

Stephan Kruse

Soot Formation of Gasoline Surrogate Components in Laminar and Turbulent Flames

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Rußbildung von Benzin-Surrogat-Komponenten in laminaren und turbulenten Flammen

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Stephan Kruse

Berichter: Universitätsprofessor Dr.-Ing. Heinz Günter Pitsch
 Professor Bassam Dally, PhD

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Zusammenfassung

Direkt-einspritzende (DI-) Ottomotoren bieten ein großes Potential hinsichtlich der Effizienzsteigerung, allerdings stellen hierbei die verschärften Rußemissionsgrenzwerte eine enorme Herausforderung dar. Um die Rußemissionen von DI-Ottomotoren zu minimieren, sind ein detailliertes Verständnis und genaue Modelle der Rußbildung von Benzinkraftstoffen essentiell. Gegenstand dieser Arbeit ist daher die grundlegende Untersuchung der Rußbildung von Benzinsurrogat-Komponenten.

In einer Gegenstromkonfiguration wurde der Rußvolumenbruch in Ethylen-, n-Heptan-, iso-Oktan- und Toluolflammen mittels Laser-induzierter Inkandescenz (LII) gemessen. Um die Messgenauigkeit der LII-Technik in Gegenstromflammen zu erhöhen, wurde ein Verfahren entwickelt, das es ermöglicht, die Auswirkungen von lokalen Schwankungen der Laserenergiedichte signifikant zu reduzieren.

Die Rußvolumenbrüche wurden daraufhin im Gegenstrombrenner für verschiedene Streckungsraten und Mischungsbrüche experimentell bestimmt. Dabei weisen die einzelnen Kraftstoffe signifikante Unterschiede in den Sensitivitäten des Rußvolumenbruchs bezüglich der Streckungsrate und des stöchiometrischen Mischungsbruchs auf. Eindimensionale Simulationen zeigen, dass die Rußbildung in Ethylenflammen von verschiedenen Modellen sehr gut vorhersagt wird, während die berechneten Rußemissionen der Benzinkraftstoffkomponenten bei allen Modellen deutlich über den experimentellen Werten liegen. Analysen der Spezieskonzentrationen in einer iso-Oktan- und einer Ethylenflamme zeigen, dass die Konzentrationen der Rußvorläufer in den iso-Oktanflammen im Vergleich zu den Ethylenflammen erheblich erhöht sind und aus den Pyrolyseprodukten des Kraftstoffs stammen.

Schließlich wurde in einer turbulenten Toluolflamme mittels verschiedener Lasermesstechniken Rußvolumenbruch, Rußpartikelgröße, Temperatur und Reaktionszone simultan bestimmt. Kombinierte Statistiken der charakteristischen Ruß- und Flammengrößen zeigen, dass Ruß aufgrund differentieller Diffusion in Gebieten niedrigerer Temperatur auf der brennstoffreichen Seite des OH-Gebiets präsent ist. Das Eindringen von Ruß in das OH-Gebiet legt nahe, dass entgegen häufiger Annahme die Rußoxidationsraten begrenzt sind und Ruß beim Eintritt in das Reaktionsgebiet nicht unmittelbar oxidiert.

Abstract

Stringent emission regulations aim to restrict soot emissions by particle mass and particle number in the transport sector. Whilst the concept of direct-injection gasoline (GDI) engines is very promising with respect to efficiency, complying with the legislative soot emission limits is challenging for GDI-engines. A sufficient reduction of soot emissions in GDI engines requires a detailed understanding and accurate modeling of soot formation processes of gasoline fuels in engines. This study provides fundamental investigations of the soot formation process of common gasoline surrogate components.

First, soot volume fraction measurements in ethylene, n-heptane, iso-octane, and toluene counterflow flames were performed by means of laser-induced incandescence (LII). This study demonstrates that large gradients in temperature and local gas composition, which are inherently present in counterflow flames, induce substantial beam steering effects. An approach is developed to ensure accurate LII measurements based on tailoring the mean laser fluence. The collected LII signal in the counterflow flames is thus nearly independent of beam steering effects.

The soot volume fraction profiles in the counterflow flames were then determined for a wide range of strain rates and stoichiometric mixture fractions. Sensitivities of soot formation on strain rate and stoichiometric mixture fraction were determined for each fuel, which are found to be strongly fuel-dependent. Simulations of the flames reveal that applied models are capable of predicting the soot volume fraction with remarkable accuracy for ethylene. For the gasoline surrogate components, however, the overall soot volume fractions are overpredicted. A reaction pathway analysis suggests that, in these flames, more soot precursors are formed via the reaction pathways involving fuel pyrolysis products.

Finally, flame structure, local gas temperature, local soot volume fraction, and primary soot particle diameter were simultaneously detected by means of optical diagnostics in turbulent toluene flames. Joint statistics of flame and soot properties indicate that, due to differential diffusion of soot, high soot concentrations are present at conditions of low temperatures and low OH concentrations. In the soot oxidation region, the presence of large particles suggests that oxidation is not sufficiently fast to burn soot completely.

Publications

This thesis is mainly based on the following journal publications.

- S. Kruse, A. Wick, P. Medwell, A. Attili, J. Beeckmann, and H. Pitsch. Experimental and numerical study of soot formation in counterflow diffusion flames of gasoline surrogate components. Submitted to *Combustion and Flame*
- S. Kruse, P. Medwell, J. Beeckmann, and H. Pitsch. The significance of beam steering on laser-induced incandescence measurements in laminar counterflow flames. *Applied Physics B*, doi.org/10.1007/s00340-018-7072-0, 2018
- S. Kruse, J. Ye, Z. Sun, A. Attili, B. Dally, P. Medwell, and H. Pitsch. Experimental investigation of soot evolution in a turbulent non-premixed prevaporized toluene flame. *Proceedings of the Combustion Institute*, doi.org/10.1016/j.proci.2018.05.075, 2018
- S. Kruse, M. S. Mansour, A. M. Elbaz, E. Varea, G. Grünefeld, J. Beeckmann, and H. Pitsch. Evaluation of partially premixed turbulent flame stability from mixture fraction statistics in a slot burner. *Combustion Science and Technology*, doi:10.1080/00102202.2018.1452393, 2018
- S. Kruse, B. Kerschgens, L. Berger, E. Varea, and H. Pitsch. Experimental and numerical study of MILD combustion for gas turbine applications. *Applied Energy*, 148:456-465, 2015

Contents

Title	i
Acknowledgments	iii
Zusammenfassung	iv
Abstract	v
Publications	vi
Contents	vii
1 Introduction and Motivation	1
2 Fundamental aspects	5
2.1 Counterflow diffusion flames	5
2.2 Physics and chemistry of soot formation	9
2.3 Energy and mass balance of laser-heated particles	11
3 Burner setups	18
3.1 Counterflow burner	18
3.1.1 Design of counterflow burner	20
3.1.2 Uncertainty quantification of strain rate and fuel mass fraction	23
3.1.3 Counterflow setup verification	27
3.2 McKenna type burner	29
3.3 Jet in hot coflow burner	30

4	Optical diagnostics	34
4.1	Non-intrusive soot diagnostics in laminar flames	34
4.1.1	Optical setup for LII measurements	35
4.1.2	Approach for LII signal quantification	37
4.1.3	Verification of LII setup in a target flame	39
4.2	Effect of beam steering on LII in counterflow flames	40
4.2.1	Beam steering in premixed flat-flames	42
4.2.2	Beam steering in counterflow flames	45
4.2.3	Compensation of beam steering in counterflow flames	50
4.2.4	Evaluation of LII accuracy in counterflow flames	55
4.3	Simultaneous measurements in turbulent flames	58
4.3.1	Optical setup for soot measurements in the JHC configuration	59
5	Application	63
5.1	Soot measurements in laminar counterflow flames	64
5.1.1	Experimental and numerical approach	66
5.1.2	Soot profiles in counterflow flames	68
5.1.3	Dependence of soot formation on strain rate and mixture fraction	72
5.1.4	Measured and computed soot formation in ethylene flames	79
5.1.5	Measured and computed soot formation of surrogate components	81
5.1.6	Analysis of soot formation in ethylene and iso-octane flames	83
5.2	Soot formation and oxidation in turbulent flames	88
5.2.1	Mean soot distribution and flame structure of turbulent toluene flames	89
5.2.2	Statistical analysis of soot-OH interaction in turbulent flame	92
6	Concluding remarks	99
	Bibliography	103