INTERACTION OF TURBULENT PREMIXED FLAMES WITH COMBUSTION PRODUCTS: ROLE OF STOICHIOMETRY

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Abstract

Stabilization methods of turbulent flames often involve mixing of reactants with hot products of combustion. The stabilizing effect of combustion product enthalpy has been long recognized, but the role played by the chemical composition of the product gases is typically overlooked. We employ a counterflow system to pinpoint the effects of the combustion product stoichiometry on the structure of turbulent premixed flames under conditions of both stable burning and local extinction. To that end, a turbulent jet of lean-to-rich, $\text{CH}_4/\text{O}_2/\text{N}_2$ -premixed reactants at a turbulent Reynolds number of 1050 were opposed to a stream of hot products of combustion that were generated in a preburner. While the combustion product stream temperature was kept constant, its stoichiometry was varied independently from that of the reactant stream, leading to reactant-to-product stratification of relevance to practical combustion systems. The detailed structure of the turbulent flame front was analyzed in two series of experiments using laser-induced fluorescence (LIF): joint CH_2O LIF and OH LIF measurements and joint CO LIF and OH LIF measurements. Results revealed that a decrease in local $\text{CH}_2\text{O}+\text{OH}$ and $\text{CO}+\text{OH}$ reaction rates coincide with the depletion of OH radicals in the vicinity of the combustion product stream. These critical combustion reaction rates were more readily quenched in the presence of products of combustion from a stoichiometric flame, whereas they were favored by lean combustion products. As a result, stoichiometric combustion products contributed to a greater occurrence of local extinction. Furthermore, they limited the capacity of premixed reactants to ignite and of the turbulent premixed flames to stabilize. In contrast, lean and rich combustion products facilitated flame ignition and stability and reduced the rate of local extinction. The influence of the combustion product stream on the turbulent flame front was limited to a zone of approximately two millimeters from the gas mixing layer interface (GMLI) of the product stream. Flame fronts that were separated from the GMLI by larger distances were unaffected by the product stream stoichiometry.

Keywords: turbulent counterflow, premixed flames, stratification, extinction

Nomenclature

Lowercase roman

c	Progress variable
d_{nozzle}	Nozzle separation distance
$f_i(T)$	LIF signal temperature dependence for species i (i.e., OH, CO, CH ₂ O)
k_{CH_2O+OH}	CH ₂ O + OH reaction rate
$k_{CH_2O+OH}^{LIF}$	Product of CH ₂ O LIF and OH LIF signals
$k_{CH_2O+OH}^{max}$	Peak CH ₂ O + OH reaction rate
k_{CO+OH}	CO + OH reaction rate
k_{CO+OH}^{LIF}	Product of CO LIF and OH LIF signals
k_{CO+OH}^{max}	Peak CO + OH reaction rate
l'	Integral lengthscale
l_F	Unstrained laminar flame thickness
v'	RMS axial velocity
x_n	Flame-normal coordinate
z	Axial burner coordinates

Uppercase roman

CH_2O_{max}	Peak CH ₂ O LIF signal
CO_{max}	Peak CO LIF signal
GMLI	Gas mixing layer interface

Ka_t	Turbulent Karlovitz number
K_{bulk}	Bulk strain rate
K_{ext}	Extinction strain rate
K_{ign}	Ignition strain rate
OH_o	OH LIF signal for $x_n = 0$
Re_t	Turbulent Reynolds number
S_L	Unstrained laminar flame speed
SR	Strain rate for laminar flame calculation
T	Temperature
T_{ad}	Adiabatic flame temperature
T_b	Product stream temperature
V_{bulk}	Cold-inlet bulk velocity

Lowercase Greek

φ_b	Product stream equivalence ratio
φ_u	Fresh reactant equivalence ratio

Uppercase Greek

Δ	Axial distance from GMLI
Δ_f	Flame-front axial coordinate with respect to GMLI
$/\nabla OH _{max}$	OH LIF signal maximum gradient

1. Introduction

Lean premixed combustion is a straightforward strategy for reducing emissions and increasing efficiency but is difficult to implement in highly turbulent devices because of instabilities and the increased risk of blowoff [1]. Mixture stratification is an effective approach to improve overall-lean combustion stability where lean and weak flames are supported by regions of richer and more robust burning. In some systems, achieving homogeneous mixing is technically not possible and stratification, if not deliberate, is simply unavoidable. The practical relevance of stratified-premixed combustion has motivated an increasing body of work in fundamental combustion research [2]. Effects of mixture stratification are essentially twofold, depending on whether they are associated with either local or large-scale inhomogeneity of the reactant stoichiometry [3]. Studies of local reactant inhomogeneity in different laboratory configurations [4-12] have shown that stratification increases the turbulent flame speed as a result of larger turbulent flame density and curvature relative to the corresponding homogeneously premixed cases. Additionally, as flame fronts propagate across equivalence ratio gradients, front- and back-support can enhance the burning rates and extend the flammability limits [13-18]. These effects, however, seem to prevail at low turbulence intensities [19-20] and become dominated by turbulent transport as the turbulence intensity is increased [11, 21-22] or by flame instabilities for certain types of fuels and mixtures [23].

Under moderately turbulent conditions, Mastorakos [24] introduced a counterflow burner where a stream of reactants was opposed to a second stream of combustion products diluted by air. In previous studies [25-26], by introducing a new turbulence generation scheme [27], we succeeded in raising the turbulent Reynolds number to values on the order of one thousand and examined the interaction of turbulent premixed flames with hot combustion products in the same counterflow configuration. The equivalence ratio of the flame generating the products was maintained at unity while parameters controlling the fluid dynamics, such as the bulk strain rate and the turbulent Reynolds number, and the chemistry, such as the reactant equivalence ratios and the temperature of the product stream, were varied independently.

The present study focuses on large-scale stratification effects pertaining to the interaction of premixed flames with hot combustion products that originate from a flame of different stoichiometry. Figure 1 illustrates such a situation relevant to systems involving large recirculation zones such as swirl-stabilized combustors, stratified-premixed gas turbines [28-31] and trapped vortex combustors [32-35].

Combustion products from flame fronts of local equivalence ratio ϕ_b may be advected towards and interact with flame fronts of a different equivalence ratio, ϕ_u . In this study, except for some global ignition experiments, the temperature of the products was kept constant while the equivalence ratios of the reactant and product streams were varied independently. Regime diagrams of turbulent premixed combustion [36] provide a conceptual framework to systematize the turbulence-flame interactions but rely on oversimplified assumptions and neglect the effects of mixture stratification.

The role of product stoichiometry is often overlooked relative to the product temperature even in laminar flame studies. The enhancing effect of heat fluxes on heat release and extinction limits has been long recognized [37] and validated [17, 38-40]. In an adiabatic system, stoichiometric combustion products are expected to provide stronger support than either lean or rich flames at lower temperature. With a new focus on the effects of product stoichiometry, the quasi-steady extinction of strained laminar CH₄/air-premixed flames was investigated numerically using detailed chemistry and transport mechanisms in Ref. [41]. For product temperatures exceeding 1600 K, two modes of extinction occurred depending on the product stoichiometry: abrupt extinction, when the products were nearly stoichiometric and smooth extinction when the products came from a lean flame. The presence of oxygen in lean counterflowing combustion products at a mole fraction as low as 10^{-2} prevented abrupt extinction. The present study investigates these scenarios under broader conditions, including a rich product stoichiometry and turbulent flow.

Laser diagnostics were employed for simultaneous laser-induced fluorescence (LIF) imaging of either CH₂O and OH or CO and OH. For appropriate choices of the LIF excitation and detection schemes, the product of the LIF images yields in the first case a quantity proportional to the forward reaction rate of $\text{CH}_2\text{O} + \text{OH} \rightarrow \text{HCO} + \text{H}_2\text{O}$, and in the second case a quantity proportional to the forward reaction rate of $\text{CO} + \text{OH} \rightarrow \text{CO}_2 + \text{H}$. The peak reaction rate of the former reaction is correlated with the primary peak of heat release for lean-to-stoichiometric CH₄/air premixed flames [42-44]. The latter reaction is a critical step for the conversion of CO to CO₂ that accounts for a third of the heat release as the dominant contribution in the oxidation layer of premixed flames.

The following sections begin with an introduction of the flow conditions, burner and experimental methodologies. The results are divided into three sections. After discussing the global stability and ignition limits of the flames, we examine the flame structure, address the influence of the combustion products and the nature of the flame front topology for different product stoichiometries.

2. Experimental methods

2.1 Turbulent counterflow burner and experimental conditions

The counterflow configuration (Fig. 2a) consisted of a turbulent stream of premixed reactants issuing from the upper nozzle and flowing against a stream of combustion products in thermodynamic equilibrium issuing from the lower nozzle. As depicted in Fig. 2b, turbulent premixed flames ignite and stabilize on top of counterflowing combustion products. The burner design [25-26] allowed for varying independently the bulk strain rate, K_{bulk} , the turbulent Reynolds number, Re_t , the temperature of the hot product stream, T_b , and the equivalence ratios of the reactant and product streams, ϕ_u and ϕ_b , respectively. In this study, reactant and product equivalence ratios were varied using the flow conditions given in Table 1. Simultaneous $\text{CH}_2\text{O}/\text{OH}$ LIF and simultaneous CO/OH LIF measurements were performed for lean-to-stoichiometric reactants and products. Additional OH LIF measurements were acquired for rich reactants as well as rich products.

The flow rate emerging from the upper nozzle was 85 LPM, resulting in a bulk jet exit velocity of $V_{bulk} = 11.2$ m/s. The bulk strain rate, defined as $K_{bulk} = 2V_{bulk}/d_{nozzle}$ was varied from 1400/s to 2240/s by changing the nozzle separation, d_{nozzle} , from 16 mm to 10 mm. Fully developed turbulence of the reactant stream was generated by a high-blockage ratio turbulence generator plate housed inside the upper nozzle [45-46]. Turbulence properties were determined at the nozzle exit using hot-wire anemometry. The axial velocity fluctuations, v' , and the integral length scale, l' , were $v' = 3.9$ m/s and $l' = 4.1$ mm. The turbulent Reynolds number, defined as $Re_t = v'l'/\nu$, was 1,050.

For the $\text{CH}_4/\text{O}_2/\text{N}_2$ reactant mixture, the equivalence ratio, ϕ_u , was varied from 0.5 to 1.2 and the O_2/N_2 molar ratio was 30/70. The adiabatic flame temperature, T_{ad} , the laminar flame speed, S_L , and the laminar flame thickness, l_F , of the premixed flames were computed using ChemkinPREMIX [47] and the UCSD chemical kinetic mechanism [48] and are listed in Table 2. The laminar flame thickness was defined based on the maximum temperature gradient as $l_F = (T_{ad} - T_u)/(\nabla T)_{max}$ where T_u is the unburned gas temperature of 294 K. The Karlovitz number, defined as $Ka_t = (u'/S_L)^{3/2}(l_F/l')^{1/2}$, was varied by one order of magnitude from 2.7 to 29. All the premixed flames nominally belonged to the thin reaction zone regime [49-50] (Fig. 3), but with some degree of localized extinction. In the conventional regime diagram of turbulent premixed combustion, no provisions are made for the role of the product stoichiometry, and thus the flames listed in Table 2 have the same coordinates in the regime diagram of Fig. 3 regardless of the equivalence ratio of the product stream.

The stream of combustion products was produced by a $\text{CH}_4/\text{O}_2/\text{N}_2$ premixed flame that was stabilized on a preburner within the bottom nozzle and operated at an equivalence ratio, ϕ_b , of either 0.7, 1.0 or 1.2. Although the resulting hot product stream was quasi-laminar, velocity fluctuations at the bottom nozzle inlet were induced by the upper turbulent jet [26]. To maintain a constant product stream temperature, T_b , at 1850 K and vary the equivalence ratio ϕ_b , we varied the O_2/N_2 molar ratio of the preburner flame from 38/62 for $\phi_b = 0.7$, to 26/74 for $\phi_b = 1.0$ and 29/71 for $\phi_b = 1.2$. The uncertainties in the equivalence ratios and dilution are estimated to be between 1-2%.

2.2 CO + OH reaction rate imaging

Simultaneous imaging of single-photon OH LIF and two-photon CO LIF was used to measure the relative forward reaction rate of the $\text{CO} + \text{OH} \rightarrow \text{CO}_2 + \text{H}$ reaction. This diagnostic approach enabled us to probe the CO to CO_2 conversion rate in lean-to-stoichiometric $\text{CH}_4/\text{O}_2/\text{N}_2$ flames. A diagram of the experimental system, consisting of two lasers, three cameras, and the axisymmetric counterflow burner, is shown in Fig. 4a. The reaction rate imaging technique was previously described in detail elsewhere [51-53]. The forward reaction-rate, $k_{\text{CO}+\text{OH}}$, is given by $k_{\text{CO}+\text{OH}} = k(T)[\text{CO}][\text{OH}]$, where $k(T)$ is the temperature-dependent rate constant. The product of the LIF signals from CO and OH can be approximated by $k_{\text{CO}+\text{OH}}^{\text{LIF}} = f_{\text{CO}}(T)f_{\text{OH}}(T)[\text{CO}][\text{OH}]$, where the temperature dependence of the LIF signals is represented by $f_i(T)$. The pump/detection scheme determines the temperature dependence of the LIF signals and can be selected such that $f_{\text{CO}}(T)f_{\text{OH}}(T) \propto k(T)$. When this relationship is achieved, the pixel-by-pixel product of the OH LIF and CO LIF signals is proportional to the forward reaction rate, i.e. $k_{\text{CO}+\text{OH}}^{\text{LIF}} \propto k_{\text{CO}+\text{OH}}$. The fidelity of the reaction rate imaging scheme is evaluated in Section 2.4.

Laser-induced fluorescence of OH was excited by the frequency doubled output of a Nd:YAG-pumped dye laser tuned to the $Q_1(13)$ line ($\lambda = 285.88$ nm) of the $A_2\Sigma^+ \leftarrow X_2\Pi^+$ ($v' = 1, v'' = 0$) band of OH. An intensified CCD camera (512×512 pixels) with a UV camera lens ($f.l. = 45$ mm, $f/1.8$) was used to record the OH LIF signal with a spatial resolution of approximately $100 \mu\text{m}/\text{px}$. The image intensifier was gated for 500 ns bracketing the dye laser pulse. The combination of a UG11 color glass filter and a high-transmission bandpass filter ($T > 80\%$, $\lambda = 305\text{-}325$ nm) transmitted the OH fluorescence from the (0,0) and (1,1) bands and blocked out-of-band interferences. The OH LIF images were corrected for the mean spatial variations in the laser sheet and the throughput of the imaging system using OHLIF measurements in the downstream of a burner-stabilized flat premixed flame to record the beam profile.

Shot-to-shot fluctuations of a few percent were negligible for the subsection of the dye-laser beam that was used for LIF imaging. Each OH LIF image was also corrected for variations in laser energy along the direction of beam propagation resulting from absorption. For this correction, the path length that the laser beam traversed through the flame was determined from the OH LIF signal in each image, and the resulting absorption was calculated at each location along the direction of beam propagation. We verified the absorption correction by checking that the average of the corrected single-shot OH LIF images was symmetric about the burner axis. The extent of the absorption correction varied on a shot-to-shot basis depending on the topology of the turbulent flame. The maximum absorption resulted in a decrease in laser energy of 15% across the probe volume. The OH LIF signals were not corrected for local variations in quenching rates and the OH ground-state Boltzmann distribution.

Two-photon CO LIF was excited by pumping overlapped transitions in the $B_1\Sigma^+ \leftarrow X_1\Sigma^+$ ($v' = 0, v'' = 0$) Hopfield-Birge system of CO using the frequency-doubled outputs from two separate 10 Hz Nd:YAG-pumped optical parametric oscillators (OPO) (14 mJ and 16 mJ/pulse at $\lambda = 230.1$ nm), while operating in the sub-quadratic COLIF regime. A cylindrical lens was used to combine the laser beams into a 14-mm-high laser sheet. The CO fluorescence was imaged onto an intensified CCD camera (512×512 pixels) using a combination of a medium-format camera lens ($f.l. = 80$ mm, $f/2$) and a 35-mm format lens ($f.l. = 58$ mm, $f/1.2$). The imaging system included a dual-band interference filter ($\Delta\lambda = 10$ nm per band) that blocked out-of-band interference and transmitted fluorescence from the $B_1\Sigma^+ \rightarrow A_1\Pi$ ($v' = 0, v'' = 1, 2$) transitions of CO at $\lambda = 483.5$ nm and 519.8 nm. For the lean and stoichiometric flames considered here, the dual-band filter provided greater signal collection than a single-band filter with negligible interference from laser-generated C_2 Swan band emission. The spatial resolution was approximately 100 $\mu\text{m}/\text{px}$. The image intensifier was gated for 500 ns, bracketing the OPO laser pulses. Shot-to-shot variations in the beam profile were recorded by splitting off portions of the 230-nm beams after the cylindrical lens and focusing them into a cell with quartz windows containing CO and N_2 at atmospheric pressure and ambient temperature. The CO LIF signal from the cell was recorded on an un-intensified CCD camera, and the resulting images were used to correct the CO LIF measurements in the counterflow flames for beam profile variations and signal intensity variations associated with the non-uniformity of the beam thickness across the probe volume. Additional uncorrected variations in the beam profile arise from beam steering as a result of the laser beam traversing the index of refraction gradients in the turbulent flames.

The two-dimensional reaction rate measurements required subpixel matching of the camera imaging planes for OH LIF and CO LIF, which was achieved using fine dotted targets and a bilinear warping algorithm for image registration. To insure that the OH LIF and CO LIF measurement planes were coincident, we carefully overlapped the two laser beams over distances that were significantly larger than the field of view for LIF imaging.

2.3 $\text{CH}_2\text{O} + \text{OH}$ reaction rate imaging

The relative forward reaction rate of the $\text{CH}_2\text{O} + \text{OH} \rightarrow \text{HCO} + \text{H}_2\text{O}$ reaction, $k_{\text{CH}_2\text{O}+\text{OH}}$, was measured using simultaneous OH LIF and CH_2O LIF imaging in the same series of lean-to-stoichiometric $\text{CH}_4/\text{O}_2/\text{N}_2$ flames as the CO/OH LIF measurements. A diagram of the experimental system is shown in Fig. 4b. Similarly to the CO + OH reaction rate imaging, the LIF pump/detection scheme for measurement of $k_{\text{CH}_2\text{O}+\text{OH}}$ was selected such that the pixel-by-pixel product of the OH LIF and CH_2O LIF images yields a quantity proportional to $k_{\text{CH}_2\text{O}+\text{OH}}$, i.e. $k_{\text{CH}_2\text{O}+\text{OH}}^{\text{LIF}} \propto k_{\text{CH}_2\text{O}+\text{OH}}$. Simultaneous imaging of CH_2O LIF and OH LIF has previously been used to probe the primary peak of heat release rate in laminar [42-44] and turbulent premixed flames [54-58].

The OH LIF excitation and detection system was similar to that used for the CO/OH LIF measurements. The same laser system was tuned to the $Q_1(6)$ line ($\lambda = 283.01$ nm) of the $A_2\Sigma^+ \leftarrow X_2\Pi^+$ ($v' = 1, v'' = 0$) band of OH. The use of a different OH rotational level improved the proportionality between the product of the CH_2O and OH LIF signals and the corresponding reaction rate. The OH LIF measurements were corrected in the same manner as described Section 2.2.

Laser-induced fluorescence of formaldehyde was excited by the third harmonic of an injection seeded Nd:YAG laser. The diode seed laser was temperature tuned to $\lambda = 354.83$ nm to excite overlapping transitions in the 4_0^1 band of the $\tilde{A}^1A_2 \leftarrow \tilde{X}^1A_1$ system. The 355 nm laser beam was formed into a sheet that overlapped the 283 nm beam within the probe volume. The CH_2O fluorescence emission was imaged onto an intensified CCD camera using a medium-format camera lens, an AR-coated singlet ($f.l. = -400$ mm), and a 35-mm format lens. A color glass filter (GG375) in the imaging system blocked elastic scattering from the laser and transmitted the CH_2O LIF signal. In addition, the laser beam was horizontally polarized to eliminate interferences from Rayleigh or Raman scattering.

Passive, single-cavity pulse stretchers were employed in the CH_2O and OH LIF systems to approximately double the effective duration of the UV laser pulses. The longer laser pulses enabled the

use of greater laser energy, viz. 1.2 mJ/pulse for OH LIF and 42 mJ/pulse for CH₂O LIF, while operating in the linear LIF regime.

2.4 Fidelity of the LIF and reaction rate measurements

In the analysis of the turbulent counterflow flames, we use the OH/CO/CH₂O LIF signals directly as well as the pixel-by-pixel product of LIF signals to determine quantities proportional to reaction rates. Conversion of the LIF signals into species mole fraction would have required additional simultaneous measurements of temperature and other major species. The discrepancies between profiles of scaled LIF signals and species mole fractions were estimated using temperature and species data from laminar flame calculations to simulate LIF signal profiles across a laminar counterflow flame. The simulations of LIF signals were limited by the availability of detailed data on energy transfer processes, such as collisional quenching, rotational energy transfer (RET), and vibrational energy transfer (VET). The LIF simulations included calculations of variations in the Boltzmann distributions and LIF quenching rates for the excited transitions. Quenching rates for OH LIF and CO LIF were computed using the quenching cross sections from Tamura *et al.* [59] and Settersten *et al.* [60], respectively. For CO LIF, the ionization rate was estimated using the ionization cross section from Di Rosa and Farrow [61]. For simulation of CH₂O LIF signals, the temperature dependence of the Boltzmann distribution of the lower states of the 4^1_0 band of the $\tilde{A}^1A_2 \leftarrow \tilde{X}^1A_1$ system was calculated using Asyrot spectral simulation software [62]. The availability of relevant data on the temperature dependent quenching rates for CH₂O LIF is very limited. As a result, the temperature dependence was modeled using an assumed $T^{-0.5}$ dependence, as in previous work [63]. Although this assumption involves uncertainty, its use does not appreciably alter the conclusions of our study. Alternative possibilities for temperature dependence models changed the proportionality constant between the reaction rate and the product of the LIF signals, but the two quantities maintained the proportionality relationship over the relevant temperature range.

Computed profiles of OH, CO and CH₂O mole fractions for a lean, freely propagating laminar premixed flame ($\phi_u = 0.7$) are plotted in Fig. 5a. The mole fraction profiles are normalized by their respective peak values and compared to the corresponding peak-normalized simulated LIF signals in Fig. 5b. The abscissa, x_n , is centered at the location of the maximum gradient of OH mole fraction in Fig. 5a and of the computed OH LIF signal in Fig. 5b. Overall, the normalized profiles of the LIF signals are in good agreement with the corresponding mole fraction profiles. The main discrepancies are: the positions of the peak CO and CH₂O LIF signals are slightly shifted upstream toward lower temperatures

relative to the corresponding peak mole fractions; the CO LIF profile is broader than the CO mole fraction profile; and, toward the flame products, the OH LIF signal decays slightly faster than OH mole fraction. If a different temperature dependence is used for modeling the CH₂O LIF collisional quenching, the peak of the simulated CH₂O LIF signal shifts slightly.

This laminar flame calculation was also used to evaluate the fidelity of the relative reaction rate profile measurements for the LIF excitation and detection schemes described in Sections 2.2 and 2.3. The forward reaction rates, k_{CH_2O+OH} and k_{CO+OH} , are plotted in Fig. 5a, and the quantities resulting from the products of the simulated LIF signals, $k_{CH_2O+OH}^{LIF}$ and k_{CO+OH}^{LIF} , are plotted in Fig. 5b. Over the temperature range of significant CO + OH reaction (1600-2400 K), the correspondence between the k_{CO+OH}^{LIF} and k_{CO+OH} profiles is excellent. Similar agreement was observed for the other lean-to-stoichiometric flames considered here with an estimated deviation on the order of $\pm 5\%$, which is within the uncertainty of the estimate. The profile of $k_{CH_2O+OH}^{LIF}$ is also in good agreement with the CH₂O + OH reaction rate, despite a small amount of broadening and a shift of the peak location towards the reactants. In the remainder of the article, the measured reaction rates for CO + OH and CH₂O + OH will be directly denoted k_{CO+OH} and k_{CH_2O+OH} , respectively.

3. Results and discussion

3.1 Extinction/Ignition of Strained Laminar Premixed Flames

To help understand the effects of counterflowing products on turbulent flames, we first use strained laminar premixed flame calculations [47-48] to analyze the extinction and ignition of laminar flames with different product and reactant stream equivalence ratios. These calculations include counterflowing streams of premixed reactants with equivalence ratios ϕ_u equal to 0.7, 1.0 and 1.2 and combustion products with equivalence ratios ϕ_b equal to 0.7, 1.0 and 1.2 in thermochemical equilibrium at a temperature of $T_b = 1850$ K. The O₂/N₂ molar ratios were prescribed as discussed in Section 2.1. In Fig. 6, we compare the variations of peak mole fractions of OH, CO and CH₂O with those of peak heat release rates in the lean, rich and stoichiometric premixed flames as a function of the strain rate for $\phi_b = 0.7$. The strain rate is defined as $SR = 2V_{reactant}/d_{nozzle}$, where $V_{reactant}$ is the imposed velocity at the cold inlet. The ordinates are normalized by the corresponding peak values in an unstrained laminar premixed flame at the same equivalence ratios. As the strain rate increases, the peak heat release rates of all three premixed flames decrease monotonically. Peak mole fractions of OH, CO and CH₂O generally follow the same trend, but with a more gradual decay than the peak heat release rate, except for a small

overshoot of the OH mole fraction in the rich premixed flame. For the three premixed flames, CH_2O is the slowest of the three species to be depleted as the strain rate increases.

In Fig. 7, normalized peak heat release rates of the lean, rich and stoichiometric premixed flames are plotted as a function of their distance, Δ_f , from the gas stagnation plane where the axial velocity is zero. As the strain rate increases, the flame front is invariably displaced closer to the gas stagnation plane [41]. For Δ_f greater than 1.75 mm, peak heat release rates decrease very modestly for decreasing Δ_f and are relatively insensitive to variations in strain rate and product stoichiometry. The “zone of influence” of the counterflowing products can be identified for Δ_f less than approximately 1.75 mm where peak heat release rates are affected by mixing of the reaction zone with the counterflowing combustion products and the flames approach extinction differently depending on the product stream stoichiometry. For strain rates as low as 500/s, the premixed flames are displaced into the region of $\Delta_f < 1.75$ mm and, therefore, are easily affected by the counterflowing stream of combustion products.

Two modes of extinction are identified depending on the composition of the product stream: 1) abrupt extinction, corresponding to a sudden drop of peak heat release rate following an incremental increase in strain rate, and 2) smooth extinction, where the heat release rate decays monotonically as the strain rate is increased, as shown in Fig. 6. For stoichiometric and rich counterflowing combustion products, the premixed flames extinguish abruptly with extinction strain rates displayed in Fig. 8. Rich products make the flame more robust with larger extinction strain rates than the flames with stoichiometric products. This enhancement is most pronounced for the lean reactant mixture ($\phi_u = 0.7$) in which large amounts of CO from the rich product stream are readily consumed by excess O_2 in the lean reactants. The maximum strain rates at which the reactant mixture will ignite by mixing with the product stream are also shown in Fig. 8. The maximum ignition strain rates are higher for the rich products than for the stoichiometric products, indicating that ignition is hindered by stoichiometric products. In the presence of lean counterflowing products ($\phi_b = 0.7$), extinction and re-ignition occur in a smooth transition with the peak heat release rate varying monotonically as a function of the strain rate, as shown in Fig. 6. In this case, we define the extinction/ignition point as the condition where the peak heat release rate is 10% of its value in the corresponding unstrained laminar flame. Figure 8 shows that the extinction and ignition strain rates for the lean product stream are on the order of 3500/s, regardless of the reactant equivalence ratio, indicating that the back-support provided by the lean product stream is quite significant. As discussed in [41], abrupt extinction of the premixed flames result from the rapid depletion of the radical pool composed of H, OH and O. In the presence of lean products, smooth

extinction is permitted by the slower decay of the OH radical concentration. Previous numerical simulations have shown that the transition of the extinction/ignition mode from abrupt to smooth occurs for a small departure of the product equivalence ratio from stoichiometric to lean [41].

These results conclusively show that the premixed flame burning rate and extinction/ignition cannot be solely characterized in terms of strain rate. The displacement of the flame front in the vicinity of the gas stagnation plane combined with the temperature and composition of the counterflowing stream play critical roles in determining the mode of extinction and ignition, i.e. abrupt or smooth, and the onset of extinction/ignition. These results from laminar flame calculations provide a framework for investigating the effects of stratification in turbulent premixed flames.

3.2 Conditions of turbulent flame global ignition and stability

Before we examine in detail the structure of turbulent premixed flames, we first address the important role of the stoichiometry globally by considering the ignitability of turbulent premixed reactants. In Ref. [26], ignitability of reactants by stoichiometric combustion products and combustion stability were shown to increase at higher product temperatures and lower bulk strain rates. Extending these results, we consider the influence of the product stoichiometry on global ignition and stability of turbulent premixed flames. In Fig. 9, the measured ignition and stability limits of lean ($\phi_u = 0.7$), stoichiometric ($\phi_u = 1.0$) and rich ($\phi_u = 1.2$) premixed reactants are shown for counterflowing product stream temperatures, T_b , from 1500K to 2000K, and equivalence ratios, ϕ_b , from 0.7 to 1.2. For each combination of T_b, ϕ_b and K_{bulk} , we determined whether or not premixed reactants with equivalence ratio ϕ_u ignited solely by mixing with the stream of combustion products and without the support of an external ignition source. Ignition is possible for product temperatures, T_b , above the dashed/solid limiting curves. Following ignition, combustion is either pulsing (domain between dashed lines and solid lines) or stable (domain above solid lines). In the pulsing regime, combustion switches on and off spontaneously and repeatedly at frequencies between approximately 1Hz and 10Hz. This unstable regime is characterized by strong acoustic pulses as the flame brush sequentially extinguishes and reignites. In the stable regime, however, the flame brush burns steadily without intermittent blow-off.

The rationale for global flame ignition and stability is defined by the competition between the ignitability of the reactants by the counterflowing products and the ability of the turbulent flame to sustain itself [26]. As the product temperature increases, reactant ignitability increases and the global stability of the flame improves. In Fig. 9a, the global ignition and stability of lean, rich, and

stoichiometric premixed flames are compared for $K_{bulk} = 1400/s$. For the three turbulent flames, ignition and stability are most facilitated by lean combustion products and most limited in the presence of stoichiometric products. For $\phi_b = 0.7$, the stoichiometric flame is the most stable of the three flames. As ϕ_b is increased to unity, ignition of the rich turbulent flame is hampered the most. For $\phi_b > 1.0$, ignition of the rich flame requires higher product temperatures than the lean and stoichiometric premixed flames. Unlike the stoichiometric flame, the lean and rich flames operate in the pulsing regime at reduced temperatures. These turbulent flames interact more with the product stream than the stoichiometric flame. Their stability, therefore, depends on the product stream temperature and composition to a greater extent. These global flame behaviors generally follow the extinction/ignition trends observed for laminar flame calculations in Section 3.1.

In Ref. [26], for $K_{bulk} = 1400/s$, the stoichiometric premixed flame was shown to be virtually unaffected by the temperature of the stoichiometric product stream because the flame brush was located sufficiently far from the combustion products. When K_{bulk} is raised to 2240/s (Fig. 9b), ignition of a stable stoichiometric premixed flame requires noticeably higher product temperatures or leaner/richer counterflowing products. Combustion remains most limited in the presence of a stoichiometric product stream. The global stability of the stoichiometric premixed flame becomes strongly dependent on T_b and ϕ_b . The pulsing combustion limit, which does not exist for the lower strain rate case, lies approximately 100K below the stable regime limit. For $T_b - \phi_b$ conditions that are below the pulsing regime limit, ignition of the stoichiometric premixed reactants by the counterflowing products is no longer possible and the flame cannot be ignited, not even by using an external ignition source.

These experiments provide qualitative evidence that product stream stoichiometry affects the survivability of turbulent premixed flames. One would expect that detailed measurements of the flame structure should confirm this finding.

3.3 Turbulent premixed flame structure

The influence of counterflowing products on turbulent premixed flame structure and reaction rates were investigated using two independent series of CH_2O/OH LIF and CO/OH LIF measurements. For these measurements, only the equivalence ratios of the reactant and product streams were varied and the other experimental parameters were fixed, with $K_{bulk} = 1400/s$, $Re_t = 1050$ and $T_b = 1850K$ (Table 1). Under these conditions, the turbulent flames were globally stable (Fig. 9).

As shown by the laminar flame calculations in Section 3.1, monitoring the distance separating the flame fronts from the product stream is of paramount importance. To that end, we define the equivalent of the gas stagnation plane under turbulent conditions as the Gas Mixing Layer Interface (GMLI), separating the turbulent opposed streams. The new definition is required since the GMLI is not fixed in space and time, unlike the gas stagnation plane, but fluctuates as a result of turbulence. The GMLI was identified using the OH LIF images in each of the two series of LIF measurements using the technique developed in previous work [26] and described here for completeness. The GMLI was determined in each single-shot OH LIF image using the procedure illustrated in Fig. 10. In this OH LIF snapshot, cold reactants and turbulent flames are located on top of the GMLI, and the counterflowing products occupy the space below. Regions occupied by reactants, turbulent flames and products are readily identified. In the cold premixed reactants, the OH LIF signal is near zero. The combustion products issuing from the bottom nozzle contain small amounts of OH radicals and, since the counterflowing products are in thermodynamic equilibrium, the concentration of OH radicals present in the product stream is fairly homogeneous. As a result, the OH LIF signal in the product stream is relatively low and uniform. In the turbulent premixed flames, the OH LIF signal is significantly higher than in the product stream.

The location of the GMLI contour was determined from the magnitude of the gradient of the OH LIF signal. For each column of pixels in the OH LIF image, the GMLI position was identified as the first peak in the magnitude of the gradient of the OH LIF signal starting from the bottom of the image. Any subsequent peaks in the OH LIF signal gradient corresponded to flame fronts. This procedure is demonstrated in the three plots in Fig. 10 where profiles of the OH LIF signal and the magnitude of its gradient are plotted along each of the three pixel columns labelled (1), (2) and (3) in the OH LIF snapshot. The GMLI and the flame front positions are successfully identified in column (1) where the premixed flame is attached to the GMLI, in column (2) where the small region of partially extinguished premixed flame is attached to the GMLI and in column (3) where the premixed flame is detached from the GMLI and the product stream transitions directly into the reactants. Because of the convoluted flame morphology, there can be more than one flame front per pixel column, e.g. columns (2) and (3). The distance that separates the turbulent flame fronts from the GMLI is denoted as Δ_f , following the notation for the laminar flame calculations in Section 3.1

Examples of simultaneous $\text{CH}_2\text{O}/\text{OH}$ LIF and simultaneous CO/OH LIF measurements are displayed in Figs. 11a and 11b, respectively, for a lean ($\phi_u = 0.7$) turbulent premixed flame opposed to

either lean ($\phi_b = 0.7$) or stoichiometric ($\phi_b = 1.0$) combustion products. The premixed flames stabilize on top of the counterflowing combustion products and away from the nozzles, which are located at $z = \pm 8$ mm. Contours of the GMLI were determined by applying the procedure described above to each of the single-shot OH LIF images. The GMLI is unsteady and moderately corrugated as a result of the interaction between the turbulent reactant stream and the nearly laminar product stream and is typically located between $z = -4$ mm and $z = 2$ mm [26]. In laminar premixed flames, the concentration of OH rises sharply in the fuel consumption layer and decays more gradually in the oxidation layer. In turbulent flames, the flame front leading edge corresponds to the boundary of the OH LIF layer. The gradual decay of OH in the flame is not systematically observed because of the unsteady nature of the flow. The regions of high CH₂O LIF and CO LIF signals form layers aligned with the flame front contours. The thickness of these layers and the magnitude of the CH₂O and CO LIF signals are not uniform along the leading edge of the flame. The forward reaction rates are also very non-uniform along the flame front contour. Regions of high CO LIF and CH₂O LIF signals do not necessarily result in large reaction rates due to a lack of overlap with the OH LIF structures. A lower CO LIF signal is also detected in the products of the turbulent flames where the OH LIF signal is high, resulting in a low rate of oxidation of CO by OH in these regions (Fig. 11b).

To evaluate the turbulent flame front structure in more detail, we analyze one-dimensional profiles of the OH, CH₂O, CO LIF signals, as well as the values of k_{CH_2O+OH} and k_{CO+OH} that are sampled along the lines labeled (1), (2) and (3) for the case of $\phi_b = 0.7$ in Figs. 11a and 11b. These 2.5 mm-long lines are centered on the leading edges of the flames at the location of the maximum magnitude of the OH LIF signal gradient, denoted as $|\nabla OH|_{max}$, and oriented normal to the flame surface. The data along each line are interpolated between pixels, and the results are plotted in Fig. 12. The reaction rates and the LIF signals were normalized with respect to their respective ensemble-averaged values computed for $\Delta_f \geq 1.75$ mm. The abscissa of each plot, x_n , is centered at the flame leading edge and is negative toward reactants and positive toward flame products. Although Figs. 11a and 11b were obtained from separate single-shot measurements, the location of each of the three lines was chosen to sample similar structures within each image. Along line (1), the turbulent flame structure is comparable to that of a laminar premixed flame with high reaction rates. For line (2), the peaks of CH₂O and CO are located further upstream of the OH layer than in the profiles along line (1). As a result, the peak reaction rates along line (2) are approximately 50% lower than those along line (1), even though the peaks of the CH₂O LIF and CO LIF signals along line (2) are comparable to those along line (1). The reduced reaction rates

in regions near line (2) are the result of misalignments of the CH_2O and CO layers with the flame leading edge. These species misalignments originate from unsteady turbulence–flame interactions and result in partial quenching of the peak heat release rate and the $\text{CO} + \text{OH}$ reaction. These effects would not be predicted by a flamelet model based on strained laminar premixed flames such as those presented in Section 3.1 [64–65]. These observations are also affected by the three-dimensional orientation of the flame front with respect to the imaging plane, resulting in seemingly thicker flame structures at locations where the flame-normal vector is not parallel to the imaging plane.

The red arrows in the images of Figs. 11a and 11b indicate regions where the amount of OH radicals is substantially reduced near the GMLI. For the stoichiometric product stream ($\phi_b = 1.0$), OH radicals can be entirely depleted above of the GMLI. In these regions, even though some amounts of CH_2O and CO remain, the $\text{CO} + \text{OH}$ and $\text{CH}_2\text{O} + \text{OH}$ reactions are quenched as a consequence of the absence of OH . For a lean product stream ($\phi_b = 0.7$), however, a thin layer of low OH LIF signal remains present along the GMLI. In Fig. 12, the flame-normal profiles sampled along line (3) in Fig. 11 show that the measured species and reaction rates are diminished in these regions along the GMLI where the flame is partially extinguished. We deduce that the primary heat release rate and the oxidation of CO by OH are significantly reduced. These differences in flame extinction in Fig. 11 can be explained with the results of the laminar flame calculations presented in Section 3.1: flames exposed to stoichiometric products are expected to extinguish abruptly whereas the reaction rates in flames exposed to the lean products ($\phi_b = 0.7$) are expected to decay smoothly with increasing strain rate. The effects of the product stream on localized extinction will be further treated in Section 3.4.

To generalize the observations made on the particular cases examined in Fig. 11, we analyze statistics of the turbulent flame structure computed using thousands of flame-front samples from a minimum of 800 single-shot images. Figure 13a shows the mean flame-normal profiles of OH , CO and CH_2O LIF signals and associated reaction rates. Each profile was normalized by its peak value. The two mean OH LIF profiles are from the separate data sets that were acquired in combination with the CH_2O and CO LIF measurements. The close agreement between these OH LIF profiles indicates that the two sets of LIF measurements are statistically consistent and can be compared. The mean turbulent flame structure is similar to that of the freely propagating laminar premixed flame computed for an equivalence ratio of 0.7 that was presented in Fig. 5. The main difference is that all mean profiles extend over a much broader spatial domain as a result of turbulent transport and flame-front folding. The mean reaction rate profiles in Fig. 13a show that the peak reaction rate of CH_2O and OH is located closer to the reactants

than the $\text{CO} + \text{OH}$ reaction, which is the primary step intervening in the relatively slow CO oxidation. This result from the turbulent flame is consistent with the computed laminar flame profiles in Fig. 5, except that the $\text{CH}_2\text{O} + \text{OH}$ reaction in the turbulent flame persists significantly farther into the product region.

The effects of turbulence on the premixed flame structure are further analyzed in terms of the peak values of the flame-normal profiles of CO LIF, CH_2O LIF and the $\text{CO} + \text{OH}$ and $\text{CH}_2\text{O} + \text{OH}$ reaction rates, denoted as $\text{CO}_{\max}$, $\text{CH}_2\text{O}_{\max}$, $k_{\text{CO}+\text{OH}}^{\max}$ and $k_{\text{CH}_2\text{O}+\text{OH}}^{\max}$, respectively. Probability density functions (PDFs) of the positions of these peaks along the flame-normal coordinate are plotted in Fig. 13b. In analyzing Fig. 13b, one must keep in mind the small shifts of the peak CO LIF and CH_2O LIF signals toward lower temperatures relative to the peak CO and CH_2O mole fractions (Fig. 5). The PDF for $\text{CH}_2\text{O}_{\max}$ indicates that most of the CH_2O peaks are located within 0.5 mm upstream of maximum OH gradients. In contrast, the CO peak positions spread out more broadly and span approximately 0.5 mm on either side of the maximum OH gradient. In rare occurrences, $\text{CO}_{\max}$ and $\text{CH}_2\text{O}_{\max}$ can be located as far as $x_n = -0.8$ mm. The peak $\text{CH}_2\text{O} + \text{OH}$ reaction rates are located in an approximately 0.5 mm band on the reactant side of the peak OH gradients, whereas the peak $\text{CO} + \text{OH}$ reaction rates are slightly more spread out on the product side of the maximum OH gradient. The most probable position of these peaks closely coincides with the corresponding peaks in the laminar flame (Fig. 5). However, the spreading of the PDF of $k_{\text{CO}+\text{OH}}^{\max}$ relative to that of $k_{\text{CH}_2\text{O}+\text{OH}}^{\max}$ in Fig. 13b indicates that the turbulent fluctuations perturb the slow CO oxidation more than the faster reaction rates occurring in the fuel consumption layer and associated with primary heat release. Biases associated with out-of-plane flame orientation effects would not account for these differences. Overall, the turbulent reaction zones are structurally similar to laminar flames but thickened as a result of the turbulent transport, which can result in large fluctuations of the reaction rates as analyzed in the following section.

3.4 Zone of influence of the product stream on turbulent flame reaction rates

This section aims at evaluating the zone of influence of the counterflowing product stream on turbulent premixed flames by establishing relationships between the turbulent flame structure and the distance between the flame fronts and the GMLI by analogy with the discussion presented in connection with Fig. 7 on the basis of laminar flame calculations. Figures 14a and 14b show scatter plots of the maximum $\text{CH}_2\text{O} + \text{OH}$ and $\text{CO} + \text{OH}$ reaction rates, $k_{\text{CH}_2\text{O}+\text{OH}}^{\max}$ and $k_{\text{CO}+\text{OH}}^{\max}$, respectively, as a function of the flame front distance from the GMLI, Δ_f , for the lean ($\phi_u = 0.7$) turbulent premixed flame opposed

to a stream of lean ($\phi_b = 0.7$) combustion products. The reaction rates and the LIF signals are normalized by their respective ensemble averages computed for flame fronts that are well separated from the combustion products ($\Delta_f \geq 1.75$ mm). Turbulence–flame interactions cause large variations of the forward reaction rates. However, $k_{CH_2O+OH}^{max}$ and k_{CO+OH}^{max} are preferentially lower in the vicinity of the GMLI, as indicated by the decay in the conditional mean curves shown in each plot. Two regions of the flow can be differentiated depending on the flame front distance from the GMLI. For Δ_f greater than approximately 1.75 mm, the conditional averages of k_{CO+OH}^{max} and $k_{CH_2O+OH}^{max}$ are relatively constant as a function of Δ_f . However, for $\Delta_f < 1.75$ mm, the conditional averages decrease rapidly as the flame fronts approach the GMLI. As in the laminar premixed flames (Fig. 7), the counterflowing products affect burning rates of turbulent flame fronts within approximately 1.75 mm of the GMLI. Turbulent transport has seemingly no effect on the extent of the zone of influence of the product stream. In Ref. [25], strained laminar flame calculations demonstrated that this zone varies more significantly as a function of the thickness of the flame oxidation layer.

The peak locations of the CO LIF and CH₂O LIF flame-normal profiles were also monitored as a function of Δ_f for the same lean flame condition ($\phi_u = \phi_b = 0.7$). However, the flame-normal OH LIF profiles did not always have a clearly defined peak because, in some instances, the OH LIF profiles monotonically increased or reached a plateau instead of a well-defined maximum, as shown in Fig. 12. Instead of using the OH LIF signal maximum, we analyzed the OH LIF signal at $x_n = 0$, denoted OH_o , which corresponds to the location of the maximum OH LIF gradient magnitude, as a function of the distance of the flame fronts from the GMLI. In Figs. 14a and 14b, the conditional averages of OH_o decrease significantly near the GMLI. Both sets of OH LIF measurements show this decay, although there is an approximately 20% discrepancy in the conditional mean values of OH_o near the GMLI as a result of incomplete statistical convergence. In contrast to this substantial decay of OH_o , the conditional average of CO_{max} remains nearly constant and CH_2O_{max} decays only slightly near the GMLI. As a result, the depletion of OH radicals is the primary cause for the overall decrease in these reaction rates near the GMLI. This result was also observed in the single-shot images of Fig. 11 where the regions along the GMLI that were completely depleted of OH remained filled with CO and CH₂O. The lack of a significant decay in CH₂O and CO as the turbulent flame fronts approach extinction is likely the result of turbulent transport of these intermediates from neighboring regions of the flame. The lack of OH radicals, and possibly H and O, preclude further oxidation of CH₂O and CO.

The inherent temperature dependence of the LIF signals can also contribute to this persistence of larger CH_2O and CO LIF signals within the zone of influence of the product stream. In the laminar flame calculations for the lean premixed flame in Fig. 5, we showed that the peak values of the CH_2O and CO LIF signals are shifted towards lower temperatures relative to the locations of the corresponding peak mole fractions. This inherent low-temperature bias of the LIF signals may temper their rates of decay as the flame approaches the GMLI and the peak temperature decreases. We evaluate the extent of this effect by comparing the decays of the peak values of OH, CH_2O and CO mole fractions with those of the corresponding peak LIF signals in a series of strained laminar flame calculations for $\phi_u = \phi_b = 0.7$ and $T_b = 1850\text{K}$. The LIF signals are simulated using the same approach described in Section 2.4. As the strain rate increases, the flame front enters the zone of influence of the product stream for $\Delta_f < 1.75\text{ mm}$ and the peak values decay. Figure 15 shows the calculations of the normalized peak OH, CH_2O and CO mole fractions and corresponding LIF signals as a function of the distance of the flame fronts from the gas stagnation plane. All of these quantities are normalized with respect to the corresponding values in an unstrained laminar flame. The decay of the peak OH LIF signal with decreasing Δ_f in Fig. 15 is in good agreement with that of the computed peak OH mole fraction and is similar to the decay of the conditional average of OH_o in Fig. 14. For CO and CH_2O , the peak mole fractions decay faster than the respective LIF signals. This effect is significantly more pronounced for CH_2O LIF than for CO LIF because of the stronger temperature dependence of the CH_2O LIF signal. Although the LIF temperature dependence contributes somewhat to the observed slower decay of the CH_2O and CO LIF signals in the zone of influence of the product stream, it does not explain the extensive persistence of the CH_2O and CO LIF signals observed in Fig. 14 as OH disappears and the turbulent flame extinguishes near the GMLI.

This analysis supports the conclusion that much of the excess CH_2O and CO LIF signals near the GMLI arise from turbulent transport of these species. In strained laminar premixed flames (Fig. 6), on the other hand, the CH_2O and CO mole fractions decrease along with the OH mole fraction as the flames approach extinction, although CH_2O decays less rapidly than CO and OH. Thus, the comparison of these profiles between laminar and turbulent conditions highlights a case in which the similarity of flame structure is violated.

The response of the turbulent flames within the zone of influence of the product stream depends on the stoichiometry of the reactant and product streams. We analyze the effects of stoichiometry on the response of the $\text{CO} + \text{OH}$ reaction rate in the near ($\Delta_f < 1.75$) and far ($\Delta_f \geq 1.75\text{ mm}$) zones with respect

to the separation distance of the flame front from the GMLI. For conciseness, the analysis is limited to the CO+OH reaction rate, but similar results are obtained for the CH₂O+OH reaction rate since the effect of OH radical depletion is common to both reactions, as shown in Figs. 14a and 14b. Figure 16 shows PDFs of k_{CO+OH}^{max} for three turbulent premixed flames ($\phi_u = 0.6, 0.8$ and 1.0) and two product stream equivalence ratios ($\phi_b = 0.7$ and 1.0). The values of k_{CO+OH}^{max} are normalized with respect to the corresponding ensemble-averaged value for $\Delta_f \geq 1.75$ mm. Figures 16a and 16b compare PDFs of k_{CO+OH}^{max} conditional on Δ_f being greater and smaller than 1.75 mm, respectively. For $\Delta_f \geq 1.75$ mm (Fig. 16a), the distributions of k_{CO+OH}^{max} for lean and stoichiometric products are approximately the same. This result confirms that the reaction rates in the far zone are not affected by the counterflowing combustion products since the flame front is sufficiently far from the GMLI. In addition, the distribution of k_{CO+OH}^{max} broadens in the leaner flames, indicating that turbulence has a stronger effect on the reaction rates of leaner flames, as expected on the basis of their larger Karlovitz numbers. The PDFs of k_{CO+OH}^{max} in Figure 16b show that the distributions of k_{CO+OH}^{max} conditioned on $\Delta_f < 1.75$ mm are shifted toward smaller values than the corresponding PDFs for the far zone in Fig. 16a. This shift increases for leaner reactant mixtures and counterflowing products. Partial-extinction characterized by low values of k_{CO+OH}^{max} is more likely in the presence of a lean product stream. Although localized extinction also occurs near the GMLI for a stoichiometric product stream, the PDFs for $\phi_b = 1.0$ in Fig. 16b do not show as large a shift towards low values of k_{CO+OH}^{max} . This result indicates that there is a relatively low probability of partial extinction and the localized extinction is abrupt for $\phi_b = 1.0$. The effect of ϕ_b on k_{CO+OH}^{max} is more important in the leanest ($\phi_u = 0.6$) of the three turbulent premixed flames in Fig. 16b, suggesting that the turbulent flame brush of such a flame is the most exposed to the counterflowing products, as further elaborated in Section 3.5.

The overall effects of the reactant and product stream stoichiometries on the turbulent flame are very sensitive to the proximity of the flame front to the GMLI. If the flame front is in the far zone, the interaction with the product stream is minimal and the reaction rates are relatively insensitive to variations in the equivalence ratio. However, the stratification between reactant and product streams becomes significant when the flame fronts cross into the zone of influence near the GMLI. In this case, the equivalence ratios determine the extinction mode (abrupt vs. smooth) as well as the probability of localized extinction.

3.5 Role of product stoichiometry on the reaction progress variable

The effects of reactant-product stratification on the mean reaction progress variable of the turbulent flames are investigated with respect to variations in the stoichiometries of the reactant and product streams. The analysis was performed for the flow conditions and the range of reactant and product equivalence ratios reported in Table 1. Following the same approach as in Ref. [26], we define a binary progress variable, c , to be zero in the reactants and unity in regions containing products from the turbulent flame. The progress variable is determined using the OH LIF images to identify the boundary between reactants and flame products, as demonstrated in Fig. 10. In Fig. 17, mean progress variable profiles for a lean ($\phi_u = 0.6$) turbulent premixed flame are contrasted with those of the stoichiometric premixed flame for lean-to-rich ($\phi_b = 0.7$ -to-1.2) streams of combustion products. They are plotted as a function of the axial distance from the GMLI, Δ (Fig. 2b). The mean progress variable represents the probability that turbulent flame products are located at a distance Δ from the GMLI. It is indicative of the extent of separation of the turbulent flame brush from the GMLI and thus from the product stream. Additionally, as demonstrated in Section 3.4, depletion of the OH LIF signal along the GMLI correlates with localized extinction and, so, the OH LIF-based progress variable, c , can also be employed to quantify the amount of localized extinction at the GMLI. For intermittent localized extinction along the GMLI, the conditional mean progress variable is less than unity and the probability of extinction at the GMLI is given by its complement to unity, $1 - \langle c | \Delta = 0 \rangle$.

The lean premixed flame ($\phi_u = 0.6$) strongly interacts with the stream of combustion products and is therefore very sensitive to the composition of the product stream. Near the GMLI, the amount of localized extinction varies quite substantially as a function of ϕ_b : from nearly no extinction when $\phi_b = 1.2$ to a probability of localized extinction of approximately 75% when $\phi_b = 1.0$. The flame is significantly enhanced by both lean and rich combustion products, with rich products having a stronger supporting effect on the burning rate than lean products. The effects of the product stream stoichiometry are the strongest within a couple of millimeters from the GMLI where the profiles of the mean progress variable vary the most as a function of ϕ_b . The mean progress variable of the stoichiometric premixed flame, on the other hand, shows very little sensitivity to variations in the stoichiometry of the product stream. Accordingly, the mean progress variable profiles for stoichiometric flame are only shown for $\phi_b = 0.7$, 1.0 and 1.2. Near the GMLI, variations are small and, away from the GMLI, these three profiles are identical, suggesting that the flame is self-sustained. The stoichiometric turbulent premixed flame stands sufficiently far away from the GMLI because of the larger flame speed and, as a result, it is not significantly affected by the composition of the product stream.

The contrasting behavior of the lean and stoichiometric flames in Fig. 17 depends on both the probability of localized extinction and the ignitability of the reactants by the hot product stream. Note that no attempt was made to force the ignition of the premixed flames with an external source of heat; ignition of the flames was only permitted by mixing of the reactants with the product stream. The effect of product stream temperature on the reactant ignitability was previously studied for a single product stream composition of $\phi_b = 1.0$ [26]. However, the effect of product stream stoichiometry on the ignitability must also be considered. In Sections 3.1 and 3.2, we showed that the ignition of premixed flames by the combustion product stream is enhanced for product equivalence ratios away from stoichiometric conditions. These observations are consistent with the behavior of the lean ($\phi_u = 0.6$) premixed flame in the vicinity of the GMLI in Fig. 17: the lower ignitability of lean reactants by a stoichiometric product stream contributes to the low probability of lean flame fronts near the GMLI. In contrast, the rich and lean product streams increase the ignitability of the lean reactants and thereby support the presence of a lean flame front near the GMLI. The presence of flame fronts away from the GMLI and outside the zone of influence of the product stream remains restrained by the ability of the premixed flames to compete with the strong adverse bulk strain rate.

These effects are further analyzed for a broader range of reactant stoichiometries in Fig. 18 where the mean progress variable profiles of lean-to-rich turbulent premixed flames ($\phi_u = 0.5, 0.7, 1.0, 1.1$ and 1.2) are compared for lean ($\phi_b = 0.7$), stoichiometric ($\phi_b = 1.0$) and rich ($\phi_b = 1.2$) streams of combustion products. Relative to the stoichiometric product stream case ($\phi_b = 1.0$), the lean ($\phi_b = 0.7$) and rich ($\phi_b = 1.2$) products enhance the combustion of all the turbulent premixed flames such that the probability of localized extinction near the GMLI decreases and the turbulent flame brush extends further from the GMLI. For $T_b = 1850\text{K}$, lean and rich products increase the ignitability of reactants for all stoichiometries (Fig. 8). The mean progress variable of the stoichiometric flame is always the largest, indicating that none of the lean and rich premixed flames extend as far from the GMLI as the stoichiometric flame. The mean progress variable profiles for the lean and rich premixed flames have a much stronger dependence on the product stream stoichiometry than that of the stoichiometric flame. Lean combustion products strongly enhance the rich premixed flames, which are much weaker in the presence of a stoichiometric product stream. For example, the entire mean progress variable profile for the rich premixed flame ($\phi_u = 1.2$) is near zero for $\phi_b = 1.0$, indicating that the flame is mostly extinguished. However, combustion of these rich reactants is supported by the lean product stream ($\phi_b = 0.7$), resulting in a robust flame with almost no extinction, as indicated by the near unity

value of the progress variable at the GMLI in the left plot of Fig. 18. Reciprocally, lean premixed flames are well sustained by a rich product stream, as seen in the right plot of Fig. 18. The combined results in Figs. 17 and 18 demonstrate conclusively the substantial sensitivity of turbulent premixed combustion to the stoichiometry of the product stream, with the laminar computations helping to explain the observed flame behaviors in terms of their ignition and extinction limits.

4. Conclusion and implications

The interaction of a turbulent stream of premixed reactants at ambient temperature with high-temperature combustion products was studied in the counterflow configuration. The focus was on the effects of the stoichiometry, which was varied in either stream independently of the other to mimic stratification effects occurring in practical systems. The principal conclusions follow:

- 1) Qualitatively, the global ignitability and stability of turbulent flames, regardless of the stoichiometry of the reactant mixture, are facilitated strongly by lean combustion products, mildly by rich products and most limited in the presence of stoichiometric products.
- 2) These results are in part confirmed by detailed measurements of the structure of the turbulent flames using $\text{CH}_2\text{O} + \text{OH}$ and $\text{CO} + \text{OH}$ reaction rates. These measurements showed the influence of the stoichiometry of the product stream when the turbulent brush is positioned within a “zone of influence” that extends approximately a couple of millimeters from the Gas Mixing Layer Interface (GMLI) between the two streams.
- 3) In particular, a decrease in local $\text{CH}_2\text{O} + \text{OH}$ and $\text{CO} + \text{OH}$ reaction rates coincided with the depletion of OH radicals in the vicinity of the combustion product stream. In regions where partial or complete localized extinction occurred, neither CH_2O nor CO decayed appreciably as a result of turbulent transport of these intermediates from neighboring regions of the flame. The $\text{CH}_2\text{O} + \text{OH}$ and $\text{CO} + \text{OH}$ reactions appeared to be more readily quenched in the presence of products of combustion from a stoichiometric flame whereas they were favored by lean combustion products.
- 4) The effect of the product stream composition is confirmed by calculations on laminar flames of the same composition, showing that the product stream composition affects the extinction and its mode (smooth or abrupt) within a zone of influence close to the gas stagnation plane comparable to that found in turbulent flames. As a result, laminar flame calculations provide remarkable

guidance in the interpretation of the present results with the fundamental structure of a laminar flame not being disrupted by the turbulence even in the presence of localized extinction.

- 5) Even though under some conditions the turbulent flame is locally extinguished and has a complex morphology in instances of flame folding, its structure is not fundamentally altered with respect to the laminar counterpart, except for the inevitable broadening of profiles as a result of turbulent transport.

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Tables

Table 1 – Experimental conditions.

K_{bulk}	1400 s ⁻¹
Re_t	1050
T_b	1850K
ϕ_u	0.5 – 1.2
ϕ_b	0.7, 1.0, 1.2

Table 2. Turbulent $CH_4/O_2/N_2$ premixed flame properties for $Re_t = 1050$ ($u' = 3.9$ m/s and $l' = 4.1$ mm).

For all the flames, the O_2/N_2 molar ratio is 30/70.

ϕ_u	T_{ad} (K)	S_L (g/hr)	S_L (cm/s)	u'/S_L	l_F (mm)	l'/l_F	Ka_t
1.20	2522	302	74.4	5.2	0.27	15.3	3.1
1.10	2549	322	79.1	4.9	0.25	16.0	2.7
1.00	2534	324	79.1	4.9	0.25	16.1	2.7
0.90	2482	303	73.5	5.3	0.26	15.6	3.1
0.80	2390	268	64.7	6.0	0.28	14.5	3.9
0.70	2255	217	52.1	7.5	0.32	12.8	5.7
0.60	2076	156	37.3	10.5	0.39	14.0	10.5
0.50	1854	92	21.7	18.0	0.57	7.1	28.5

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Figure 16- PDFs of k_{CO+OH}^{max} for different combinations of reactant and product streams conditioned on the proximity of the turbulent flame fronts to the GMLI: a) $\Delta_f \geq 1.75$ mm, b) $\Delta_f < 1.75$ mm. For each flame condition, values of k_{CO+OH}^{max} are normalized with respect to its ensemble-average for $\Delta_f \geq 1.75$ mm.

Figure 17 - Effect of productstream equivalence ratio ϕ_b on the mean progress variable of turbulent premixed flames of equivalence ratios $\phi_u = 0.6$ (solid lines) and $\phi_u = 1.0$ (Dashed lines). Vertical red dashed line indicates the approximate boundary of the zone of influence of the product stream on the reaction rates.

Figure 18 - Effect of product stream equivalence ratio ϕ_b on the OH-based mean progress variable of turbulent premixed flames of equivalence ratios ϕ_u equal to 0.5, 0.7, 1.0, 1.1 and 1.2. Vertical red dashed line indicates the approximate boundary of the zone of influence of the product stream on the reaction rates.