

# Studies of Evaporating Sprays; Issues & Diagnostic Tools

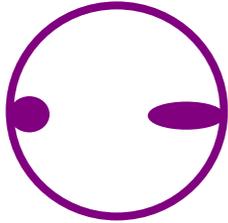
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***IEA Task Leaders Meeting, August 2006***

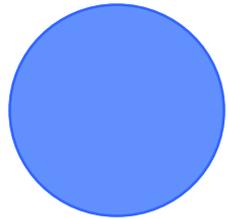
# Why Evaporating Sprays?

- Our motivation was initially driven by the need to understand the behaviour of the latest technology Gasoline Direct Injection fuel injection into realist temperatures and pressures (typically up-to ~15 bar and ~650K).
- The latest trend, similar to diesel engines, is to have spray-guided mixing where the injected fluid initially comprises of high momentum small droplets (~ 10  $\mu\text{m}$  dia).
- Such sprays are typically very dense and limit the application of conventional methods such as PDA; further PDA is poor at measuring the liquid volume fraction.
- We explore the application of combined Mie-LIF to these sprays to measure average SMD diameters using the LSD approach.

# Laser Sheet Dropsizing



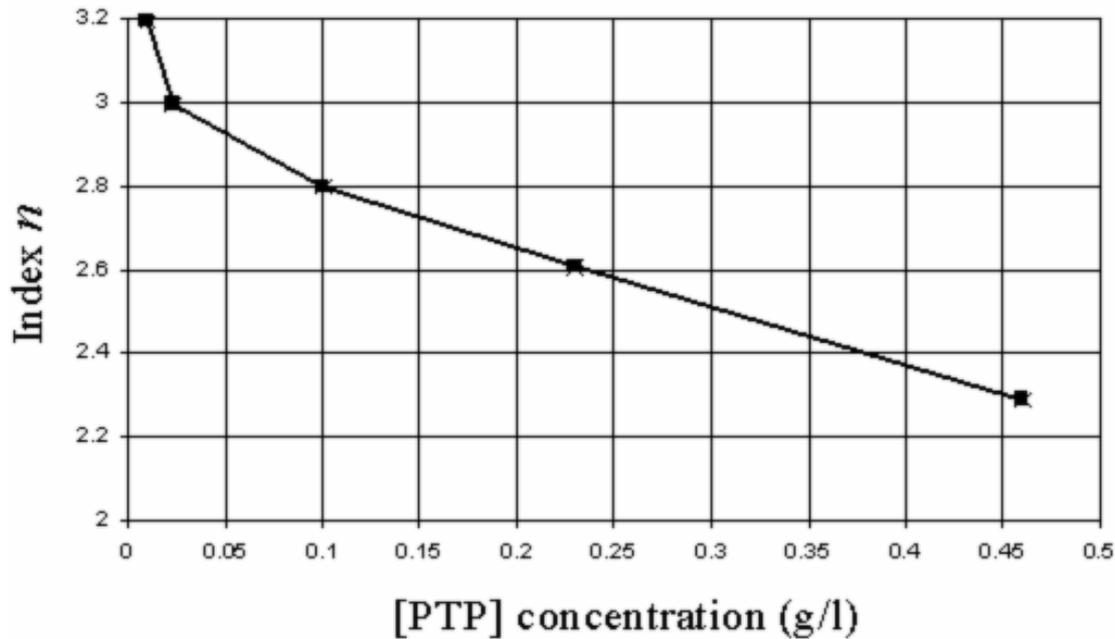
**Mie Scattering** →  $d^2$ .



**LIF from optimum fluorophor conc**

→  $d^3$ .

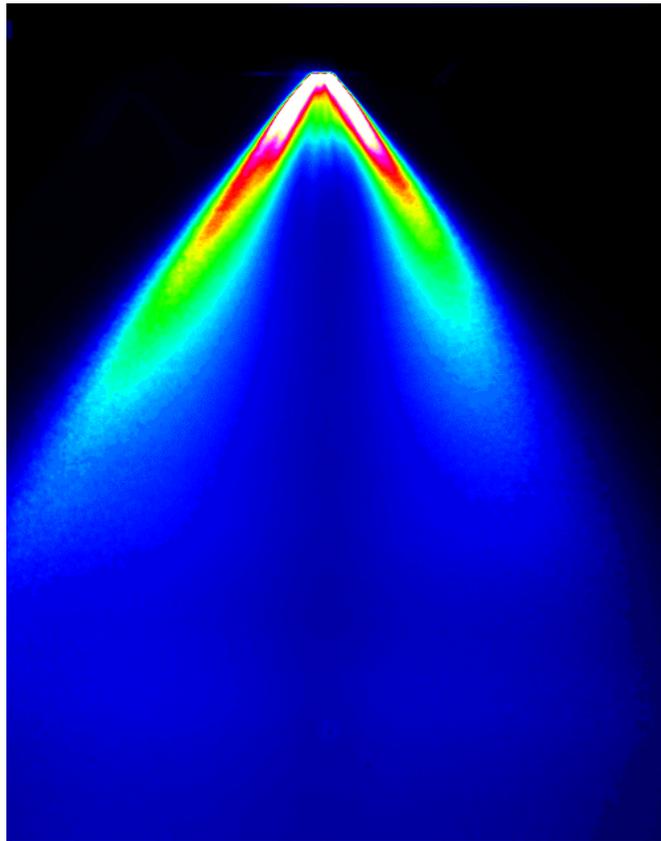
$$S_{LSD} = \frac{S_{LIF}}{S_{Mie}} = \frac{C_{LIF} \sum_i d_i^3}{C_{Mie} \sum_i d_i^2} \propto \frac{\sum_i d_i^3}{\sum_i d_i^2} \equiv d_{32}, \text{ Sauter Mean Diameter.}$$



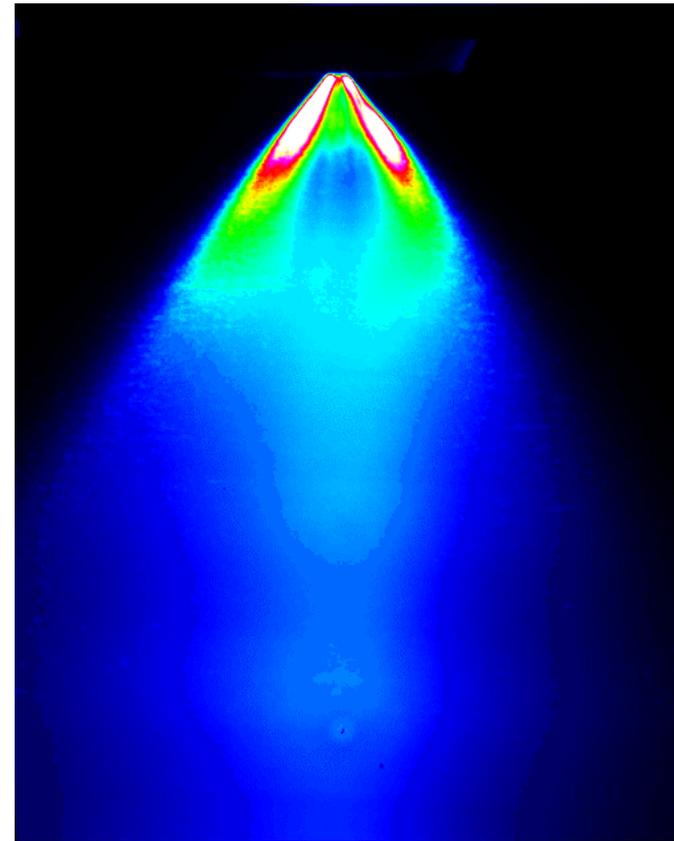
$$S_{LIF} \propto d_p^n$$

- Droplet LIF varies with the fluorescent agent concentration.
- High concentrations result in laser absorption mainly in the front surface of the droplet and therefore  $\Rightarrow$  “ $d_p^2$ ” dependence.
- Low concentrations (optically thin) should give a “ $d_p^3$ ” dependence, **but** “high gain” dyes give superadiant emission(ASE) and this generates more emission from larger droplets resulting in a “ $d_p^{3+\delta}$ ” dependence [N.B. very small droplets can vary from this behaviour].

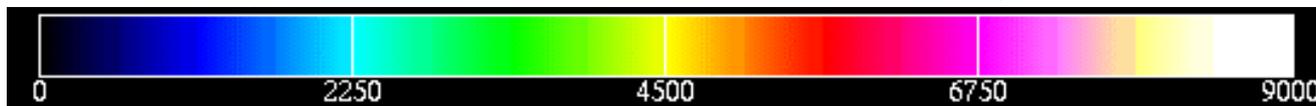
## Mean LIF & Mie images from a Delavan pressure swirl atomiser.



Mean LIF (3,000 laser shots)

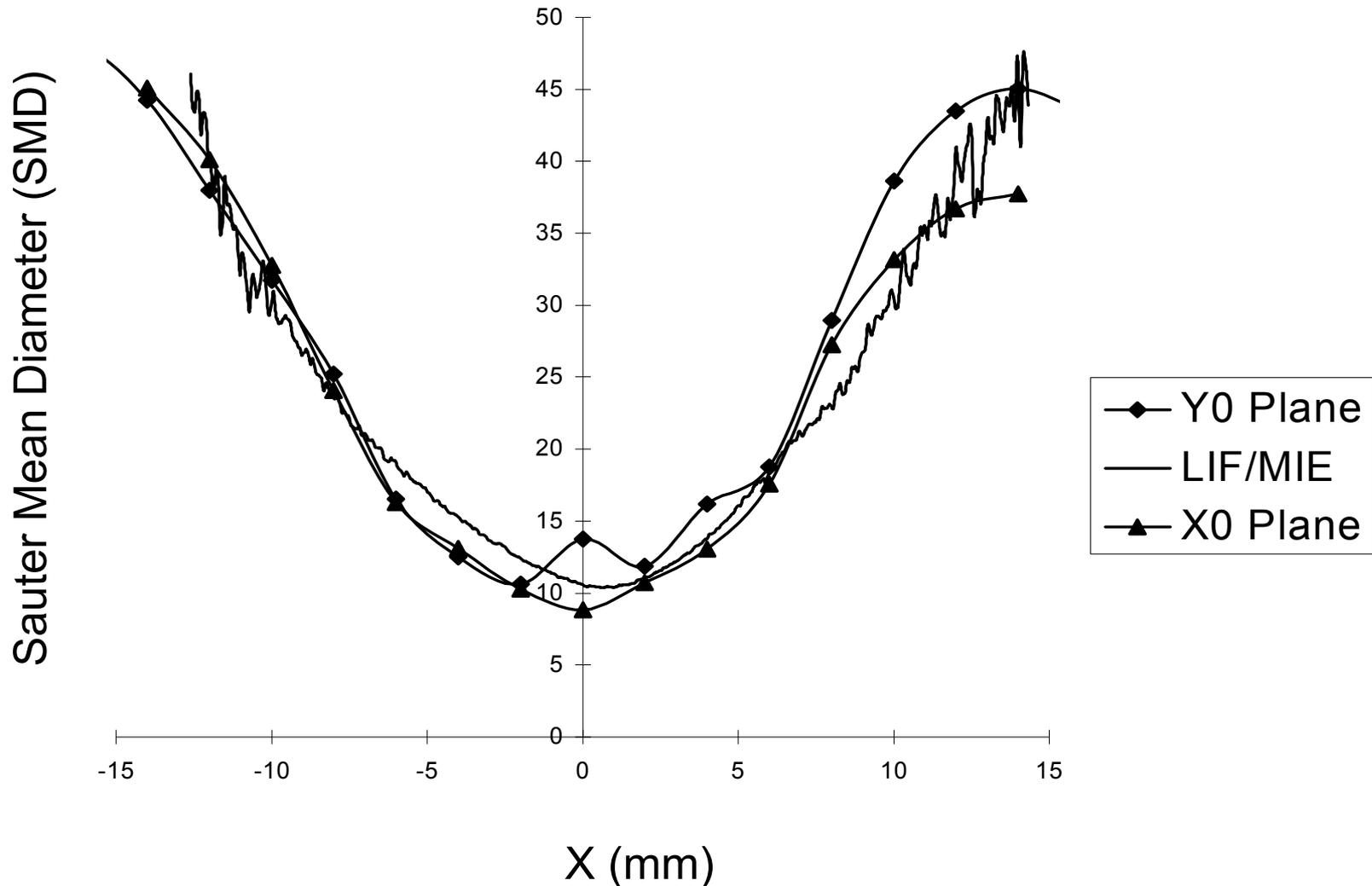


Mean Mie (3,000 laser shots)

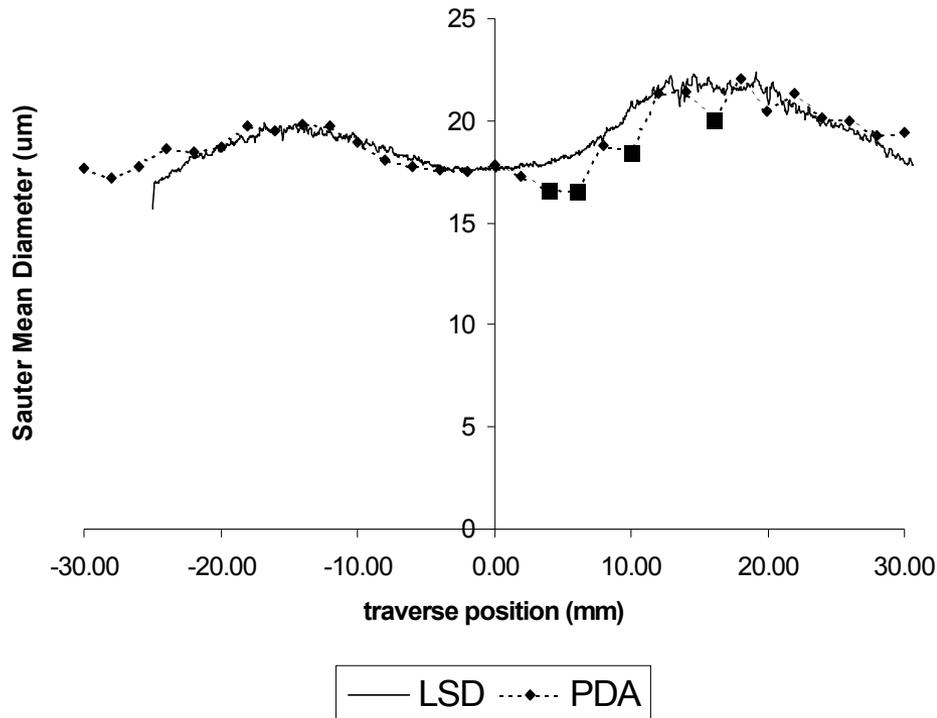


(False colours scale in counts)

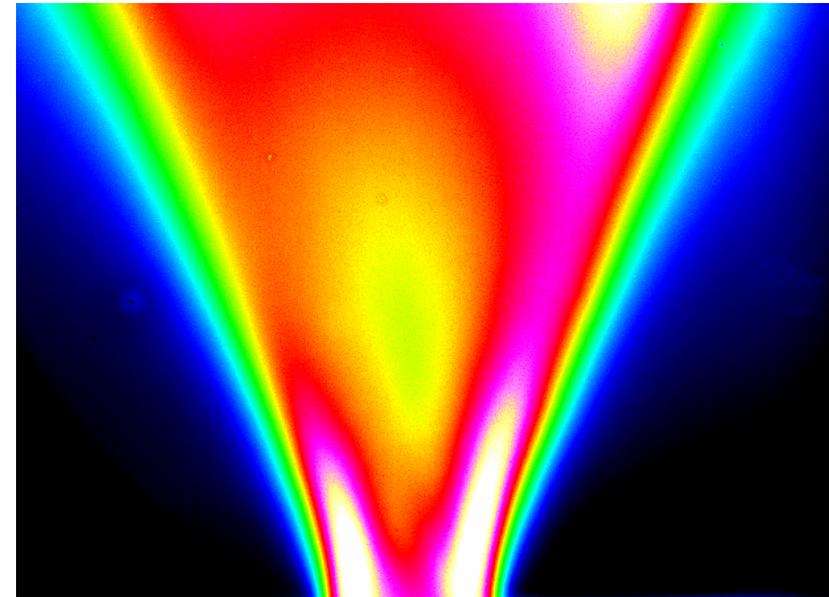
## SMD comparison mean LSD versus PDA for a Delavan pressure swirl atomiser at $z = 20\text{mm}$ downstream of the injector.



# LSD on an “dense” Air-Spray Fuel Injector – an example of PDA issues.



Comparison of LSD & PDA.



Mean LIF Image - also a Map of Fuel Volume Fraction.

## Extending LSD to evaporating sprays.

- PLIF is now widely used for in-cylinder mapping of fuel to estimate AFR or equivalence ratio.
- Tracers with good fluorescence properties and similar physical & chemical properties are used to mark the fuel.
- The temperature and pressure dependence of the tracers in the **vapour phase** are well established
- **How can we accommodate the P & T variations in a calibration?**
- **What is the fidelity of the tracers (e.g. do they co-evaporate & how sensitive to co-evaporation are we)?**

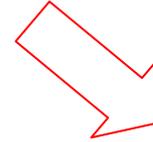
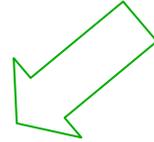
# Tracer requirements

- The tracer SHOULD co-evaporate but we should also ask how important this is.
- Tracer LIF should be independent of T & P – through a careful selection of ketones this can achieve this to ~95%.
- Tracer should match the properties of the fuel – for this exercise we have restricted ourselves to the tractable problem of iso-octane.
- Tracer must give good  $d^3$  response.

## What do we know about droplet evaporation & mass transport ?

- In most systems the droplets do not survive for more than 2 msec and often less!
- In the previous data we see the relative droplet surface area decrease by nearly 100 in about  $10^\circ$  of c.a. at 1,000 rpm. This gives a reduction of 10 in droplet size or **1,000 in fuel mass fraction!**
- Typical mass evaporation rates are of  $\sim 10^{-6}$  m<sup>2</sup>/sec whereas typical vapour mass diffusion rates are  $\sim 10^{-9}$  m<sup>2</sup>/sec. This large difference will limit the distillation behaviour of droplets. This suggest flash evaporation will be important **unless** the air pressure is high.

# Two extreme evaporation regimes

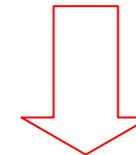
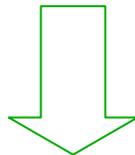


Slow / Distillation  
Evaporation

Rapid / Flash  
Evaporation

diffusion  
rate  $\gg$  evaporation  
rate

diffusion  
rate  $\ll$  evaporation  
rate



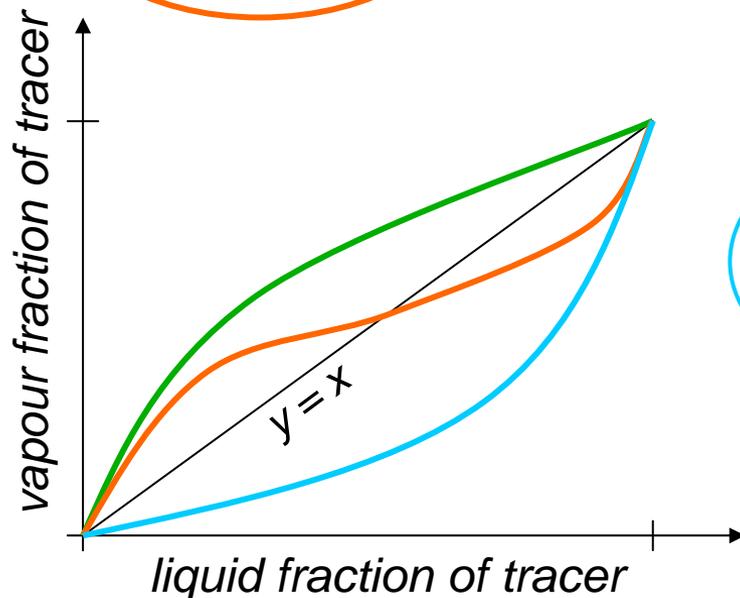
Differential evaporation

Homogeneous evaporation

# Slow / Distillation Evapouration for a « fuel + tracer » mixture

Azotropic  
mixtures

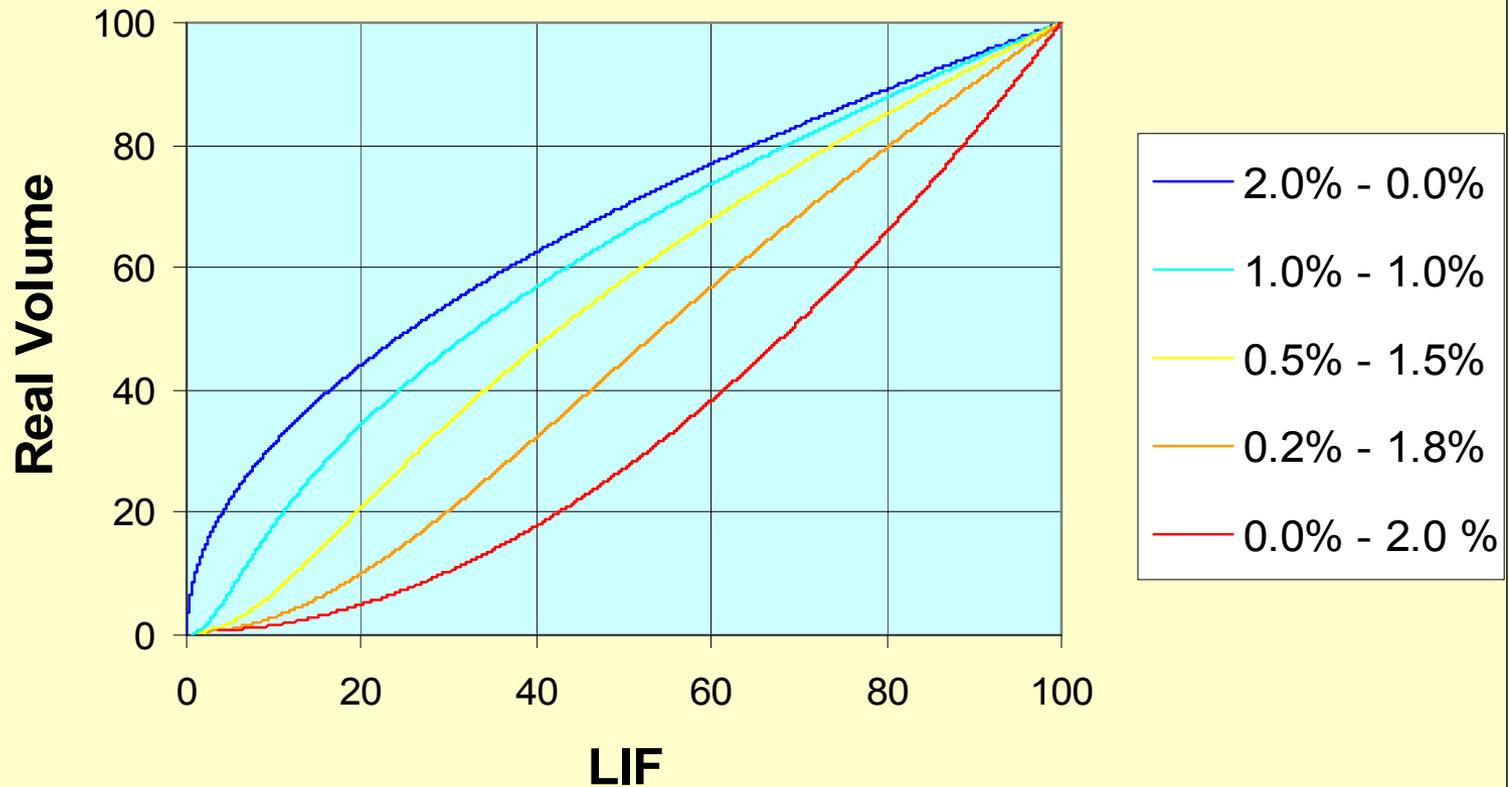
Non-Azotropic  
mixtures



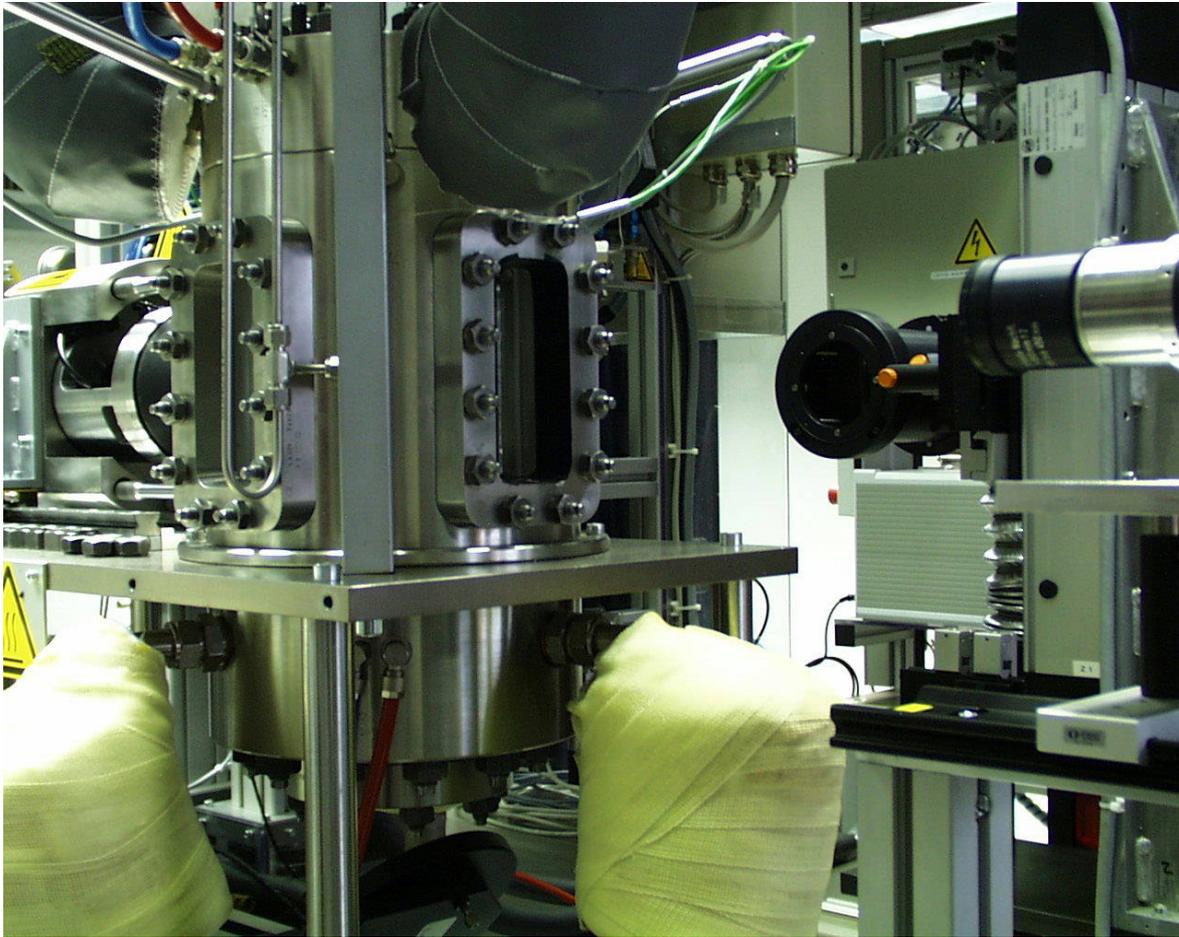
Preferential  
evapouration of  
the fuel

Preferential  
evapouration of  
the tracer

## Pentanone / Hexanone in Iso-Octane



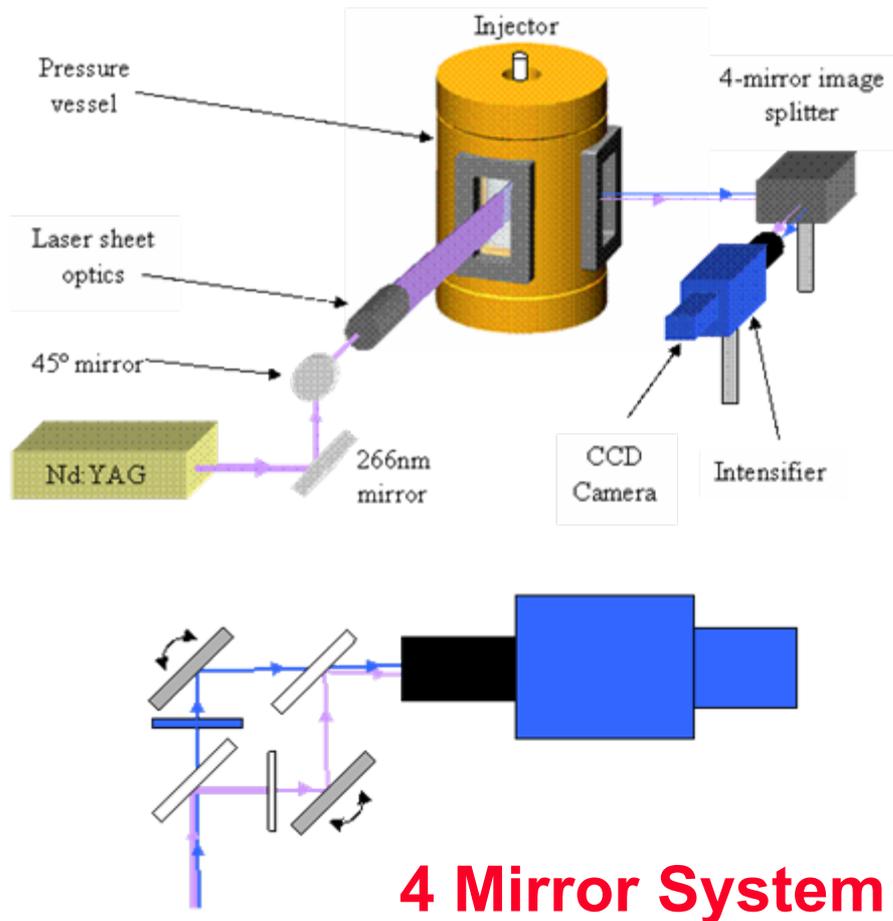
## BMW Optical Bomb for Injector Testing



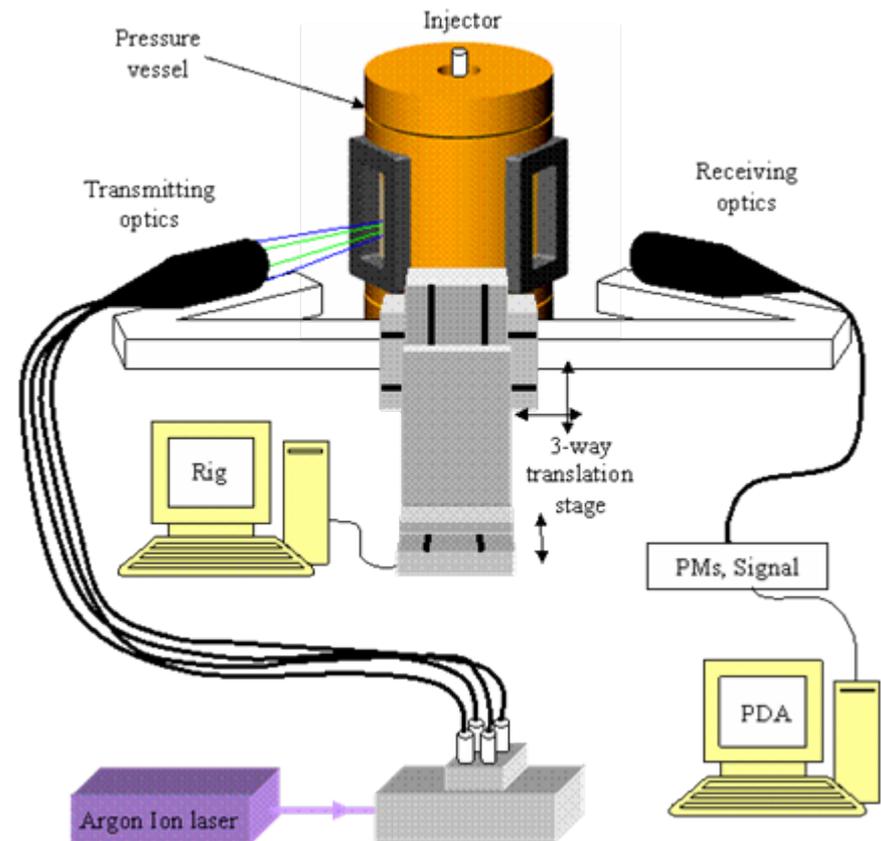
- Operates on pure  $N_2$  flow, P to 20 bar & T to 700K.
- Continuous flow therefore allows multiple repeat injections.
- Fuel pressure to 200 bar.

# Experimental Details – 2 Nearly Identical Pressure Vessels

## LSD – Mie - LIF

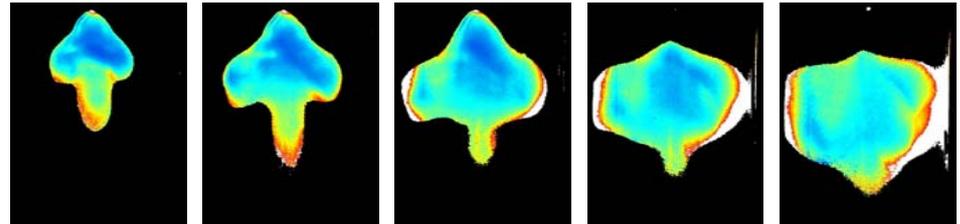


## PDA

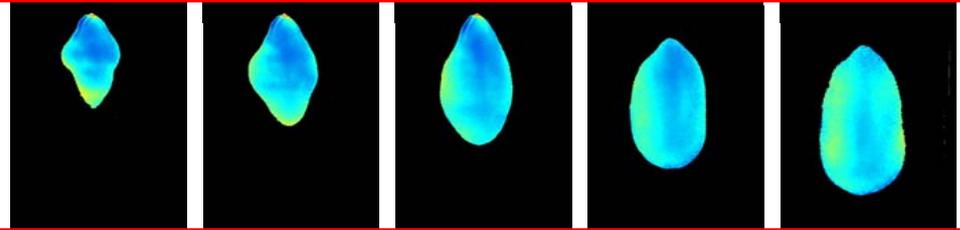


## LSD Images

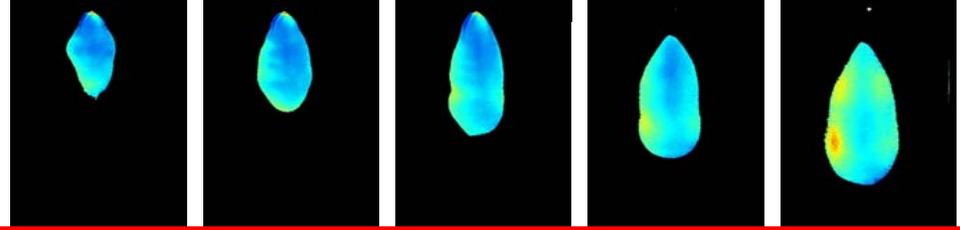
1bar - 25°C



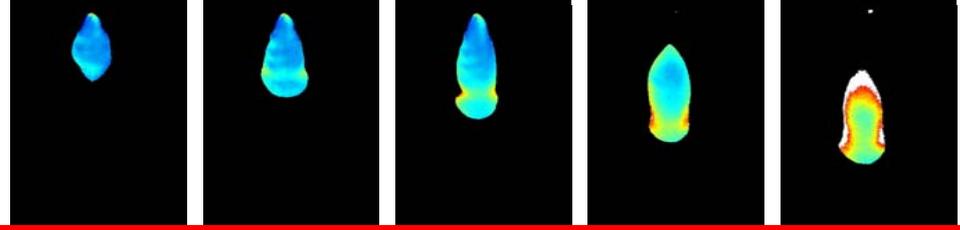
3bar - 135°C



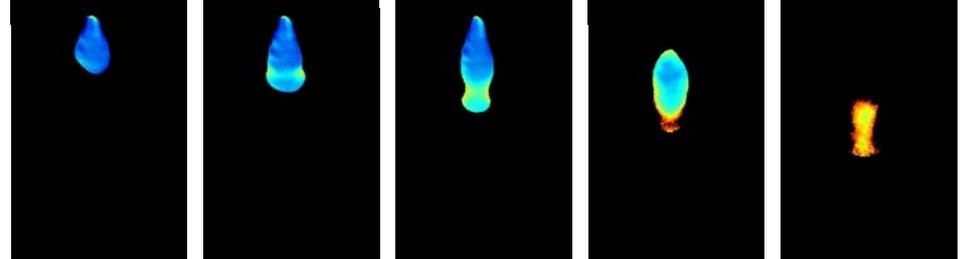
5bar - 197°C



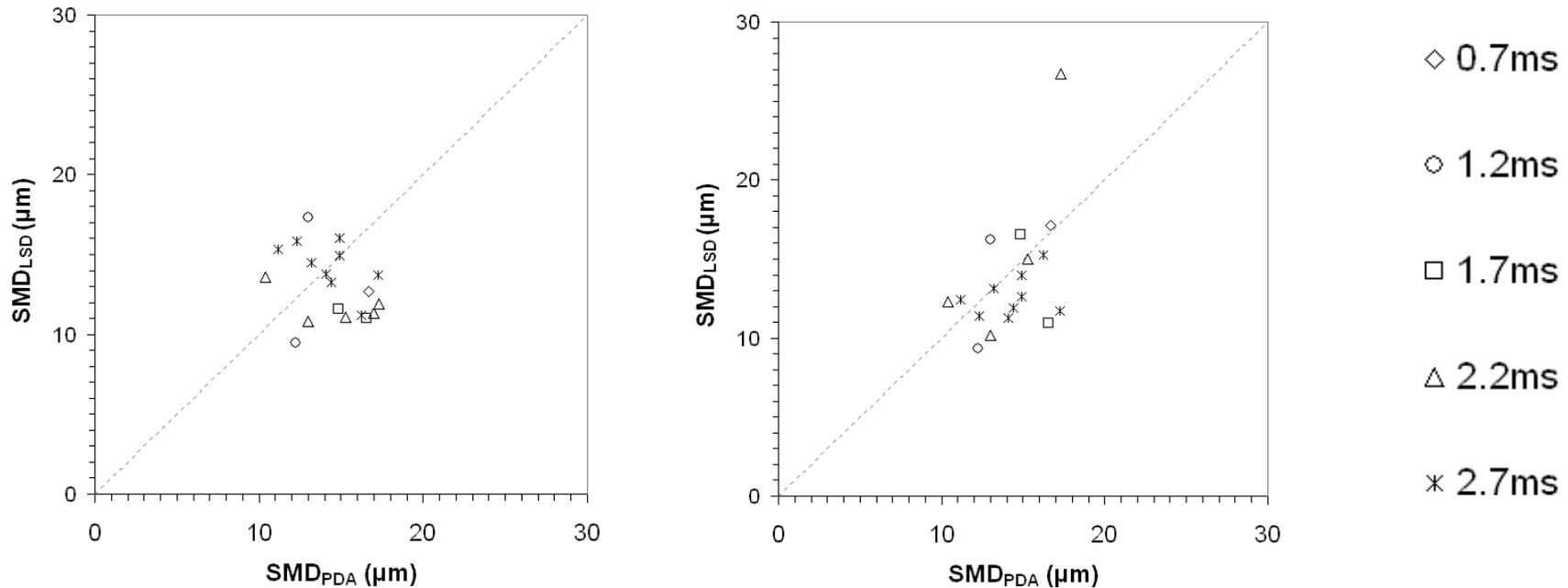
10bar - 297°C



15bar - 365°C

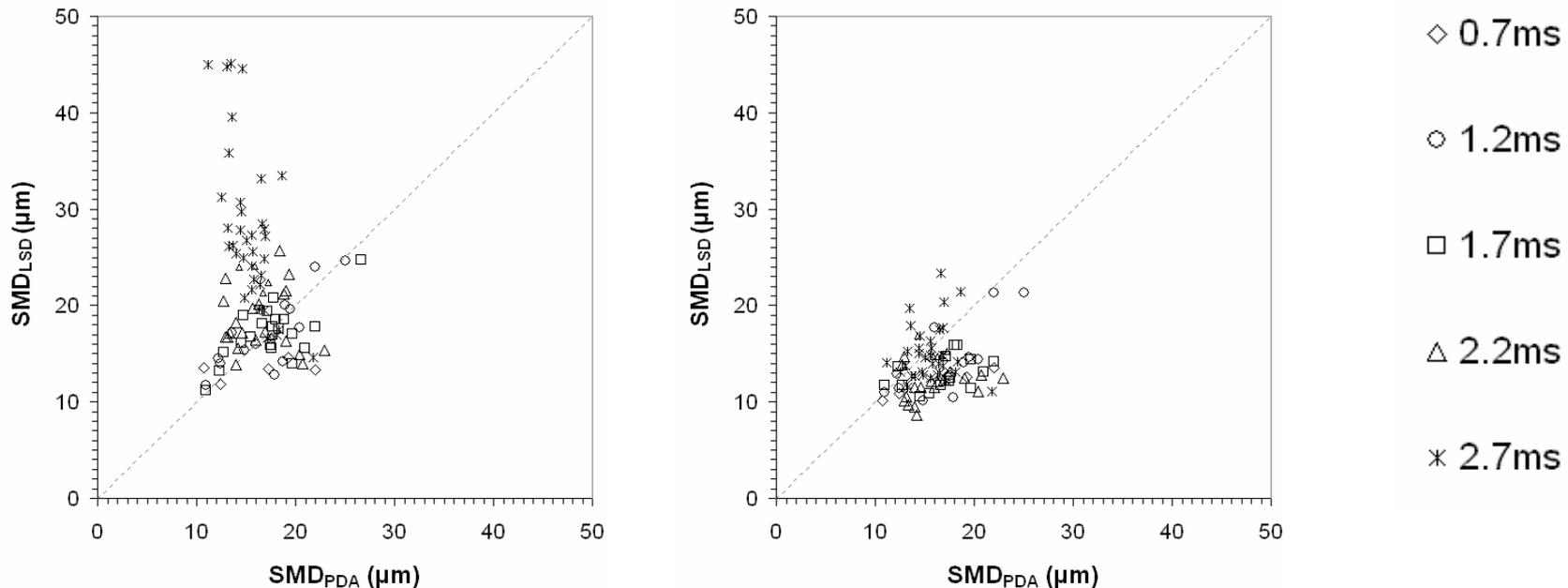


# LSD & PDA Comparison @ 1bar - 25°C



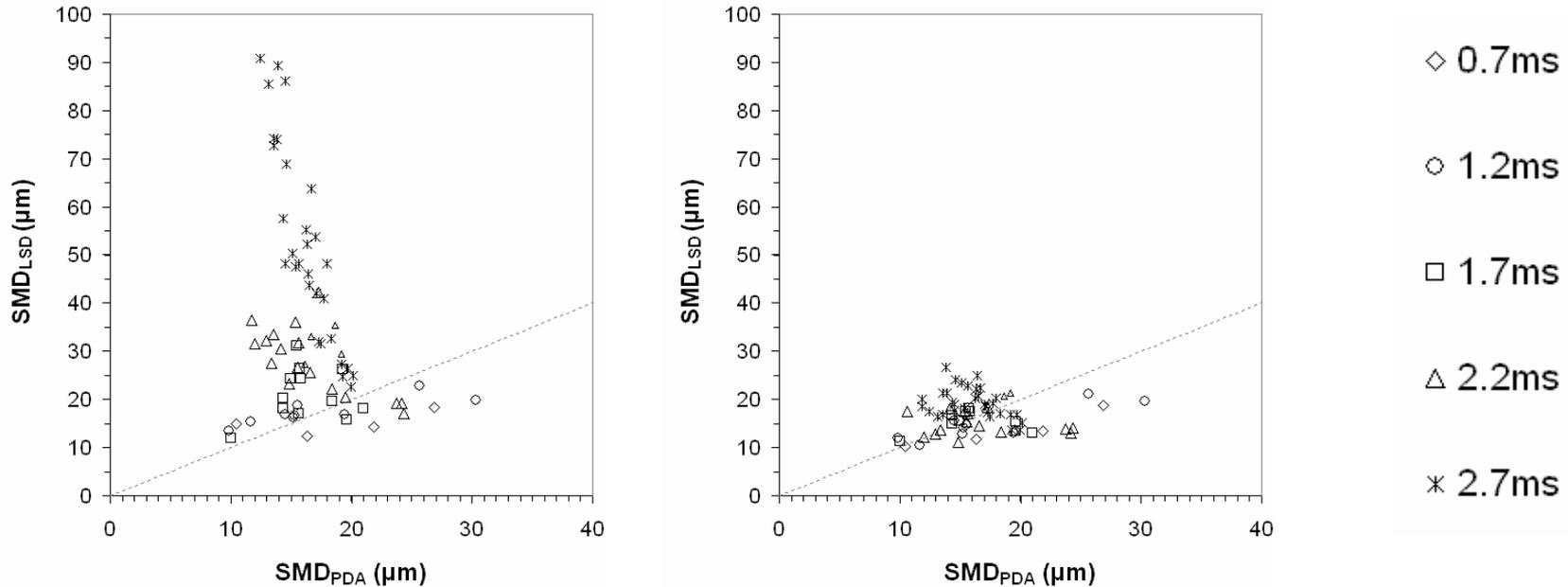
**Various times after start of injection - 2 / 0 / 98  
(%pentanone/%hexanone/%iso-octane) (left)  
& 0.5 / 1.5 / 98 mixtures (right)**

# LSD & PDA Comparison @ 3bar - 135°C



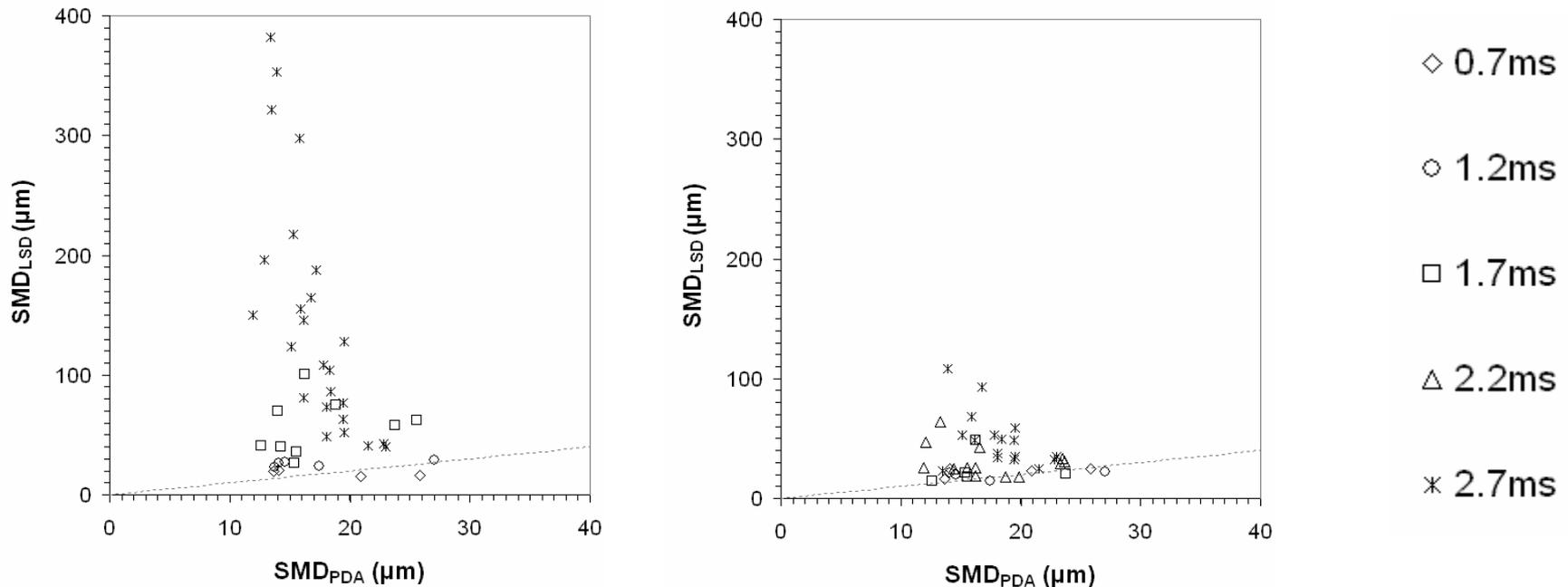
**Various times after start of injection - 2 / 0 / 98  
(%pentanone/%hexanone/%iso-octane) (left)  
& 0.5 / 1.5 / 98 mixtures (right)**

# LSD & PDA Comparison @ 5bar - 195°C



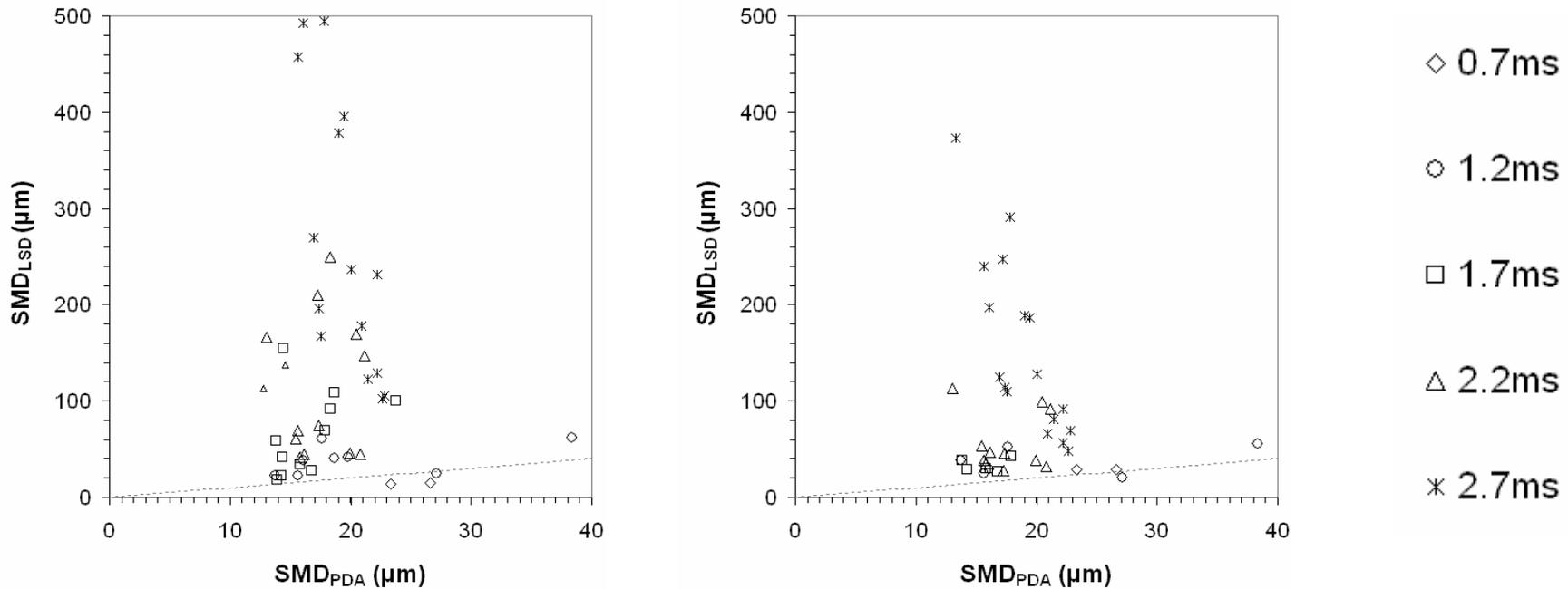
**Various times after start of injection - 2 / 0 / 98  
(%pentanone/%hexanone/%iso-octane) (left)  
& 0.5 / 1.5 / 98 mixtures (right)**

# LSD & PDA Comparison @ 10 bar - 295°C



**Various times after start of injection - 2 / 0 / 98  
(%pentanone/%hexanone/%iso-octane) (left)  
& 0.5 / 1.5 / 98 mixtures (right)**

# LSD & PDA Comparison @ 15 bar - 360°C



**Various times after start of injection - 2 / 0 / 98  
(%pentanone/%hexanone/%iso-octane) (left)  
& 0.5 / 1.5 / 98 mixtures (right)**

# Conclusions:

- PDA struggles with high density, small droplet size sprays and is not suitable as a routine diagnostic as the measurements times (hours) and data processing (days) is too long.
- We have found a combination of LIF tracers suitable for iso-octane sprays operating up to modest evaporation rates.
- Our comparison with real sprays show that the LSD technique is good to  $\sim 5$  bar and  $200^{\circ}\text{C}$ .
- At high evaporation rates LIF of vapour confuses the issue.
- At high evaporation rates LIF directly measures the liquid volume fraction and is arguably the best diagnostic. At these conditions large increases in the LIF/Mie ration can be interpreted as indicators of rapid evaporation.
- The next challenges are to:
  - Look at diesel sprays.
  - To understand sprays undergoing super-critical evaporation.