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# NUMERICAL STUDY OF TURBULENT NORMAL DIFFUSION FLAME CH<sub>4</sub>-AIR STABILIZED BY COAXIAL BURNER

#### by

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The practical combustion systems such as combustion furnaces, gas turbine, engines, etc., employ non-premixed combustion due to its better flame stability, safety, and wide operating range as compared to premixed combustion. The present numerical study characterizes the turbulent flame of methane-air in a coaxial burner in order to determine the effect of airflow on the distribution of temperature, on gas consumption, and on the emission of  $NO_x$ . The results in this study are obtained by simulation on FLUENT code. The results demonstrate the influence of different parameters on the flame structure, temperature distribution and gas emissions, such as turbulence, fuel jet velocity, air jet velocity, equivalence ratio, and mixture fraction. The lift-off height for a fixed fuel jet velocity is observed to increase monotonically with air jet velocity. Temperature and  $NO_x$  emission decrease of important values with the equivalence ratio, it is maximum about the unity.

Key words: non-premixed combustion, coaxial burner, turbulent flame, lift-off height, equivalence ratio, mixture fraction, NO<sub>x</sub> emission

## Introduction

Generally, there are three main types of the flame: the premixed flame, diffusion flame (non-premixed flame), and partially premixed flame. The diffusion flame used in the most industrial applications, such as industrial furnaces, diesel engines, aeronautics (rocket engine), *etc.* due to its better flame stability and safety as compared to premixed combustion. In this process fuel and oxidizer are injected into the flame by two streams, one for gas other for air or oxygen. The two different configurations of the diffusion flame are: normal diffusion flame (NDF), in a simple coaxial burner, the central gas jet enveloped by annular air jet. Inverse diffusion flame (IDF) is a kind of flame with an inner air jet is surrounded by outer fuel jet. The research of non-premixed combustion started in 1928 by Burke and Schumann on the flame properties such flame length, lift-off from the burner, temperature, and mass fractions of combustion products. These properties depend upon mixture fraction. The principal objectives of the research in the field of non-premixed combustion are: reducing atmospheric pollution and maintain a more stable flame, therefore a good performance of combustion are: the reduction of atmospheric pollution and the improvement of flame stability, therefore a good performance of combustion system.

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Several researchers reported the effect of air-fuel velocity ratio on the characteristics of NDF and IDF. The first detailed investigation was performed by Mahesh and Mishra [1, 2] for experimental study characterizing the turbulent liquefied petroleum gas (LPG) for inverse diffusion flame stabilized in a back-step burner in terms of visible flame length, dual flame structure, centerline temperature distribution and oxygen concentration. Sze, et al. [3] carried out experiments on the appearance, temperature distribution, and NO<sub>x</sub> emission of two inverse diffusion flames, one with circumferentially arranged ports and the other with co-axial jets, both burning liquefied petroleum gas. Sobiesiak and Wenzell [4] reported the effect of air and fuel jet diameters and the ratio of inner air jet and outer fuel jet velocities on temperature distribution, flame length, and stability limits on inverse diffusion flame. Fernandez et al. [5] carried the numerical study of Liftoff and blow out of methane jet diffusion flame between parallel streams of methane diluted with nitrogen and air. Santos and Costa [6] characterized jet diffusion flames for three fuels such as methane, ethylene, and propane gases experimentally in terms of flame lift-off height, flame length, and NO, emission. Meunier et al. [7] investigated on NO, emissions from turbulent propane diffusion flames. The study includes both experimental measurements and numerical 2-D simulations and aims to provide a better understanding of the dominant physical effects associated with the NO, production in turbulent diffusion flames. Montgomery et al. [8] reported the effect of co-flow velocity on a lifted methane-air jet diffusion flame. The lift-off height increases with jet exit velocity and with the air co-flow velocity. Many researchers [9-14] carried experimental and numerical out the effect of hydrogen jet on stability, distribution of temperature, and NO<sub>x</sub> emission. Oh et al. [9, 13] investigated the effect of nitrogen dilution and acoustic excitation on the stabilization and NO<sub>y</sub> emission of turbulent diffusion flames. They concluded that the turbulent flame propagation velocity increased as the mole fraction of nitrogen diluents gas decreased. However, the dilution of nitrogen is characterized by lower mixture fraction and reduced burning velocity. This premixing effect resulted in a lower flame temperature and lower NO, production. Flame length decreased with acoustic excitation, NO, emission was reduced and minimized at a resonance frequency. Mishra and Kumar [10, 11] investigated the effect of H<sub>2</sub> addition on the flame length, soot free length fraction, radiant fraction, gas temperature, and emission NOx in the LPG composed of 69% C3H8 and 30% C4H10 jet diffusion flame. El-Ghafour [12] performed an experimental investigation on turbulent natural gas-hydrogen jet diffusion flame and observed the effect of hydrogen addition on flame stability, flame length, flame structure, exhaust species concentration and pollutant emissions. Other experimental studies [15-17] on the characteristics of high temperature diluted air combustion showed that it yields very low NO<sub>x</sub> emission, low luminosity, higher flame stability, large flame volume and a more uniform temperature. Several studies reported the control of the flame by inclined oxygen jets [18-20]. This idea enhanced the premixed fuel-oxidant, therefore reduces emission of NO<sub>r</sub>.

In this context, the present work provides a numerical investigation of the turbulent methane-air jet in a coaxial burner. The aim is to examinine the effects of air jet velocity on the flame stability, distribution of temperature, and gas emission with fixed fuel jet velocity.

## Numerical modeling

## Mathematical formulation

The numerical model solves the steady equations for conservation of mass, momentum, energy, and species for a coaxial burner in ambient air. The inner fuel stream injected the pure methane, is introduced into the flow field through the nozzle with diameter of 6 mm and with fixed

mean velocity of 10 m/s. This leads to a fuel exit Reynolds number of 5217. The stoichiometric mixture fraction  $\xi$  is 0.055. Through a surrounding nozzle having diameter of 18 mm, air is injected with different velocities, fig. 1. The modelisa- tion of this configuration is obtained by FLUENT. We used the *k*- $\varepsilon$  turbulent model. The calculations have been performed in axisymmetric co-ordinates (x, y), 250 mm for x and 1000 mm for y.

The numerical model solves the independent time equations for conservation of mass, momentum, energy, and individual species:

$$\nabla(\rho \mathbf{V}) = 0$$

$$\vec{\nabla}(\rho \vec{V} \vec{V}) = -\nabla P + \nabla(\tau) + \rho \vec{g}$$
(2)

$$\rho \frac{\partial h}{\partial t} + \rho \vec{\mathbf{V}} \nabla h = \frac{\partial P}{\partial t} + \vec{\mathbf{V}} \nabla P - \nabla j_{\mathbf{q}} + q_{\mathbf{r}}$$
(3)





$$\rho V \nabla Y_{t} - \nabla \rho D \nabla Y_{t} = \omega_{t} \tag{4}$$

(1)

$$\xi = \frac{m_1}{\dot{m}_1 + \dot{m}_2} \tag{5}$$

Here, the stress tensor, heat flux  $j_q$  and diffusive flux  $j_i$  are defined as:

$$\boldsymbol{\tau} = \boldsymbol{\mu} [ (\nabla \vec{\mathbf{V}} + \nabla \vec{\mathbf{V}}^T) - \frac{2}{3} \nabla \vec{\mathbf{V}} I ]$$
(6)

$$j_{q} = -\lambda \nabla T + \sum_{i=1}^{n} h_{i} j_{i}$$
<sup>(7)</sup>

$$j_i = -\rho D_i \nabla Y_i \tag{8}$$

The heat radiation  $q_r$  is defined by:

$$q_r = -4\sigma \sum_{i=1}^{n} (p_i a_{p,i}) (T^4 - T_{amb}^4)$$
(9)

where  $\sigma$  is the Stefan-Boltzmann constant and  $p_i$  the partial pressure of species *i*,  $a_{p,i}$  – the Plank mean absorption coefficients for radiating species *i* [21], *P* – the static pressure, g – the term of gravity,  $Y_i$  – the mass fraction,  $D_i$  – the thermal diffusivity, *T* – the temperature,  $\rho$  – the density, *h* – the specific enthalpy,  $\omega_i$  – the chemical production rate,  $m_i$  – the mass flux of fuel and air, and  $\xi$  – the mixture fraction.

To complete the problem formulation, one needs to provide an expression for the methane consumption rate. Here, we shall consider an overall irreversible reaction where the gas reacts with the air to produce combustion products and heat according to:

$$CH_4 + 9.5238(0.21O_2 + 0.79N_2) \rightarrow CO_2 + 2H_2O + 7.5238N_2 + Q$$
(10)

## Turbulence modeling

The *k*- $\varepsilon$  turbulence model presented by Launder and Spalding [22] is used to model the turbulent kinetic energy *k* and the dissipation rate of kinetic energy  $\varepsilon$ . The transport equations for *k* and  $\varepsilon$  expressed by:

$$\nabla(\rho Vk) = \nabla \left(\mu + \frac{\mu_{t}}{\sigma_{k}}\right) \nabla k + P_{k} - \rho \varepsilon - Dt$$
(11)

and

$$\nabla(\rho V\varepsilon) = \nabla \left(\mu + \frac{\mu_{t}}{\sigma_{\varepsilon}}\right) \nabla \varepsilon + C_{\varepsilon 1} \frac{\varepsilon}{k} P_{k} - C_{\varepsilon 2} \rho \frac{\varepsilon^{2}}{k}$$
(12)

where

$$\mu_{t} = \rho C_{\mu} \frac{k^{2}}{\varepsilon}$$
(13)

For the ideal gas

$$Dt = 2\rho \frac{\varepsilon k}{\gamma RT}$$
(14)

The constants:  $C\varepsilon 1 = 1.44$ ;  $C\varepsilon 2 = 1.92$ ,  $\sigma_k = 1$ ;  $\sigma_{\varepsilon} = 1.3$ , and  $C\mu = 0.09$ .  $P_k$  represents the generation of the kinetic energy caused by velocity gradient and volume forces.

## **Combustion modeling**

For the combustion reaction model, we will use the generalized finite-rate chemistry model to analyze the methane-air combustion system. The combustion will be modeled using a global one-step reaction mechanism as depicted the eq. (15):

$$CH_4 + 9.5238(0.21O_2 + 0.79N_2) \rightarrow CO_2 + 2H_2O + 7.5238N_2$$
 (15)

The eddy dissipation model is applied to model the heat release and also to take into account the turbulence-reaction interactions. The eddy-dissipation model computes the rate of reaction under the assumption that chemical kinetics are fast compared to the rate at which reactants are mixed by turbulent fluctuation. In the flame zone, the eddy dissipation rate is generally smaller than the Arrhenius rate, and reactions are limited by the extent of mixing. Based on the assumption that the reactant rate is limited by both the mixing of the reactants and the heating of the reactants through the mixing with hot reaction products, the net rate of production of species *i* due to reaction,  $R_i$ , is given by the smaller of the reactant mixing rate and product mixing rate [23]:

$$R_{i} = \min\left[\nu_{i}^{\prime}M_{w,i}A\rho\frac{\varepsilon}{k}\min_{R}\left(\frac{Y_{R}}{\nu_{R}^{\prime}M_{w,R}}\right),\nu_{i}^{\prime}M_{w,i}AB\rho\frac{\varepsilon}{k}\frac{\Sigma PY_{P}}{\sum_{j}^{N}\nu_{j}^{\prime\prime}M_{w,j}}\right]$$
(16)

where A and B are model constants (A = 4 and B = 0.5),  $v'_i$  and  $v''_j$  are the stoichiometric coefficient for reactant *i* and product *j* in reaction,  $M_w$  is the molecular weight,  $Y_R$  and  $Y_P$  are mass fraction of any reactant and product species, respectively.

## **CFD modeling**

#### Discretization procedure

In the present study, the commercial CFD package FLUENT6.3 was used to model combustion, heat transfer, and  $NO_x$  emission in a diffusion flame methane-air with a coaxial burner. A finite volume approach is employed to solve the governing equations including mass, momentum, energy, and species in addition to the turbulence transport and combustion model. The governing equations were discretized in the whole domain using the second-order schemes. In addition, the segregated solution method and the simple pressure-velocity coupling algorithm were used to solve the governing equations. The Standard *k-\varepsilon* turbulence model, the eddy dissipation model for turbulent reacting flows, the thermal, prompt and N<sub>2</sub>O-intermediate models for NO<sub>x</sub> emission were introduced to the CFD code. The convergence criterion requirement is set to be 10<sup>-6</sup> for energy and about 10<sup>-3</sup> for the other terms of the transport equations.

#### Computational domain and boundary conditions

The axisymmetric computational domain (half of the flame) was extended 100 cm in the axial direction and 25 cm in the radial direction and had a coaxial burner whose the radius of methane injector equal 3 mm and the radius of annular jet equal to 9 mm. A total number of 16200 ( $200 \times 81$ ) quadrilateral cells were generated using non-uniform grid spacing to provide an adequate resolution near the jet axis and close to the burner where gradients were large. The grid spacing increased in the radial and axial directions since gradients were small in the far-field. The computational domain was bounded by the axis of symmetry, velocity inlet for methane, velocity inlet for air, pressure inlet for ambient air and pressure outlet to the upper boundary condition (ex-



Figure 2. Mesh geometry

haust), fig. 2. In this study, pure methane ( $\rho_g = 0.61517 \text{ g/l}$ ) (mole fraction XCH<sub>4</sub> = 1) flows through the inner stream ( $d_f = 6 \text{ mm}$ ) at 10 m/s, while air ( $\rho_a = 1.177 \text{ g/l}$ ) flows through the outer stream ( $d_a = 18 \text{ mm}$ ) at different velocity (variation inlet of air velocity: 0 m/s to 20 m/s). Air is composed by 0.21 mass fraction of O<sub>2</sub> and 0.79 mass fraction of N<sub>2</sub>. Gas and air temperatures were both 25 °C, and the ambient pressure was 101325 Pa.

#### **Results and discussion**

In this study, we investigate the effect of air velocity variation on the turbulent diffusion flame. For this reason, with a constant velocity of methane flow 10 m/s, we varied the air flow velocity between 0 and 20 m/s, as shown in tab. 1. The mixture fraction and equivalence ratio, two important parameters in a problem of non-premixed combustion, are given in (tab. 1). Our aim is to have the influence of air flow on the temperature distribution, on gas consumption and pollutant emissions.

Case No.	d <sub>g</sub> [mm]	d <sub>a</sub> [mm]	Gas flow $Q_{\rm g}$ [lmin <sup>-1</sup> ]	Air flow $Q_{\rm a}$ [lmin <sup>-1</sup> ]	Gas velocity $V_{\rm g}  [{\rm ms}^{-1}]$	Air velocity $V_{\rm a}  [{\rm ms}^{-1}]$	Reg	$V_{\rm a}/V_{\rm g}$	ξ	φ
1	6	18	16.956	0	10	0	5217	0	1	-
2	6	18	16.956	94.92	10	7	5217	7/10	0.0853	1.6
3	6	18	16.956	135.6	10	10	5217	1	0.0613	1.12
4	6	18	16.956	203.46	10	15	5217	3/2	0.0417	0.74
5	6	18	16.956	271.2	10	20	5217	2	0.0316	10.56

Table 1. Geometrical and physical properties of the problem

#### Model validation

In order to validate the CFD results, the predicted radial temperature gradients are compared to flame temperature measurements reported by [24]. Figure 3(a-c) shows the radial temperature distribution from experimental measurement and numerical prediction at different position from the burner. The radial temperature gradients, in the present flame, represent the width of the reaction zone of the flame. The predicted radial temperature profile shows satisfactory with the experimental measurements in terms of both trend and values. The numerical sim-

ulation shows the predicted radial gradients of temperature imply a same reaction zone that corresponds to the experimental measurements. The temperature values on the burner axis given by the numerical simulation are less than the values corresponding to the experimental measurement. For example, as shown in fig. 3(b), the predicted and measured locations of maximum temperature are 20.87 and 19.37 mm, while the predicted and measured values of maximum temperature are 1789.83 and 1670.73 K. This difference is a consequence of heat transfer between the flame and the surrounding environment.



#### Temperature distribution

For a fixed methane flow rate and increasing values of air flow, the flame temperature distribution, the consumption of methane, the distribution of oxygen and emission NO<sub>x</sub> are determined numericaly. Figure 4(a-e) represents the spatial temperature distribution as a function of the variation of velocity ratio. We note, for a low air flow, the flame temperature is low and extended into a large area in a computational domain. But, the flame becomes thinner and higher when we increase the air flow. For example, in the absence of airflow (velocity ratio  $V_a/V_g = 0$ ), first the maximum temperature does not exceed 1590 °C, secondly, the flame starts the burner exit (no lifted flame), fig. 4(a). The maximum flame temperature varied from 1779 °C for  $V_a/V_g = 2$ . The flame lifts fronts on the injector nozzle of the burner with increasing air jet velocity, as shown in figs. 4(b-e).

The radial variation of temperature at different heights and for different velocity ratio is shown in fig. 5. For a low velocity ratio  $(V_a/V_g = 0)$ , fig. 5(a), the flame starts from the burner nozzle and the temperature of the flame front is low, because the combustion reaction occurs in the absence of air. Flame temperature maximum is important when increasing velocity ratio, but appear far from the burner nozzle at a height to be near 20 cm as shown in fig. 5(b-e). The temperature is maximum at the flame front where the fuel and air are well premixed. The maximum temperatures represent the stoichiometric line, fig. 6. Figure 6 shows the distribution of maxi-



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mum temperature in the computational domain so it shows the width of flame. The increase of velocity ratio leads to a decrease in the flame width and favors the temperature of flame front, therefore favors an increase in  $NO_x$  emission.

If the velocity ratio increases, the temperature on the burner axis increases. For example, at a height of 50 cm from the nozzle burner and for a low velocity ratio, the temperature is 846 °C, the maximum temperature is 1043 °C, 1097 °C, 1296 °C, and 1457 °C corresponding, respectively, to 7/10, 1, 3/2, and 2 velocity ratio.

# Mole fraction of $CH_4$ and $O_2$

## Effect of velocity ratio on the consumption of methane

Figure 7(a-c) represent the radial variation of methane at different positions of the flame and for different velocity ratio. We selected three positions downstream the burner (10 cm, 20 cm, and 50 cm) and four velocity ratios and show their effect on the CH<sub>4</sub> consumption along the flame. Figure 7(a) shows the radial variation of CH<sub>4</sub> mole fraction near the nozzle of the burner (y = 10 cm) for different velocity ratio, the mole fraction of methane presents a maximum on the burner axis and decreases when the radial position x increases and extends towards zero in



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the ambient zone. First, the maximum mole fraction decreases when the velocity ratio increases, This indicating that the amount of oxygen entrained into the jet fuel, increases with increasing air velocity. Secondly, the flame is thin when the velocity ratio increases. For example, for a low-velocity ratio ( $V_a/V_g = 7/10$ ), in a position near the burner nozzle (y = 10 cm), the mole fraction of gas equal to 0.70 on the burner axis and extend to zero at 2.1 cm radial distance. In a same position and for high velocity ratio ( $V_a/V_g = 2$ ) the CH<sub>4</sub> mole fraction equals to 0.344 and the amount of methane is consumed at a 1.63 cm in the radial axis. Two figures, fig. 7(b) and 7(c). show the

variation of CH<sub>4</sub> mole fraction for different velocity ratio at the position *y* equals to 20 cm and 50 cm along the axis of the burner. The amount of methane, consumed by the combustion reaction, depends upon the height along the burner axis and the velocity ratio, so its decreases with the height *y* as the velocity ratio increases. At a height of 50 cm along the burner axis, fig. 7(c), the maximum mole fraction equals to 0.37 for a velocity ratio ( $V_a/V_g = 7/10$ ) and equals to 0.133 for a velocity ratio ( $V_a/V_g = 2$ ). This means that at the distance of 50 cm



Figure 6. Line of maximum temperature at different velocity ratio

downstream the burner, the gas is consumed completely by the combustion reaction. For a fixed velocity ratio, for example  $(V_a/V_g = 3/2)$  CH<sub>4</sub> mole fraction varies from 0.48 at position of 10 cm to 0.21 at y = 50 cm.



#### Distribution of oxygen in the reaction zone

Figure 8 represents the radial variation of the mole fraction of oxygen at 10 cm, fig. 8(a), 20 cm, fig. 8(b), and 50 cm, fig. 8(c) downstream the burner nozzle (the oxidizer air is composed of 21% oxygen and 79% nitrogen). The distribution of oxygen in the flame indicates the quality of mixture between the fuel and air. In the non-reactive zone and near the burner, fig. 8(a), the variation of mole fraction of oxygen represents a minimum on the axis burner and increases rapidly to the maximum 0.21. In an intermediate section, fig. 8(b), the curves of variation of mole fraction of oxygen represent a minimum in the flame front, where the mixing between methane and air is near the stoichiometry. Near the stoichiometric conditions, the quantity of gas and oxygen injected into the reaction is completely consumed by combustion re-

action. The radial position of the minimum decreases when the velocity ratio increases, for example, for a ratio  $(V_a/V_g = 7/10)$  the minimum mole fraction of oxygen located at a radial position of 4 cm, against for ratio  $(V/V_g = 2)$  this minimum is located at 2.8 cm radially. So the flame becomes thinner when the air velocity increases. In the upper part of the flame corresponding y = 50 cm, fig. 8(c), the mole fraction of oxygen varies between 0 and 0.05 in the flame zone, indicating that the amount of oxygen injected by the burner is completely consumed by combustion reaction.



#### NO<sub>x</sub> emission

The presence of nitrogen in the combustion reaction at a high temperature promotes the  $NO_x$  production. This nitrogen is found with a low amount in the fuel and a large quantity in the air. The NO is formed mainly by the Thermal route, Prompt NO, Fuel NO and N<sub>2</sub>O intermediate.

The formation of thermal NO is determined by direct oxidation of nitrogen in the oxidizer and was described by Zeldovich [25]. It is depends on the following reactions:

$$N_2 + O \iff NO + N$$
 (R1)

$$N + O_2 \quad \leftrightarrow \quad NO + O$$
 (R2)

The overall reaction is given by:

$$N_2 + O_2 \leftrightarrow 2NO$$
 (R3)

These reactions are slow and require high temperatures. Another reaction that may be added to the mechanism of thermal NO formation is defined by (R4):

$$N + OH \leftrightarrow NO + H$$
 (R4)

The Prompt NO mechanism is responsible for NO formation in the flame front. This mechanism is through by the rapid reaction of hydrocarbon radicals with molecular nitrogen:

$$CH + N_2 \leftrightarrow HCN + N$$
 (R5)

The Fuel NO mechanism is responsible for the formation of  $NO_x$  from the nitrogen content in the fuel. This mechanism depends mainly on the nitrogen content of the fuel. For certain fuels this mechanism is responsible for much of the formation of  $NO_x$ .

Generally  $N_2O$  emission in the flame is low and the mechanism of their formation is given by Faivre [20] by the reactions:

$$NCO + NO \iff N_2O + CO$$
 (R6)

$$NH + NO \leftrightarrow N_2O + H$$
 (R7)

More N<sub>2</sub>O reacts with H atoms by the reaction:

$$N_2O + H \leftrightarrow N_2 + OH$$
 (R8)

The amount of NO emission is conventionally quantified by the emission index  $EINO_x$ , which is generally defined as the mass of pollutat emitted per unit mass of fuel consumed, and can be obtained using the equation [15]:

$$EI_{\rm NO_x} = \frac{M_{\rm NO_x} \int_V \omega_{\rm NO_x} dV}{M_{\rm CH_4} \int_V \omega_{\rm CH_4} dV} 10^3$$
(11)

Figure 9 represents the variation of the index emission NO<sub>x</sub> as a function of velocity ratio (depends air flow) for a coaxial burner. The four curve of fig. 9 shows, respectively, the formation of thermal NO, prompt NO, N<sub>2</sub>O intermediate, and total NO<sub>x</sub>. The N<sub>2</sub>O intermediate, emission in this case is negligible compared to the other two routes. NO prompt is the basic mechanism of NO<sub>x</sub> formation in our case, by cons, when using pure oxygen as an oxidizer, the NO thermal becomes the basic mechanism of NO formation since the flame front temperature is very high. If the velocity ratio increases of  $(V_a/V_g = 7/10)$  to  $(V_a/V_g = 3/2)$ , the NO prompt formation decreases and the NO thermal emission increases, but the total  $EI_{NO_x}$  formation decreases. The increase in  $EI_{NO_x}$  in the condition  $(V_a/V_g = 7/10)$  and  $V_a/V_g = 1)$  can be attributed to the increase in flame temperature due to the enhanced gas-air mixing leading to higher a NO<sub>x</sub> level.

The excess of oxygen and nitrogen in the flame  $(V_a/V_g = 2)$  causes a decrease in the NO thermal formation since the temperature of the flame front starts to decrease, and increased NO prompt formation as the important amount of nitrogen in the flame.

## Conclusions

The turbulent normal diffusion flame (NDF) stabilized on a coaxial burner is investigated in the present study for different velocities ratio. The temperature distribution, meth-



Figure 9. Emission index EINO<sub>x</sub> at different velocity ratio

ane and oxygen profiles in the flame and  $NO_x$  emission index are compared for different velocities ratio. The following results were obtained.

- With a gas flow rate constant (16.956 l/min), the variation of air flow in a coaxial burner • configuration, causes a change in the flame structure and temperature distribution. The increasing of velocity ratio, due to increased air flow, causes an increase on the flame temperature, lift-off flame, and lends to a thinner flame.
- The increase on the air velocity promotes the combustion reaction due to the enhanced gas-air mixing. If the air velocity increases, the methane is rapidly consumed by the combustion reaction, almost totally methane is consumed for a velocity ratio equal to 2 at a height of 50 cm.
- If the oxidant is air, the flame characterized by a lower temperatures and high NO<sub>y</sub> emission, compared to the use of pure oxygen. So for ensure the combustion reaction in a stoichiometric conditions, it requires very large air flow, which favors the lift flame. The amount of oxygen is low on the flame front due to the enhanced gas-air mixing. At a fixed axial position; this amount is increased when the velocity ratio increases.
- The increase in the NO formation by the prompt mechanism, as the main contributor, • demonstrated the observed increase in the  $NO_x$  emission index. The  $NO_x$  formation depends on the type of oxidizer (pure oxygen or air), the type of fuel (contains nitrogen or no), the velocity ratio and depends upon the flame temperature.

## Nomenclature

- diffusion coefficient of species i  $D_i$
- diameter of air jet, [mm]  $d_{a}$
- $d_{\rm g}$ diameter of gas jet, [mm]
- acceleration of gravity,  $[ms^{-2}]$ g
- enthalpy of species i, [J]  $h_i$ \_
- diffusive flux [molm<sup>-2</sup>s<sup>-</sup>  $J_i$
- $J_a$ heat flow caused by the diffusive flux,  $[Jm^{-2}s^{-1}]$
- k - turbulent kinetic energy,  $[m^2 s^{-2}]$
- $M_{w}$ molecular weight, [kgmol<sup>-1</sup>]
- Р - absolute pressure, [Pa]
- $Q_a$
- air flow, [lmin<sup>-1</sup>]
  gas flow, [lmin<sup>-1</sup>]  $Q_{\rm g}$
- heat radiation, [J]  $q_{\rm r}$
- universal gas constant, [JKg<sup>-1</sup>mol<sup>-1</sup>K<sup>-1</sup>] R
- Reg Reynolds number for methane
- $(=\rho V_{\rm g} d_{\rm g}/\mu)$ Т - temperature, [K]
- v
- overall velocity vector, [ms<sup>-1</sup>]  $V_{a}$ - air velocity, [ms<sup>-1</sup>]

- $V_{\rm g}$ - gas velocity, [ms<sup>-1</sup>]
- x - radial co-ordinates, [mm] v
  - axial co-ordinates, [mm]
- $Y_i$ - mass fraction of species *i*

Greek symbols

ε

λ

- dissipation rate of kinetic energy,  $[m^2s^{-3}]$
- thermal conductivity,  $[Wm^{-1}K^{-1}]$
- dynamic viscosity,  $[kgm^{-1}s^{-1}]$ μ
- stoichiometric coefficient for reactant *i*  $v'_i$ product j
- stoichiometric coefficient for product j
- $v_j''$ - mixture fraction
- density, [kgm<sup>-3</sup>] ρ
- Stefan-Boltzmann constant  $\sigma$ 
  - $(= 5.6704 \cdot 10^{-8}) [Wm^{-2}K^{-4}]$
- τ stress tensor
- equivalence ratio φ
- production rate of species *i*,  $[molm^3 s^{-1}]$  $\omega_i$

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