







Module 5: Premixed Flame

Lecture 25: Effect of Oxygen Concentration on S_L

The Lecture Contains:

-  [Effect of Oxygen Concentration on \$S_L\$](#)
-  [Effect of Initial Pressure and Temperature on \$S_L\$](#)
-  [Effect of Inert Additives](#)
-  [Flame Extinction](#)
-  [Flame Quenching](#)
-  [Simplified Analysis for Quenching Diameter](#)

 **Previous** **Next** 

Module 5: Premixed Flame

Lecture 25: Effect of Oxygen Concentration on S_L Effect of Oxygen Concentration on S_L

For various combinations of O_2 and N_2 , there is a drastic increase in S_L .

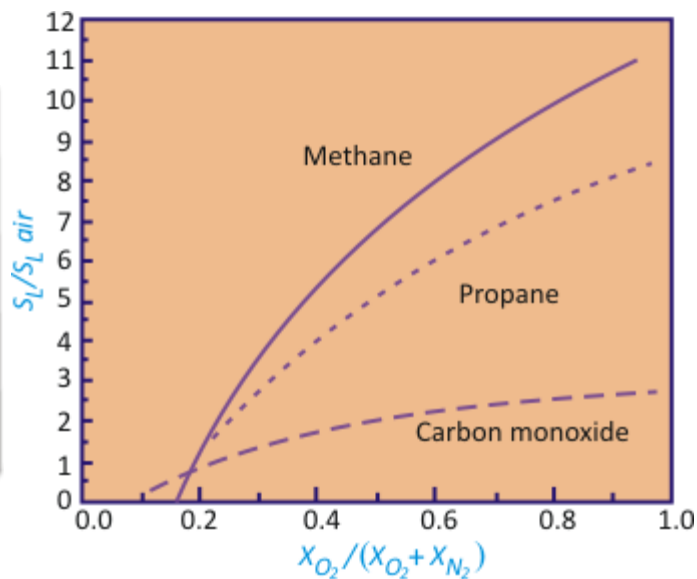
For methane, S_L is increased by 10 times

For propane, S_L is increased by 7.5 times

For CO, S_L is increased by 2.4 times.

What is the reason for this change?

Higher the oxygen level, higher will be the adiabatic flame temperature
Thus higher burning velocity



(Figure 25.1)

◀ Previous Next ▶

Module 5: Premixed Flame

Lecture 25: Effect of Oxygen Concentration on S_L Effect of Initial Pressure and Temperature on S_L

$$S_L \propto p^{(n-2)/2}$$

This relation is supported by experimental data

n : Overall order of global chemical reaction

$n < 2$ for HC flames with $S_L < 50 \text{ cm/s}$

$n = 2$ for HC flames with

$S_L \approx 50 - 100 \text{ cm/s}$

$n > 2$ for HC flames with

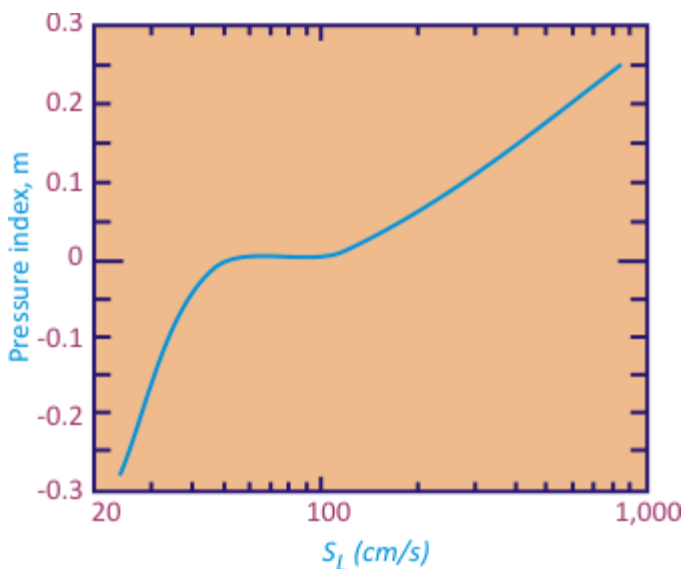
$S_L > 100 \text{ cm/s}$

Pressure index, $m = (n - 2)/2$

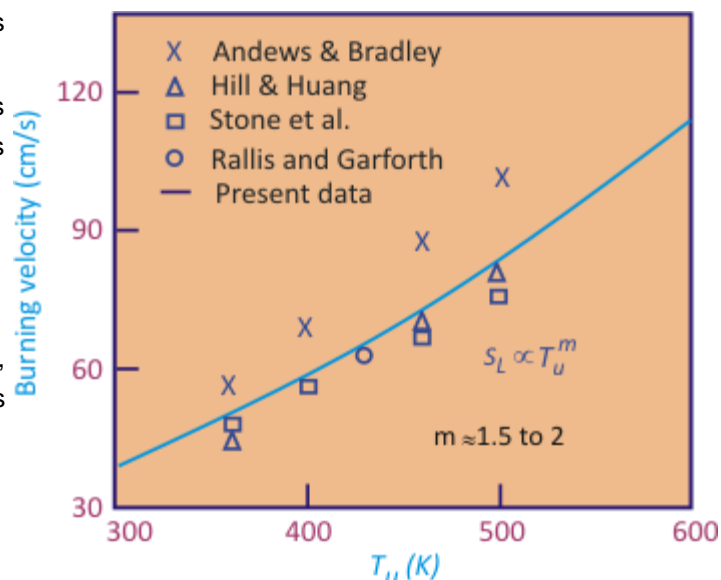
For $S_L < 50 \text{ cm/s}$, m is negative, indicating burning velocity increases with decreasing pressure

For $S_L \approx 50 - 100 \text{ cm/s}$, m is constant, indicating burning velocity is constant

For $S_L > 100 \text{ cm/s}$, m is positive, indicating burning velocity decreases with decrease in initial pressure



(Figure 25.2)



(Figure 25.3)

◀ Previous Next ▶

Module 5: Premixed Flame

Lecture 25: Effect of Oxygen Concentration on S_L

Effect of Inert Additives

Inert gas additives produce the following effects

1. Reduction of burning velocity
2. Narrowing of flammability limits
3. Shifting of S_L peak

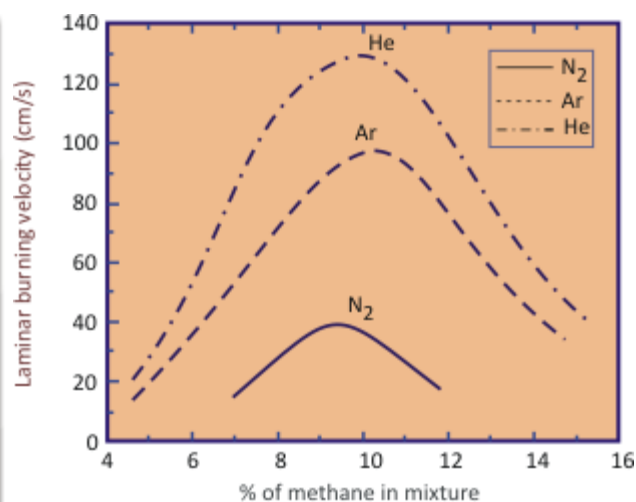
Why there is a shift in S_L peak ?

Addition of inert gas changes the ratio of thermal conductivity and sp. heat of the mixture, resulting in change in S_L .

Burning velocity with He addition is higher than Ar and N_2 . Why ?

Lower MW will lead to higher burning velocity
MW of He is much lower than Ar and hence higher S_L

Ar and N_2 have almost same thermal diffusivity
Hence for same heat release, Ar attains higher flame temperature \rightarrow higher S_L .



Clingman et al 1995

(Figure 25.4)

◀ Previous Next ▶

Module 5: Premixed Flame

Lecture 25: Effect of Oxygen Concentration on S_L

Flame Extinction

◀◀ Previous Next ▶▶

Flame Quenching

Fuel	Oxidizer	SL (cm/s)	dq (mm)
CH ₄	Air	40	2.5
CH ₄	O ₂		0.3
C ₃ H ₈	Air	45	2.0
C ₃ H ₈	O ₂		0.25
C ₂ H ₂	Air	140	0.8
C ₂ H ₂	O ₂		0.2
CO	Air		2.8
H ₂	Air	210	0.5
H ₂	O ₂		0.2

Quenching diameter for various stoichiometric fuel-oxidizer ratio.

Module 5: Premixed Flame

Lecture 25: Effect of Oxygen Concentration on S_L

Simplified Analysis for Quenching Diameter

Condition for flame propagation

Heat generated due to chemical reaction >
heat lost due to conduction

Rate of heat generated per unit volume

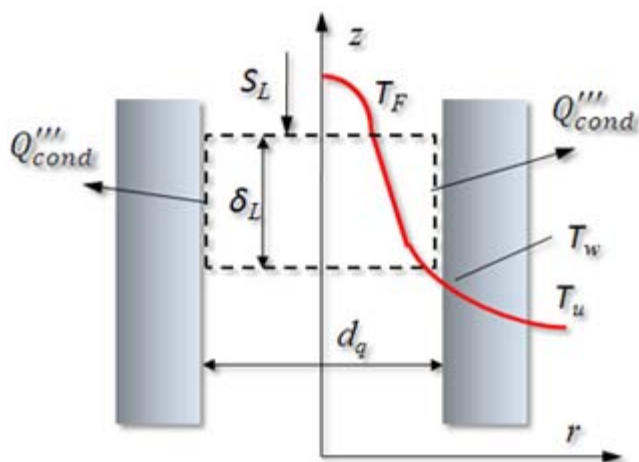
$$\dot{Q}''' = \bar{m}_F''' \Delta \hat{H}_c = \dot{Q}_{cond}$$

Heat generated in flame volume

$$\dot{Q} = \bar{m}_F''' \Delta \hat{H}_c \frac{\pi}{4} d_q^2 \delta_L$$

Heat loss rate due to wall conduction

$$\dot{Q}_{cond} = k_g \pi d_q \delta_L \frac{dT}{dr}$$



(Figure 25.5)

Assuming linear temperature distribution in flame,

$$\frac{dT}{dr} = \frac{T_F - T_u}{d_q/C}$$

Quenching diameter

$$d_q = \sqrt{\frac{4Ck_g}{\bar{m}_F''' \Delta \hat{H}_c} \frac{T_F - T_u}{C}}$$

After simplification,

$$d_q = \sqrt{8C} \cdot \delta_L$$

◀ Previous Next ▶