

CHARACTERIZATION OF NANOSECOND MICRO-MIXERS

S. Mitic¹, J. van Nieuwkesteele², A. van den Berg², S. de Vries¹

¹Section Enzymology – Department of Biotechnology – Delft University of Technology, Julianalaan 67, 2628 BC Delft, The Netherlands

²MESA+ Research Institute - University of Twente, 7500 AE Enschede, Postbus 217, The Netherlands

Characterization of a four-jet tangential stainless steel micro-mixer

The novel micro-mixer contains four channels (\varnothing 70 μ m) ending in a 'premixing' chamber of 120x120x70 μ m (fig. 1).

After the 'premixing' chamber, the fluids enter a small 'mixing' chamber and are mixed within \sim 2 μ s (fig. 2). The liquids leave at an orifice (\varnothing 20 μ m) travelling further as a free jet (max. linear jet flow \sim 200 ms⁻¹, dynamic pressure \sim 400 bar) [1].

The mixer is employed to study enzyme reactions in a freeze-quench set up yielding an instrumental dead time of 60 μ s, 100 fold shorter than any other device.

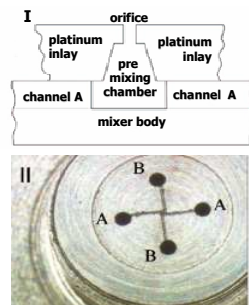


Fig. 1: (I) Geometry of the mixer; (II) Channels and the premixing chamber of dismounted mixer

Mixing efficiency

To characterize the mixing efficiency of the micro-mixer, the fluorescence quenching of o-hydroxycinnamic acid, HCA was monitored in a jet (fig. 2).

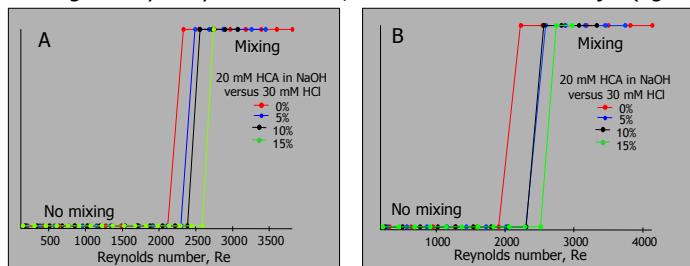


Fig. 2: Mixing at various concentrations of glycerol. The mixing was followed by the quenching of HCA(alk) vs HCl, (A) 1:1 flow rate ratio: HCA(alk) vs HCl; (B) 2:1 flow rate ratio: HCA(alk) vs HCl.

For a 1:1 and 2:1 flow rate ratio between HCA(alk) and HCl mixing occurs in a transient regime at $Re > 2300$ and $Re > 2200$, respectively. With increasing glycerol concentrations in both solutions, the Re stays the same \sim 2600, in agreement with turbulent mixing theory.

Characterization of four-jet tangential glass-silicon micro-mixer chips

To determine where mixing occurs, in the 'premixing' or in the 'mixing' chamber a few generations of novel chips for optical studies are designed (fig. 3). The opposite channels etched in 360 μ m thick Si are offset with respect to each other by $\frac{1}{2}$ width of the channel. The channels are covered by 1.1 mm thick glass. The orifice is in the Si part of the mixer (\varnothing 25 μ m – 70 μ m), providing a mixing volume of the small chamber between 16-127 pL.

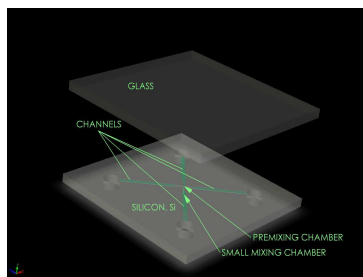


Fig. 3: Glass-silicon four-jet micro-mixer chip

To determine the percentage of premixing in the micro-mixers, fluorescence appearance during deprotonation of 8-hydroxypyrene 1,3,6-trisulfonic acid, trisodium salt, HPTS was monitored in the premixing chamber using a fluorescence microscope. When NaOH was 1.5, 10 or 30 fold in excess over HCl, it was determined that 6, 14 or 16.5% / 2.5, 8.8 or 11.4% / 2.5, 10 or 13.5% and 2, 8.3 or 10% of premixing occurred in the premixing chamber of the chips with 70/50/30 and 25 μ m orifice diameter, respectively (fig. 4 and fig. 5).

[1] A. Cherepanov and S. de Vries. *BBA*, 1656, 1-31, **2004**.

[2] F. Wiertz, *et al. FEBS Letters*, 575, 127-130, **2004**.

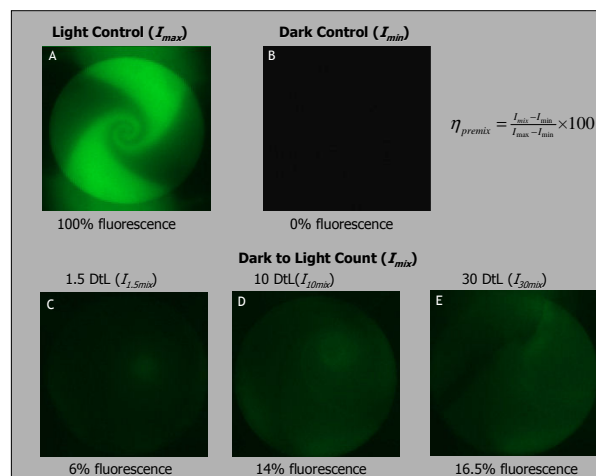


Fig. 4: Fluorescence appearance of HPTS dye in the 'premixing' chamber of the chip with 70 μ m orifice diameter and at the flow rate of 10 mL.min⁻¹: (A) HPTS(alk) in NaCl vs NaOH in NaCl -> light control; (B) HPTS(ac) in NaCl vs HCl in NaCl -> dark control; (C, D, E) HPTS(alk) in NaCl vs HCl in NaCl when NaOH is 1.5; 10 and 30 fold in excess over HCl, respectively -> dark to light count - percentage of premixing in the premixing chamber

Values of calculated Re in the mixing chamber for the chips (\varnothing 25 μ m-70 μ m), ($Re \gg 2000$ transient/turbulent flow) are respectively, 1.9/2.7/4.5 and 5.4 fold larger than their values of Re in the premixing chamber, ($Re < 2000$, laminar flow).

Since at $Re < 2200$ no mixing occurs, all ($> 85\%$) of the mixing must occur in the small chamber. The calculated mixing time is 760/560/310 and 360 ns for the chips with 70/50/30 and 25 μ m orifice diameter, respectively.

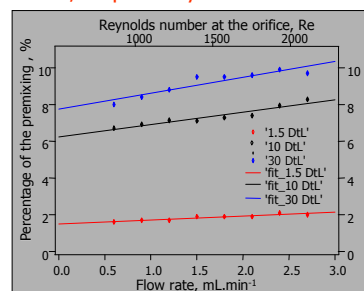


Fig. 5: Percentage of premixing in the premixing chamber during deprotonation of HPTS when NaOH was 1.5; 10 or 30 fold in excess over HPTS in HCl as a function of the flow rate in the system and calculated values of the Re at the orifice at each flow rate for the chips with 25 μ m orifice diameter.

Conclusion

The time of mixing can be even shorter \sim 200 ns if the same experiment would be successfully done with the (\varnothing 20 μ m) chip, as in the stainless steel mixer. This mixer would then be 100 fold faster than any described in the literature.

Outlook

For optical studies a glass cylinder will be used in stead of a free jet. The cylinder (25-50 mm diameter path length) is positioned under an inverted fluorescence microscope, connected to a monochromator and a Hg light source with collimating optics.

Using both mixing systems, the time course of chemical and enzymatic reactions, applying UV-Vis absorbance and fluorescence spectroscopy will be studied with a time resolution in the nanoseconds.