

What's a fire ?

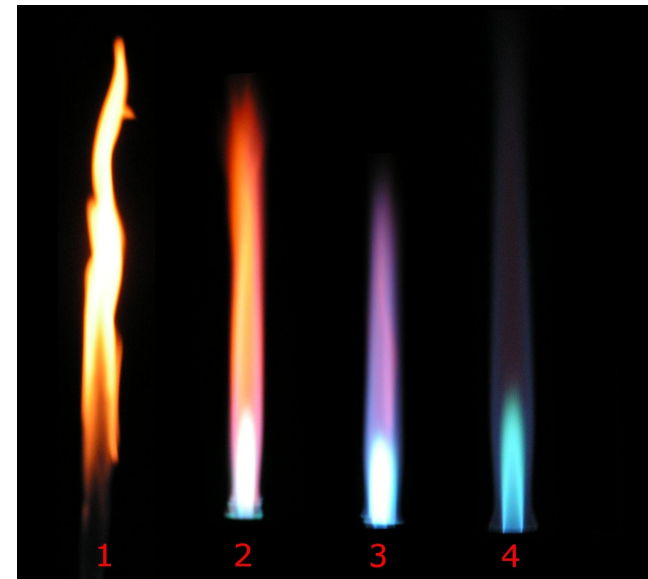
The rapid oxidation (combustion) of a material (the fuel) releasing heat, light and various reaction products (*adapted from standard definitions*)

Contrary to many combustion devices (engines, bunsen burner,...) :

- fire is a self-sustained process,
- often uncontrolled.



Fire spreading in a grass fuel



Bunsen Burner (combustion device)

Common fuels

All include carbon in any form (hydrocarbons, alcohols, carbohydrates)

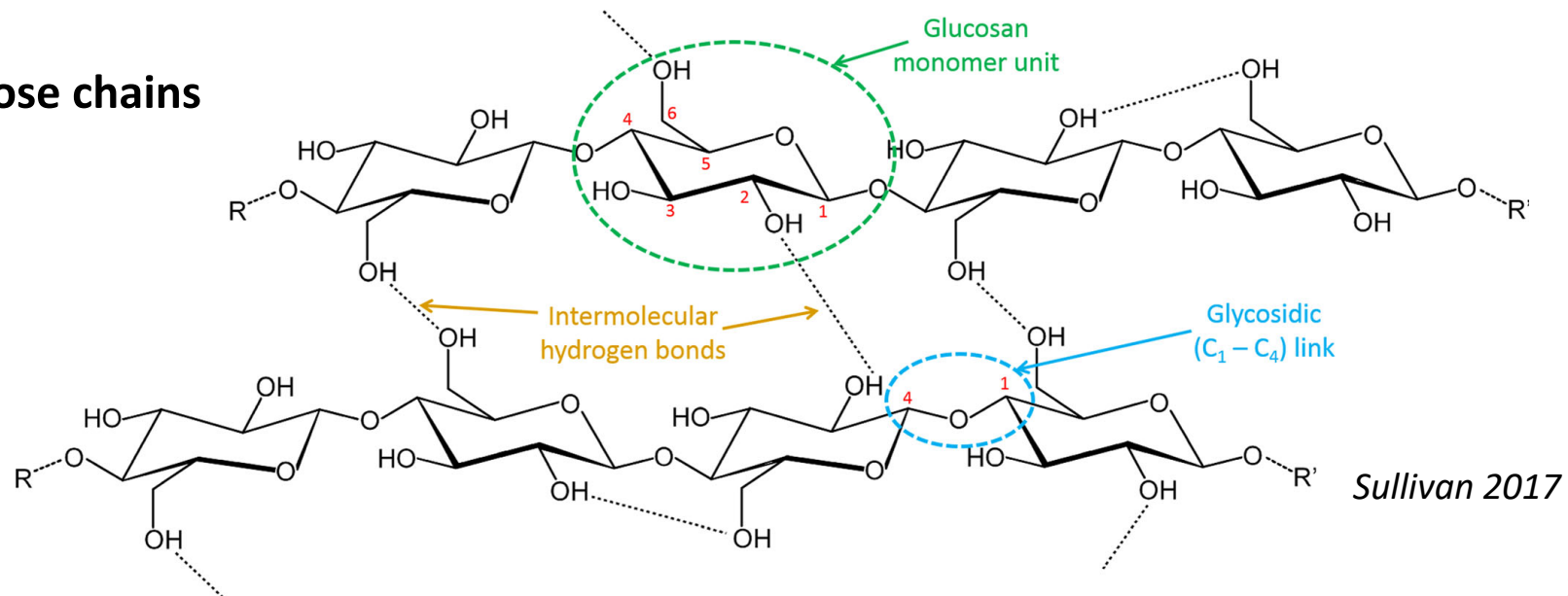
Gaseous : methane (natural gas), propane, ethylen, ...

Liquid : alcohols, gasoline

Solid : coal, synthetic polymers, wood and other natural fibers

Wood and other biomass fuels are mostly composed of natural polymers of carbohydrates ($C_m(H_2O)_n$) : cellulose, hemicelluloses and lignin.

Cellulose chains



Biomass fuels

Table 2 Approximate analysis of some biomass species taken from Shafizadeh [52], Mok and Antal [61] and Demirbaş [62, 63].
Source: modified from [41] with permission from Elsevier

Sample	Cellulose (%)	Hemicelluloses (%)	Lignins (%)	Other ^a (%)
Shafizadeh [52]				
Softwood	41.0	24.0	27.8	7.2
Hardwood	39.0	35.0	19.5	6.5
Wheat straw	39.9	28.2	16.7	15.2
Rice straw	30.2	24.5	11.9	33.4
Bagasse	38.1	38.5	20.2	3.2
Mok and Antal [61]				
<i>Eucalyptus saligna</i>	45	15	25	15
<i>Eucalyptus gummifera</i>	38	16	37	9
Sweet sorghum	36	18	16	30
Sugar cane bagasse	36	17	17	30
<i>Populus deltoides</i>	39	21	26	14
Demirbaş [62, 63]				
Softwood (av.)	45.8	24.4	28.0	1.7
Hardwood (av.)	45.2	31.3	21.7	2.7
Wood bark	24.8	29.8	43.8	1.6
Wheat straw	28.8	39.1	18.6	13.5
Tobacco stalk	42.4	28.2	27.0	2.4
Tobacco leaf	36.3	34.4	12.1	17.2
Spruce wood	50.8	21.2	27.5	0.5
Beech wood	45.8	31.8	21.9	0.4
Ailanthus wood	46.7	26.6	26.2	0.5

^a Other can consist of organic compounds such as starch or inorganic material such as salts, minerals, water and extractives

Cellulose, hemicelluloses and lignin are non-soluble

Extractives (i.e. soluble) include waxes, resin, simple sugars, starches, proteins, ..., and volatile organic compounds (VOCs, such as terpenes)

Fuels in 'natural' fires

All live and dead elements of vegetation + Soil organic material

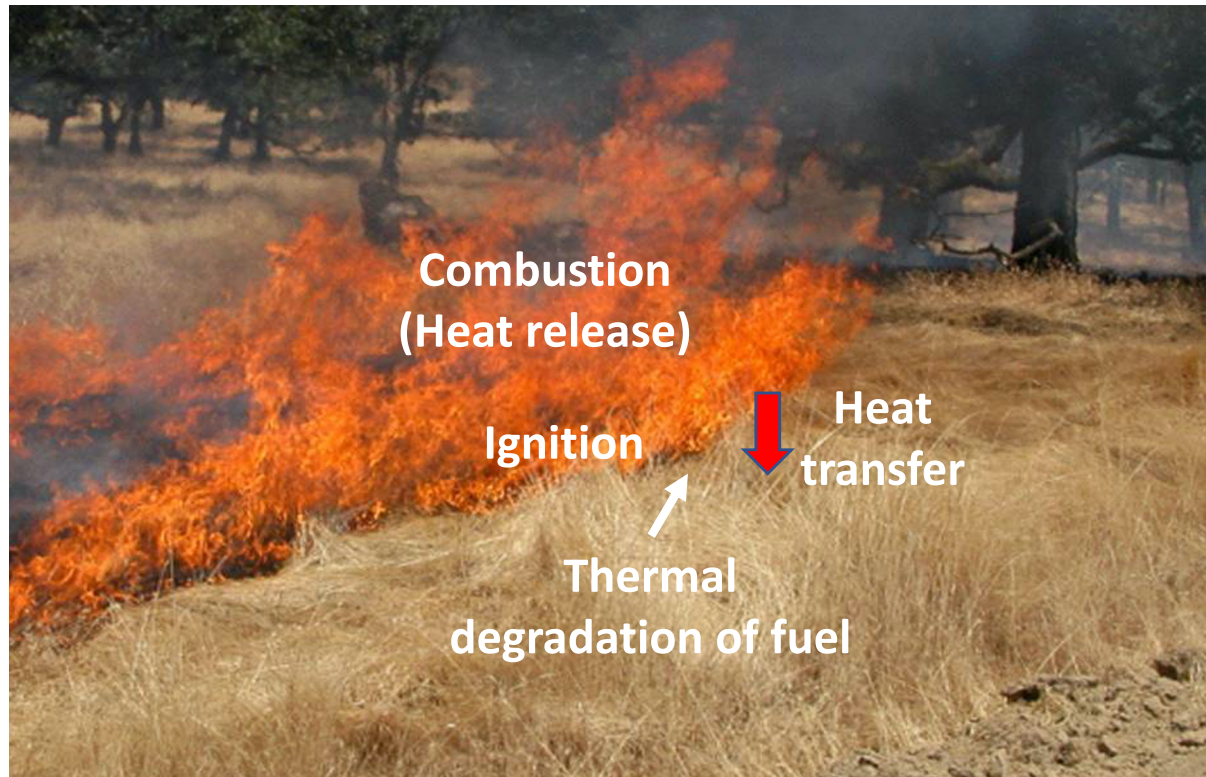
Fine elements (< 6 mm) drive fire spread

Larger fuels, even trunks, may burn (at least partially)

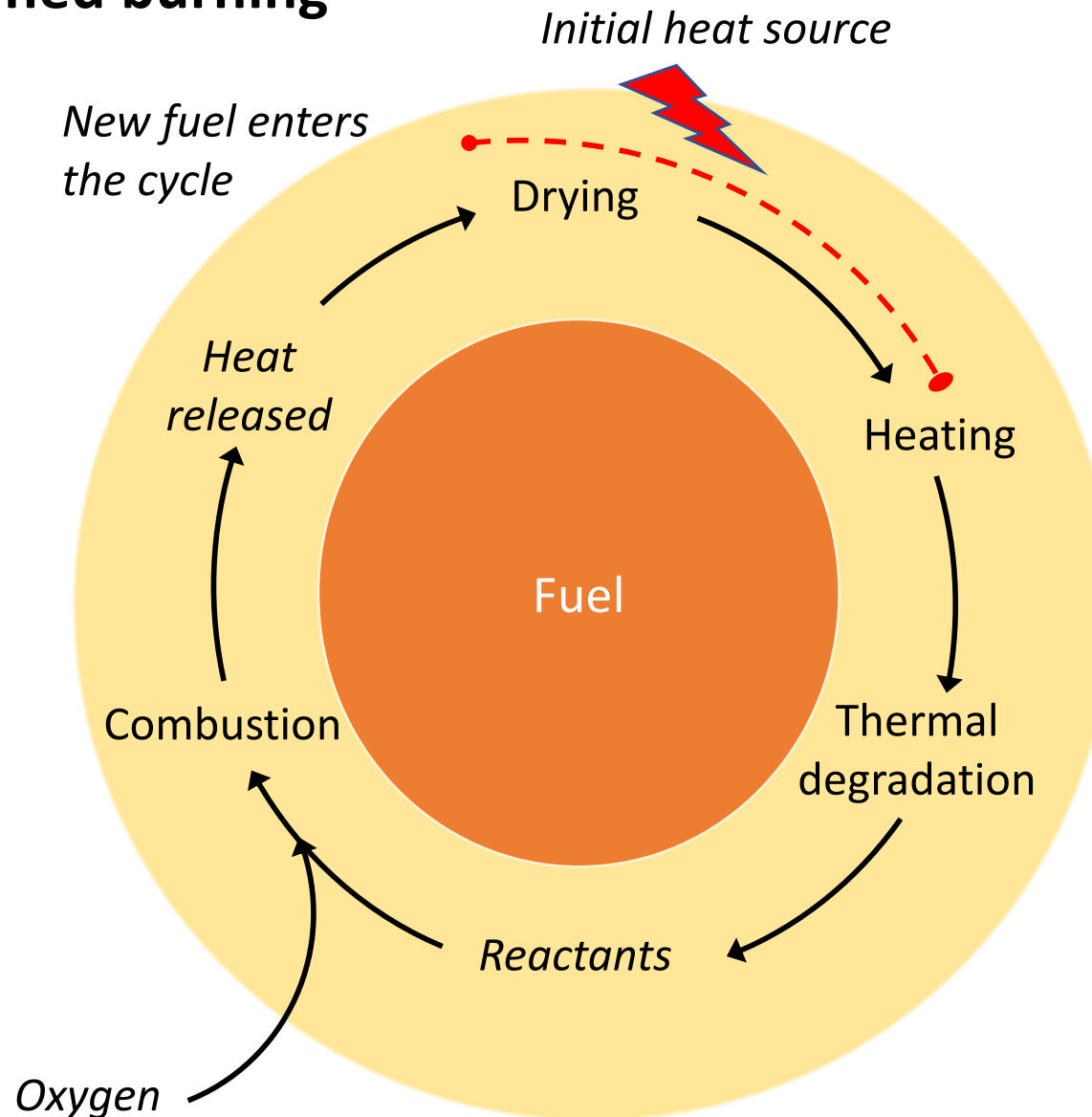


Self-sustained burning

Fundamental processes

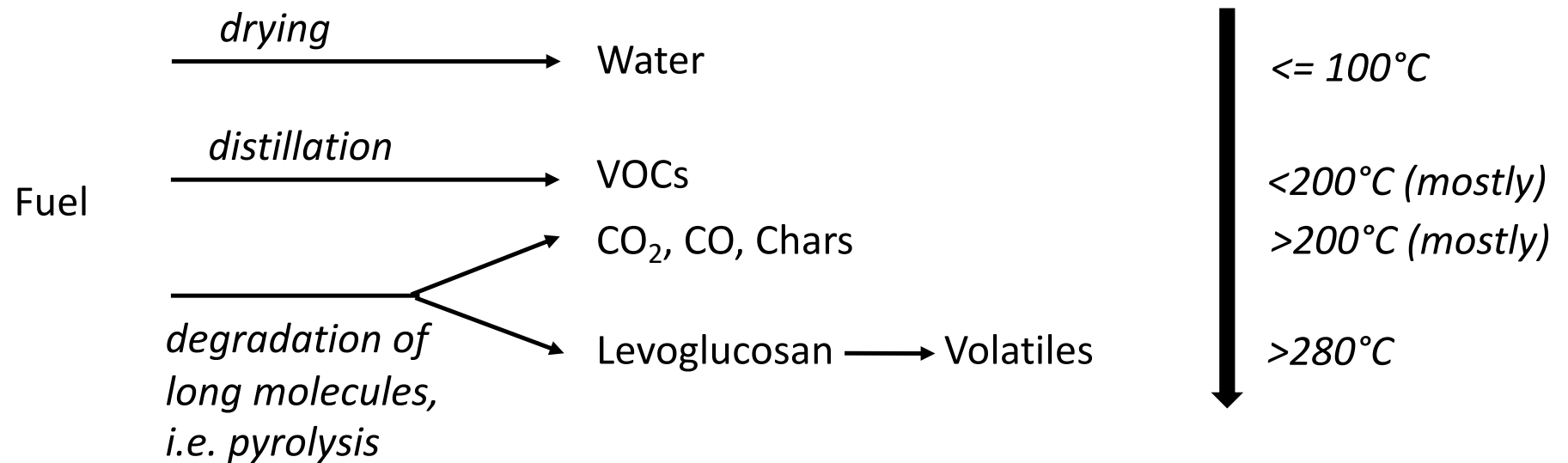


Self-sustained burning



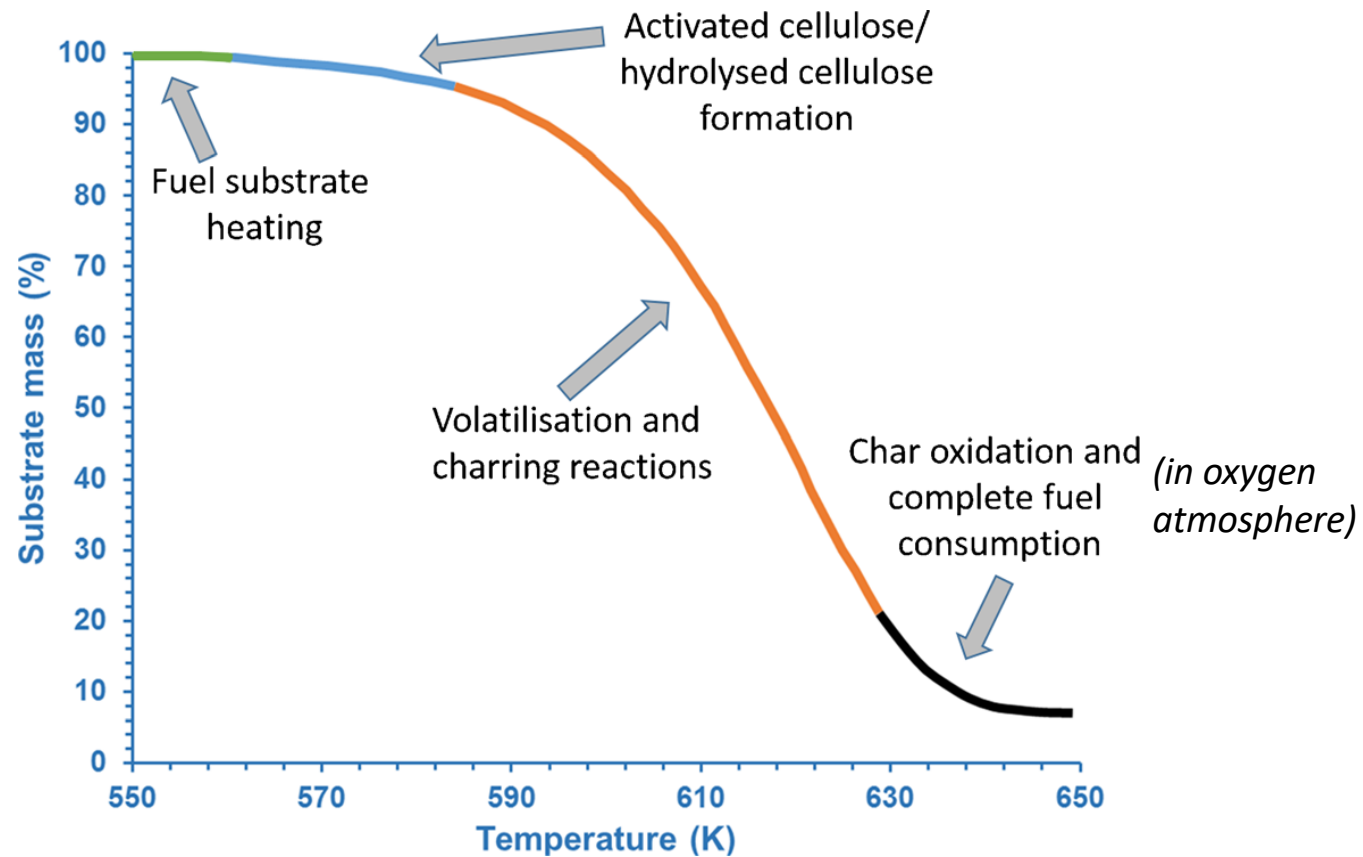
Thermal degradation

Under heating, fuels undergo **drying, distillation and pyrolysis**



Thermal degradation

Thermo-gravimetric analysis



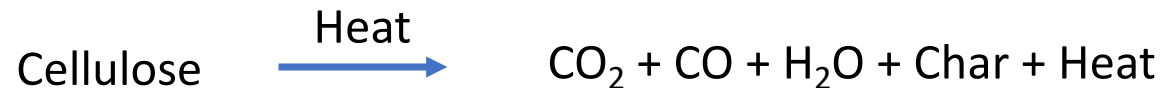
Sullivan 2017

Thermal degradation (details on pyrolysis of cellulose)

Cellulose (usually 40 to 50% of material) has been studied extensively

Two main pathways : char formation and volatilisation

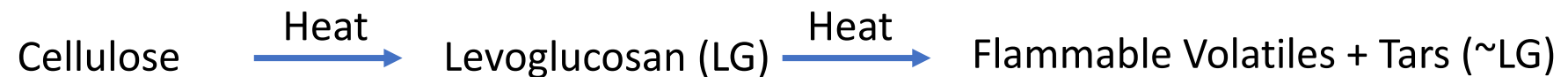
Char formation at low temperature (< 280°C) :



Char formation at high temperature :



Volatilisation (high temperature):



Char formation is slightly exothermic

Volatilisation is slightly endothermic

Ignition

Ignition is the appearance of combustion, generally accompanied by a flame.

Needs **fuel, heat and oxygen**, but the conditions are more drastic :

1- Most biomass fuels do not burn when **O₂ concentration** is below 15%
(atmospheric air normally contains 21% oxygen in volume)

2- Gaseous flammable products (released by distillation and pyrolysis of solid fuel) have lower and upper **flammability limits**, i.e. they don't ignite outside these limits

3- **Fuel heating must be enough, and fast enough**, to get a sufficient flow of flammable products to exceed the lower flammability limit when mixed with the air

In practice, ignition may happen if the woody fuel is "rapidly" heated to a critical temperature:

- 300-350°C with a pilot flame (**pilot-ignition temperature**)
- a higher threshold (600°C) with no pilot (spontaneous ignition temperature)

Pre-heating of the solid fuel elements

Prior to ignition, fuel must be pre-heated to ignition temperature T_{ig}

Both the dry material and the water of the fuel must be heated from ambient temperature T_a (300 K) to T_{ig} (~600 K)

Dry material : $Q_{dig} = C_{pd} (T_{ig} - T_a) = 390 \text{ J/g}$ (g of dry material)

Liquid water must be heated up to 100°C ($T_{boil} = 373 \text{ K}$) and then vaporized:

$$Q_w = C_{pw}(T_{boil} - T_a) + L_v = 2560 \text{ J/g} \text{ (g of water)}$$

C_{pd} is the specific heat of dry matter (1.3 J/K g)

C_{pw} is the specific heat of liquid water (4.18 J/K g)

L_v is the latent heat of vaporization of water (2257 J/g)

If the water content of fuel is **FMC** :

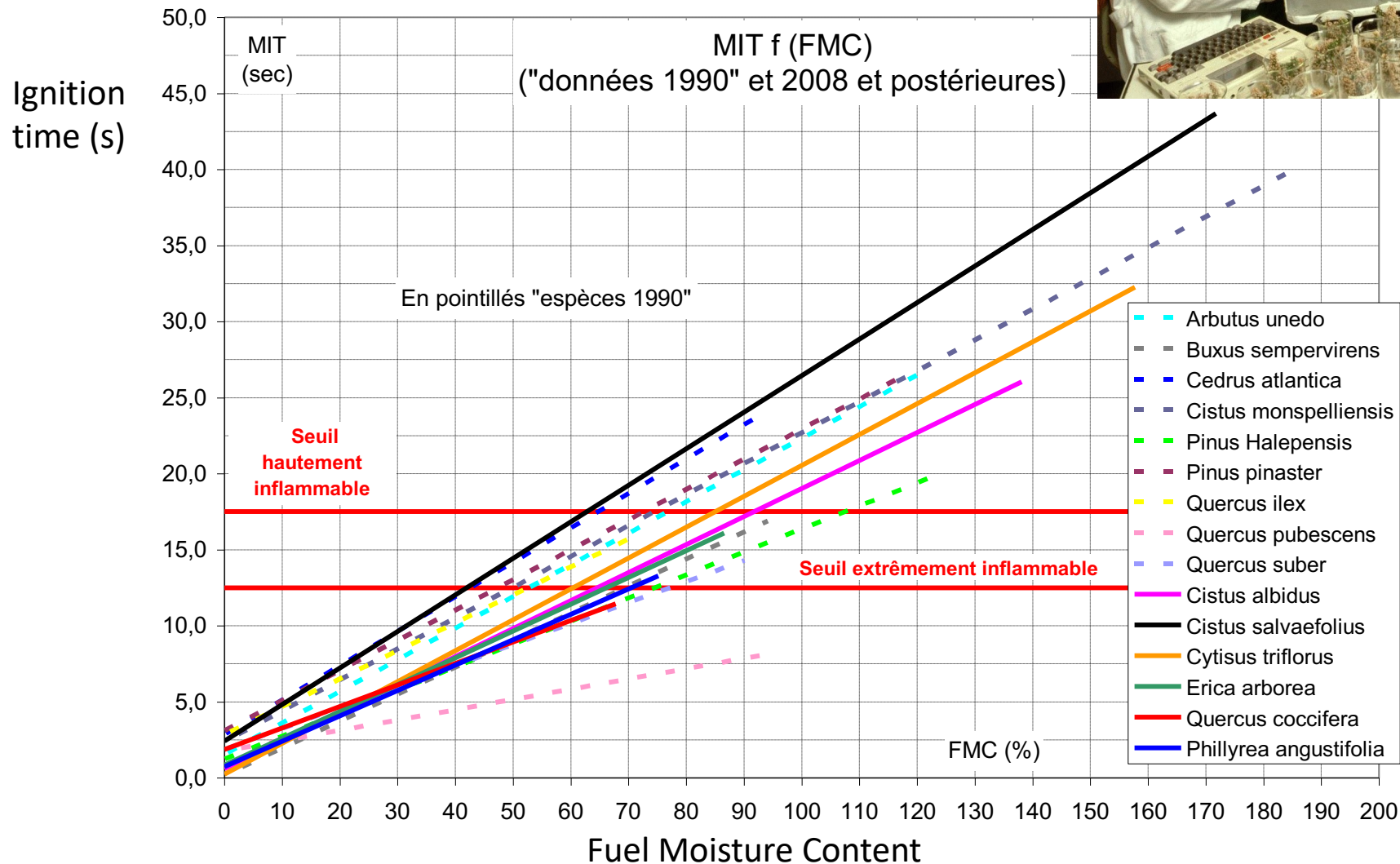
Fuel Moisture Content = mass of water / mass of dry fuel

Then the **heat of pre-ignition of the fuel** is :

$$Q_{ig} = Q_{dig} + FMC Q_w$$

Ignition tests

Compare species samples
Assess moisture effect



Combustion

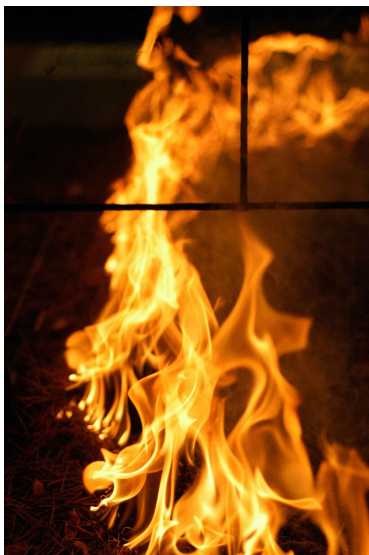
Flaming combustion :

- Products of volatilisation (gaseous fuels) react with oxygen in the air
- Fast reaction controlled by mixing of gaseous fuel and oxygen

Glowing or smouldering (low temperature) combustion :

- Oxidation of chars
- Slower reaction controlled by oxygen diffusion to char surface
- Incomplete combustion, especially smouldering

Flaming



Glowing



Smouldering



Combustion products

Complete combustion (theoretical):

- CO₂ and H₂O are the only final products
- O₂ consumption may be computed if fuel composition is known

Combustion is largely incomplete in wildland fires, which is measurable thanks to the Equivalent Oxygen to Fuel Ratio (EOFR):

$$\text{EOFR} = \frac{\text{moles of oxygen actually consumed}}{\text{moles of oxygen consumed in complete combustion}}$$

Typical emission factors of wood combustion (g per 1000 g of fuel)

95% of carbon released in CO₂, CO and CH₄

Type of combustion :	Complete	Flaming	Smoldering
EOFR	1.00	0.93	0.80
Water (H ₂ O)	559	546	523
Carbon dioxide (CO ₂)	1821	1632	1283
Carbon monoxide (CO)	0	90	257
Methane (CH ₄)	0	3	9
Other hydrocarbons	0	2	6
Particulate matter (PM)	0	9	25

In *Fire Science*, Springer, 2021

Combustion products

Significance : impacts to the atmosphere, air pollution

CO₂ is the most abundant species by far, but relatively inert

Other, less abundant, products raise specific concerns :

- CO: toxicity
- CH₄: high global warming potential
- NMOCs* , Nitrogen molecules, Sulfur dioxide (SO₂) : toxicity, impacts on atmospheric chemistry
- PM : toxicity, impacts on radiative forcing

*NMOCs : Non-methane organic compounds, a number of molecules each representing very low contribution

Combustion products

Modified Combustion Efficiency (MCE) and emission factors (g/kg) for different fire types

Data from airborne or tower-based measurements above fires (Urbanski 2014)

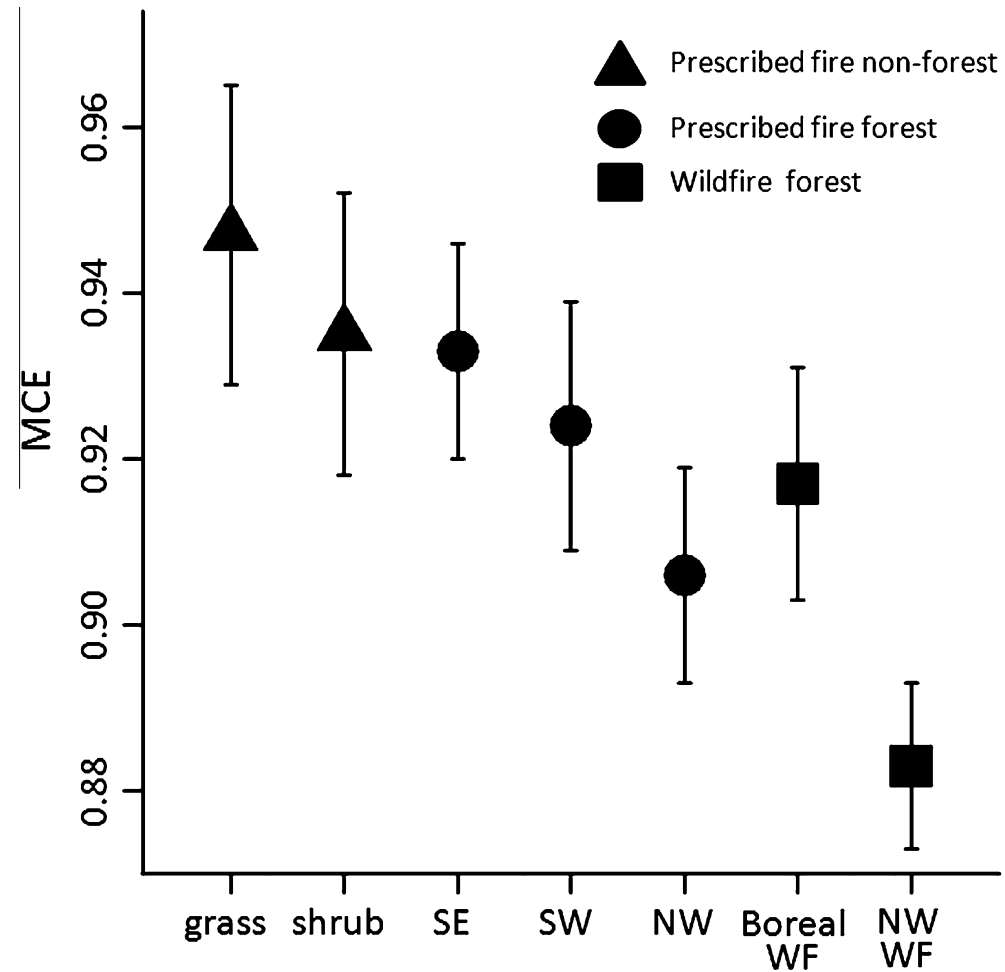
	Prescribed fires			Wildfires	
	Northwest conifer forest	Western Shrubland	Grassland	Northwest conifer forest	Boreal forest
MCE	0.906	0.935	0.947	0.883	0.917
Carbon dioxide (CO ₂)	1598	1674	1705	1600	1641
Carbon monoxide (CO)	105	74	61	135	95
Methane (CH ₄)	4.86	3.69	1.95	7.32	3.38
NMOCs	47.3	24.6	23.9	59.6	38.3
PM < 2.5 μm	17.6	7.06	8.51	23.2	21.5
Nitrogen molecules	3.75	3.93	3.68	3.66	2.2

$MCE = CO_2 / (CO_2 + CO)$, an indicator of smoldering vs flaming activity

Orders of magnitude are similar among fire types, but significant differences appear (uncertainties reported in Urbanski 2014) :

e.g. more efficient combustion in grasslands and shrublands (see next Figure)

Combustion products



MCE = $\text{CO}_2 / (\text{CO}_2 + \text{CO})$ in products, an indicator of smoldering vs flaming activity

Combustion products

Modified Combustion Efficiency (MCE) and emission factors (g/kg) for smoldering fuels

Data from ground-based measurements (Urbanski 2014)

	Stumps and Logs	Temperate forest duff/soil	Boreal forest duff/soil
MCE	0.796	0.752	0.790
Carbon dioxide (CO ₂)	1408	1305	1436
Carbon monoxide (CO)	229	271	244
Methane (CH ₄)	13.9	7.47	8.42
NMOCs	84.9	247	183
PM < 2.5 μm	33	50	20.6
Nitrogen molecules	0.48	3.34	3.34

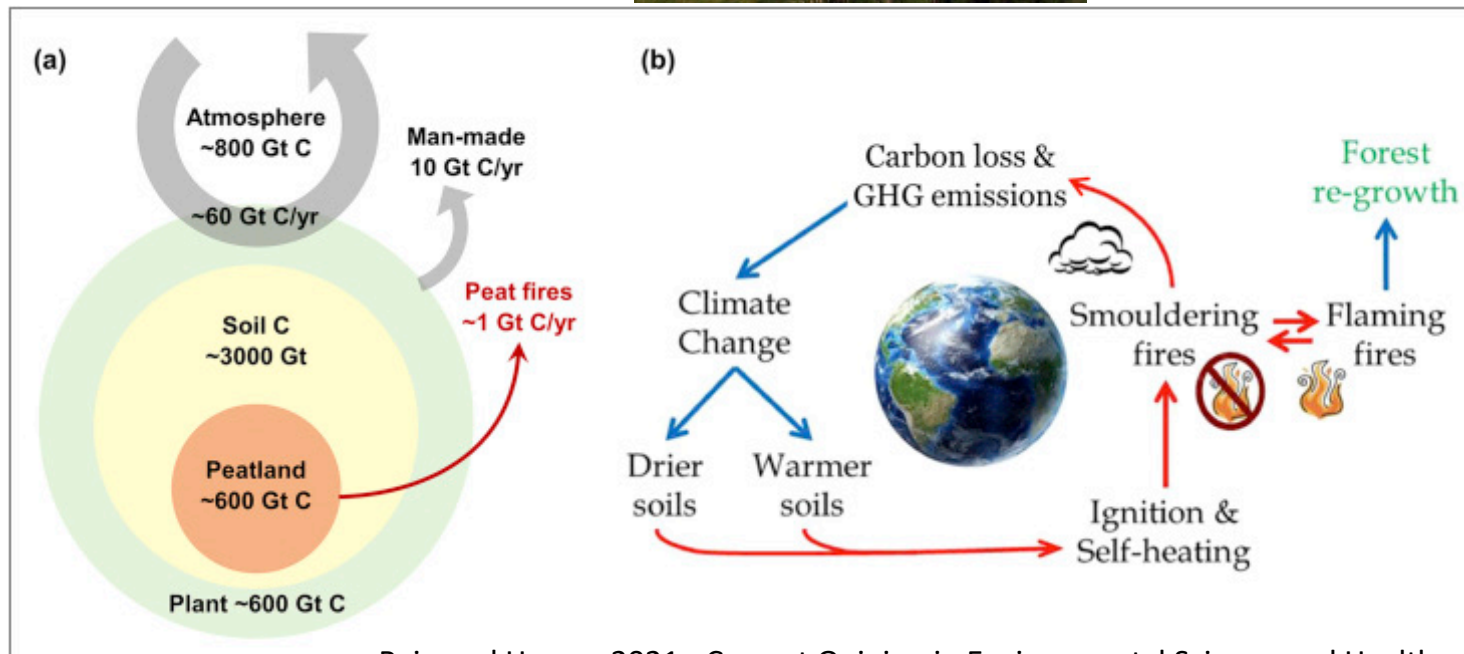
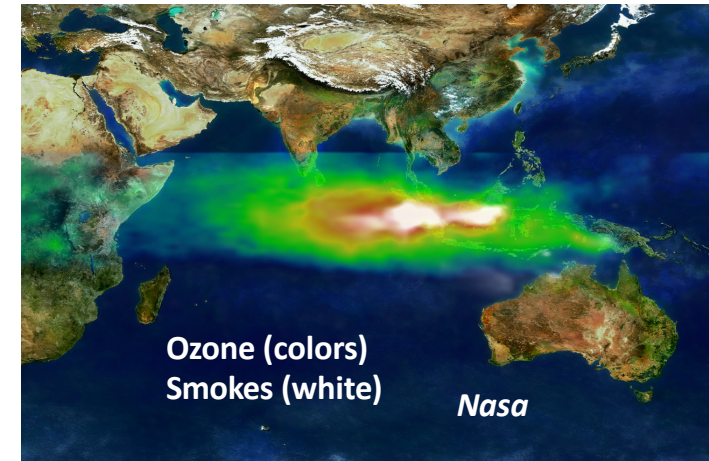
As expected, pollutants are much more represented over these smoldering fuels than above flaming fires with a significant convection column transporting smokes to the upper atmosphere.

Combustion products

Peat fires release huge amount of carbon and pollutants, with significant impacts to the carbon cycle (feedback) and the human health (haze)



Peat fires, 1997, Indonesia



Combustion heat release

The heat released by combustion may be computed when reactants and products are known. By convention, energy release (exothermicity) is negative

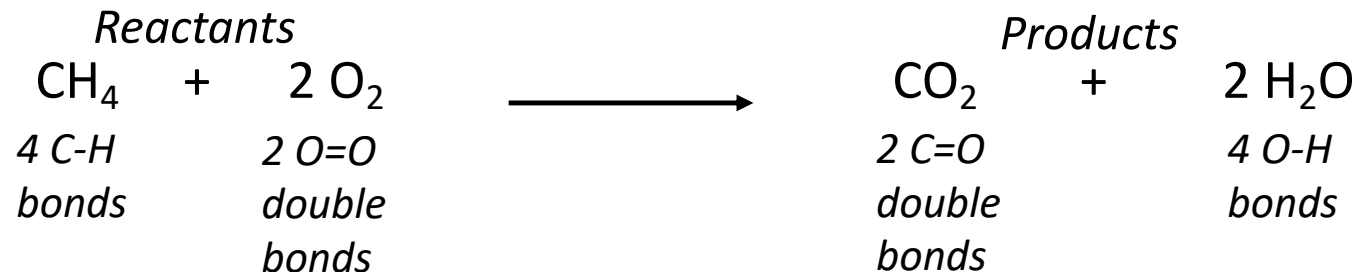
Net energy release (ΔH) =

Energy to form the bonds of products + Energy to break the bonds of reactants

\swarrow < 0
 \longrightarrow = - "bond energy"

\swarrow > 0
 \longrightarrow = "bond energy"

Example : combustion of methane



$$\Delta H = - 2 \times 804 - 4 \times 460 + 4 \times 413 + 2 \times 497 = - 802 \text{ kJ/mol} = - 50.1 \text{ kJ/g}$$

\uparrow
C=O

\uparrow
O-H

\uparrow
C-H

\uparrow
O=O

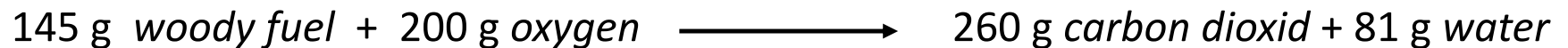
Another way to compute the heat of combustion is to use the enthalpies of formation of products and reactants.

Combustion heat release

The heat of combustion of woody fuels can be computed considering the following reaction, where $C_6H_9O_4$ represents an average composition of the dry fuel :



In mass:



High heat of combustion : $\Delta H_{fuel} = 20.1 \text{ kJ/g}$, when the final state of *water* is liquid

Low heat of combustion : $\Delta H_{fuel} = 20.1 - 1.4 = 18.7 \text{ kJ/g}$, when the final state of *water* is vapor

Incomplete combustion releases less heat.

But it is better to measure the heat of combustion, as the exact composition of the material is usually unknown

Combustion heat release

The *heat of combustion* of a burning material depends on the material (composition) and on the fire conditions (temperature, oxygen supply, water content).

Indeed, both influence the proportion of pyrolysis products, which have different heats of combustion

Pyrolysis product	High heat of combustion ΔH_H (kJ/g)
Chars (C)	32
CO	10
CH ₄	50
Levoglucosan	17

Combustion heat release

The *heat of combustion* of woody fuels varies with their composition and the proportion of char produced (data from Rothermel 1976)

Substance	Proportion of char produced	Higher heat of combustion $\Delta H H$ (kJ/g)
Cellulose and hemicellulose (50-75%)	0.092*	16.1
Lignin (15-35%)	0.624	24.5
Extractives (0.2-15%)	0.285	32.3

*the amount of char produced increases when silica-free minerals are present.

Extractives include volatile organic compounds, which are highly flammable substances

Minerals tend to inhibit flaming combustion, promoting char formation.

Combustion heat release

Combustion characteristics of different fuel elements (from Susott 1982)

Fuel type	Ash content (%)	Fraction of char (%)	Higher heat of combustion (kJ/g)	Energy for volatiles (kJ/g)	Energy for char (kJ/g)
Grasses	6.5-9.5	22-25	19.4-20.2	12.0-12.2	7.1-8.2
Foliage	1.5-7.1	25-34	20.6-23.3	10.9-15.8	7.5-10.6
(Small) Stems	2.2-6.1	22-28	20.0-22.4	10.9-15.2	7.2-9.1
Wood	0.2-0.6	15-24	19.6-21.0	12.6-14.6	5.0-7.6
Rotten wood	0.2-0.2	21-41	20.3-23.1	10.4-13.6	6.8-12.6
Bark	0.5-17.7	28-47	21.5-24.0	7.7-12.8	8.9-14.3
Duff	31.2-34.1	35-39	20.3-23.3	8.9-11.1	11.4-12.2

Note that the heat of combustion is not so variable

Combustion heat release

Combustion characteristics of
Mediterranean fuels (from Madrigal
et al 2011)

Measurements :

GHC (MJ/kg) : Gross heat of combustion (high heat of
combustion measured with bomb calorimeter)

PHRR (kW/m²) : peak heat release rate

THR (MJ/m²) : total heat release

TTI (s) : time to ignition of the sample

FD (s) : Flaming duration

MLR (g/s) : peak mass loss rate

Note :

The initial sample load is 10g/100 cm², i.e. 1kg/m².

Thus, THR can also be expressed as MJ/kg and
compared with GHC.

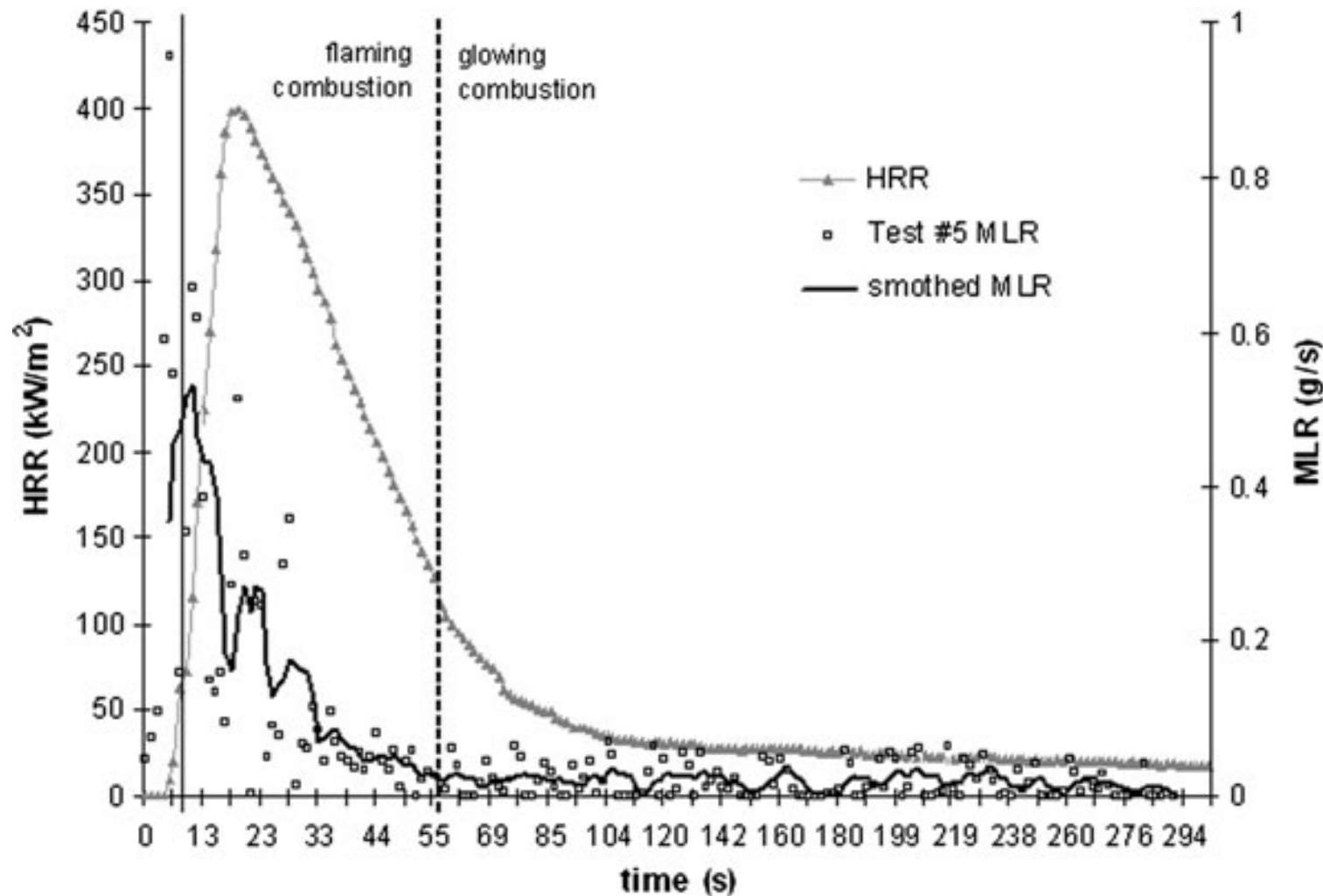


Figure 1. a General view of the MLC device. b Detail of the thermopile. c Detail of methane burner used to calibrate thermopiles. d Porous holder with *Pinus pinaster* dead litter, immediately before a test.

Combustion heat release

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Combustion heat release

Combustion characteristics of Mediterranean fuels (from Madrigal et al 2011)

Species	Growth form	Plant parts collected	Bomb calorimeter GHC (MJ/kg)	Mass loss calorimeter				
				PHRR (kW/m ²)	THR (MJ/m ²)	TTI (s)	FD (s)	MLR (g/s)
Live fuels								
<i>Aparagus acutifolius</i>	Herb, perennial	Green stalks and leaves	20.25	326	18.54	3	57	0.056
<i>Cistus ladanifer</i>	Shrub	Green twigs and leaves	21.24	390	23.93	12	65	0.054
<i>Cistus laurifolius</i>	Shrub	Green twigs and leaves	21.44	303	19.48	5	68	0.06
<i>Crataegus monogyna</i>	Tree, shrub	Green twigs and leaves	19.96	284	16.84	3	29	0.06
<i>Cynodon dactylon</i>	Graminoid, annual	Green leaves	17.66	238	14.47	2	40	0.059
<i>Cytisus scoparius</i>	Shrub	Green twigs and leaves	21.05	426	21.85	12	49	0.046
<i>Daphne gnidium</i>	Shrub	Green twigs and leaves	19.93	353	25.87	3	82	0.061
<i>Erica arborea</i>	Shrub	Green twigs and leaves	22.86	359	19.19	7	47	0.06
<i>Eucalyptus camaldulensis</i>	Tree	Green twigs and leaves	19.87	349	20.01	2	46	0.059
<i>Eucalyptus globulus</i>	Tree	Green twigs and leaves	22.42	397	20.33	4	48	0.06
<i>Juniperus oxycedrus</i>	Tree, shrub	Green twigs and leaves	20.34	367	19.63	4	57	0.063
<i>Lavandula stoechas</i>	Shrub	Green twigs and leaves	20.99	343	17.81	4	62	0.06
<i>Quercus coccifera</i>	Tree, shrub	Green twigs and leaves	19.95	353	17.73	11	38	0.062
<i>Quercus ilex</i>	Tree, shrub	Green twigs and leaves	19.81	438	23.94	17	35	0.061
<i>Rubus ulmifolius</i>	Shrub	Green twigs and leaves	19.14	327	17.82	11	64	0.058
<i>Ulex europaeus</i>	Shrub	Green twigs and leaves	21.43	507	25.98	11	94	0.059
<i>Pinus halepensis</i>	Tree	Green twigs and leaves	21.34	356	23.18	4	65	0.062
<i>Pinus pinaster</i>	Tree	Green twigs and leaves	21.37	389	15.81	13	61	0.049
<i>Pinus pinea</i>	Tree	Green twigs and leaves	20.37	331	12.81	9	81	0.063
<i>Pinus sylvestris</i>	Tree	Green twigs and leaves	21.68	375	20.39	6	62	0.06
Dead fuels								
<i>Brachypodium retusum</i>	Herb, perennial	Dead stalks and leaves	19.59	264	16.86	3	98	0.065
<i>Pinus halepensis</i>	Tree	Needle litter	22.5	495	25.8	4	60	0.072
<i>Pinus pinaster</i>	Tree	Needle litter	21.54	472	26.07	8	53	0.064
<i>Pinus pinea</i>	Tree	Needle litter	20.47	375	19.65	6	68	0.064
<i>Ulex europaeus</i>	Shrub	Twigs and leaves litter	20.23	260	16.89	4	54	0.057
<i>Ulex europaeus</i>	Shrub	Aerial dead twigs and leaves	22.09	512	27.39	8	98	0.065

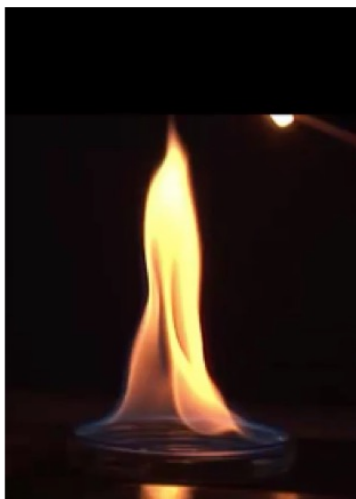
Flame - mixing

The locations where the combustion of volatiles occurs (reaction zone)

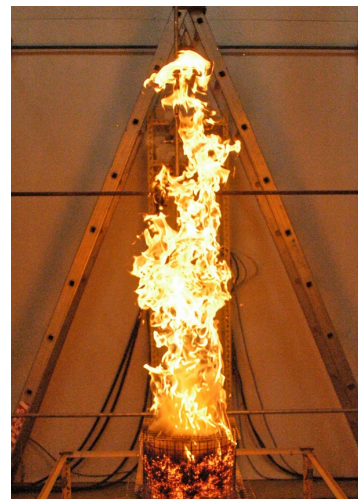
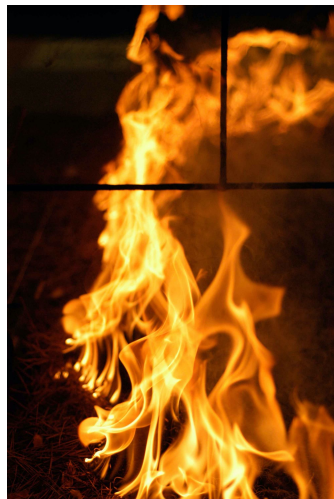
It is visible in fires thanks to the formation of soot particles radiating in the visible wavelengths (yellow color) at high temperatures

Flames in fires are diffusion flames, i.e. mixing of oxygen and flammable gas realized thanks to molecular and turbulent diffusion

Turbulence increase



Laminar



Highly turbulent 28

Flame - mixing

Molecular diffusion : driven by concentration gradients of oxygen and gaseous fuel

Turbulent diffusion : driven by eddy chaotic motion

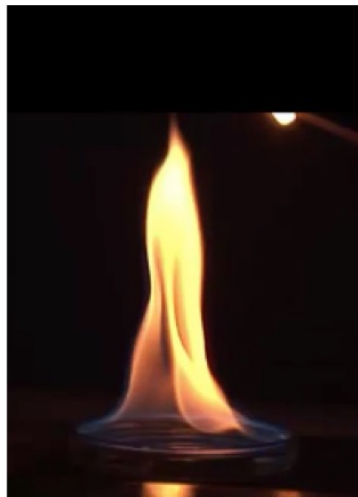
-> much more efficient for mixing than molecular diffusion

Mixing (which ensures oxygen supply) is fundamental in wildland or natural fires, as it controls the rate of combustion (the speed at which fuel or O₂ is consumed)

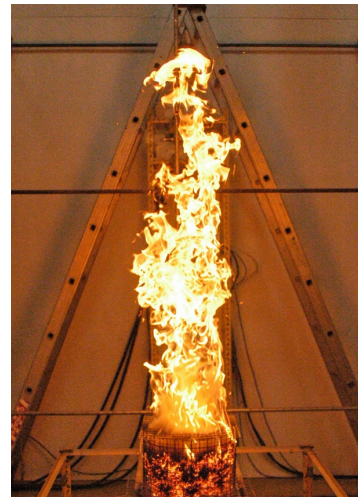
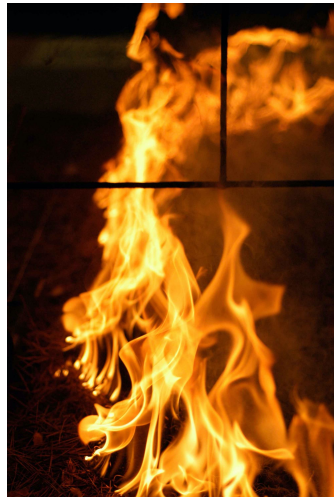
Only
molecular diffusion

Turbulence increase

Dominated by
turbulent diffusion



Laminar



Highly turbulent 29

Flame - temperature

Theoretical (adiabatic) flame temperatures are above 2000°C

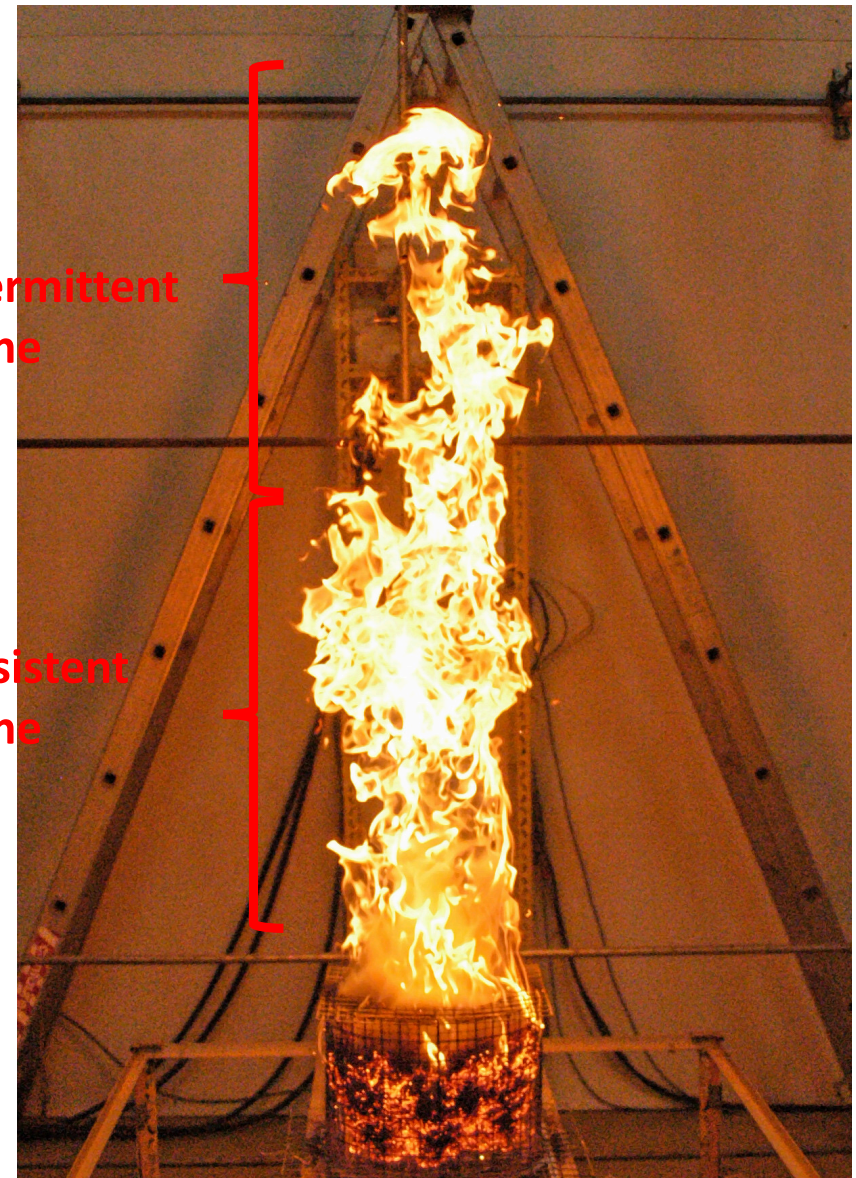
But actual temperature are much lower due to heat losses :

typically 800- 1000°C in average in the core of the flames

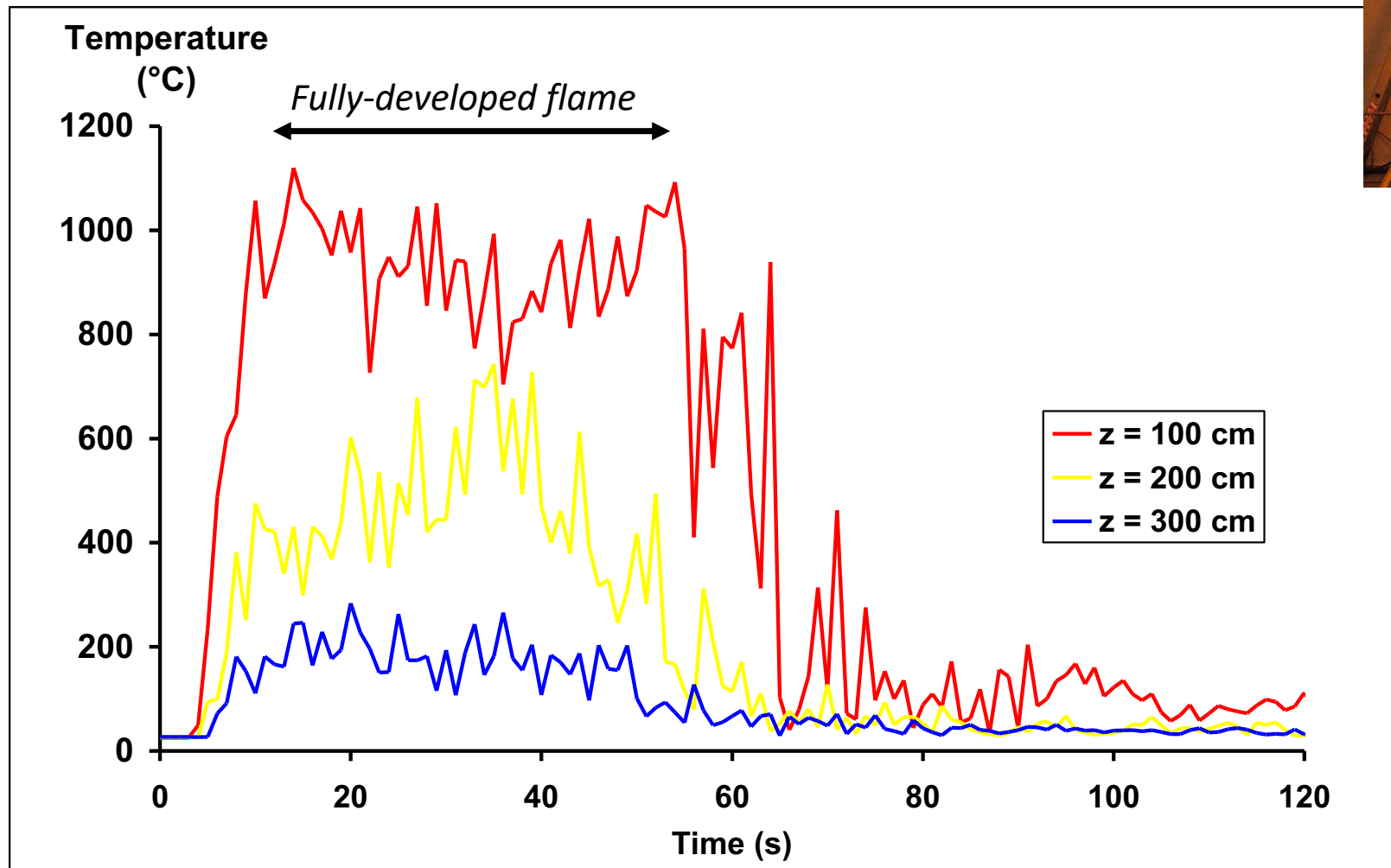
$$Z = \frac{z}{H}$$

Local, instantaneous values fluctuate a lot

Average temperature decreases above the persistent flame, and radially

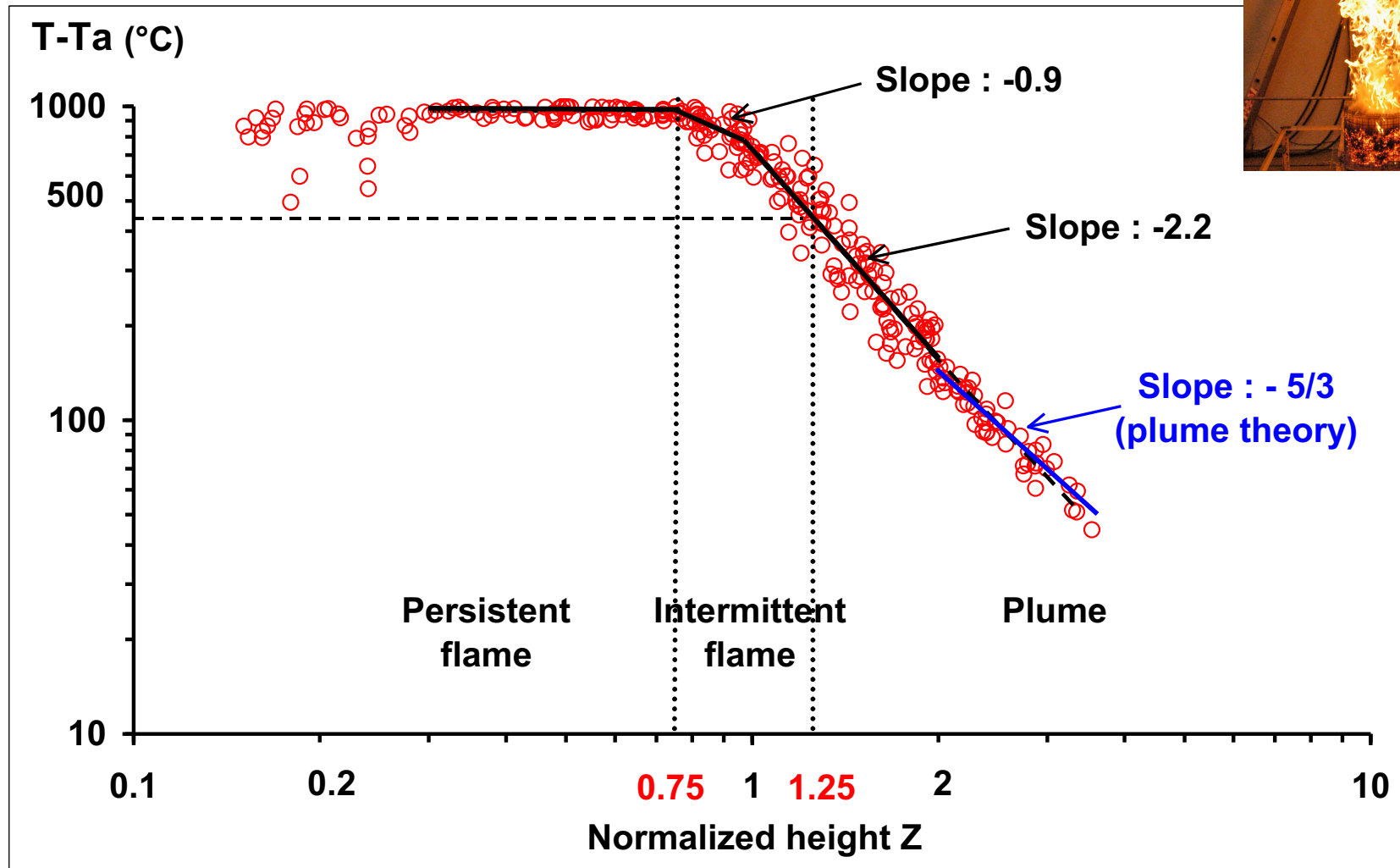
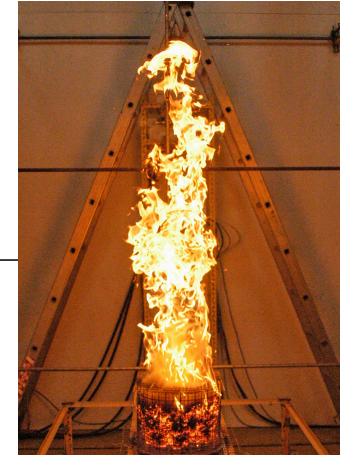


Flame - temperature



Flame - temperature

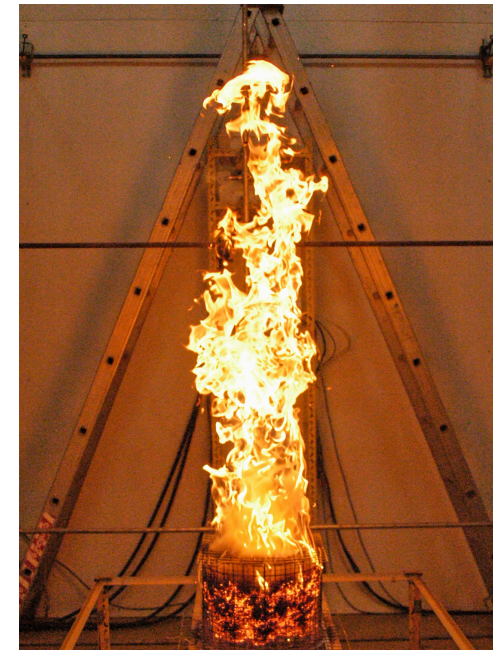
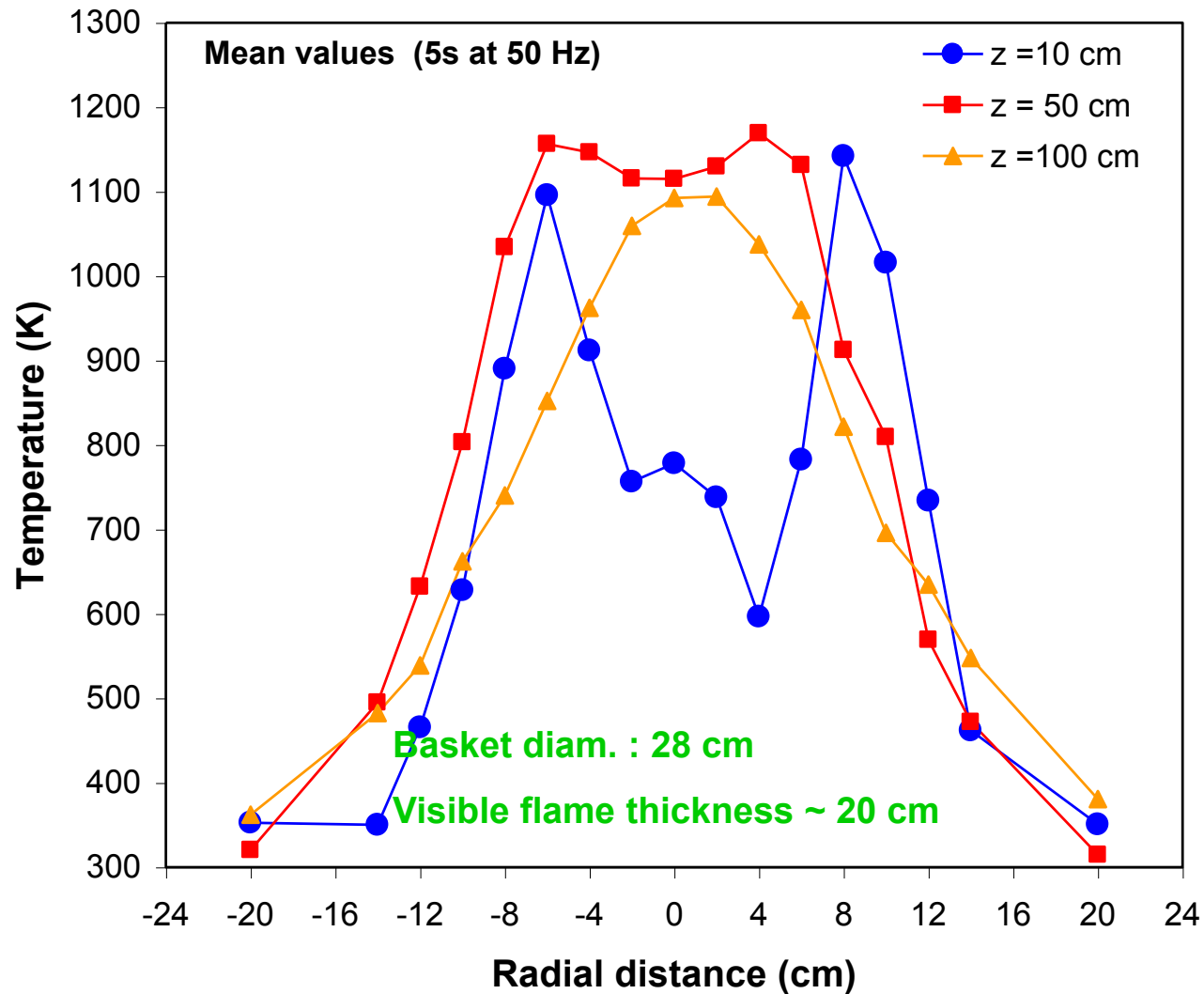
Vertical profile of time-averaged temperature



$$Z = \frac{z}{H} \quad H \text{ is mean flame height}$$

Flame - temperature

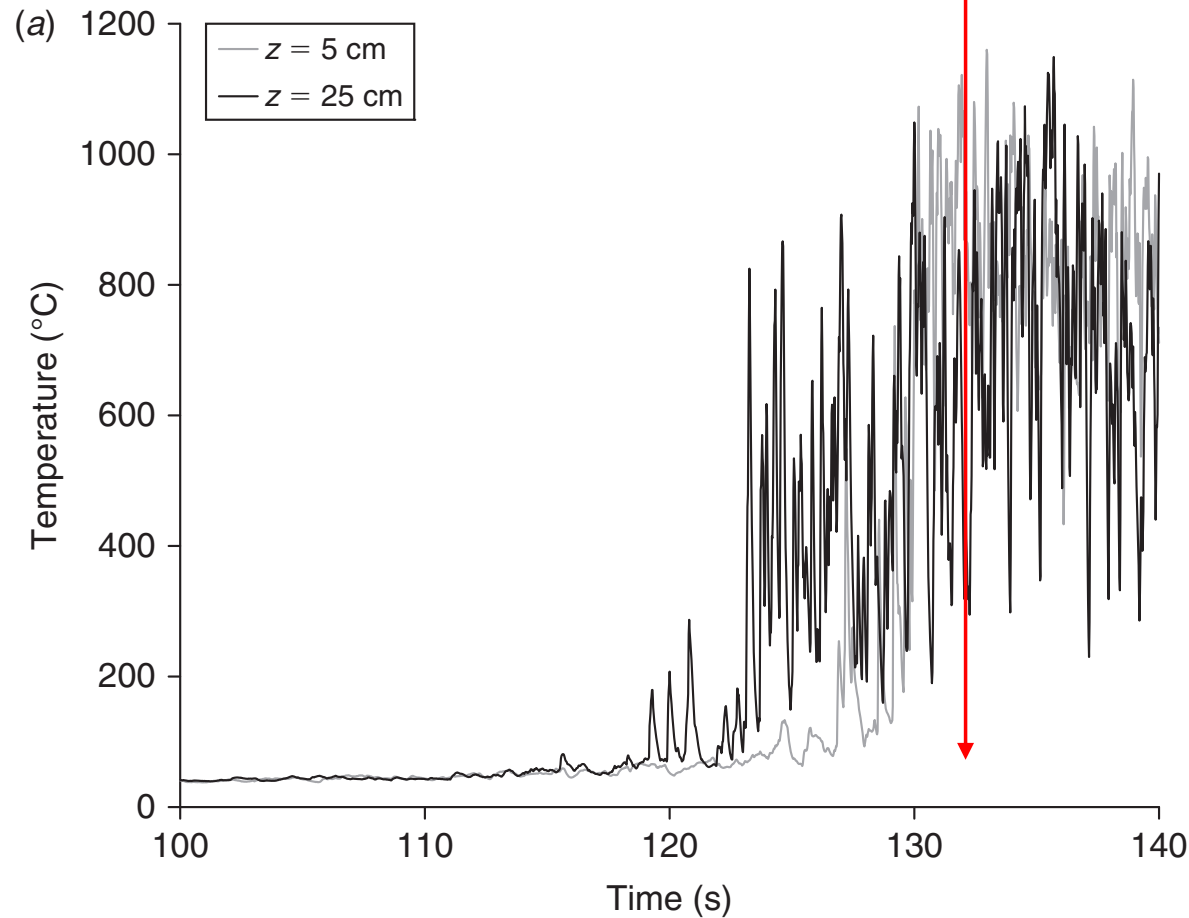
Radial profile of time averaged temperature



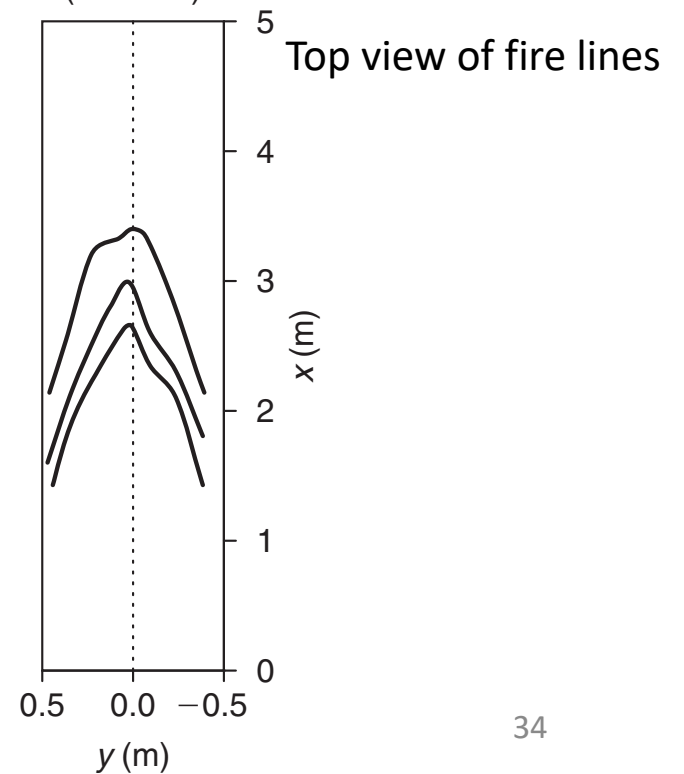
Flame - temperature

Temperature recordings in a spreading fire

Head fire (flame) reaches the thermocouple



30°, 1 m
(test #44)

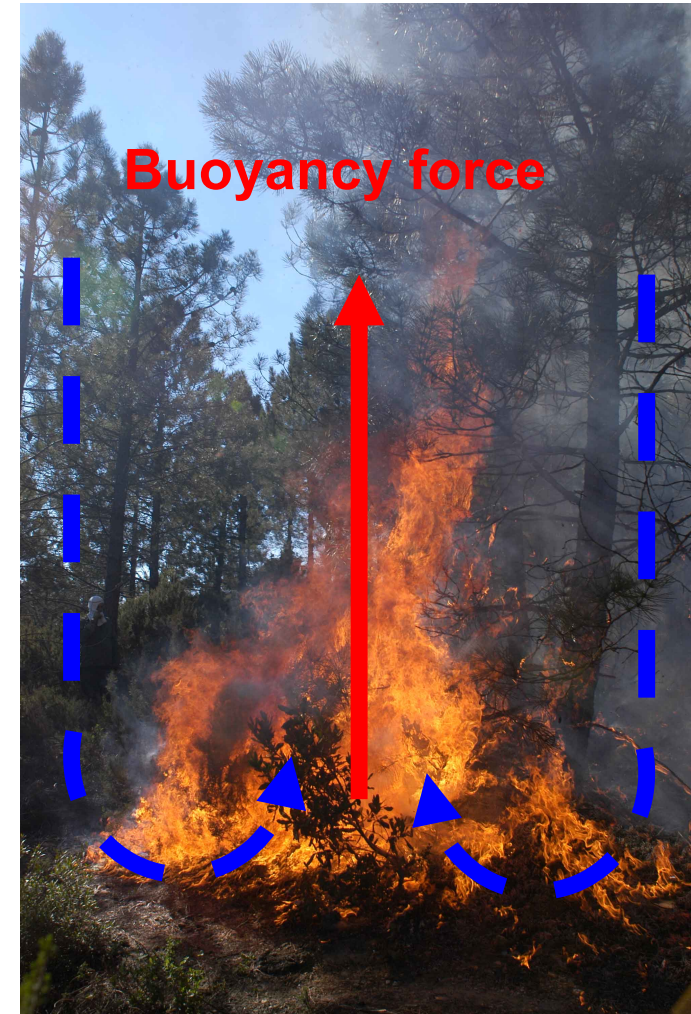


Buoyancy and fire-induced flow

Hot air is much less dense than fresh air (ambient a), generating a buoyancy force :

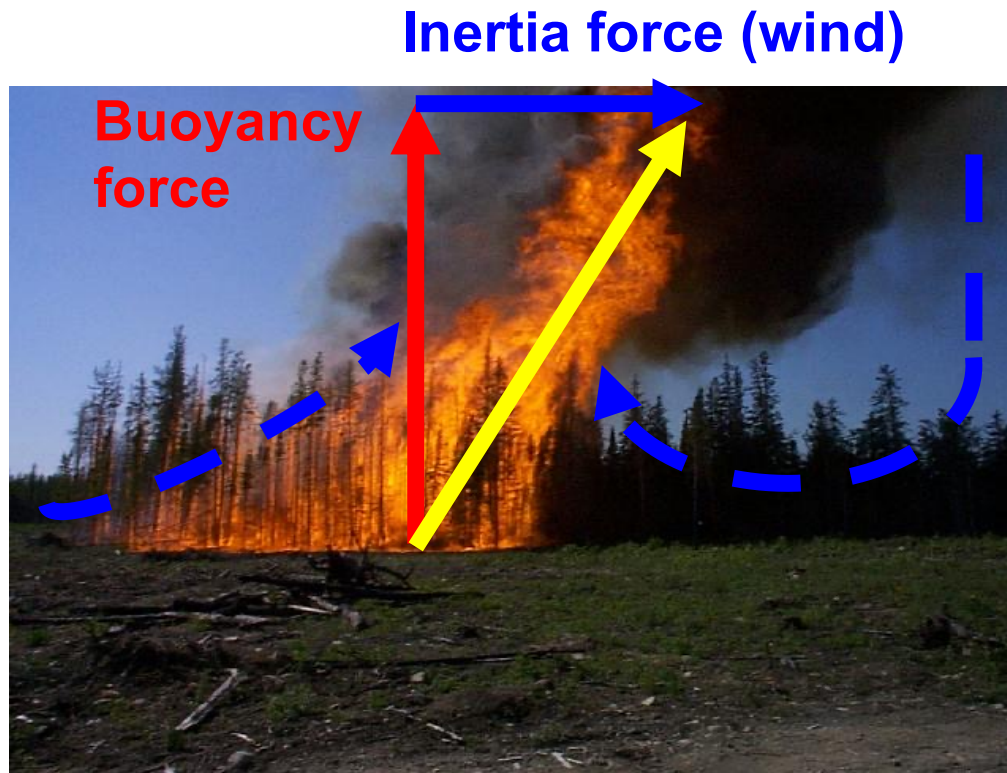
$F = g (\rho_a - \rho)$, g is gravity and ρ is density
(F expressed as forced per unit volume of air)

Rising hot air and gas must be replaced by fresh air

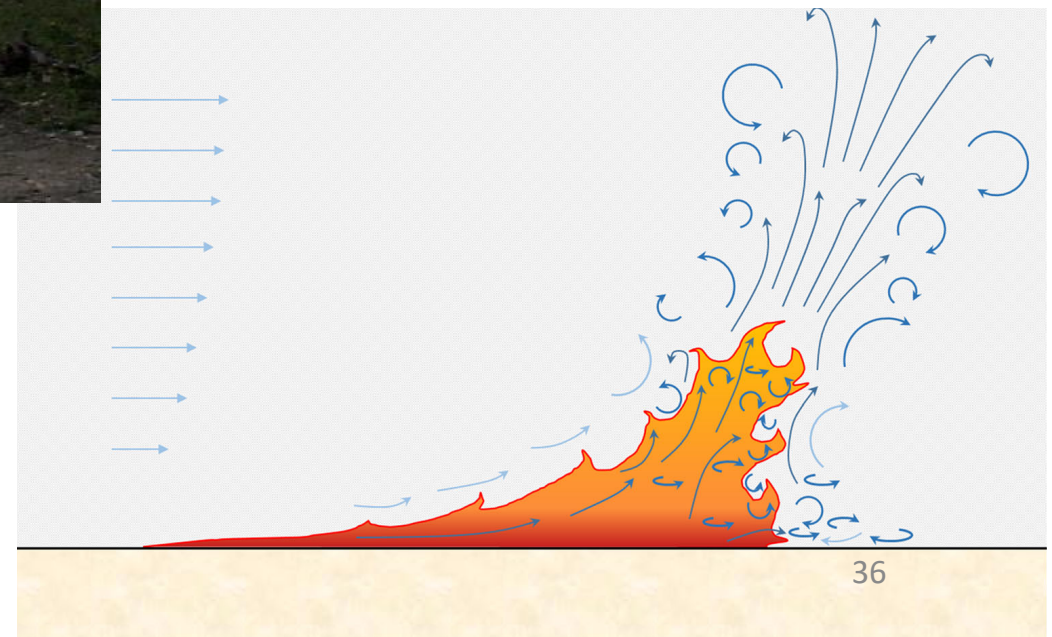


Fire-induced downdraft

Buoyancy and wind interactions



Fire-induced drafts

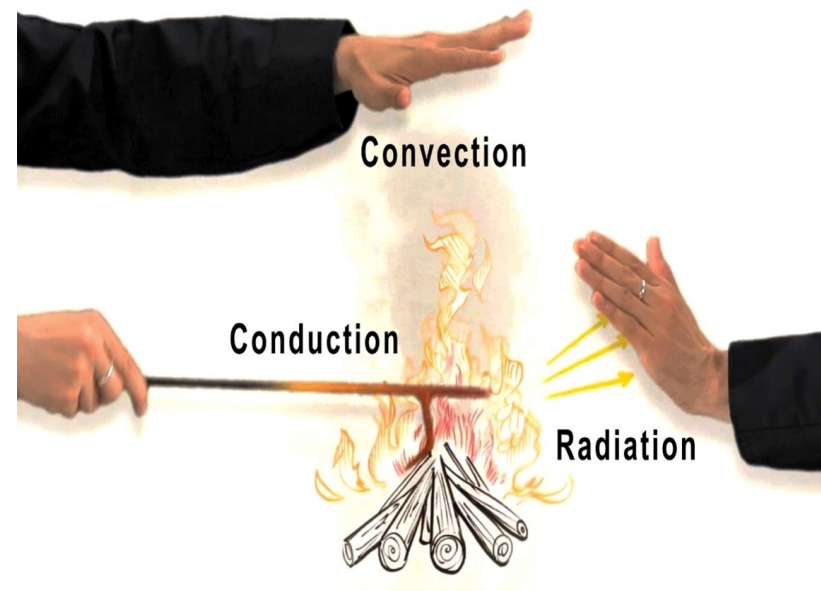


Heat transfer

The process by which heat (thermal energy) is exchanged between two objects or two parcels of matter of different temperatures

3 modes :

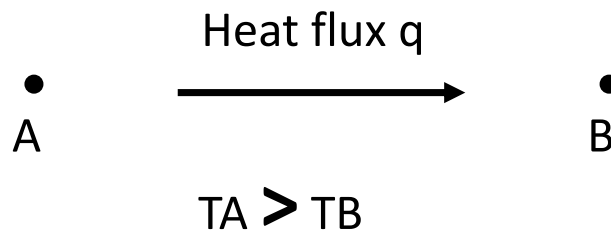
- conduction
- thermal radiation
- convection



Conduction

Conduction is the transfer of heat from one molecule of matter to another by direct contact

In fires : relatively slow transfer, operating within solid fuels and soil layers



Fourier law

$$q = -\lambda \frac{\partial T}{\partial x} \quad (\text{kW/m}^2)$$

Thermal conductivity λ :

- copper : 400 W/m K
- dry wood : 0.1-0.2 W / m K
- dry air : 0.02-0.03 W / m K

Thermal radiation

Thermal radiation is the transfer of heat energy by electromagnetic waves from a heat source to an absorbing material

Semi-transparent bodies (the air) allow the waves to pass, so the transfer

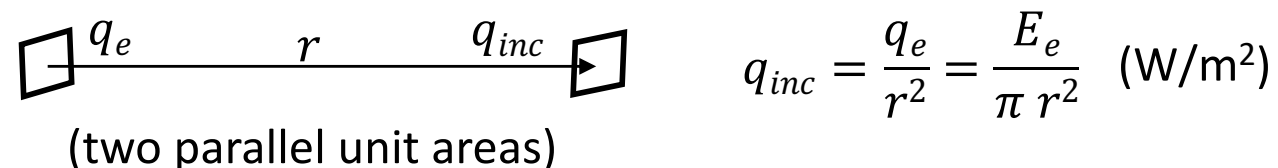
Any body emits radiant energy due to its temperature T (Stefan-Boltzman law):

$$E_e = \varepsilon \sigma T^4 \quad (\text{Emissive power of grey bodies, W/m}^2)$$

ε is body emissivity (=1 for a black body, 0 for a non emitting surface)

σ the Stefan-Boltzman constant

The heat flux decreases with the square of the distance to the source



(two parallel unit areas)

$$q_{inc} = \frac{q_e}{r^2} = \frac{E_e}{\pi r^2} \quad (\text{W/m}^2)$$

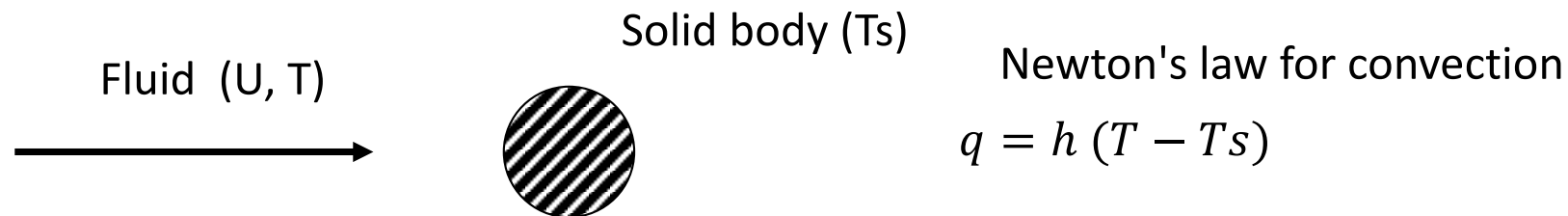
Any non-transparent body absorbs incident radiant energy according to

$$q_a = \alpha q_{inc} \quad (\text{absorptivity } \alpha = \varepsilon \text{ for diffuse - grey bodies})$$

Convection (convective heat transfer)

Convection is the transfer of heat resulting from the motion of a fluid.
In fires : the fluid is air or hot gases (flame, smoke).

The fluid transports heat and exchanges it (by conduction) with the solid body, namely, in fires, the fuel elements



The convective heat transfer coefficient h depends on U (fluid velocity), T (fluid temperature), and other fluid properties, as well as on body geometry

This is a very efficient way of heating, especially when flame is in contact with fuel.

Note : the term *convection* is also widely used to designate the natural movement of fluid due to fluid density differences, for example above a fire ...

Dominant heat transfer in horizontal fire spread

Head Fire

Wind and/or slope affect fire spread with radiant and convective heat.

Backing Fire

Conduction/radiation within fuel bed is dominant factor in fire spread. Much less dependent on wind and slope.

Wind



Dominant heat transfer in vertical fire spread

Convection above the fire heats and may ignite tree foliage



Dominant heat transfer in glowing embers

Radiation, and conduction within charring material

