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Thermal ignition in buoyancy-driven boundary layer flows along inclined hot plates

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1. INTRODUCTION

The criterion of ignition of combustible boundary-layer flow over a heated surface is a fundamental problem of combustion theory, tackled by experiment, series expansion, numerical simulation or asymptotic analysis in the limit of large activation energy. Because the behavior of most fires on earth is governed by buoyancy-induced convection, ignition in boundary-layer flow under natural convection is related not only to ignition phenomena influenced by natural convection on the academic side, but also to the onset of accidental fire on the practical side.

Laminar combustible boundary-layer flow along a vertical plate at a high-temperature wall serves as a prototypical problem to assess the effect of natural convection on ignition. Ignition phenomena of this buoyant boundary-layer flow were investigated numerically and experimentally [1, 2]. Ono *et al.* [1] proposed a numerical correlation $\ln[\sqrt{x_1}P^{n-1}] = E/RT_w + \text{constant to evaluate ignition dis$ tances for selected system parameters. The results predicted from this numerical correlation agreed satisfactorily with their experimental data. However, the constant in the numerical correlation was undetermined. Obviously this constant is a function of other system parameters, e.g. reactant concentrations, Pr number, etc. No complete ignition criterion for such a typical problem, to the authors' knowledge, is published, even though the method of large activation energy asymptotics is well established to obtain a theoretical ignition criterion. An asymptotic analysis is superior to numerical methods because asymptotic solutions correlate all system parameters into a single equation, and thereby facilitate physical interpretations.

The ignition phenomena of combustible boundary-layer flow under purely natural convection at various angles of inclination θ are of our interest (Fig. 1). The objective of the present work is to obtain an explicit criterion of ignition, which is a function of θ , the *Pr* number, the temperature at the wall and other properties of the combustible gas, by means of an asymptotic analysis in the limit of large activation energy.

2. FORMULATION

The approximations of a steady, two-dimensional and laminar boundary-layer flow are applicable. The other assumptions are as follows: the specific heats at constant pressure of the various species are equal to a constant; radiative heat transfer, Soret and Dufour effects are neglected; the flow is the ideal gas mixture with constant values of $\rho\mu$, $\rho\lambda$ and $\rho^2 D_i$. For thermal ignition, we assume a global onestep irreversible chemical reaction $(F + v_0 O \rightarrow \text{ products})$ with $\omega = BT^{\alpha} (\rho Y_F / W_F)^{n_F} (\rho Y_0 / W_0)^{n_0} \exp(-E/RT)$ [1-3].

With the above assumptions, the conservation equations are

$$\frac{\partial(\rho u)}{\partial x} + \frac{\partial(\rho v)}{\partial y} = 0 \tag{1}$$

$$\rho\left(u\frac{\partial u}{\partial x} + v\frac{\partial u}{\partial y}\right) = g(\rho_{\infty} - \rho)\cos\theta + \frac{\partial}{\partial y}\left(\mu\frac{\partial u}{\partial y}\right)$$
(2)

$$\rho c_{\rm P} \left(u \frac{\partial T}{\partial x} + v \frac{\partial T}{\partial y} \right) = \frac{\partial}{\partial y} \left(\lambda \frac{\partial T}{\partial y} \right) + q W_{\rm F} \omega \tag{3}$$

	NOME	NCLATUR	E
A	defined in equation (27)	β	${ ilde T}_{ m w} - { ilde T}_{ m v}$
В	frequency factor	$\dot{\Delta}$	defined in equation (29)
с	defined in equation (32)	ű.	dimensionless streamwise coordinate
$C_{\rm P}$	specific heat at constant pressure		defined in equation (18)
Ċ	defined in equation (25)	η	dimensionless transverse coordinate
D	mass diffusivity		defined in equation (9)
E	activation energy	θ	angle of inclination
f	dimensionless stream function	i.	thermal conductivity coefficient
g	acceleration of gravity	μ	viscosity coefficient
H	$=(4v_{x}^{2}/g)^{1/3}$	v	stoichiometric coefficient
n	total reaction order, $n_{\rm F} + n_{\rm O}$	v ,	kinematic viscosity outside boundary layer
n_i	reaction order of species <i>i</i>	ű.	transformed transverse coordinate defined
Р	pressure		in equation (24)
Pr	Prandtl number	ρ	density
q	specific heat of combustion	ψ	stream function
R	gas constant	0	specific reaction rate.
\$	dimensionless streamwise coordinate		
	defined in equation (9)	Subscripts	
Sc	Schmidt number		
Т	temperature	L L	fuel
$T_{\rm a}$	dimensionless activation energy, $c_{\rm P} E/Rq$	Г ;	index for species
и	streamwise velocity	1 [index for species
r	transverse velocity		avidant
W	molecular weight	0	oxidant at wall
X	coordinate along wall	w	at wan outside boundary laver
r	coordinate normal to wall	1.	outside boundary layer.
Y	mass fraction.		
		Superso	cripts
Greek symbols		~	dimensionless quantity
χ	temperature exponent		differentiation with respect to η .

$$\rho\left(u\frac{\partial Y_{\rm F}}{\partial x} + v\frac{\partial Y_{\rm F}}{\partial y}\right) = \frac{\partial}{\partial y}\left(\rho D_{\rm F}\frac{\partial Y_{\rm F}}{\partial y}\right) - W_{\rm F}c_2 \tag{4}$$

$$\rho\left(u\frac{\partial Y_{\rm O}}{\partial x} + v\frac{\partial Y_{\rm O}}{\partial y}\right) = \frac{\partial}{\partial y}\left(\rho D_{\rm O}\frac{\partial Y_{\rm O}}{\partial y}\right) - v_{\rm O}W_{\rm O}co \qquad (5)$$
$$P = \rho RT \qquad (6)$$

$$P = \rho RT$$
 (

subject to the boundary conditions

$$u = v = 0, \quad T = T_{w}, \quad \partial Y_{i}/\partial y = 0 \quad \text{at } y = 0$$
 (7)

$$u \to 0, \quad T \to T_{i}, \quad Y_{i} \to Y_{i}, \quad \text{as } y \to \infty.$$
 (8)

Dimensionless coordinates and variables are introduced as follows



Fig. 1. $x_0/x_{1,\theta=0}$ vs angle of inclination θ (radian) for varied \tilde{T}_w .

$$s = x/H, \quad \eta = s^{-1/4} \int_0^{y/H} \rho(x, y') / \rho_\infty dy'$$
 (9)

and

$$\tilde{T} = c_{\rm p} T/q, \quad \tilde{Y}_{\rm F} = Y_{\rm F},$$

$$\tilde{Y}_{\rm O} = Y_{\rm O}/(\nu_{\rm O} W_{\rm O}/W_{\rm F}), \quad T_{\rm a} = c_{\rm P} E/Rq \tag{10}$$

with $H = (4v_{\infty}^2/g)^{1/3}$. The dimensionless stream function f is defined as

$$f(s,\eta) = \psi(x,y)/4v_{x}s^{3/4},$$
 (11)

where ψ satisfies

$$\partial \psi / \partial x = -\rho v / \rho_{\infty}$$
 and $\partial \psi / \partial y = \rho u / \rho_{\infty}$. (12)

Let $f' \equiv \partial f / \partial \eta$, $Pr \equiv Pr_{\infty}$ and $Sc_i \equiv Sc_{i\infty}$. The appropriate dimensionless equations are

$$\frac{\partial^2 f'}{\partial \eta^2} + 3f \frac{\partial f'}{\partial \eta} + 2\zeta \left(\frac{\partial f}{\partial \zeta} \frac{\partial f'}{\partial \eta} - f' \frac{\partial f'}{\partial \zeta} \right) = 2f'^2 + \left(1 - \frac{\tilde{T}}{\tilde{T}_x} \right) \cos \theta$$
(13)

$$Pr^{-1}\frac{\partial^2 \tilde{T}}{\partial \eta^2} + 3f\frac{\partial \tilde{T}}{\partial \eta} + 2\zeta \left(\frac{\partial f}{\partial \zeta}\frac{\partial \tilde{T}}{\partial \eta} - f'\frac{\partial \tilde{T}}{\partial \zeta}\right) = -\zeta \tilde{\omega} \quad (14)$$

$$Sc_{i}^{-1}\frac{\partial^{2}\tilde{Y}_{i}}{\partial\eta^{2}} + 3f\frac{\partial\tilde{Y}_{i}}{\partial\eta} + 2\zeta\left(\frac{\partial f}{\partial\zeta}\frac{\partial\tilde{Y}_{i}}{\partial\eta} - f'\frac{\partial\tilde{Y}_{i}}{\partial\zeta}\right) = \zeta\tilde{\omega}, \quad i = F,O$$
(15)

associated with the boundary conditions

$$f' = 0, \quad v = 0, \quad \tilde{T} = \tilde{T}_{w}, \quad \partial \tilde{Y}_{i} / \partial \eta = 0 \quad \text{at} \quad \eta = 0 \quad (16)$$
$$f' \to 0, \quad \tilde{T} \to \tilde{T}_{\infty}, \quad \tilde{Y}_{i} \to \tilde{Y}_{i\infty} \quad \text{as} \quad \eta \to \infty, \quad (17)$$

where the dimensionless streamwise coordinate

$$x = 2 D(x + 1/2) D(x + 1) = 1 = 1 = 1 = 1 = 1 = 1 = 1$$

$$\zeta = 2B(x/g)^{1/2} (P/R)^{n-1} v_{O}^{n_{O}} W_{F}^{1-n} (q/c_{P})^{1+\alpha-n}$$
(18)

and the reaction rate

$$\tilde{\omega} = \tilde{T}^{1+\alpha-n} \tilde{Y}_{F}^{n_{F}} \tilde{Y}_{O}^{n_{O}} \exp\left(-T_{a}/\tilde{T}\right).$$
(19)

3. IGNITION ANALYSIS

As a standard procedure of asymptotic analysis [3], we seek first the solutions in the chemically frozen state. In the absence of a chemical reaction as $T_a \rightarrow \infty$, this problem admits similar solutions that depend on η only. Thus we have

$$f''' + 3ff'' = 2f'^{2} + (1 - \tilde{T}/\tilde{T}_{\infty})\cos\theta$$
(20)

$$Pr^{-1}\tilde{T}'' + 3fT' = 0 \tag{21}$$

$$Sc_i^{-1} \tilde{Y}''_i + 3f \tilde{Y}'_i = 0, \quad i = F, O$$
 (22)

with boundary conditions (16) and (17). Moreover, the boundary condition v = 0 at $\eta = 0$ (16) can be reduced to f = 0 at $\eta = 0$. This is a standard two-point boundary-value problem that is solved numerically using a computer-library subroutine (BVPFD from IMSL User's Manual, 1989) [4]. In particular, the solutions of species are simply

$$\tilde{Y}_{i,f} = \tilde{Y}_{ix}, \quad i = F, O.$$
⁽²³⁾

The location of ignition is expected to be near the wall because the temperature there is the greatest. In the limit of large activation energy, the variation of temperature from the chemically frozen state to the weakly reactive state at ignition is expected to be quite small $[O(\varepsilon)]$ [3]. As a result, the stream function f and zeroth-order concentrations of reactants are assumed to be the same as those in the chemi-

cally frozen state, when we solve the energy equation with a weakly chemical reaction term.

A new coordinate transformation is introduced as

$$\xi = \int_0^{\eta} \exp\left(-3Pr \int_0^{\eta'} f(\eta'') \,\mathrm{d}\eta''\right) \mathrm{d}\eta'/C \qquad (24)$$

with

$$C(Pr,\theta,\tilde{T}_{w},\tilde{T}_{\omega}) = \int_{0}^{\infty} \exp\left(-3Pr\int_{0}^{\eta'} f(\eta'') \,\mathrm{d}\eta''\right) \mathrm{d}\eta' \qquad (25)$$

In terms of this new transverse coordinate, equation (14) is rewritten as

$$\frac{\partial^2 \tilde{T}}{\partial \xi^2} - 2\zeta f' \operatorname{Pr} A(\eta) \frac{\partial \tilde{T}}{\partial \zeta} = -\zeta \tilde{\omega} \operatorname{Pr} A(\eta), \qquad (26)$$

where

$$A(\eta) = \left(C \exp\left(3Pr \int_0^{\eta} f(\eta') \,\mathrm{d}\eta'\right)\right)^2 \tag{27}$$

and the solution of temperature in the chemically frozen limit becomes recast into

$$\tilde{T}_{\rm f} = \tilde{T}_{\rm w} - \beta \xi \tag{28}$$

with $\beta = \tilde{T}_{w} - \tilde{T}_{\infty}$.

Equation (26) is similar to that of the previous work [3]. The detailed procedure of asymptotic analysis can be found and is not reproduced here. The key steps, which are quite standard in the asymptotic analysis of ignition, are illustrated as follows. The solution in the inner diffusive-reactive region near the wall is locally similar after introducing an inner expansion and a stretched coordinate. Besides the isothermal condition at the wall, the second boundary condition is obtained through matching with the outer diffusive-convective solution which is chemically frozen to all orders. A relevant Damköhler number is defined as

$$\Delta = 2\varepsilon \operatorname{Pr} C^2 \beta^{-2} \zeta_1 \tilde{T}_{w}^{1+\alpha-n} \tilde{Y}_{F\infty}^{n_F} \tilde{Y}_{O\infty}^{n_o} \exp\left(-T_a/\tilde{T}_w\right)$$
(29)

with a small expansion parameter $\varepsilon = \hat{T}_{w}^{2}/\theta$. Finally, this problem admits double solutions for $\Delta < 1$ but no solutions for $\Delta > 1$; thereby the Damköhler at ignition is $\Delta_{I} = 1$ [3].

According to equation (29), the corresponding dimensional ignition distance is

$$x_{1} = \left[\left(\frac{g^{1/2} \beta^{2}}{4\varepsilon BPrC^{2}} \right) \left(\frac{W_{F}^{n_{v}-1} W_{O}^{n_{o}}}{Y_{F\infty}^{n_{o}} Y_{O\infty}^{n_{o}}} \right) \\ \times \left(\frac{P}{R} \right)^{1-n} T_{w}^{n-x-1} \exp\left(\frac{E}{RT_{w}} \right) \right]^{2}.$$
(30)

4. RESULTS AND CONCLUSIONS

According to the above equation, the ignition distance x_1 becomes recast into

$$\ln\left[\sqrt{x_{\rm I}}P^{n-1}\right] = E/RT_{\rm w} + c \tag{31}$$

with the constant c as

$$c = \ln \left[\left(\frac{g^{1/2} \beta^2}{4\varepsilon B \operatorname{Pr} C^2} \right) \left(\frac{W_{\mathrm{F}}^{n_{\mathrm{F}}-1} W_{\mathrm{O}}^{n_{\mathrm{O}}}}{Y_{\mathrm{F}^{n_{\mathrm{F}}}}^{n_{\mathrm{F}}} Y_{\mathrm{O}^{n_{\mathrm{O}}}}^{n_{\mathrm{O}}} \right) R^{n-1} T_{\mathrm{w}}^{n-n-1} \right].$$
(32)

Equation (31) is identical with the experimentally verified correlation proposed by Ono *et al.* [1] for a vertical plate. Here we not only analytically determine this constant c, but also extend the application of equation (31) to all inclined plates.



Fig. 2. $x_{1/x_{1,\theta}=0}$ vs angle of inclination θ (radian) for varied *Pr* number.

For quantitative presentations, we adopted the input system parameters $n_{\rm F} = 0.15$, $n_{\rm O} = 1.65$, $\alpha = 0.0$, $\tilde{T}_{\tau} = 0.0075$, $T_{\rm a} = 0.4$ and $\tilde{Y}_{\rm Fx} = \tilde{Y}_{\rm Ox} = 0.06$. A plot of $x_{\rm I}/x_{\rm I,\theta=0}$ vs θ for varied \tilde{T}_{w} is presented in Fig. 1. Here $x_{1,\theta=0}$ is the ignition distance for a vertical plate (Pr = 1 and $\tilde{T}_{w} = 0.023$). Obviously a slightly increased wall temperature results in a substantially decreased ignition distance for a large activation energy. For fixed \widetilde{T}_{w} , the magnitude of $x_{1}/x_{1,\theta=0}$ decreases with increasing angle of inclination θ . This fact signifies that a weaker streamwise flow is induced for a greater value of θ . The value of $x_1/x_{1,\theta=0}$ vs θ for some selected *Pr* numbers is shown in Fig. 2. According to this figure, the ignition distance for a larger Pr number is smaller than that for a smaller Pr number. The Pr number is defined as a ratio of momentum diffusivity to thermal diffusivity. From the physical point of view, the thermal energy generated from the chemical reaction near the high-temperature wall is transported away from it at a smaller rate for a greater Pr number; thereby the ignition criterion is satisfied more readily and then the magnitude of ignition distance decreases.

The dependence of the ignition distance (x_i) on the accel-

eration of gravity (g), the ambient pressure (P) and the concentrations of reactants (Y_{Fz} and Y_{0z}) can be readily determined from equation (30). The result shows $x_1 \sim g P^{2(1-n)} Y_F z^{n_1} Y_O z^{n_0}$. For common hydrocarbon fuels, the total reaction order *n* exceeds unity [5]. Thus the growth of the ambient pressure results in decreasing x_1 .

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