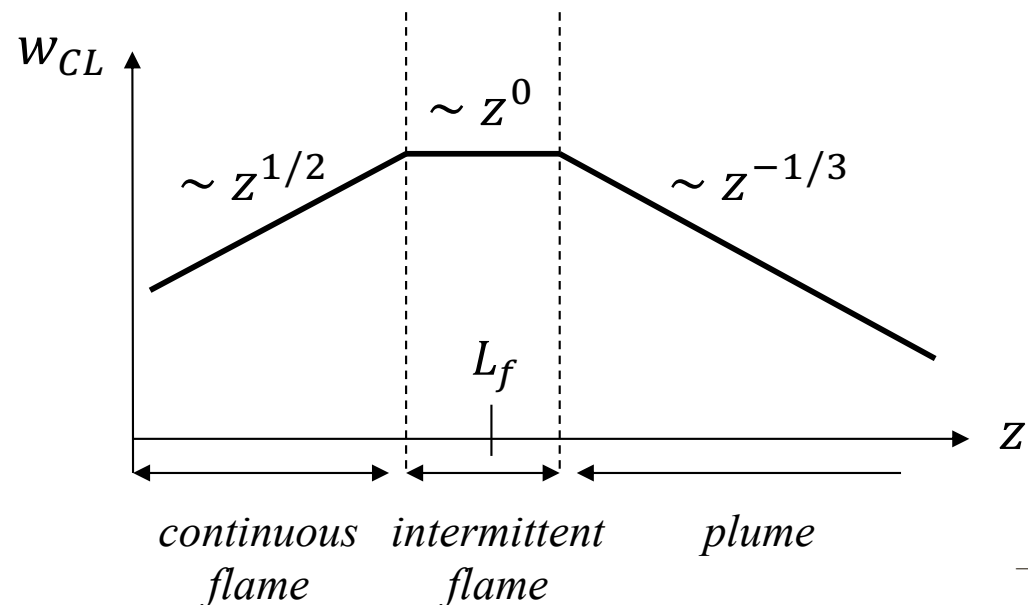
Plume Structure

■ McCaffrey correlations (pool fire config.)

- Description of near-field/flame region as well as far-field/plume region

$$\left| \begin{array}{ll} w_{CL}(z) = 6.8 \times z^{1/2}, & z < (0.08 \times \dot{Q}_{fire}^{2/5}) \\ w_{CL}(z) = 1.9 \times \dot{Q}_{fire}^{1/5}, & (0.08 \times \dot{Q}_{fire}^{2/5}) \leq z \leq (0.2 \times \dot{Q}_{fire}^{2/5}) \\ w_{CL}(z) = 1.1 \times \dot{Q}_{fire}^{1/3} \times z^{-1/3}, & (0.2 \times \dot{Q}_{fire}^{2/5}) < z \end{array} \right.$$

(\dot{Q}_{fire} in kW)





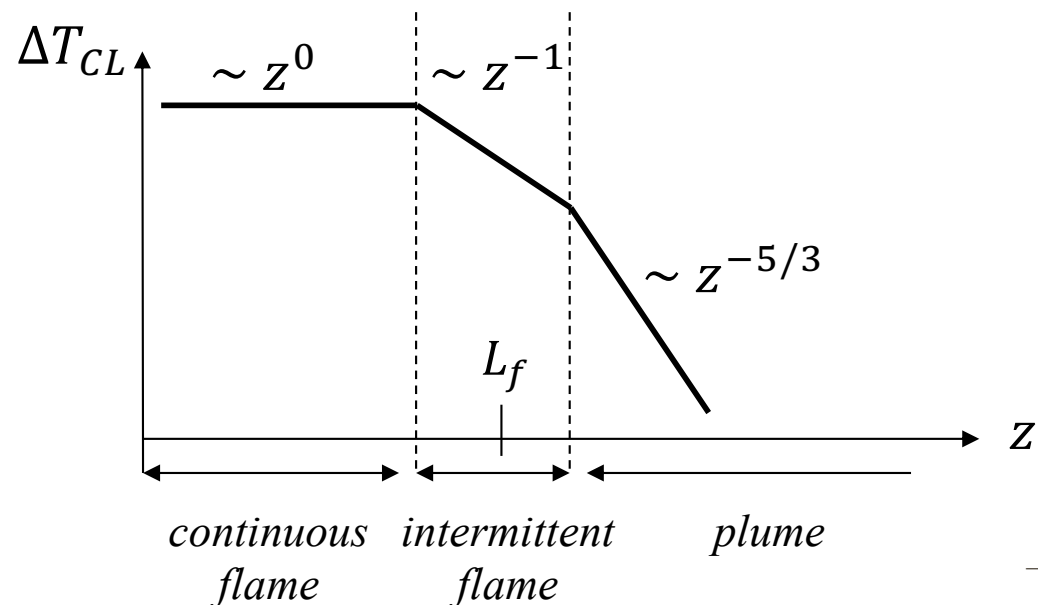
Plume Structure

■ McCaffrey correlations (pool fire config.)

- Description of near-field/flame region as well as far-field/plume region

$$\left\{ \begin{array}{l} \Delta T_{CL}(z) = T_{CL}(z) - T_{air} = 850, \quad z < (0.08 \times \dot{Q}_{fire}^{2/5}) \\ \Delta T_{CL}(z) = T_{CL}(z) - T_{air} = 67 \times \dot{Q}_{fire}^{2/5} \times z^{-1}, \quad (0.08 \times \dot{Q}_{fire}^{2/5}) \leq z \leq (0.2 \times \dot{Q}_{fire}^{2/5}) \\ \Delta T_{CL}(z) = T_{CL}(z) - T_{air} = 22 \times \dot{Q}_{fire}^{2/3} \times z^{-5/3}, \quad (0.2 \times \dot{Q}_{fire}^{2/5}) < z \end{array} \right.$$

(T_{CL} in K, \dot{Q}_{fire} in kW)





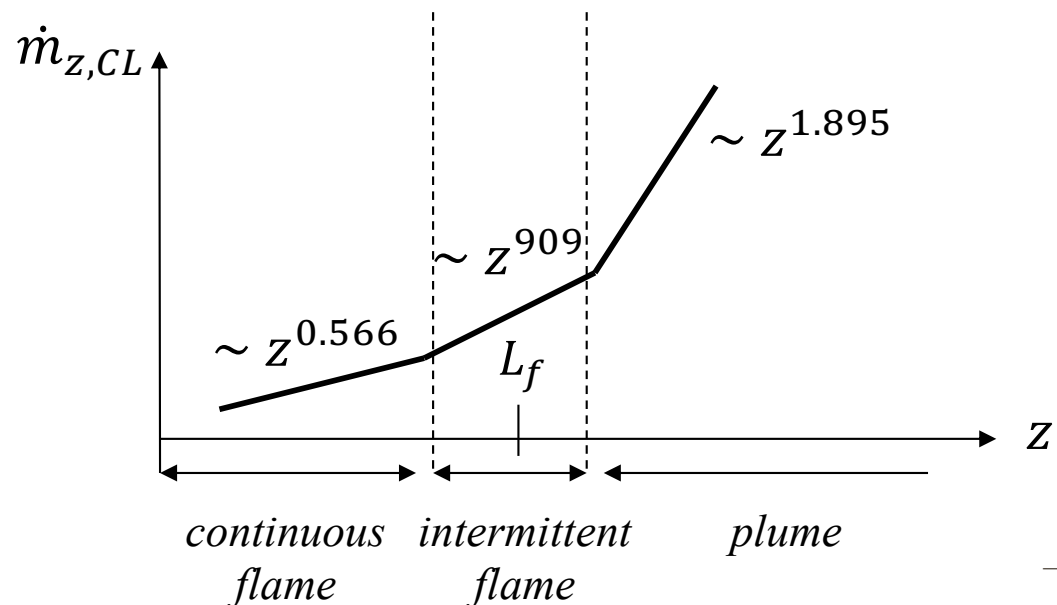
Plume Structure

■ McCaffrey correlations (pool fire config.)

- Description of near-field/flame region as well as far-field/plume region

$$\left| \begin{array}{ll} \dot{m}_{z,CL}(z) = 0.011 \times \dot{Q}_{fire} \times (z/\dot{Q}_{fire}^{2/5})^{0.566}, & z < (0.08 \times \dot{Q}_{fire}^{2/5}) \\ \dot{m}_{z,CL}(z) = 0.026 \times \dot{Q}_{fire} \times (z/\dot{Q}_{fire}^{2/5})^{0.909}, & (0.08 \times \dot{Q}_{fire}^{2/5}) \leq z \leq (0.2 \times \dot{Q}_{fire}^{2/5}) \\ \dot{m}_{z,CL}(z) = 0.124 \times \dot{Q}_{fire} \times (z/\dot{Q}_{fire}^{2/5})^{1.895}, & (0.2 \times \dot{Q}_{fire}^{2/5}) < z \end{array} \right.$$

(\dot{Q}_{fire} in kW)





Flame Structure

■ Mean flame height (line fire configuration)

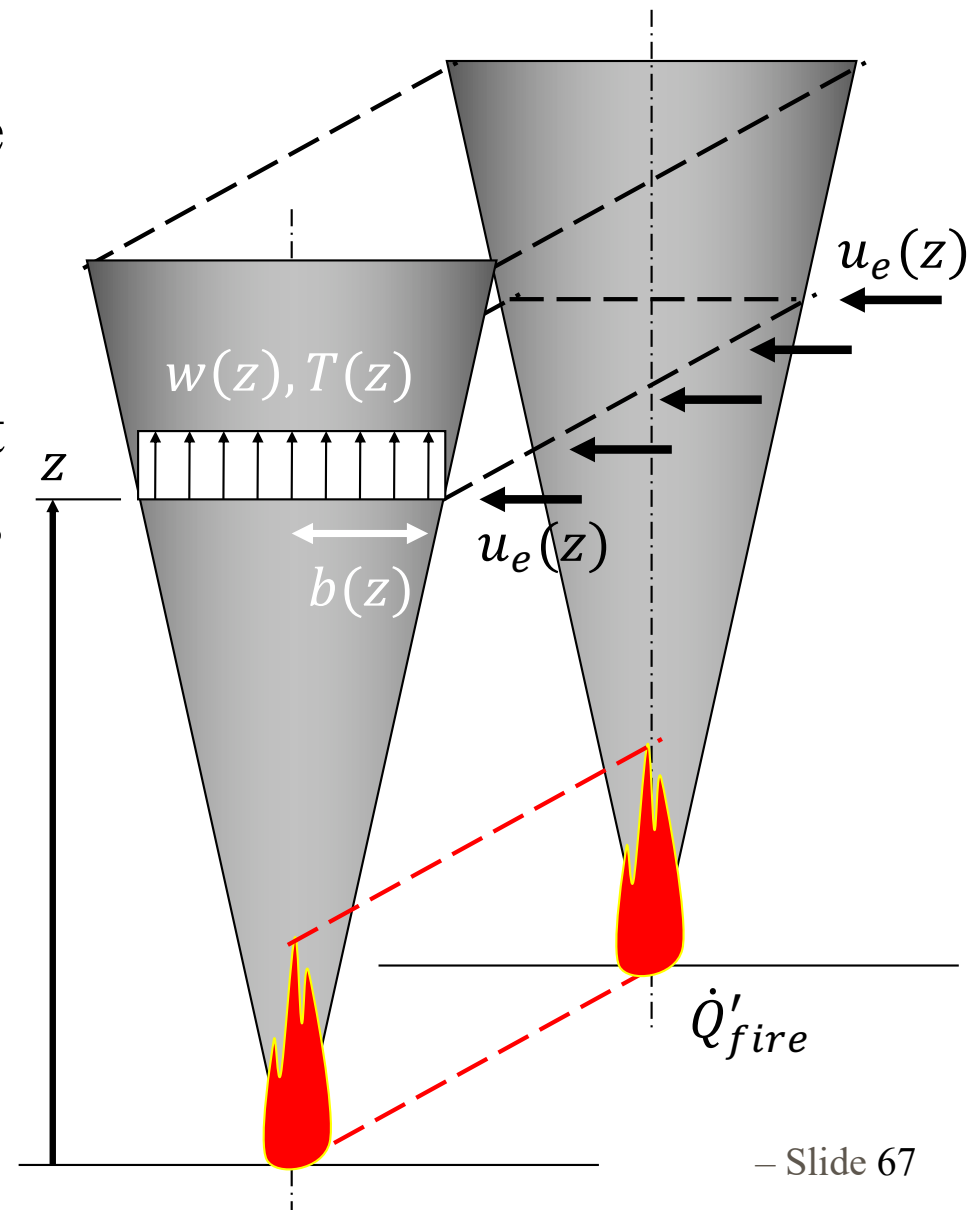
- 2D plane geometry
- Fire intensity: HRR per unit length of fireline [W/m], \dot{Q}'_{fire}
- Buoyant flames

(Byram 1959) (Yuan & Cox 1996)

$$L_f = 0.0775 \times (\dot{Q}'_{fire})^{0.46}$$

$$L_f = 0.034 \times (\dot{Q}'_{fire})^{2/3}$$

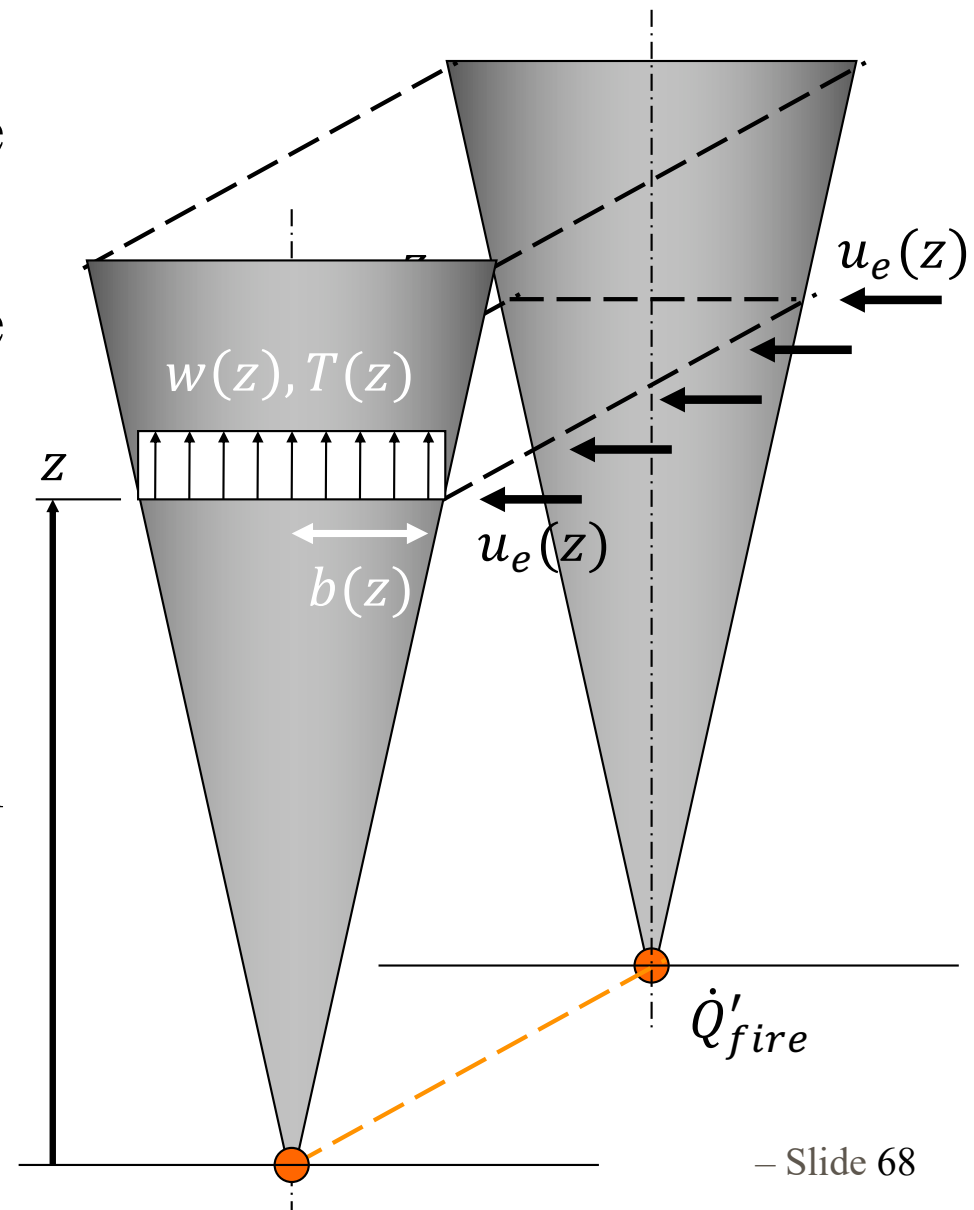
(\dot{Q}'_{fire} in kW/m)





Plume Structure

- Ideal buoyant plume (line fire configuration)
 - Two-dimensional plane geometry
 - Line source release of heat (far field plume theory)
 - “Fire induced wind”: $u_e(z) = \alpha \times w(z)$, empirical entrainment factor α ($\alpha \sim 0.1-0.2$)





Plume Structure

■ Ideal buoyant plume (line fire configuration)

$$b(z) = C_b \times z$$

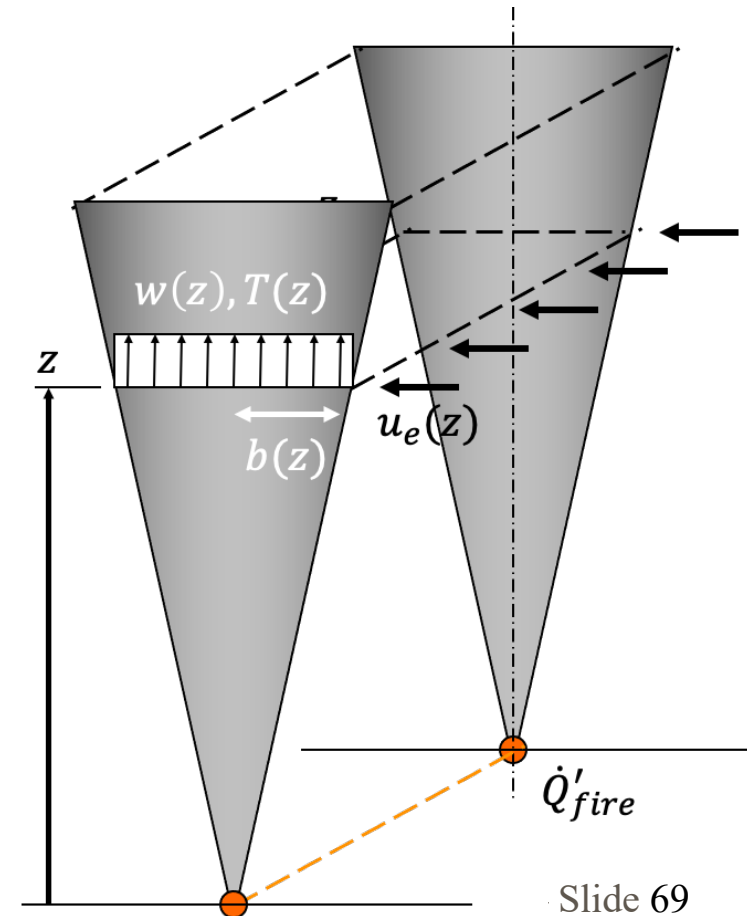
$$w(z) = C_w \times (\dot{Q}'_{fire})^{1/3}$$

$$\frac{T(z) - T_{air}}{T_{air}} = C_T \times (\dot{Q}'_{fire})^{2/3} \times z^{-1}$$

$$\dot{m}'_z(z) = \rho(2b)w = C_{\dot{m}'} \times (\dot{Q}'_{fire})^{1/3} \times z$$

$$C_b = \alpha; \quad C_w = \left(\frac{1}{2\alpha} \frac{g}{(\rho c_p T)_{air}} \right)^{1/3}$$

$$C_T = \frac{1}{2C_b C_w (\rho c_p T)_{air}}; \quad C_{\dot{m}'} = \frac{1}{C_T (c_p T)_{air}}$$





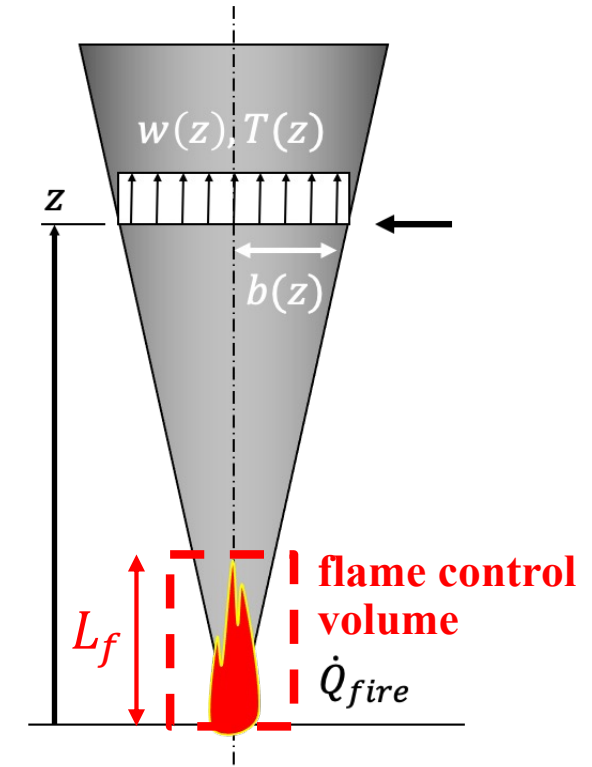
Flame Structure

- Importance of air entrainment, $u_e(z)$ (fire-induced/external wind)
 - Pool fire configuration
 - Global equivalence ratio for the flame region

$$\phi = \frac{(m_F^i / m_{O_2}^i)}{(MW_F / (v_{O_2} MW_{O_2}))} = r_S \left(\frac{m_F^i}{m_{O_2}^i} \right)$$

$$\Rightarrow \phi \approx r_S \times \left(\frac{\dot{m}_{fuel}}{\dot{m}_{air} \times Y_{O_2,air}} \right)$$

$$\Rightarrow \phi \approx r_S \times \left(\frac{\dot{m}_{fuel}}{\int_0^{L_f} (\rho u_e \times 2\pi b) dz \times Y_{O_2,air}} \right) \rightarrow$$



$$\phi \approx 0.1 \quad \text{over-ventilated fire}$$



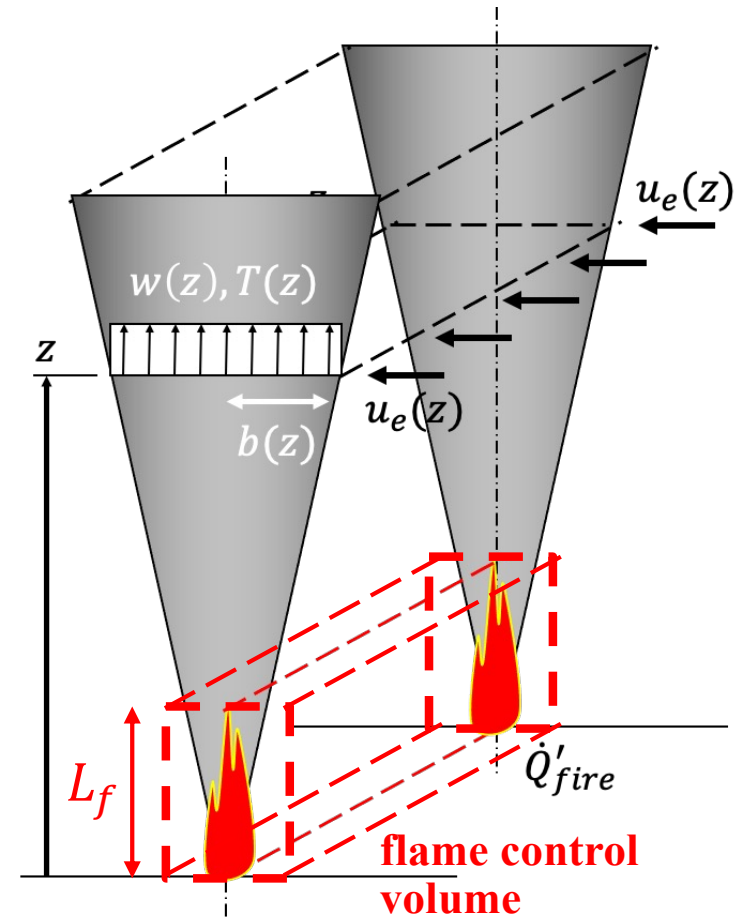
Flame Structure

- Importance of air entrainment, $u_e(z)$ (fire-induced/external wind)
 - Line fire configuration
 - Global equivalence ratio for the flame region

$$\phi = \frac{(m_F^i / m_{O_2}^i)}{(MW_F / (v_{O_2} MW_{O_2}))} = r_S \left(\frac{m_F^i}{m_{O_2}^i} \right)$$

$$\Rightarrow \phi \approx r_S \times \left(\frac{\dot{m}_{fuel}}{\dot{m}_{air} \times Y_{O_2,air}} \right)$$

$$\Rightarrow \phi \approx r_S \times \left(\frac{\dot{m}'_{fuel}}{\int_0^{L_f} (\rho u_e \times 2) dz \times Y_{O_2,air}} \right)$$



$$\phi \approx 0.1 \text{ over-ventilated fire}$$



Flame Structure

■ Importance of air entrainment

- Fuel package experiments: wood cribs (S. McAllister & M. Finney, *Fire Technology* 2015)



Under-ventilated wood crib
(flame stabilized on the outer edge of the crib, $\phi > 1$)



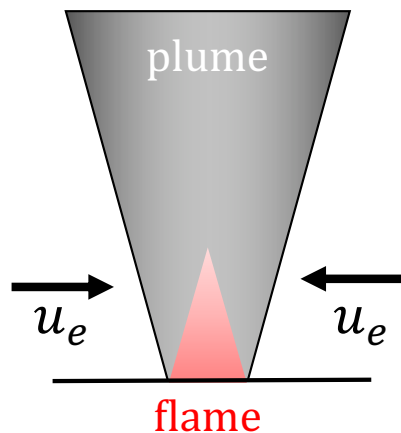
Over-ventilated wood crib
(flames stabilized around individual sticks, $\phi < 1$)



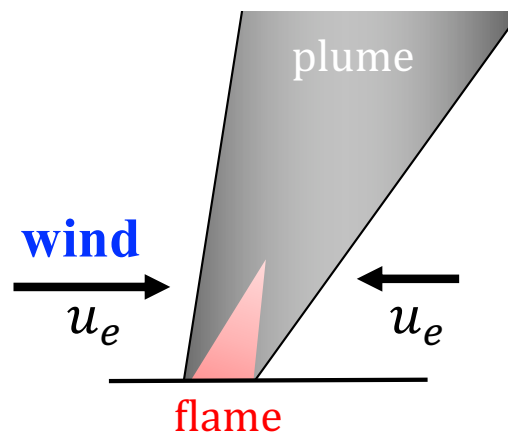
Flame Structure

■ Importance of air entrainment

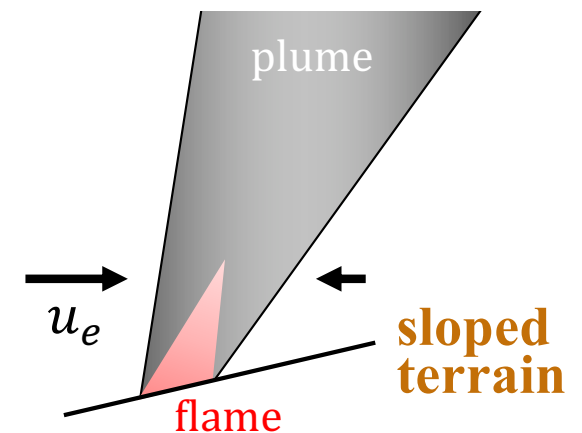
- Asymmetry in the entrained air flow will change the flame geometry



Symmetric



Asymmetric

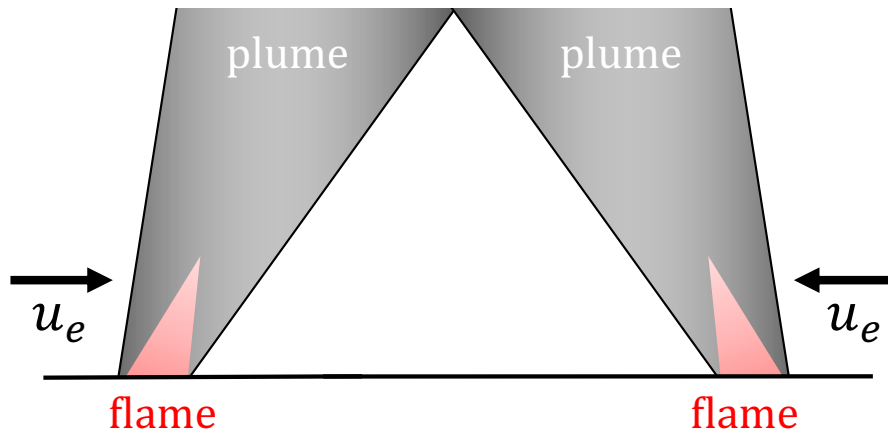


Asymmetric



Flame Structure

- Importance of air entrainment
 - Asymmetry in the entrained air flow will change the flame



Merging of fire fronts





Outline

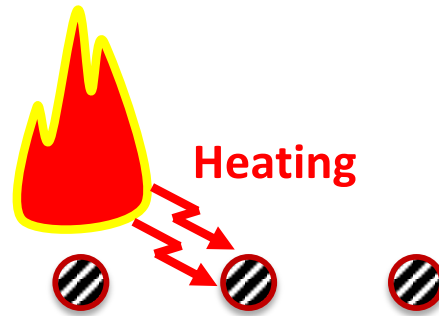
- Introduction
- Pyrolysis and Combustion
- Combustion: the Thermodynamics Viewpoint
- Combustion Chemistry
- Flame Structure
- **Flame Effects**



Flame Effects

■ Hazards

- Transport of combustion heat and mechanisms for further flame spread



- Radiative heat transfer – flame radiation and/or particle-to-particle radiation
- Convective heat transfer – flame/plume contact
- Transport of firebrands – spotting



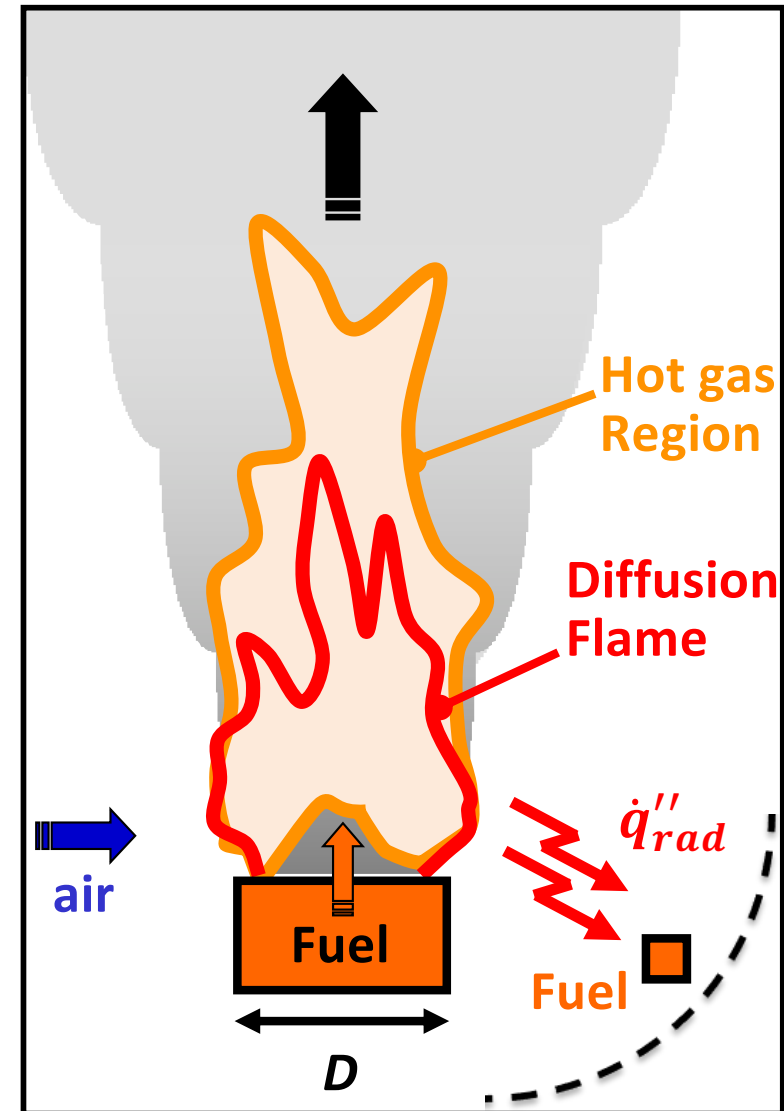
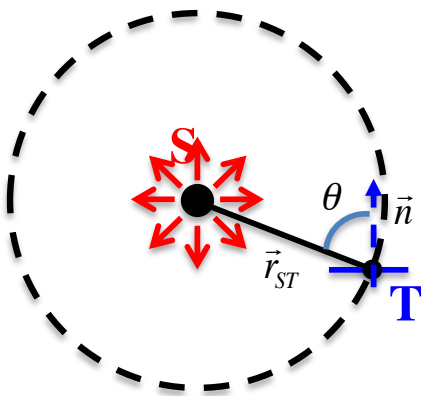
Flame Effects

■ Hazards

- Mechanisms for flame spread
- Radiative heat transfer
 - **Critical exposure conditions for spread:** $\dot{q}''_{rad} > CHF$ to achieve ignition conditions
 - Point source model

$$\dot{q}''_{rad} = \frac{(\chi_{rad} \times \dot{Q}_{fire})}{(4\pi d^2)} \times \cos(\theta)$$

$$\chi_{rad} = (\dot{Q}_{rad} / \dot{Q}_{fire}) \sim 0.3$$

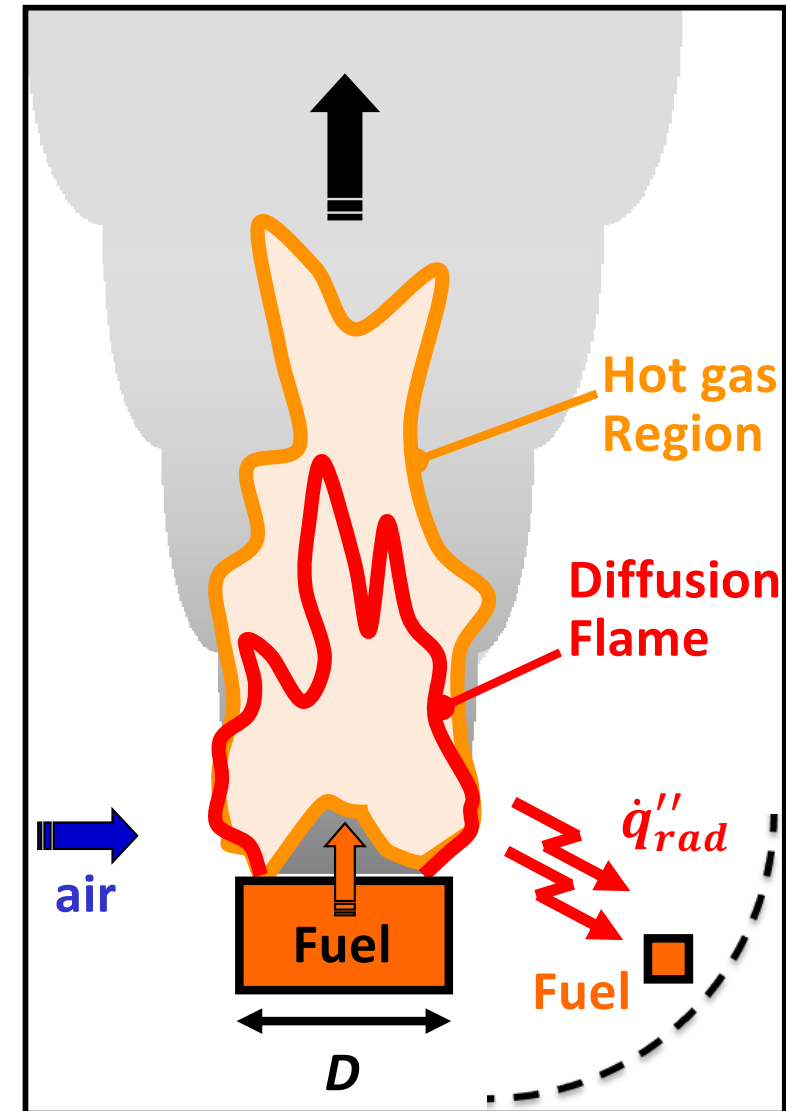
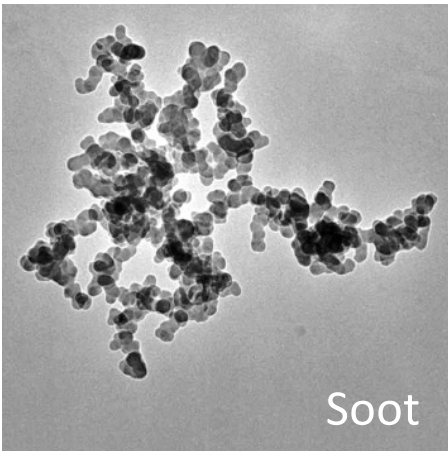




Flame Effects

■ Hazards

- Mechanisms for flame spread
- Radiative heat transfer
 - Radiative power of the flame/plume (i.e. value of χ_{rad}) determined by chemical composition of the gas: CO_2 , H_2O , and soot particles



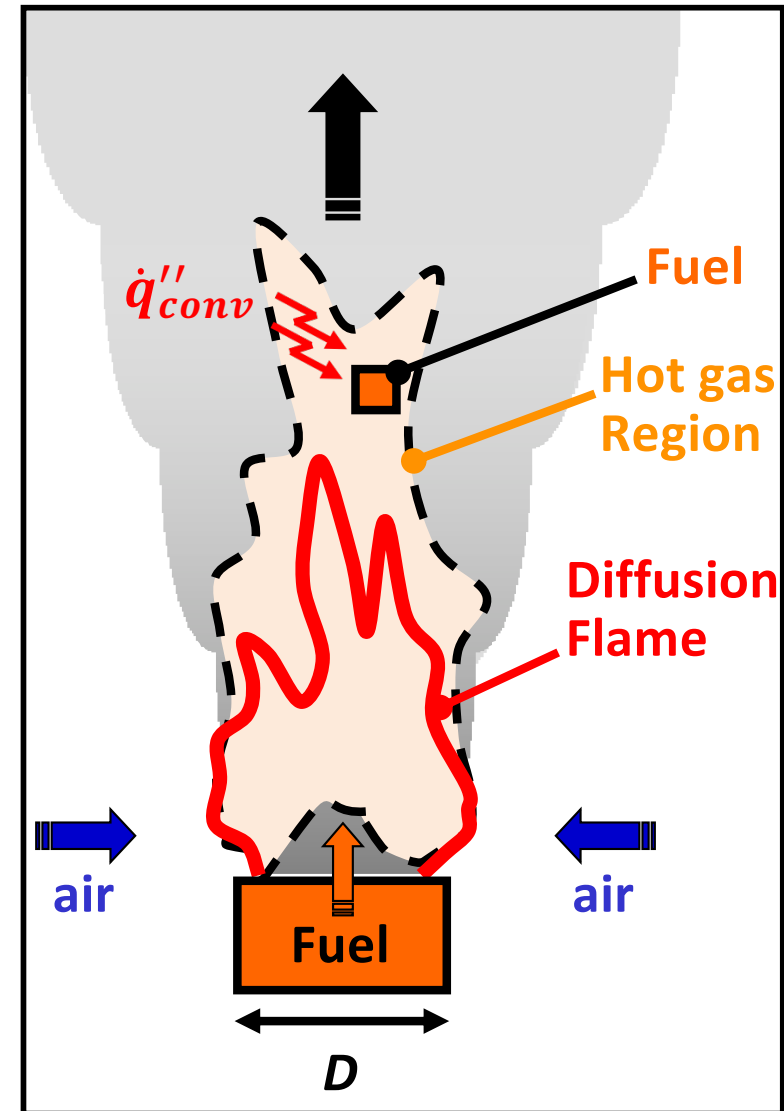


Flame Effects

■ Hazards

- Mechanisms for flame spread
- Convective heat transfer

➤ **Critical exposure conditions for spread:** $T > T_{pyro} \sim 600 \text{ K}$ to achieve ignition conditions

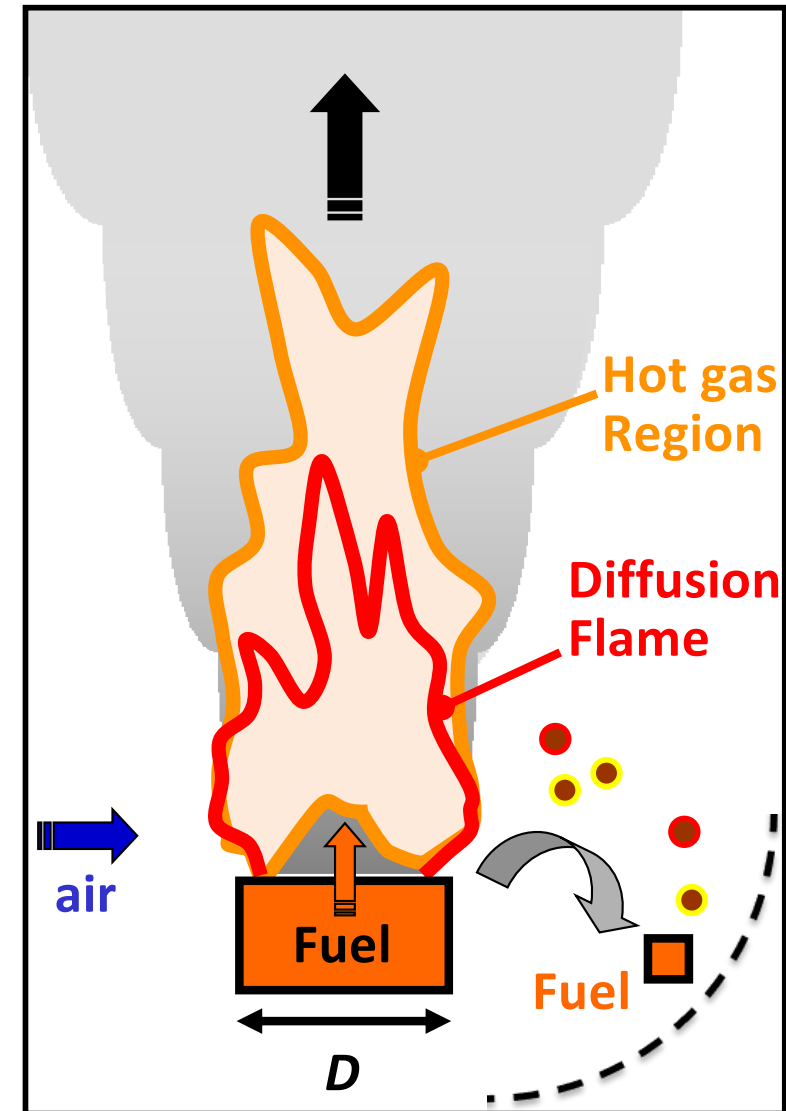




Flame Effects

■ Hazards

- Mechanisms for flame spread
- Transport of firebrands
 - **Critical exposure conditions for spread:** $\dot{m}''_{embers} > threshold$ to achieve ignition conditions

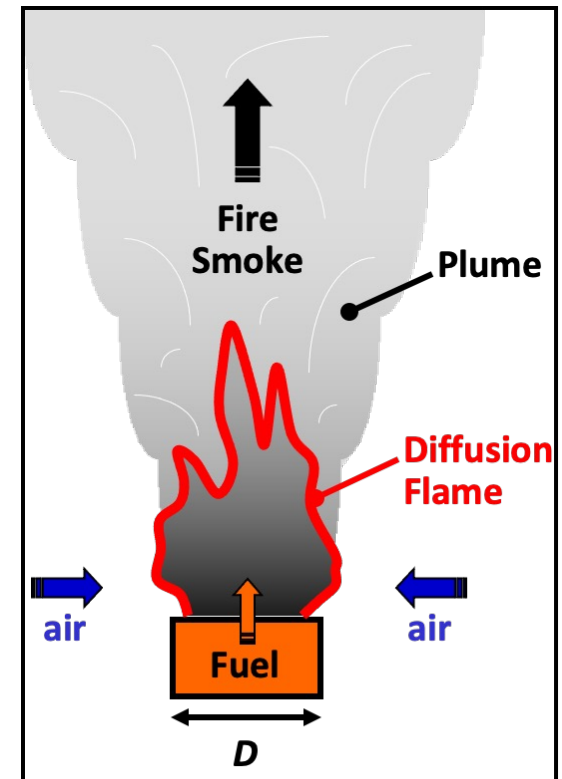




Flame Effects

■ Hazards

- Emissions of chemical species that may have a negative impact on human/environment/climate systems
 - Fire smoke contains toxic chemical species: *CO*, *HCN*, fine particulate matter (PM2.5), *etc*
 - Fire smoke contains atmospheric pollutants: *CO*, *NO/NO₂*, *PAHs*, particulate matter, *etc*
 - Fire smoke contains greenhouse gases: *CO₂*, *CH₄*, *N₂O*, *etc*

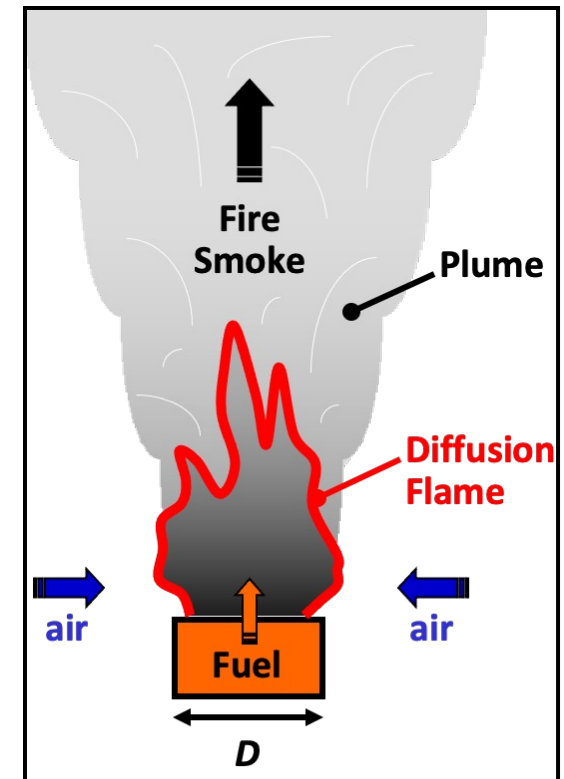




Flame Effects

■ Hazards

- Emissions of chemical species that may have a negative impact on human/environmental health
 - **Fire-smoke-exposed region:** not well understood
 - High-temperature combustion chemistry
 - Plume dynamics
 - Low-temperature atmospheric chemistry





Conclusion

- The basic physical concepts used to describe combustion phenomena have been introduced
 - Scope of discussion
 - Solid fuel (vegetation biomass)
 - Flaming combustion
 - Non-premixed flame
 - Buoyancy-driven flame
 - Turbulent flame
 - No wind and flat terrain conditions
 - Non-spreading flame



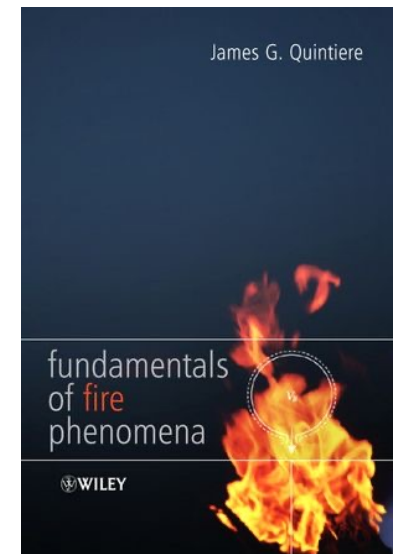
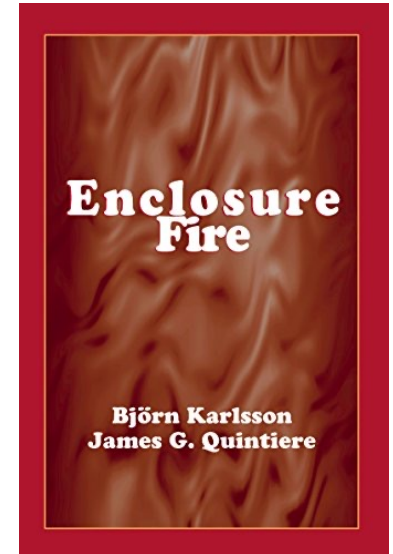
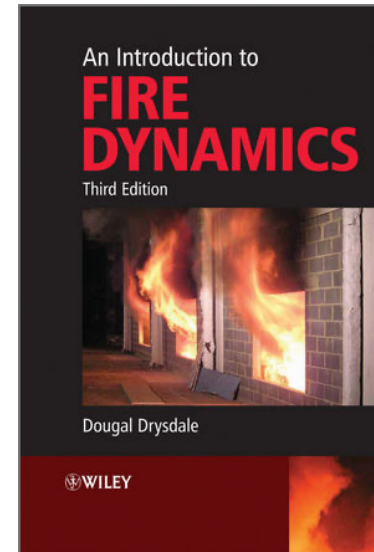
Conclusion

- Flame structure
 - Global combustion equation
 - Over-ventilated vs under-ventilated fire conditions
 - Combustion chemistry (Arrhenius model); infinitely fast chemistry assumption; rate of combustion controlled by mixing of $Fuel/O_2$; flame structure described by mixture fraction variable
 - Flame height as the characteristic length scale of the combustion region; correlations for flame height
 - Mechanisms for flame spread: convective heat transfer; radiative heat transfer (importance of soot particles); firebrands
 - Emissions of chemical species: toxic chemical species, atmospheric pollutants, greenhouse gases



Further Reading

- Fundamentals of compartment fire dynamics
 - Karlsson B., Quintiere J.G., *Enclosure Fire Dynamics*, CRC Press LLC, 2000
 - Drysdale D., *An Introduction to Fire Dynamics*, 3rd, Wiley, 2011
 - Quintiere J.G., *Fundamentals of Fire Phenomena*, Wiley, 2006





Further Reading

- Fundamentals of wildland fire behavior
 - Finney M., McAllister S., Grumstrup T., Forthofer, J., *Wildland Fire Behavior - Dynamics, Principles and Processes*, CSIRO Publishing, 2021

