

Effect of Propane-Methanol Blending Ratio on the Stretched and Unstretched Flame Speeds at Elevated Initial Temperatures

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The unstretched flame speed of the premixed propane-methanol/ air flames has been studied experimentally in a constant volume combustion chamber with central ignition. The experiments were done at atmospheric pressure and stoichiometric air/fuel ratio. Various blending ratios of methanol (0%, 20%, 40%, 60%, 80%, 100%) by volume, and different elevated initial temperatures (348 K, 373 K, and 398 K) were used in this study. In general, the results indicated that the unstretched flame speed increased with increasing both methanol blending ratio and initial temperature. For M60 the increment value of unstretched flame speed at $T_i=398K$ was about 9% compared with that of pure propane and by elevating the initial temperature for the same blend ratio (M60) from 348 K to 398 K the increment value was about 8.8%. It is also noticed that Markstein length decreased with increasing both initial temperature and blending ratio, which is indicate that flame instability increased with increasing these parameters.

Keywords: *Blending, Constant volume chamber, Initial temperature, Markstein length, Un-stretched flame speed*

1. INTRODUCTION

Increasing the concern of fossil fuels depleting and air pollution lead to that improving engine fuel economy and reducing exhaust emissions had become the major topics in combustion researches and engines manufacture [1-7]. Many studies focused to search for alternative fuels in addition to enhance the combustion process since the 1970s [8]. Some of the alternative fuels reported in the literature which are used for spark ignition engines were alcohols, compressed natural gas (CNG), hydrogen, liquid petroleum gas (LPG), dimethyl ether (DME), and others that used in various internal combustion engines [9].

Methanol is considered as one of the important alternative fuels due to its advantages of environmental friendly, can be prepared from multiple sources, and having high octane number compared with that of fossil fuels [10]. However, it also has some problems due to its low energy density and high latent heat of vaporization. It is reported that methanol fuel having a high specific fuel consumption when used in internal combustion engines compared with conventional fuels such as gasoline or diesel fuels [10]. Therefore, methanol is widely used as an additive into other fuels. In addition to that recent increased the use of propane as a fuel in transport vehicles [11] led the propane to be a key fuel candidate of study.

Laminar flame speed is a necessary parameter

describing the physical and chemical properties in propagation of flame, and to validate the chemical kinetic model. It is essential for studying turbulent flame [12]. Constant Volume Chamber (CVC) method is one of several methods used in the field of flame speed, which is broadly applied as one of the non-stationary methods to measure burning velocity (S_u) and flame speed (S_l) with reliable experimental results [13]. This method can be applied at a wide range of initial pressure (0.1 – 60 bar) compared with other methods [14]. Several factors influence on the accuracy of experiments in the CVC method, including the effect of spark ignition, chamber confinement, radiation heat loss, flame stretch, and flame instability [11,13,14]. To get precise measurements of laminar flame speed, the effect of the above-mentioned factors should be accurately minimized.

The internal cavity of the CVC can be designed with cylindrical or spherical shapes at different dimensions. Salih and Chaichan, [17] conducted their experiments in a cylindrical chamber with a length and inner diameter of 305 mm. Galmiche et al. [18] used a cylindrical vessel made from stainless steel with 24.32 L inner volume. Hinton and Stone [19] utilized 160 mm inner diameter of spherical vessel, while Mannaa et al. [20] used a spherical stainless steel combustion chamber with inner diameter of 330 mm. Other study conducted by Yasiry and Shahad [21] with 190 mm inner diameter and 250 mm length cylindrical vessel. Cylindrical cavity with an inner diameter of 80 mm and a length of 268 mm used by Yao et al. [22]. Bao et al. [23] carried out their experiments in a spherical vessel with 350 mm inner diameter. Lhuillier et al. [24] done their experiments in a spherical stainless steel combustion chamber with a volume of 4.2l. Linteris and Babushok

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[25] used a spherical vessel with volume of 3.05 L. A stainless-steel cylindrical combustion chamber with an internal diameter and length of 305 mm utilized by Duva et al. [26].

Flame stretch is a quantity which measures the stretch amount of the surface of flame due to the curvature and the outer velocity field strain. It is required to studying the effect of the stretch on the burning velocity. The rate of flame stretch is $K = 2S_n/r_f$ for the spherical propagation flame. The un-stretched flame speed (S_u) and Markstein length of burned gas, (L_b) can be determined by extrapolate the S_n to zero stretches.

Initial temperature considered as the most significant factor influencing flame speed and burning velocity with direct proportional relation. It allows a more stable flame propagation. The following equation is the most common correlation expressing the effect of initial pressure (P_i) and temperature (T_i):

$$S_u = S_{u,o} \left(\frac{T_i}{T_{i,o}} \right)^\gamma \left(\frac{P_i}{P_{i,o}} \right)^\beta \quad (1)$$

where (o) represent the reference conditions ($T_{i,o} = 298$ K, $P_{i,o} = 1$ bar). The exponent power for pressure and temperature (β , and γ) were discussed broadly by Konnov et al. [27]. It differs as a function of several factors such as: equivalence ratio, fuel type, and the oxidizer. Varghese and Kumar [28] examined laminar flame speed for mixtures of $H_2/CH_4/CO_2/CO/N_2$ /air at high initial pressures and temperatures. They suggested a power-law correlation, as shown below:

$$S_u = S_{u,o} \left(\frac{T_i}{T_{i,o}} \right)^{\alpha_o + \alpha_1} \left(1 - \frac{P_i}{P_{i,o}} \right) \left(\frac{P_i}{P_{i,o}} \right)^{\beta_o + \beta_1} \left(1 - \frac{T_i}{T_{i,o}} \right) \quad (2)$$

Han et al. [29] investigated the S_u for natural gas at an initial pressure of two bars under different initial temperatures. They showed that laminar flame speed increased with increasing the temperature for all mixtures examined.

Many researchers investigate S_u and S_f for methanol blending fuels. Metghalchi and Keck [30] proposed empirical equations for S_u of the mixtures of methanol, indolene and isooctane at different initial temperatures and pressures. Beeckmann et al. [31] and Zhang et al. [32] studied the laminar flame speed of methanol-isooctane blends at different initial temperatures and atmospheric pressure. They found that the blends have S_u values higher than isooctane but lower than methanol. Liu et al. [33] investigated laminar flame speed for hydrogen-methanol blends, and suggested correlation for it.

According to the literature review, there are a few studies reported data for the blends of methanol and propane. Therefore, the aim of this study is to investigate the effect of methanol-propane blending ratios and the effect of initial temperature (that have not been used in other previous studies) on the unstretched flame speed and Markstein length in a cylindrical constant volume combustion chamber (CVCC) using schlieren technique. The study was conducted under atmospheric pressure and at stoichiometric ratio, for

different initial temperatures (348 K, 373 K, and 398 K) and various blending ratios:

- 0% methanol-100% propane (M0)
- 20% methanol-80% propane (M20)
- 40% methanol-60% propane (M40)
- 60% methanol-40% propane (M60)
- 80% methanol-20% propane (M80)
- 100% methanol-0% propane (M100)

2. EXPERIMENTAL SETUP

In this study, the Constant Volume Combustion Chamber (CVCC) method and schlieren photography technique are used to measure the flame speed. The complete setup of the rig and the schematic layout are shown in figures (1) and (2), respectively. The parts of the rig are fabricated and assembled in the laboratories of Mechanical Engineering Department at the University of Technology / IRAQ.

The combustion chamber is built from a stainless-steel type 316 cylinder with 305 mm inner diameter, and 400 mm length with a volume of 29 L. Two pressure resisting high transparency windows with 120 mm net diameter and thickness of 20 mm are fixed on the opposite sides of the cylinder to allow the optical access for combustion process. Seven sockets are welded to the cylinder, two of them placed at opposed, co-linear position used to insert and fixed two spark ignition electrodes, which are connected to the ignition system. The others used for liquid fuel injection, admitting gaseous fuel and air, exhaust gases, and inserting thermocouple type PT-100, and pressure transmitter. A robust stainless-steel stand was used to hold the combustion chamber.

The fuel injection unit, (figure 3) is used to induct the gaseous fuel (propane)/air mixture into the combustion chamber, while the liquid fuel (methanol) was injected directly to the combustion chamber using liquid fuel injection valve. An electrical control board unit is fabricated and utilized to facilitate and regulate conducting the experiments accurately. The unit supplies 12 V DC to the solenoid valves and regulates and controls the voltage entering to the ignition burner transformer. An electronic circuit unit is constructed to supply the required powerful spark to the electrodes and to control its timing. A schlieren technique was used to visualize the process of flame propagation with a high-speed camera (AOS -Q PRI) using a light source and plano-convex lenses as shown in figure (4).

The thermocouple and the pressure transmitter were calibrated regularly over different temperature and pressure ranges at the Central Organization for Standardization and Quality Control (COSQC) of Iraq.

3. TEST PROCEDURE

The first step started with vacuum process to scavenge the combustion chamber from any previous gases until its pressure reaches approximately -0.94 bar, then flushing process takes place by injecting air inside the combustion chamber for 3-4 minutes. This step is repeated twice to ensure that the combustion chamber is flushed completely from previous test. The vacuum

process is repeated for the three times to prepare the CC for injecting fuels and air according to the mixing ratio which was calculated previously. After that, turning on the heating system to heat up the combustion chamber to the desired temperature. In the next step, propane fuel is injected to the combustion chamber by fuel injection unit, while methanol fuel is injected by the liquid fuel injection valve according to its partial pressure. The mixing process is based on Gibbs-Dalton law of partial pressures for each mixture component. Subsequently, the air is admitted to the combustion chamber until

reaches the required total pressure. The mixture is left for 5-6 minutes to ensure perfect mixing before igniting the mixture. The spark duration is set at 10 ms. Then the ignition switch and the camera trigger with 600×600 pixels, 3536 fps are pushed simultaneously to initiate recording and combustion processes. The results are recorded for each condition. In addition, the photographs of flame propagation are taken and then analysed using Tracker 5.1.5 software.



Figure 1. Photograph of the experimental apparatus used in the study. (1) Combustion chamber, (2) Thermocouple, (3) Pressure transmitter, (4) Liquid fuel injection valve, (5) Inlet valve, (6) Exhaust valve, (7) Electrical heating tape, (8) Sparking ignition electrodes, (9) Ignition burner transformer, (10) Injection unit, (11) Vacuum pump, (12) Electrical control board unit, (13) Light Source, (14) Two lenses (15) High-speed camera, and (16) PC computer.

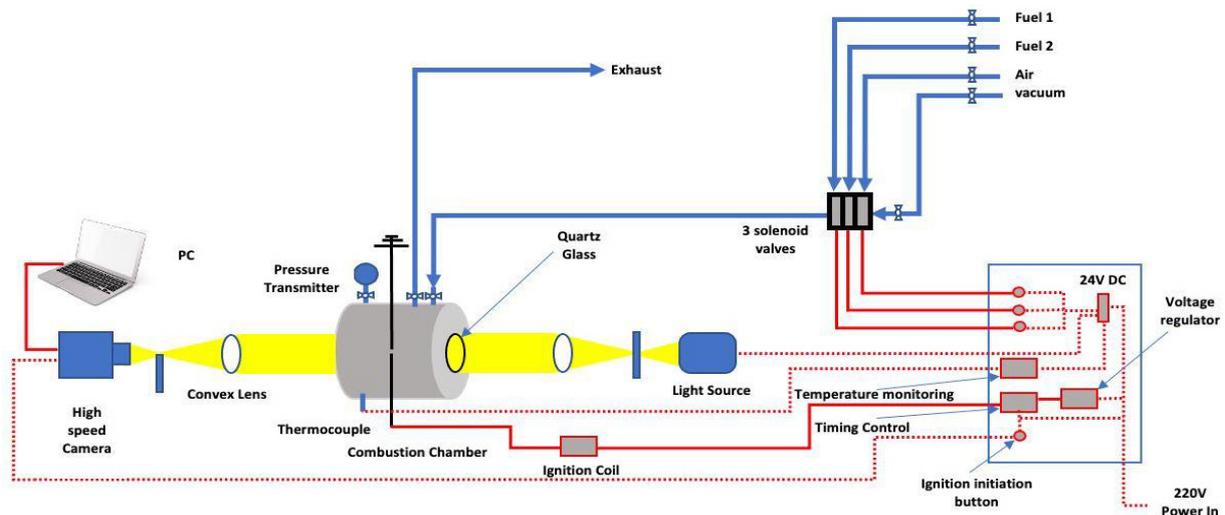


Figure 2. Schematic diagram of the experimental set up.

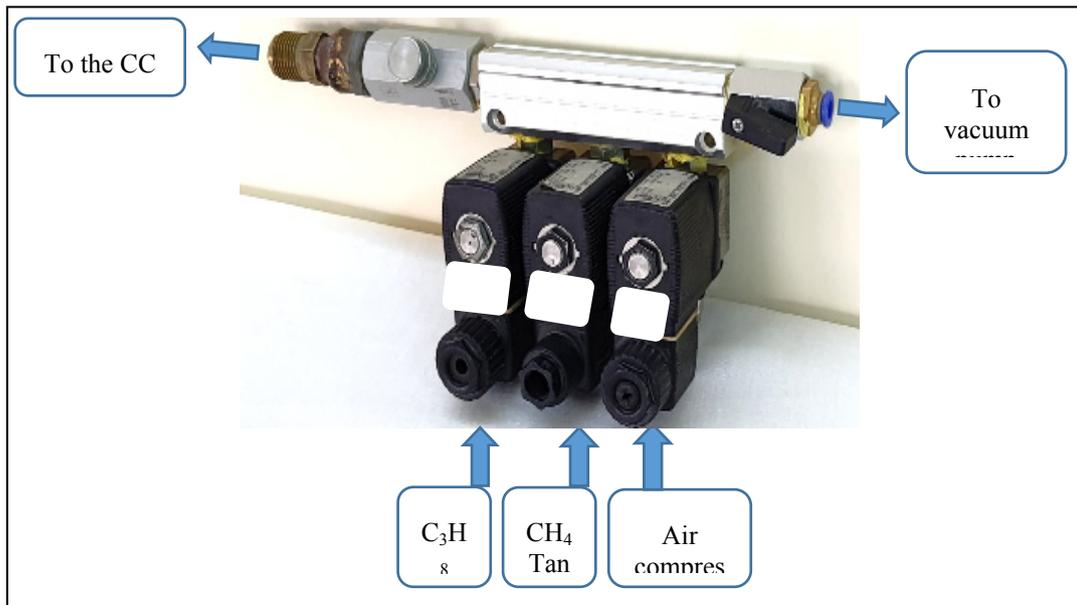


Figure 3. Fuel injection unit

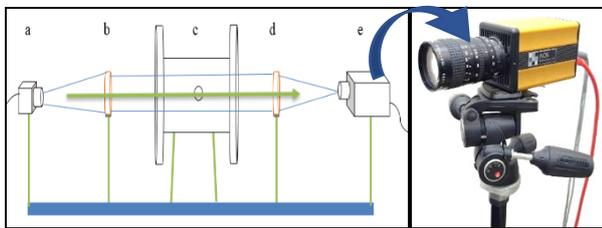


Figure 4. Schematic diagram for capturing unit a) illuminator, b) collimating lens, c) combustion chamber, d) de-collimating lens and e) high speed camera.

To test the accuracy of our system, the laminar burning velocity (S_u) for methane/air mixture at 1bar and ambient temperature for different equivalence ratios are compared with results of other researchers as shown in figure (5). Also, repeatability tests are carried out by repeated the experimental test for five times using the same fuel type (methane) and the same conditions (equivalence ratio, initial temperature, and initial pressure). Figure (6) shows that the results of repeatability are reasonable, since all the curves obtained have the same trend and the maximum percentage error at a radius of 2cm was 5.3 %.

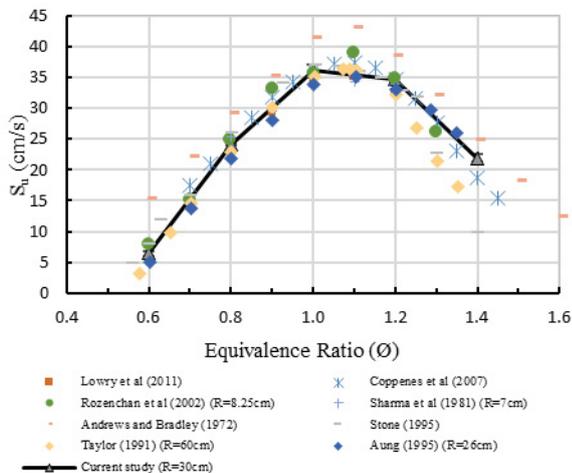


Figure 5. Comparison of experimental data for S_u of methane at standard conditions with the published results.

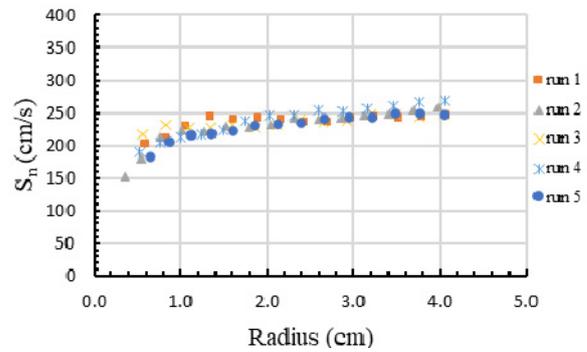


Figure 6. Stretched flame speed with flame radius for methane at $T_i=306$ K, $P_i=1$ bar, and $\phi=1$.

4. RESULTS AND DISCUSSIONS

In order to present an example of the flame propagation video recording, Figure (7) presents frames sequence of the spherical expanding flame for stoichiometric methanol/air mixture at different times and different initial temperatures. The subsequent frames are approximately every 3 ms. Tracker 5.1.5 software is applied to extract information about variation of flame radius with time to follow the flame front advancement. The instantaneous stretched flame speed (S_n) is the slope of the line segment involving two neighboring radii against the time. It can be seen that the flame propagated faster when the unburned mixture temperature is higher.

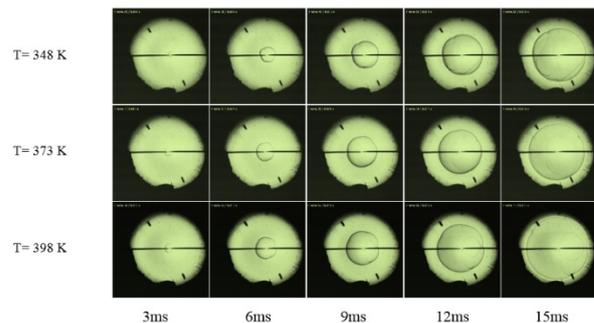


Figure 7. Photographs of flame propagation for M40 at $P_0= 1.0$ bar, and different initial temperatures.

4.1 Stretched laminar flame speed

The stretched laminar flame speed (S_n) has been extracted from the video recordings. Figure (8) shows the stretched laminar flame speed versus flame radius for different blending ratios and initial temperatures. The results show that the S_n increased with increasing initial temperature, since high temperatures boost the dissociation reactions that produce free radicals. These radicals initiate combustion reaction, which is why the velocity of flame propagation increased [41]. Likewise, with the increase in initial temperature, the adiabatic flame temperature increased and reaction rates become faster.

The data is extracted from the pre-pressure zone at the flame radius extending between 7–40 mm which show approximately constant flame speed with flame radius. The stretched laminar flame speed behavior subdivided into three regions. The first region; ignition affected region which is at flame radius below 7 mm, where the excess ignition energy could affect the flame propagation. Bradley et al. [42] clarified that this excess

energy can overestimate the flame speed for the fuel/air mixtures at a flame radius below 5 mm. In this region the flame speed cannot be used to determine the S_u due to the interference of underdeveloped flame. The second region extends until the pressure starts to rise. In this region the stretched flame speed (S_n) would remain approximately constant. Prathap et al. [43] proposed that the flame radius smaller than one-third the radius of the combustion chamber was reasonable for the calculation.

The third region begin from flame radius greater than 40 mm, where the strain in flow field would significantly change flame speed in addition to the effect of combustion pressure which is no more can be assumed constant. The first and third regions are excluded from the calculation of stretch flame speed. Sahu et al. [44] used cylindrical CC and choose 10 mm and 34mm (which forms about 4.0% of the total vessel volume) as the lower and upper critical flame radii for calculating flame speed.

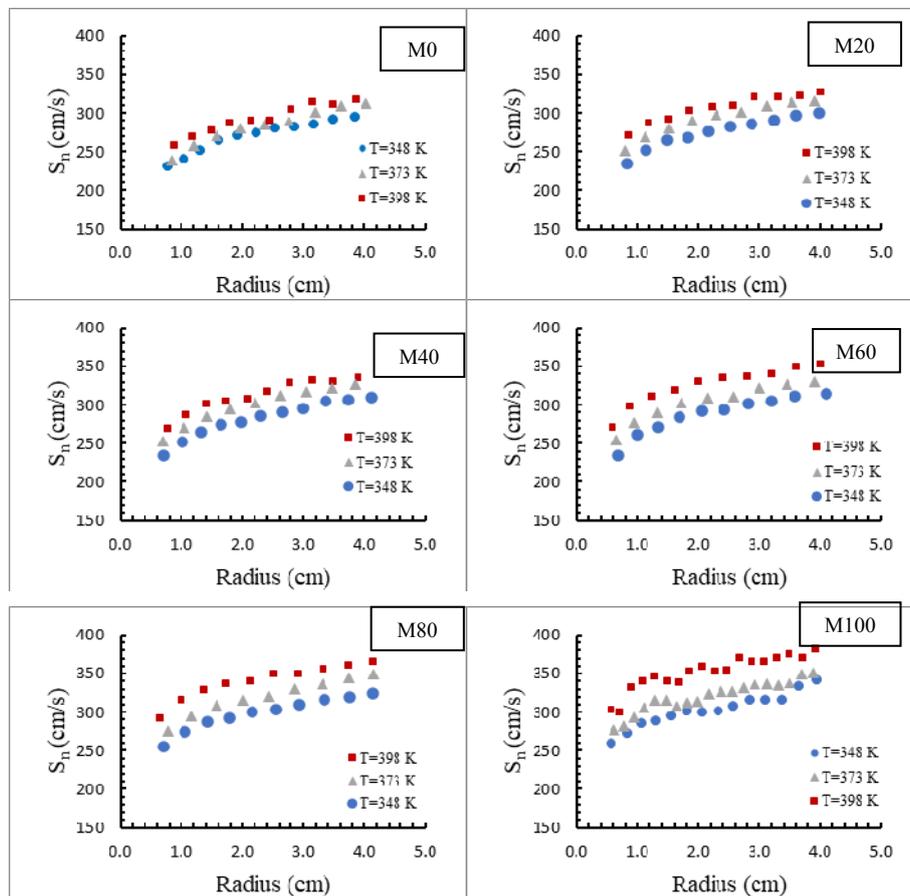


Figure 8. Variation of S_n with flame radius for different initial temperatures and blending ratios.

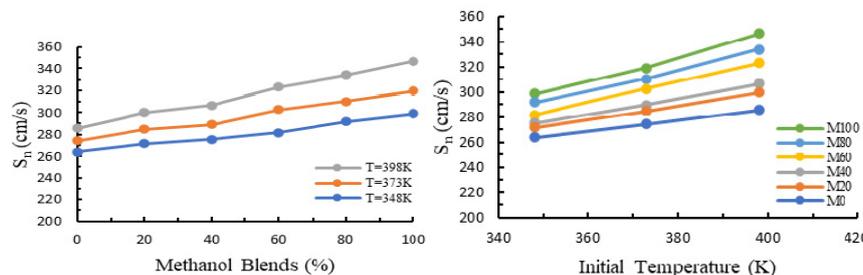


Figure 9. Variation of S_n at 20 mm flame radius versus: (a) methanol blending ratios (b) initial temperatures.

Figure (9) illustrates the variation of S_n at 20 mm flame radius with methanol blending ratio and different initial temperatures (348 K, 373 K, and 398 K). It is shown that the stretched flame speed increases with increasing methanol ratio. It is found that the value of S_n increased by about 11% for M60 at $T_i=398\text{K}$ compared with that of pure propane (M0). It is also seen that the stretched flame speed increased with increasing initial temperatures. By rising the initial temperature for M60 from 348 K to 398 K the stretched flame speed increased by 12.8%. Similar trend is found in previous study [41]. It was also noticed that the maximum stretched flame speed has the same trend as the variation of S_n with initial temperature for all blending ratios as demonstrated in figure (10).

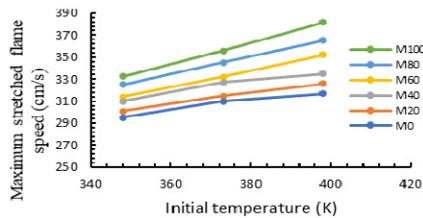


Figure 10. Variation of max. S_n versus initial temperature for different methanol blending ratios.

4.2 Un-stretched flame speed and stretched rate

When the flame-radius is small which is happened at early stage of flame propagation, the stretch rate of the flame front surface is too high due to high curvature of flame front. The stretch rate is directly proportional to the flame speed and inversely proportional with flame radius. As mentioned in the previous section the first and third region of flame propagation are excluded from the calculation and only the second region data are used to obtain the unstretched flame speed. A linear correlation between the flame stretch rate and the stretched flame propagation speed was used in this study.

The stretch rate is inversely proportional to the radius. Therefore, its effect decreases as the flame propagates due to the increases of flame radius. This would be expected on very general grounds since as increasing the flame radius, the flame shape becomes less curved and it seems as a planar, one-dimensional and hence, un-stretched flame. The stretch rate (α) has excessive values during the initial stages of flame propagation due to the significant curvature related with a small spherical surface. Flame speed is affected by flame stretch; it can be accurately determined when the stretch is negligible i.e. when $r \rightarrow \infty$.

The stretched flame propagation speeds decrease with the increase of stretch rate as shown in figure (11). It is clear that the influence of the stretch rate on flame during propagation is different for flames under different fuel blends and initial temperatures. It can be noticed from these graphs that the maximum stretch occurs at maximum initial temperature, and the stretched flame speeds decreased as the stretch rate increases. Similar trend was found in previous study [31]. The figure also shows that the gradients of S_n - α lines take the negative value, representing the positive values of Markstein-length.

Figure (12) presents the variation of Markstein length with initial temperature at various blending ratios. It can be noticed that Markstein length decreases with increasing both initial temperature and blending ratio. This indicates that flame stability decreased with increasing these parameters. It is clear from this figure that Markstein length of fuel blend is located between of their parent fuels.

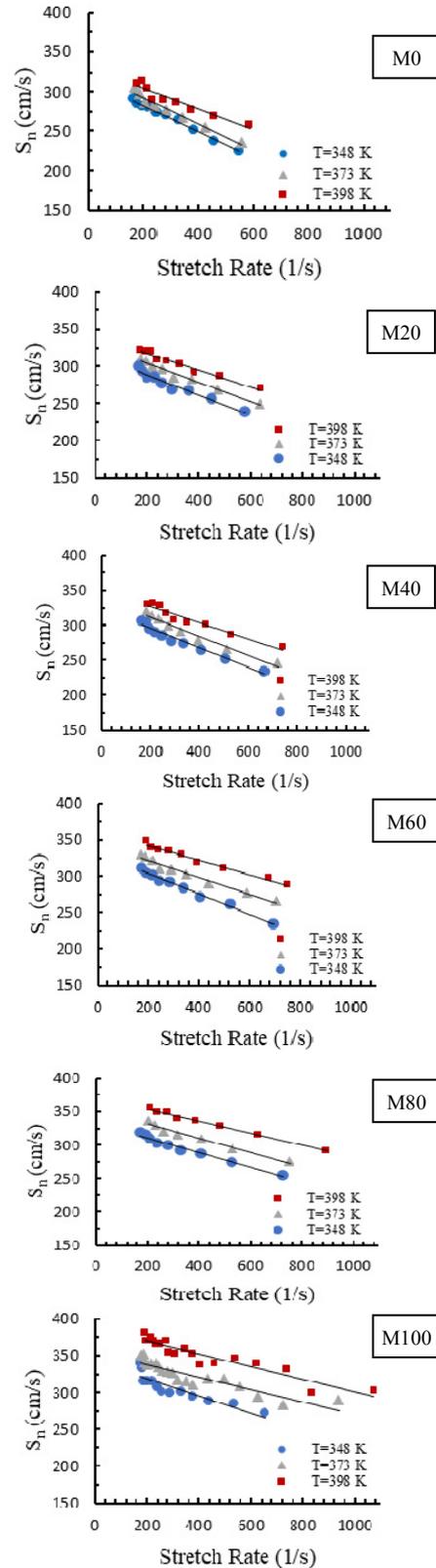


Figure 11. Stretched flame speed versus stretch rate for various initial temperatures and blending ratios

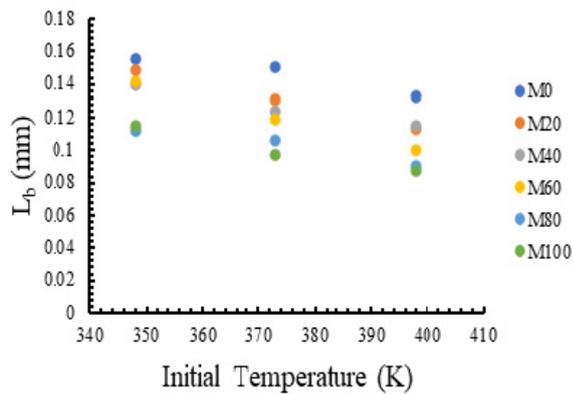


Figure 12: Markstein length versus initial temperature at different methanol blends.

The un-stretched laminar flame speed (S_i) is calculated as the intercept point of S_n at $\alpha=0$ in the plot of S_n versus α . The strain in the flow field causes a severe decrease of S_n in small stretch rate, and the cellular flame causes a severe increase of S_n in small stretch rate [21]. For reasons mentioned above the unstretched flame seed is obtained for the flame in the second stage of propagation which is at radius from 8-40 mm in the calculations.

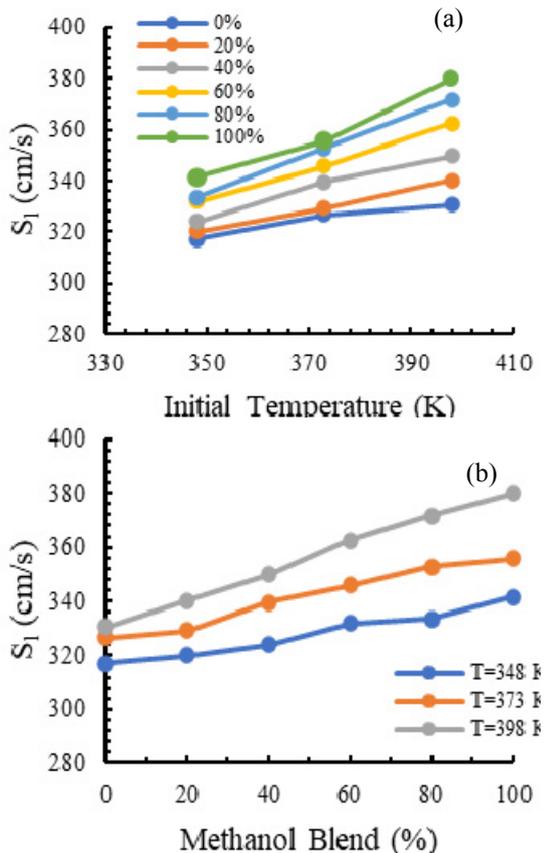


Figure 13. Unstretched flame propagation speed for stoichiometric mixtures versus (a) initial temperature (b) methanol blending ratio.

Figure (13) explains the effect initial temperature and methanol blending ratio on the un-stretched flame speed (S_i). It can be seen that the un-stretched flame speed increased with increasing the initial temperature. By elevating the initial temperature for M60 from 348 K to 398 K the value of the un-stretched flame speed (S_i) increased by about 8.8%. This trend is in agreement

with previously published results by Bao et al. [23]. It was also noticed that S_i increased with increasing methanol blending ratio. For M60 the value of S_i at $T_i=398\text{K}$ increased by about 9% compared with that of pure propane.

5. CONCLUSIONS

A constant volume combustion chamber has been design and constructed for studying the effect of propane-methanol blending ratios, and initial temperatures on stretched and un-stretch laminar flame speed. The experiments are performed for the stoichiometric mixtures of different blends. The first and third stages of flame propagation are excluded from the study. High speed photography technique is used to record the flame propagation. It can be conclude from the present study that:

- 1- Generally, the maximum stretch rate increases with increasing the initial temperature and the blending ratio.
- 2- The stretched flame speed (S_n) decreases with increasing the stretch rate. Whereas, for M60 and $T_i=373\text{ K}$ the stretch flame speed decreased from 329.6 cm/s to 265 cm/s by increasing the stretch rate from 169 s^{-1} to 702.1 s^{-1} .
- 3- The stretched flame speed (S_n) increased with increasing both initial temperature and methanol blending ratio. Raising the initial temperature of M60 from 348 K to 398 K led to increase the stretched flame speed by 12.8%. Furthermore, the value of S_n is increased by about 11% for M60 at $T_i=398\text{K}$ compared with that of pure propane (M0).
- 4- The un-stretched flame speed increased with increasing the initial temperature. By elevating the initial temperature for M60 from 348 K to 398 K the value of S_i is increased by about 8.8%.
- 5- The un-stretched flame speed increased with increasing methanol blending ratio. For M60 the value of S_i is increased by about 9% compared with that of pure propane.
- 6- The Markstein length decreased with increasing both initial temperature and blending ratio. For M60 increasing the initial temperature from 348 K to 398 K reduced the Markstein length from 0.142 mm to 0.1 mm. And for $T_i=373\text{ K}$ increasing the blending ratio from M20 to M80 reduced the Markstein length from 0.131 mm to 0.106 mm. This means that flame stability decreases with increasing these parameters. Furthermore, Markstein length of fuel blend is located between of their parent fuels.

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**УТИЦАЈ ОДНОСА МЕШАЊА ПРОПАНА И
МЕТАНОЛА НА БРЗИНЕ РАСТЕГНУТОГ И
НЕРАСТЕГНУТОГ ПЛАМЕНА НА
ПОВИШЕНИМ ПОЧЕТНИМ
ТЕМПЕРАТУРАМА**

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Брзина нерастегнутог пламена мешавине пропан-метанол/ваздух је експериментално проучавана у комори за сагоревање константне запремине са централним паљењем. Експерименти су рађени при атмосферском притиску и стехиометријском односу ваздух/гориво. У овом истраживању коришћени су различити односи мешања метанола (0%, 20%, 40%,

60%, 80%, 100%) и различите повишене почетне температуре (348 K, 373 K и 398 K). Уопштено говорећи, резултати су показали да се брзина нерас-тегнутог пламена повећава са повећањем и односа мешања метанола и почетне температуре. За М60 вредност повећања брзине пламена при $T_i = 398\text{K}$ била је око 9% у поређењу са чистом про-паном, а

повећањем почетне температуре за исти однос мешавине (М60) са 348 K на 398 K вредност прираста је била око 8,8%. Такође је примећено да се Маркштајнова дужина смањује са повећањем и почетне температуре и односа мешања, што указује да се нестабилност пламена повећава са повећањем ових параметара.