Ignition of premixed spherical spray flames

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Abstract

This study aims to build a theoretical model to describe the ignition process of spherical spray flame laden with fuel droplets under overall fuel-lean condition. Two characteristic fronts are introduced to reveal where the evaporation starts and ends in the ignition process. The results demonstrate that the flame has been strengthened with extra fuel droplet adding while the minimum ignition energy (MIE) is decreased. The evolution of the two evaporation fronts in the flame kernel and propagating branches has also be revealed coupled with the evaporation zone length.

Keywords

Ignition; Spray flame; Fuel droplet; Evaporation fronts.

Introduction and physical model

Spray combustion are widely used in propulsion system such aero-engines and rocket engines. Compared to gaseous flame, the existence of fuel/oxidizer spray in the combustion domain may bring some intriguing features in the ignition process, i.e. the liquid phase interaction with the gaseous flame kernel during the ignition period [1]. Theoretical analysis could provide a good reference on the ignition characteristics of spray flame.



Figure.1. Schematic of outwardly propagating spherical flame in liquid fuel mists: (a) heterogeneous flame, (b) homogeneous flame. Circle: fuel droplet. Yellow line: flame front (R_f) ; green line: evaporation completion front (R_c) ; blue line: evaporation onset front (R_v) . Black arrow: flame propagation direction. Red Spark (Q): forced ignition energy at the flame centre.

In this work, forced ignition of one-dimensional spherical flame in partially pre-vaporized fuel sprays will be studied. Initially, the gaseous fresh mixture is assumed to be fuel-lean and the fuel droplets are uniformly and dilute dispersed. The forced ignition is modelled as a localized energy deposition in fuel sprays. If the ignition energy is larger than MIE, a spray flame kernel would be generated and continuously propagate outwardly. During the flame development process, based on the droplet distribution relative to the reaction front, two general scenarios exist, i.e., evaporating droplets in: (1) both pre- and post-flame zones and (2) pre-flame zone only. The sketches of their physical models are shown in Figs 1(a) and 1(b), respectively.

There are three characteristic locations for liquid and gas phases, including flame front (R_f) , droplet evaporation onset (R_v) and completion (R_c) fronts. For the latter two, R_v corresponds to the location where the droplets are just heated up to boiling temperature and hence start to evaporate. For $R < R_v$, the droplet temperature remains constant and evaporation continues [2,3]. The evaporation onset front R_v is always before the flame front R_f , indicating that onset of droplet vaporization spatially precedes the gaseous combustion. Moreover, R_c denotes the location at which all the droplets are critically vaporized. When $R < R_c$, no droplets are left and hence their effects on the gas phase diminish. For brevity, hereafter, we term the first (Fig. 1a) and second (Fig. 1b) cases as heterogeneous (abbreviated as "HT") and homogeneous ("HM") flames, respectively. There are four zones in both flames. Specifically, zone 1 represents the pre-vaporization zone before R_v , for homogeneous flame. As for 3, it represents post-flame evaporation zone before R_c for heterogeneous flame, and pre-flame zone without evaporation for homogeneous flame. Meanwhile, 4 is the post-flame droplet-free zone for both flames.

Governing Equations and boundary conditions

For the gaseous flames, the well-known diffusive-thermal model [4,5] is adopted. This model has been used in numerous studies on gaseous and two-phase flames [6,7]. One-step chemistry is considered, i.e. $F + O \rightarrow P$, with F, O and P being fuel vapour, oxidizer and product, respectively. Only Fuel-lean fresh mixture (i.e. equivalence ratios of fuel vapor and fuel droplets are above unity) is studied in this work and fuel vapour *F* is the deficient species. Following previous theoretical analysis for both gaseous flames and two-phase flames with dispersed liquid droplets [6], we adopt the quasi-steady state assumption in the moving coordinate system attached to the stably propagating flame front $R_f(t)$, i.e. $\eta = r - R_f(t)$. Moreover, due to relatively dilute fuel droplet concentration, their influences on the reaction zone thickness are small and therefore gaseous combustion still dominates [3,8]. In addition, due to the kinematic and thermal equilibrium between the two phases, the droplets approximately follow the carrier gas in velocity and temperature (zone 1)[9,10]. Therefore, the non-dimensional governing equations for the gaseous temperature, fuel vapour mass fraction and droplet loading can be written as

$$-U\frac{dT}{d\eta} = \frac{1}{\left(\eta + R_f\right)^2} \frac{d}{d\eta} \left[\left(\eta + R_f\right)^2 \frac{dT}{d\eta} \right] + \omega_c - q_v \omega_v \tag{1}$$

$$-U\frac{dY_F}{d\eta} = \frac{Le}{\left(\eta + R_f\right)^2} \frac{d}{d\eta} \left[\left(\eta + R_f\right)^2 \frac{dY_F}{d\eta} \right] + \omega_c - \omega_v \tag{2}$$

$$-U\frac{dY_d}{d\eta} = -\omega_v \tag{3}$$

The term ω_v in Eqs. (1)–(3) is the non-dimensional droplet evaporation rate, i.e.

$$\omega_{\nu} = \frac{\Omega(T - T_{\nu})H(T - T_{\nu})}{q_{\nu}} \tag{4}$$

Here, H(x) is the Heaviside function, which indicates the evaporation process only occurs when the droplet temperature reaches the boiling temperature.

The non-dimensional boundary conditions for both gas phase (*T* and *Y_F*) and liquid phase (*Y_d*) at the left boundary (spherical centre, $\eta = -R_f$) and the right boundary ($\eta \rightarrow +\infty$) are [2,3,8,11,12]

$$\eta = -R_f: \left(\eta + R_f\right)^2 \frac{dT}{d\eta} = -Q, \qquad \frac{dY_F}{d\eta} = 0, \ Y_d = 0$$
(5)

$$\eta \to +\infty$$
: $T = 0$, $Y_F = 1$, $Y_d = \delta$ (6)

Here δ is the initial mass loading of the fuel droplet in the fresh mixture. In this paper, the fuel droplet is treated as adding to the gaseous mixture, and the overall fuel mass fraction 1+ δ is still under fuel-lean condition. At the evaporation onset front, $\eta = \eta_{\nu}$, the gas temperature (*T*), fuel mass fraction (*Y_F*), and fuel droplet mass loading (*Y_d*) satisfy the following jump conditions [2,3,8,11,12]

$$T = T_{\nu}, \qquad [T] = \left[\frac{dT}{d\eta}\right] = [Y_F] = \left[\frac{dY_F}{d\eta}\right] = 0, Y_d = \delta \tag{7}$$

where the square brackets, i.e. $[f] = f(\eta^+) - f(\eta^-)$, denote the difference between the variables at two sides of a location. At the evaporation completion front, $\eta = \eta_c$, the jump conditions for the gas temperature (*T*), fuel mass fraction (*Y_F*), and droplet mass loading (*Y_d*) take the following form [3]

$$\begin{cases} [T] = \left[\frac{dT}{d\eta}\right] = [Y_F] = \left[\frac{dY_F}{d\eta}\right] = 0, Y_d = 0, \qquad \eta_c > 0\\ [T] = \left[\frac{dT}{d\eta}\right] = [Y_F] = \left[\frac{dY_F}{d\eta}\right] = 0, [Y_d] = 0, \qquad -R_f < \eta_c < 0 \end{cases}$$
(8)

Large activation energy of the gas phase reaction is assumed in this study. The corresponding jump conditions are

$$T = T_f, \qquad Y_F = [Y_d] = 0 \tag{9}$$

$$-\left[\frac{dT}{d\eta}\right] = \frac{1}{Le} \left[\frac{dY_F}{d\eta}\right] = \left[\sigma + (1-\sigma)T_f\right]^2 exp\left[\frac{Z}{2}\left(\frac{T_f - 1}{\sigma + (1-\sigma)T_f}\right)\right] = \chi(T_f)$$
(10)

Analytical Solutions

Governing equations (1) – (3) with proper boundary and jump conditions (i.e. Eqs. 5 – 9) can be solved analytically. The solutions for gas temperature *T*, fuel vapour mass fraction Y_F , and droplet mass loading Y_d in four zones in both homogeneous and heterogeneous flames are derived. For heterogeneous flame, the solutions of gas temperature *T*, fuel vapor mass fraction Y_F , and droplet loading Y_d from zones 1 to 4 are (the number subscripts indicate different zones as shown in Fig. 1a)

$$\begin{cases} T_{1}(\eta) = T_{v} \frac{I(\eta, U)}{I(\eta_{v}, U)} \\ T_{2}(\eta) = T_{v} + k_{1}L_{1}(\eta) + k_{2}L_{2}(\eta) \\ T_{3}(\eta) = T_{v} + \zeta_{1}L_{1}(\eta) + \zeta_{2}L_{2}(\eta) \\ T_{4}(\eta) = Q[I(\eta_{c}, U) - I(\eta, U)] + T_{v} + \kappa_{1}L_{1}(\eta_{c}) + \kappa_{2}L_{2}(\eta_{c}) \end{cases}$$
(11)

$$\begin{cases} Y_{F,1}(\eta) = 1 - \frac{DF_1(\eta_v)}{DI(\eta_v, LeU)} I(\eta, LeU) \\ Y_{F,2}(\eta) = \frac{DF_1(\eta_v)I(\eta_v, LeU) - [F_1(\eta_v) - 1]DI(\eta_v, LeU)}{DI(\eta_v, LeU)[I(\eta_v, LeU) - I(0, LeU)]} [I(\eta, LeU) - I(0, LeU)] + F_1(\eta) \\ Y_{F,3}(\eta) = G(\eta) - G(0) \\ Y_{F,4}(\eta) = -G(0) \end{cases}$$
(12)

$$\begin{cases}
Y_{d,1}(\eta) = \delta \\
Y_{d,2}(\eta) = \delta + \frac{\Omega}{Uq_{\nu}} \int_{\eta_{\nu}}^{\eta} [T_2(\eta) - T_{\nu}] d\eta \\
Y_{d,3}(\eta) = \frac{\Omega}{Uq_{\nu}} \int_{\eta_c}^{\eta} [T_3(\eta) - T_{\nu}] d\eta \\
Y_{d,4}(\eta) = 0
\end{cases}$$
(13)

where I(x, y), $L_1(\eta)$ and $L_2(\eta)$ are

$$I(x,y) = e^{-yR_f} \int_{x}^{+\infty} (\xi + R_f)^{-2} e^{-y\xi} d\xi$$
(14)

$$L_{1}(\eta) = exp\left[-\frac{(K+U)(\eta+R_{f})}{2}\right] M\left(1+\frac{U}{K}, 2, K(\eta+R_{f})\right)$$
(15)

$$L_{2}(\eta) = exp\left[-\frac{(K+U)(\eta+R_{f})}{2}\right]N\left(1+\frac{U}{K}, 2, K(\eta+R_{f})\right)$$
(16)

Here $K = \sqrt{U^2 + 4\Omega}$, M(a, b, c) and N(a, b, c) are the Kummer confluent hypergeometric function and the Tricomi confluent hypergeometric function, respectively [13],

$$M(a, b, c) = \frac{\Gamma(b)}{\Gamma(b-a)\Gamma(a)} \int_0^1 e^{ct} t^{a-1} (1-t)^{b-a-1} dt$$
(17)

$$N(a,b,c) = \frac{1}{\Gamma(a)} \int_{0}^{+\infty} t^{a-1} e^{-ct} (1+t)^{b-a-1} d\xi$$
(18)

where $\Gamma(x) = \int_0^{+\infty} t^{x-1} e^{-t} dt$. The expressions for k_1, k_2, ζ_1 , and ζ_2 in Eq. (11) are

$$k_{1,2} = T_v \frac{DI(\eta_v, U)}{I(\eta_v, U)} \frac{L_{2,1}(\eta_v)}{L'_{1,2}(\eta_v)L_{2,1}(\eta_v) - L'_{2,1}(\eta_v)L_{1,2}(\eta_v)}$$
(19)

$$\zeta_{1,2} = \frac{(T_f - T_v)L'_{2,1}(\eta_c) + QL_{2,1}(0)DI(\eta_c, U)}{L'_{2,1}(\eta_c)L_{1,2}(0) - L'_{1,2}(\eta_c)L_{2,1}(0)}$$
(20)

where $DI(x, y) = \frac{\partial I(x, y)}{\partial x} = -e^{-y(x+R_f)} (x+R_f)^{-2}$. $F_1(\eta)$ and $G(\eta)$ hold as the following form for heterogeneous flame

$$F_{1}(\eta) = -\frac{\Omega}{q_{\nu}} \int_{0}^{\eta} \frac{e^{-LeU(x+R_{f})}}{(x+R_{f})^{2}} \int_{0}^{x} e^{LeU(y+R_{f})} [T_{2}(y) - T_{\nu}] (y+R_{f})^{2} dy dx$$
(21)

$$G(\eta) = -\frac{\Omega}{q_{\nu}} \int_{\eta_c}^{\eta} \frac{e^{-LeU(x+R_f)}}{(x+R_f)^2} \int_{\eta_c}^{x} e^{LeU(y+R_f)} [T_3(y) - T_{\nu}] (y+R_f)^2 dy dx$$
(22)

 $DF_1(\eta)$ is the first order derivative of $F_1(\eta)$.

For homogeneous flame, the solutions of gas temperature *T*, fuel vapor mass fraction Y_F , and droplet loading Y_d from zones 1 to 4 are (the number subscripts indicate different zones as shown in Fig. 1b)

$$\begin{cases} T_{1}(\eta) = T_{v} \frac{I(\eta, U)}{I(\eta_{v}, U)} \\ T_{2}(\eta) = T_{v} + k_{1}L_{1}(\eta) + k_{2}L_{2}(\eta) \\ T_{3}(\eta) = T_{v} + \kappa_{1}L_{1}(\eta) + \kappa_{2}L_{2}(\eta) \\ T_{4}(\eta) = Q[I(\eta_{c}, U) - I(\eta, U)] + T_{v} + \kappa_{1}L_{1}(\eta_{c}) + \kappa_{2}L_{2}(\eta_{c}) \end{cases}$$
(23)

$$\begin{cases}
Y_{F,1}(\eta) = 1 - \frac{DF_1(\eta_v)}{DI(\eta_v, LeU)}I(\eta, LeU) \\
Y_{F,2}(\eta) = \frac{[I(\eta, LeU) - I(0, LeU)]}{I(0, LeU)} \left[F_2(\eta_v) - 1 - \frac{DF_2(\eta_v)I(\eta_v, LeU)}{DI(\eta_v, LeU)}\right] + F_2(\eta) \\
Y_{F,3}(\eta) = \frac{[I(\eta, LeU) - I(0, LeU)]}{I(0, LeU)} \left[F_2(\eta_v) - 1 - \frac{DF_2(\eta_v)I(\eta_v, LeU)}{DI(\eta_v, LeU)}\right] \\
Y_{F,4}(\eta) = 0
\end{cases}$$
(24)

$$\begin{cases}
Y_{d,1} = \delta \\
Y_{d,2} = \delta + \frac{\Omega}{Uq_v} \int_{\eta_v}^{\eta} (T_2 - T_v) d\eta \\
Y_{d,3} = 0 \\
Y_{d,4} = 0
\end{cases}$$
(25)

 $F_2(\eta)$ holds as the following form for homogeneous flame,

$$F_{2}(\eta) = -\frac{\Omega}{q_{\nu}} \int_{\eta_{c}}^{\eta} \frac{e^{-LeU(x+R_{f})}}{(x+R_{f})^{2}} \int_{\eta_{c}}^{x} e^{LeU(y+R_{f})} [T_{2}(y) - T_{\nu}] (y+R_{f})^{2} dy dx$$
(26)

while $DF_2(\eta)$ is the first order derivative of $F_2(\eta)$.

Moreover, the correlations describing flame speed U, flame temperature T_f , evaporation onset location η_v and droplet completion location η_c under different flame radii R_f are also derived. For HT flame, the correlations are

$$\begin{cases} \zeta_{1}L_{1}'(0) + \zeta_{2}L_{2}'(0) - k_{1}L_{1}'(0) - k_{2}L_{2}'(0) = \chi(T_{f}) \\ \begin{cases} \frac{DF_{1}(\eta_{v})I(\eta_{v}, LeU) - [F_{1}(\eta_{v}) - 1]DI(\eta_{v}, LeU)}{DI(\eta_{v}, LeU) [I(\eta_{v}, LeU) - I(0, LeU)]} - DG(0) \end{cases} / Le = \chi(T_{f}) \\ T_{v} + k_{1}L_{1}(0) + k_{2}L_{2}(0) = T_{f} \\ \delta + \frac{\Omega}{Uq_{v}} \int_{\eta_{v}}^{0} (T_{2} - T_{v})d\eta = \frac{\Omega}{Uq_{v}} \int_{\eta_{c}}^{0} (T_{3} - T_{v})d\eta \end{cases}$$
(27)

where $DG(\eta)$ is the first order derivative of $G(\eta)$. For HM flame, the correlations are

$$\begin{cases} -\left[Q + \frac{k_{1}L'_{1}(\eta_{c}) + k_{2}L'_{2}(\eta_{c})}{DI(\eta_{c},U)}\right]R_{f}^{-2}e^{-UR_{f}} = \chi(T_{f})\\ \left[F_{2}(\eta_{v}) - 1 - \frac{DF_{2}(\eta_{v})I(\eta_{v},LeU)}{DI(\eta_{v},LeU)}\right]\frac{DI(0,LeU)}{LeI(0,LeU)} = \chi(T_{f})\\ T_{v} + k_{1}L_{1}(0) + k_{2}L_{2}(0) = T_{f}\\ \delta + \frac{\Omega}{Uq_{v}}\int_{\eta_{v}}^{0}(T_{2} - T_{v})d\eta = \frac{\Omega}{Uq_{v}}\int_{\eta_{c}}^{0}(T_{3} - T_{v})d\eta \end{cases}$$
(28)

Results and Discussion

Analysis on the ignition of spherical spray flame will be conducted based on the correlations derived above (Eqs. 27 and 28). *Z* and σ are set to be Z = 10 and $\sigma = 0.15$ [6,9]. T_v and q_v is assumed to be 0.15 and 0.4, respectively [14].



Figure.2. Flame propagation speed as a function of flame radius at different ignition energies: (a) $\delta = 0.1$, $\Omega = 0.2$. *Le* = 1.0 and (b) $\delta = 0$ (gaseous flame).

Fig. 2(a) shows the flame propagation speed as a function of flame radius at different ignition energies for spray flame. The Lewis number is set to be 1.0. For comparison, the results from gaseous flame is also shown in Fig. 2(b). The trajectories of spray flames are considerably affected by the ignition energy. Specifically, when Q = 0, the spray flame is initiated at a flame ball (termed as the outer flame ball, marked as R_z^+) with the radius of about 1.0, and propagate outwardly towards larger radii. With increased ignition energy Q, a new flame kernel branch emerges, besides the right flame propagating branch. The flame kernel branches start at a high propagation speed, resulting from the deposited ignition energy. However, due to insufficient energy, the flame kernel propagating speed gradually decays until a flame ball is formed (U = 0). This is the inner flame ball R_{Z}^{-} , corresponding to ignition failure as the flame kernel cannot propagate outwardly. With further increased ignition energy, the flame kernel branch and the flame propagating branch move towards each other, with approaching flame ball radii from them. When the ignition energy is equal to the MIE Q_c (e.g., 0.062475 in Fig. 2a), the twin flame ball solutions coincide, i.e., $R_Z^+ = R_Z^-$, and therefore the two branches merge. For $Q > Q_c$, the flame kernel can continuously propagate outwardly and consecutively reach the spherical flames with large radii.

Comparing Figs. 2(a) and 2(b), we can see that the flame trajectories in spray flames and gaseous flames are similar, indicating that the existence of fuel droplet on overall lean mixture has limited influence on flame ignition process. The small decrease in MIE for lean spray flame could be explained as the chemical heat release of the fuel vapor from droplet evaporation is larger than the evaporative heat loss. However, this decrease is small since the droplet loading is small, while the net heat release from evaporation product is also minimal in the flame kernel

branch compared to the ignition energy input. This also leads to a higher propagation speed for spray flame when it propagates at a large flame radius (e.g., line 4 at R_f = 10 in Figs. 2a and 2b).



Figure.3. (a) Evaporation onset location, (b) evaporation completion location and (c) evaporation zone length as functions of flame radius at different ignition energies when $\delta = 0.1$, $\Omega = 0.2$ for Le = 1.0. FF: Flame front.

The novelty of our theoretical model is that it could reveal the evaporation related fronts (onset and completion) within the flame initiation and propagation process. Here we define the length of the droplet evaporation zone, $\Delta \eta = \eta_v - \eta_c$, based on the difference between the droplet onset and completion locations. The changes of the evaporation onset and completion locations and the evaporation zone length are presented in Fig. 3. When the ignition energy Q is less than Q_c (e.g., lines 1–3), the solutions for droplet onset and completion locations have left and right branches. They respectively correspond to the flame kernel and propagating branches. Both droplet evaporation onset front monotonically increases with flame radius at the flame kernel branch. Because the effects of the ignition energy fade, and accordingly the temperature gradient in the pre-heat zone is decreased. For the evaporation completion front, it is clear that at the very beginning moment, the droplet distributed in the whole domain as $\eta_c = -R_f$. Later, with the ignition energy adding, both the flame front and evaporation completion front would move outwardly. However, as the flame propagation speed rapidly decreased due to the fading of ignition energy, the flame front left behind the evaporation completion front. Hence, in the flame kernel branch, the flame would experience a transition from HT to HM. For the flame propagation branch, both evaporation onset and completion locations decrease as the spray flame expands. This is due to the gradual enhancement of the flame.

When the ignition energy equals the MIE (e.g., line 4), the two separate branches for the evaporation onset and completion locations merge, where the inner and outer flame ball solutions coincide. In this scenario, both η_c and η_v increase due to the flame kernel decaying, and then decrease towards the spherical spray flames with large radii. If the ignition energy is further increased, this non-monotonicity gradually becomes weak since the initial effect of the ignition energy is strengthened.

Fig. 3(c) shows the length of the evaporation zone as a function of flame radius. For the flame kernel branch, the evaporation zone length $\Delta\eta$ increases and quickly decreases with the flame kernel decaying process. However, in the flame propagation branch, the evaporation zone length is increased with the flame radius. When the ignition energy is larger than MIE, the evaporation length curves of the two branches merge.

Conclusions

Initiation of spherical spray flame has been theoretically studied in this work. A theoretical model has been established to describe the flame initiation process coupled with correlations to depict the nonlinear relation between the flame propagation speed, flame temperature, evaporation onset and completion front. The results on the minimum ignition of lean spray flame laden with fuel droplets shows that the extra fuel droplet adding has decrease the level of MIE. Meanwhile, the results of the evaporation fronts revealed that the flame is initiated from heterogeneous flame to homogeneous flame in the flame kernel branch despite the amount of the ignition energy.

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Nomenclature

- U normalized flame speed
- Y_F normalized fuel mass fraction
- ω_c normalized chemical reaction rate
- *R_f* normalized flame radius
- Ω heat transfer coefficient
- *T_f* normalized flame temperature
- q_v normalized latent heat of vaporization
- η_v evaporation onset front
- *Z* Zel'dovich number

- T normalized gaseous temperature
- Y_d normalized evaporation rate
- ω_v normalized evaporation rate
- Le Lewis number of fuel
- δ initial droplet loading
- T_{v} normalized boiling temperature
- η flame front attached coordinate
- η_c evaporation completion front
- σ thermal expansion ratio

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