

# 3D magnetic resonance velocimetry for the characterization of hydrodynamics in microdevices

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1	3D Magnetic resonance velocimetry for the characterization of
2	hydrodynamics in microdevices: application to micromixers and
3	comparison with CFD simulations
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#### 12 Highlights

An NMR instrumentation is developed to measure 3D velocity fields in microdevices
MRI measurements in an arrow-type micromixer are compared to CFD simulations
Small manufacturing defects strongly affect the flow and the mixing performance
Good agreement is obtained between numerical data and experimental measurements

17 Abstract

18 3D velocity field measurements in an arrow-type micromixer are performed using nuclear 19 magnetic resonance (NMR) imaging methods. A specific device has been set up including a home-20 made radiofrequency milli-coil matching the geometry of the manufactured micromixer, allowing 21 an improvement of the signal-to-noise ratio by a factor 7 comparing to a standard commercial 22 hardware. The NMR images obtained are compared to computational fluid dynamics (CFD) 23 simulations. The very good agreement obtained between numerical data and experimental 24 measurements revealed, on the one hand, the capacity of the developed method to perform a 25 detailed and accurate study of phenomena occurring in a microstructured system and on the other 26 hand, the great impact of small manufacturing defects on the hydrodynamics in a micromixer and 27 thus on the mixing performances.

<u>Keywords</u>: computational fluid dynamics, magnetic resonance imaging, micromixers, magnetic
 resonance velocimetry, NMR instrumentation

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#### 32 **1. Introduction**

33 Microstructured reactors are attracting a lot of interest due to their intense heat and mass transfer capabilities. Their potential has been demonstrated through many applications in chemistry and 34 35 process engineering, for example for flexible and decentralized synthesis processes (Hessel et al., 36 2005; Lee & Fu, 2018). They allow fine control of selectivity in complex competitive syntheses, 37 and thus a reduction in purification costs and energetic impacts. The growing interest in these 38 microstructured systems has led to more and more innovative designs and a variety of different 39 geometries are presented in the literature from simple to complex (Bayareh et al., 2020). In the case 40 of micromixers, a simple design (T or Y shape) can be as efficient as a more complex one (Dreher 41 et al., 2009), the mixing performance being essentially related to the amount of mechanical energy 42 injected and whether it is injected at the precise location where it is needed (Falk & Commenge, 43 2010). The studies carried out on micromixers with the T and Y geometries showed that in a range 44 of Reynolds numbers between Re = 5 and Re = 600, several hydrodynamic regimes govern the 45 flow in the mixing channel and impact the mixing quality (Engler et al., 2004; Galletti et al., 2019; 46 Hsieh, 2013; Lobasov et al., 2018; Peng et al., 2022). With a high sensitivity to the local geometry 47 (channel angle and cross section), with increasing Reynolds number values, the so-called stratified 48 flow first evolves to a steady symmetric vortex flow, which becomes asymmetric, then periodic and finally unsteady, before reaching the laminar-turbulent transition around Re = 600. These 49 50 flow regimes drastically improve the mixing quality and thus manage to reconcile performance and 51 manufacturing simplicity. Experimental quantification of the mixing efficiency of micro-mixers is 52 mainly performed via chemical characterizations using selective test reactions, such as the 53 Villermaux-Dushman protocol, in which information on the mixing performance is given by the 54 spectrophotometric analysis of the reaction products (Commenge & Falk, 2011). Other 55 experimental methods use optical detection, which is generally the method of choice to study 56 microfluidic devices. Optical methods are however limited to transparent media and might be either 57 invasive as they usually require the use of tracers that can alter the properties of the fluid system, 58 or unconclusive as soon as vortical multilamination effects alter the contrast in the field of view. 59 Additional information on the evolution of the flow regime in micromixers can be obtained by 60 means of analytical methods or numerical simulations, but the latter are generally performed in 61 ideal geometries i.e. without considering potential imperfections such as geometric defects that 62 may result from the fabrication of these microscale systems.

63 Experimental methods based on the nuclear magnetic resonance (NMR) phenomenon allow 64 non-invasive analysis of flows in microdevices. In literature NMR is conventionally used as a 65 powerful analytical method in chemistry (Ionin & Ershov, 1995), biology (Wüthrich, 1990), 66 rheology (Milc et al., 2022), porous media (Ferrari et al., 2018; Perrin et al., 2022), materials 67 (Blümich Bernhard, 2000; Robert et al. 2018), in the biomedical field (Pettegrew, 2012) or in 68 medicine through magnetic resonance imaging (MRI). Although existent, the literature on the 69 application of MRI tool for the characterization of hydrodynamics (Gladden & Alexander, 1996; 70 Stapf & Han, 2006) and compared to CFD simulations in engineering is sparser (Mendez et al., 71 2017; Chung et al. 1993). Indeed, NMR gives access to quantitative 3D measurements of the flow 72 field, including in opaque devices that are inaccessible to optical velocimetry techniques. 73 Moreover, its time-resolution is sufficient for characterizing transient flow phenomena (Callaghan 74 & Xia, 1991; Oliveira et al., 2020; Seymour & Callaghan, 1997).

In practice, limitations related to measurement sensitivity may preclude the use of NMR on small scale devices handling a very low fluid volume when using standard methods and commercial hardware. Significant progress has been made over the last years in the NMR characterization of

78 such small systems (down to the nanoliter) (Lacey et al., 1999; Webb, 2012) by increasing the 79 strength of the static magnetic field (Fu et al., 2005) or by using cryoprobes (Kovacs et al., 2005). 80 Another efficient approach to improve the signal-to-noise ratio and allow direct access to the 81 internal parts of the microstructured device is to reduce the dimensions of the radio frequency (RF) 82 coil in order to cover the region of interest as closely as possible and thus maximize the so-called 83 filling factor, *i.e.* the fraction between the sample volume and the coil volume (Hoult & Richards, 84 2011; Webb, 2008, 2013). Specific three-dimensional or planar (two-dimensional) coils have been 85 designed for small volume samples (Eroglu et al., 2003; Massin et al., 2002; Olson et al., 1995; 86 Peck et al., 1995; Trumbull et al., 2000) using specific microfabrication techniques such as laser 87 processing or etching and copper printing (Klein et al., 2013a) or deposition on supports such as 88 polyimide films (Dechow et al., 2000; Meier et al., 2014; Murphree et al., 2007; Woytasik et al., 89 2006). These types of coils are incorporated into microstructured devices for microscale studies 90 such as quantitative reaction monitoring of processes (Ciobanu et al., 2003; Wensink et al., 2005), 91 dynamic studies in devices handling small volumes (Harel & Pines, 2008; Klein et al., 2013b; J. 92 Zhang & Balcom, 2010; X. Zhang & Webb, 2005) or performing velocity field mapping and 93 distributions inside micromixers (Ahola et al., 2006, 2012; Akpa et al., 2007).

In this study, an NMR instrumentation is specifically developed for 3D velocity field measurements in a micromixer of millimeter dimensions. The MRI results obtained are compared to computational fluid dynamics (CFD) simulations to study the hydrodynamics in the device highlighting the sensitivity of the flow and mixing performances to the presence of small manufacturing defects.

### 99 2. Materials and Methods

#### 100 2.1. Conception, fabrication and characterization of the micromixer

101 The micromixer investigated in this study has been machined in a block of polymethyl methacrylate 102 (PMMA) using a digital milling machine. It consists of three cylindrical channels, with an internal 103 diameter of 1 mm, arranged in an arrow configuration (Figure 1). The mixing angle,  $\theta = 30^{\circ}$ , 104 defines the angle between the axes of the outlet and one inlet. The zone of confluence of the three 105 channels will be referred throughout this paper as "mixing zone", and the point where their axes 106 are supposed to meet will be referred as "mixing point".

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Figure 1: Experimental setup. (a) Schematic description of the fabricated micromixer. (b) Front view showing the NMR coil folded on both sides of the detection zone. (c) Assembly mounted on a commercial coil head ready to be inserted into the spectrometer.







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114 As discussed later in this paper, large differences were observed between the experimental velocity measurements and the first CFD simulations, which raised doubts about the presence of 115 manufacturing defects in the prototype. In order to check this possibility, 3D images of the 116

117 micromixer were recorded by MRI and X-ray Computed Tomography (µCT) for characterizing its 118 true geometry. The MRI was performed on a Bruker Avance III 600 WB spectrometer. The 119 superconducting magnet is oriented vertically and generates a static magnetic field  $B_0$  with an intensity of 14.1 T which corresponds to a proton <sup>1</sup>H Larmor frequency of 600 MHz. The 120 121 spectrometer is equipped with a commercial Bruker MicWB40 microimaging coil. The micromixer 122 was filled with a CuSO<sub>4</sub> solution 0.01 M in order to shorten the NMR relaxation time. The acquired image, consisting of voxels of  $100 \times 100 \times 54 \,\mu\text{m}^3$ , provided a 3D mapping of the micromixer (see 123 124 Fig.S1 of the Supporting information). The analysis of the obtained data revealed a slight 125 misalignment of the three microchannels. Indeed, the two inlet axes and the outlet axis do not 126 perfectly converge toward the supposed mixing point (Fig. S2). Moreover, the images analysis 127 revealed the presence of two relatively large machining chips nearby the mixing zone, the first at 128 the end of one inlet, and the second one at the entrance of the outlet channel (Figure 2 and Fig.S3).

129 A X-ray  $\mu$ CT scan was also performed in order to characterize the micromixer with a higher 130 spatial resolution. The dataset was recorded using a Phoenix Nanotom 180 *kV* GE X-ray system 131 with a source-sample distance of 30 *mm*, an accelerating voltage of 80 *kV* and a current of 80  $\mu$ *A*. 132 The spatial resolution was set to 6.81  $\mu$ *m*/*voxel*. 2 000 projections were made (as averages of 3 133 projections of 1 000 *ms* each), resulting in a scan time of 134 *min*. Data processing was done 134 using Avizo® 2022 and VGSstudio 3.5.

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Figure 2: 3D image of the micromixer, obtained by X-ray  $\mu$ CT, showing manufacturing defects. (1): The black arrow shows a misalignment of the 3 channels. (2): Ortho-slice recorded near the mixing point: the white arrows show the presence of solid parts as observed by X-ray  $\mu$ CT (a) and MRI (b)

The raw images were segmented in order to isolate the three channels. The ortho-slice in the vicinity of the mixing point (Figure 2) confirmed the presence of the solid PMMA residues (white color, showed with the white arrows) that were not removed during the milling process. As discussed later in the results section, compared to the "ideal geometry" case, the presence of these defects leads to significant perturbations of the hydrodynamics in the micromixer.

143 **2.2**.

#### MRI instrumentation and experiments

144 2.2.1. MR velocimetry measurements

MRI allows the localization of the <sup>1</sup>H NMR signal in order to reconstruct a 1D, 2D or 3D image. The principle relies on the application of consecutive magnetic field gradient pulses that make the main magnetic field  $B_0$  vary along the three spatial directions and modifies the precession frequency of the proton <sup>1</sup>H nuclear magnetic moments according to Larmor's law (frequency encoding) as well as their phase (phase encoding). Details about the formation of MRI images are fund in textbooks (Callaghan, 1991). The schematic of the 2D velocimetry gradient-echo sequence used in this study is shown on Figure 3. In order to measure velocity components, a specific encoding of the positions of the magnetic moments is produced by two gradient pulses of amplitude  $G_{\nu}$ , duration  $\delta$  and separated by a time  $\Delta$  (Caprihan & Fukushima, 1990; Callaghan, 1991; Blümich, 2000). The relationship linking the phase  $\Phi$  of the measured NMR signal and velocity is given by the relation :

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$$\Phi = \gamma \delta \Delta \overrightarrow{G_{\nu}}. \vec{\nu}$$

157 where  $\gamma$  is the proton gyromagnetic ratio.

In practice, a phase difference image is recorded, taking as reference the image measured when  $G_{\nu} = 0$ . Since the direction in which the flow is measured is determined by the direction of the encoding gradient, the velocity vector is determined at each position in a 2D section by applying  $\overrightarrow{G_{\nu}}$  along the three orthogonal directions in space.



Figure 3: Acquisition protocol of 2D MRI velocity maps, based on the gradient echo sequence.

RF: radiofrequency pulses  $G_{ss}$ : slice-selection gradient  $G_{\omega}$ : frequency-encoding gradient  $G_{\phi}$ : phase-encoding gradient TE: echo time TR: repetition time  $G_{v}$ : velocity-encoding gradient To perform the MR velocimetry experiments, the micromixer was integrated inside a homemade specific radiofrequency (RF) coil which was used for both the RF emission and the reception of the NMR signals. The assembly was placed in a gradient imaging system delivering a maximum magnetic field gradient intensity of  $150 \ G/cm$ .

166 2.2.2. Specific MRI instrumentation

167 The developed radiofrequency milli-coil was adapted to the geometry and dimensions of the 168 micromixer (Figure 1 (b) and Figure 1(c)). The geometry chosen for this study was a Helmholtz 169 coil composed of a pair of copper loops placed face to face and 2.5 mm apart. The diameter of 170 each loop was 5 mm. The fabrication of the milli-coil was realized using an etching process on a 171 DuPont<sup>TM</sup>Pyralux<sup>®</sup>AC flexible copper/Kapton polyimide substrate. The width of the copper strip 172 etched on the polyimide substrate was 0.4 mm and its thickness was 45 µm. The fabricated milli-173 coil was connected to an electronic circuit composed of capacitors so that the resulting RLC circuit 174 resonated at the proton (<sup>1</sup>H) resonance frequency of 600 MHz (tuning) and its impedance matched 175 the characteristic impedance of the coaxial cable used for the transmission of the generated RF 176 pulses and the received NMR signal (matching). For tuning and matching, a commercial 177 exchangeable surface coil head Bruker MicWB40 was re-used. A pre-tuned circuit was added to complete the tuning with a set of fixed capacitors: ( $C_{tuning} = 2.7 \ pF$ ,  $C_{matching} = 1.2 \ pF$ ). 178

Electromagnetic simulations were carried out in order to determine the distribution of the radiofrequency field generated at 600 *MHz* by the fabricated milli-coil. The simulations were performed with COMSOL multiphysics® and allowed computing the normalized radiofrequency field intensity over a 5.7 *mm* distance in the direction of the outlet of the micromixer. The measurement of the NMR signal intensity map (Figure 4 (b)), performed on the micromixer filled with the copper sulfate solution is in very good agreement with this simulation (Figure 4 (c)). According to these results, the profiles present a plateau where the radiofrequency field is homogeneous. The deviation from homogeneity over a length of 2 *mm* is only 1.3 % on the profile obtained by the electromagnetic simulation and 3 % on the MRI profile.

For a detailed and accurate study of the hydrodynamics occurring in the micromixer, the MRI velocity measurements presented in this work were located within the homogeneity zone of the radiofrequency field produced by the milli-coil.



Figure 4: (a) Schematics of the Helmholtz NMR coil, composed of a pair of copper loops placed face to face on either side of the mixing zone. (b) Normalized 2D MR image of the CuSO<sub>4</sub> solution in the mixer. (c) Comparison of the experimental and simulated normalized NMR intensity profiles along the centerline of the outlet channel showing a homogeneous intensity plateau of about 2 mm in length.

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In order to evaluate the improvement of the signal-to-noise ratio (SNR), MRI measurements
were performed on both the milli-coil and the commercial coil Bruker MicWB40 (with a detection
volume 10 times larger) in similar conditions.

195 An axial NMR signal intensity map (measured in the plane orthogonal to the main flow 196 direction) was performed with the milli-coil considering a field of view of  $7.5 \times 7.5 \ mm^2$ , corresponding to a spatial resolution of  $30 \times 30 \ \mu m^2$  per pixel. The NMR signal was averaged 197 198 over a slice thickness of 200 µm and accumulated 8 times. The slice position considered is at 3 mm 199 from the mixing point. A similar measurement was performed with the commercial coil Bruker 200 MicWB40. The results demonstrate a ~ 7 fold improvement of the SNR, evaluated by dividing the 201 mean value of the signal (in the channel) by the mean value of the noise (outside the channel). The 202 improvement of the signal-to-noise ratio with the developed milli-coil allows, with a higher image 203 quality, a detailed characterization of the velocity field in the micromixer (Figure 5).

(a)

(b)

#### NMR intensity [a.u.]



Figure 5: comparison of the MRI measurements performed with the commercial coil and with the milli-coil showing the improvement of the signal to noise ratio and the image quality. (a) 2D NMR signal intensity maps (b) 2D velocity map of the z component (flow rate 1.4 ml/min per inlet, slice thickness 0.2 mm, slice position at 3 mm from the mixing point) obtained with the commercial coil Bruker MICW40 (25 mm ID), in-plane resolution  $30 \times 30 \,\mu\text{m}^2$  and 8 accumulations of the NMR signal) and with the developed milli-coil (in-plane resolution  $24 \times 24 \,\mu\text{m}^2$  and 4 accumulations). The distances on the axes are in millimeters.

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#### 2.2.3. Parameters and conditions for MRI velocity measurements

206 The fluid flowing through the micromixer was an aqueous CuSO<sub>4</sub> solution 0.01 *M* characterized 207 by a longitudinal relaxation time  $T_1 \sim 100 \text{ ms}$ . This concentration is adequate to significantly 208 shorten the NMR relaxation times while remaining sufficiently low to not alter the physical 209 properties of the water phase, including the self-diffusion coefficient. The water flow was achieved 210 through 1 mm ID PTFE tubings using a multi-channel peristaltic pump (Ismatec Reglo ICC Digital) used with Tygon® S3<sup>TM</sup> E-LFL, 1.52 mm ID pump tubing. The peristaltic pump ensures 211 212 that a same flow rate was injected at each inlet during all the experiments. Experiments were 213 conducted for eight different flow rates ranging from 1.4 to 4.2 ml/min at each inlet (see Table 214 1), corresponding to outflow Reynolds numbers in the range [59 - 178] (note that the outflow 215 Reynolds number is the double of that of the inflow).

The protocol described earlier (gradient echo sequence with phase encoding) was used to measure both axial (in planes orthogonal to the main flow direction) and sagittal velocity maps (in a plane orthogonal to the axial images containing all three channels of the micro-mixer). The measurements were carried out using the parameters summarized in Table 2. Each velocity component was mapped separately, the *z*-component corresponding to the main flow direction. In order to check the accuracy of the measurements, the average flow rate was calculated from the velocity images in several sections and for the different flow rates supplied by the pump. The results agree over the entire measurement range, with an error of less than 5%. The reconstruction of the velocity field was made with Paravision 5.1 software. The presentation of the maps and the comparison with the CFD results were done using Python 3.7 in Jupyter Notebook 6.0.1.

Table 1: flow conditions in the micromixer			Table 2: parameters for MRI velocity		
Flow rate per inlet	Reynolds number	Maximum velocity		Parameter	Characteristics
( <i>ml/min</i> )	(outflow)	(Poiseuille		TR	300 ms
()	()	flow) $(cm/s)$		TE	5 <i>ms</i>
1.4	59	12		Δ	2.16 ms
1.9	80	16		δ	2.05 ms
2.4	102	20		acquisition time	300 s
2.9	123	25		accumulations	4
3.2	136	27		thickness	200 μ <i>m</i>
3.4	144	28	axial	field of view	$6 \times 6 mm^2$
3.9	165	30	images	spatial resolution	$24 \times 24 \ \mu m^2$
4.2	178	32		thickness	400 u <i>m</i>
			sagittal	field of view	$1 \times 1 \ cm^2$
			images	spatial resolution	$39  imes 39 \ \mu m^2$

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#### 2.3. CFD modeling and simulation

#### 229 2.3.1. Geometry and mesh

Modeling and CFD simulations of the hydrodynamics in the micromixer were conducted using the ANSYS 2019 commercial software package (including ANSYS Fluent). A first geometry was drawn using ANSYS Design Modeler following the dimensions presented in Figure 1 (a), i.e. a diameter of 1 *mm* with inlets and outlet pipes of 10 *mm* length each. This design will be referred through the remaining of this paper as "ideal geometry". ANSYS Meshing software was used for meshing the numerical domain. Given the form of the micromixer, the mixing zone was meshed using tetrahedral cells while the remaining parts, consisting of the inlets and outlet channels, were discretized using hexahedral elements (Figure 6). A total of about  $13.4 \times 10^6$  cells was employed for ensuring a mesh-independent solution for all the cases studied.

239 As discussed later in this paper, numerical results obtained using the "ideal geometry" did not 240 allow reproducing the experimental velocity fields. Accordingly, as explained in Section 2.1, the 241 real micromixer geometry has been identified through X-ray µCT scanning. This second design 242 will be referred through the remaining of this article as "real geometry". As mentioned earlier and 243 detailed in the Supplementary material section, compared to the "ideal geometry", the real one 244 involves a slight misalignment of the inlets and outlet microchannels and includes two relatively 245 large machining chips nearby the mixing zone. The "real geometry" was first meshed via Avizo 246 software and then exported toward ANSYS Meshing where the mesh was further refined reaching a total of about  $7.9 \times 10^6$  tetrahedral elements (Figure 6). Such a large number of computational 247 248 cells is necessary for reaching mesh-independent solutions given the significant levels of numerical 249 diffusion of tetrahedral elements.



(a)

(b)



Figure 6: Mesh used for the CFD simulations corresponding to the "ideal geometry" (a) and the "real geometry" (b) of the micromixer.

#### 250 2.3.2. Hydrodynamics modeling

For both ideal and real geometries, the flow field has been determined by solving the Navier-Stokes equations assuming an incompressible and isothermal flow (and hence, uniform water properties):

$$div(\vec{u}) = 0 \tag{1}$$

255 
$$\rho \frac{\partial \vec{u}}{\partial t} + \rho \, \overline{div}(\vec{u} \otimes \vec{u}) = -\vec{\nabla}p^* + \mu \, \vec{\Delta} \, \vec{u}$$
(2)

where *t* denotes time,  $\vec{u}$  the velocity vector,  $\rho$  the water density,  $\mu$  its dynamic viscosity and  $p^*$  the driving pressure.

258 The first term in Equation 2 describes transient flow behavior. Since this study focuses on the 259 steady-state regime only, this term has been considered null and a steady-state solver has been used 260 to compute the steady-state solution. However, for the "real geometry" case under a flow rate of 261 4.2 ml/min at each inlet, convergence could not be achieved this way. Indeed, under these 262 operating conditions, the machining chips led to some low amplitude vortex shedding phenomena. Hence, for this case only, the full unsteady Navier-Stokes equations were solved until reaching the 263 264 permanent flow regime where the time-averaged solution was computed. Advancement in time was achieved through a second-order implicit scheme with time steps of  $10^{-4}s$ . 265

Simulations were conducted considering uniform velocity profiles at the inlets, uniform pressure
at the outlet and no-slip condition at the wall. The advective terms have been discretized using the
QUICK scheme and the diffusion terms were central-differenced.

269 Since MRI measurements correspond to space-averaged (namely along the depth of the sample 270 slices) velocities, in order to allow reliable comparison, CFD data were interpolated over a coarser grid corresponding to the MRI slice thickness in order to achieve the same spatial resolutionbetween experimental and numerical results (see Fig.S4).

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#### 2.3.3. CFD investigation of mixing efficiency

As detailed later in the results section, flow simulations revealed that manufacturing defects and the presence of residual machining chips have significant effects on the hydrodynamics in the micromixer, which is expected to substantially impact the mixing efficiency. Accordingly, mixing simulations were conducted using both the "ideal" and "real" geometries, and that for all of the 8 flow rates investigated in this study (

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These simulations consist in continuously injecting a tracer (i.e. a passive scalar) at one inlet where its concentration is assumed to be uniform and equal to  $C_{max}$ , whereas at the second inlet, its concentration is considered to be zero, i.e. tracer-free water is injected. It should be underlined that the value of  $C_{max}$  is arbitrarily chosen as it does not influence the mixing quality. Indeed, the tracer concentration is assumed to have no effect on the water density or physiochemical properties.

286 The concentration field is determined by solving the mass transport equation:

287 
$$\frac{\partial C}{\partial t} + div(C\vec{u}) = D\Delta C$$
(3)

where *C* is the tracer local concentration and *D* its molecular diffusivity in the liquid, taken here as  $2 \times 10^{-9} m^2 s^{-1}$  which corresponds to water self-diffusion coefficient at 20 °*C*. At the outlet section, a zero normal-concentration-gradient boundary condition is assumed. The first term in Equation 3 corresponds to temporal variations. As for the flow simulations (Section 2.3.2), it was considered null apart for the case involving the "real geometry" under a flow rate of 4.2 ml/min at each inlet where no steady-state solution could be achieved. For this case, Equations (1), (2) and (3) are simultaneously solved using a time step of  $10^{-4}$  *s* until reaching the permanent flow regime where the time-averaged solution is computed.

296 Mixing quality can be quantified by computing the concentration standard deviation at the outlet 297 or more correctly, the concentration mass-weighted uniformity index. However, because of the low 298 diffusivities of liquids, when dealing with laminar or transitional flows as in this study, numerical 299 diffusion effects become significant and may even exceed molecular diffusion effects. This is 300 particularly true when the computational domain involves tetrahedral cells. Therefore, despite the 301 large number of computational elements used (Section 2.3.1) which sufficiently reduces numerical 302 diffusion, mixing simulations will be only qualitatively analyzed in this article. No quantification 303 of the mixing quality will be presented so as to not cumbersome the paper with discussions on 304 numerical diffusion effects especially since no experimental measurements are available for 305 supporting mixing simulations results.

**306 3. Results and discussion** 

3.1.

307

# Hydrodynamics in the micromixer

308 *3.1.1. CFD simulations with the "ideal geometry"* 

309 CFD simulations carried out using the "ideal geometry" revealed similar flow behavior 310 (presented in Figure 7 and Fig.S5 to Fig.S11 in the supplementary material) for all eight flow rates 311 investigated (

312 ). The flow remains steady and preserves its double symmetry along the zy and zx planes. At 313 the entrance of the outlet channel (i.e., in the vicinity of the mixing zone), four vortices develop in 314 the cross-stream direction, and their intensity increases with increasing flow rate. Indeed, once 315 inertial effects are sufficient, secondary flow structures are known to form in the mixing channel 316 (Bothe et al., 2006; Lobasov et al., 2018). These vortices promote lateral displacement of the fluid 317 parcels and thus enhance the mixing quality. When progressing downstream toward the outlet, the 318 secondary flow intensity attenuates (and thus the cross-stream velocity components) as the flow 319 becomes increasingly axisymmetric and tends to fully develop into a classic Poiseuille profile. As 320 could be expected, the development length increases with the flow rate.

#### 321

#### 3.1.2. MRI velocity measurements vs CFD results

MRI velocity measurements along the outlet channel confirmed some of the general flow features predicted by the CFD simulations with the "ideal geometry" (Section 3.1.1) such as the presence of secondary flows. Nonetheless, major differences existed between experimental and numerical results. Indeed, MRI revealed that even for the lowest flow rates investigated, the double symmetry of the flow is not preserved (Figure 7 and Fig. S12 to Fig. S15 in the supplementary material). The secondary flows consisted of one or two asymmetric cells rather than four vortices as predicted by the simulations.

As discussed in Section 2.1, these large differences between numerical and experimental data raised doubts about the presence of manufacturing defects in the prototype. Accordingly, the real geometry of the micromixer was recorded using X-ray CT which revealed some minor defects as well as the presence of two machining chips that were not removed during the milling process as explained earlier. 334 CFD results obtained using the "real geometry" are in very good agreement with experimental 335 measurements as can be seen in Figure 7 (and Fig. S16 to Fig. S22 in the supplementary material), 336 and allow reproducing the high asymmetry of the velocity fields as revealed by the MRI, which 337 proves that even small defects may have a big impact on the hydrodynamics and hence on the 338 mixing efficiency in micromixers. We note that the velocity map presented on Figure 7a shows 339 important phase wrapping in the mixing zone ( $V_x$  component), manifested by an abrupt change in 340 velocity for small lateral displacements. We have not tried to correct this artifact because the axial 341 maps discussed in this paper are measured well above the mixing zone.

342 For the lowest flow rate investigated, CFD results are in excellent agreement, qualitatively and 343 quantitatively, with experimental data (Figure 8). As the flow rates is increased, although it remains 344 acceptable, the agreement between the two sets of results tends to decrease, and this for different 345 reasons. Indeed, with increasing flow rates, larger velocity gradients occur in the mixing zone, i.e. 346 velocity values vary more and more significantly along the axial position along the outlet channel, 347 in particular in the vicinity of the mixing zone. Accordingly, a small uncertainty in the exact 348 location of the MRI measurement section may lead to large differences with the numerical results 349 as the flow becomes very sensitive to axial position. Moreover, MRI images accuracy deteriorates 350 as the velocities are gradually increased. For instance, from a flow rate of 3.9 ml/min at each inlet 351 (corresponding to axial velocities in the mixing channel reaching about 30 cm/s, see Table 1), 352 measurements using the current experimental apparatus are no longer reliable as the signal-to-noise 353 ratio becomes very low (Fig. S23 in the supplementary material). The effect of temporal averaging 354 during the observation time  $\Delta$  should also be considered to better understand the limitations of the 355 measurements. Indeed, when the flow rate is high, the average fluid displacement during  $\Delta$  can be 356 much larger than the voxel sizes which can lead to significant image distortions. This should be taken into consideration for further comparison and to better delimit the range of velocitiesaccessible to the experiment.

In any case, in the range of the measurable velocities, the successful comparison between MRI measurements and CFD data allows validating the presence of complex flow features in the mixing channels and reveals the great impact that small manufacturing defects may have on the hydrodynamics in micromixers.

363

364

(a)







(b) slice position P1







Figure 7: 2D Velocity maps obtained by CFD simulations on ideal and real geometry and by MRI measurements are presented from left to right respectively. The flow rate on each inlet is 1.4 ml/min corresponding to  $Re_{outlet}$  of 60. (a) 2D velocity maps measured in an orthogonal plane containing the three channels. From (b) to (d): 2D axial velocity maps at slice positions P1, P2 and P3 corresponding respectively to 2 mm, 3 mm and 4 mm from the mixing point. The transverse vector  $\overrightarrow{V_{xy}}$  is represented with arrows superimposed to the velocity map of the z –component, the color of the arrows (black-and-white scale) and their width representing the norm of the vector. Velocity profiles along a centerline in the x and y directions allow quantitative comparison of MRI to CFD. The distances on the axes are in millimeters.









(c)



Figure 8: 3D velocity maps obtained by MRI measurements and by CFD simulations on real geometry for different Reynolds number on the outlet at slice positions P1 and P2 corresponding respectively to 2mm and 3mm from the mixing point. (a), (b) and (c) correspond to Re<sub>outlet</sub> of 80, 102 and 123 respectively. Velocity profiles are plotted along central lines in the x and y directions for the quantitative comparison between CFD simulations (ideal and real geometries) and MRI measurements.

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#### 3.1.3. Effects of manufacturing defects on the mixing efficiency

In the previous sections, it has been shown that manufacturing defects have a great impact on the hydrodynamics downstream the mixing zone, and thus, they are expected to significantly influence the mixing efficiency. Mixing simulations (Section 2.3.3) were conducted in order to check this eventuality.

375 Simulations revealed that the exact micromixer geometry has a great impact on the mixing 376 efficiency as can be seen from Figure 10. Indeed, in the case of the "ideal" geometry (and for all 377 eight flow rates investigated), despite the presence of secondary flows, given the double symmetry 378 of the flow, the main contact zone between the two fluid streams remains quasi-planar which 379 practically limits mixing to the sole effects of diffusion. In the case of the "real" geometry, the flow 380 perturbation induced by the manufacturing defects engenders a larger contact zone between the 381 two streams which is supposed to increase diffusive mixing (note that simulations probably 382 overestimate the thickness of the mixing front because of numerical diffusion effects). 383 Additionally, the large secondary flow cells engender engulfment of the two streams which is 384 expected to enhance the mass-transfer efficiency. Hence, it is clear that these non-voluntary 385 manufacturing defects have a positive impact on the mixing quality although they generate 386 additional mechanical energy dissipation.

When increasing the flow rate, as inertial effects becomes larger, mixing efficiency is improved in both geometries. However, while this enhancement remains insufficient with the ideal design as the main contact zone remains quasi-planar, it is much more pronounced in the case of the real geometry as the contact zone becomes larger. Moreover, as mentioned in the previous section, for a flow rate of  $4.2 \ ml/min$ , in the case of the real geometry, the presence of manufacturing chips induces some unsteadiness in the flow which is expected to further enhance the mixing quality.



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Figure 9: Concentration field at sections located 3, 4, 5, 6 and 7 mm downstream of the mixing
point: case where the flow rate at each inlet is 3.4 ml/min.

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#### 398 **4.** Conclusion

In this study, NMR imaging methods were used to study the hydrodynamics occurring in the mixing channel of a micromixer in an arrow configuration. A specific device was set up including a home-made radiofrequency coil with reduced dimensions and adapted to the geometry and the dimensions of the manufactured micromixer. In the homogeneity zone of the radiofrequency field generated by the developed milli-coil, the signal-to-noise ratio was improved by a factor 7 in comparison with a standard commercial hardware.

405 MRI measurements were performed for the imaging of the 3D velocity field in the micromixer.
406 On another hand, computational fluid dynamics (CFD) simulations of the hydrodynamics in the

407 device were carried out. The 2D velocity maps obtained by MRI measurements showed some 408 different symmetries on the fluid flow evolution and on the development of the secondary flows 409 compared to the CFD simulations performed on an "ideal" geometry of the micromixer. This 410 comparison revealed that the flow field was sensitive to the presence of small manufacturing 411 defects. Accounting for these imperfections in the numerical model (real geometry of the 412 micromixer extracted by X-ray  $\mu$ CT) allowed obtaining, in the range of the flow rates investigated, 413 a good agreement between experimental results and numerical data, hence confirming the 414 potentiality of the developed NMR instrumentation for characterizing flow phenomena in 415 millimetric devices.

As the flow field was very sensitive to the manufacturing defects, the mixing efficiency was also impacted. The simulations on the real geometry revealed that the defects engendered a larger contact zone between the two fluid streams and their engulfment. This is expected to enhance both mass-transfer and diffusive mixing.

#### 420 **References**

- Ahola, S., Casanova, F., Perlo, J., Münnemann, K., Blümich, B., & Stapf, S. (2006). Monitoring
  of fluid motion in a micromixer by dynamic NMRmicroscopy. *Lab Chip*, *6*(1), 90–95.
  https://doi.org/10.1039/B510708C
- Ahola, S., Telkki, V.-V., & Stapf, S. (2012). Velocity distributions in a micromixer measured by
  NMR imaging. *Lab on a Chip*, *12*(10), 1823. https://doi.org/10.1039/c2lc21214e
- Akpa, B. S., Matthews, S. M., Sederman, A. J., Yunus, K., Fisher, A. C., Johns, M. L., & Gladden,
  L. F. (2007). Study of Miscible and Immiscible Flows in a Microchannel Using Magnetic
  Resonance Imaging. *Analytical Chemistry*, *79*(16), 6128–6134.
  https://doi.org/10.1021/ac070364a

- Bayareh, M., Ashani, M. N., & Usefian, A. (2020). Active and passive micromixers: A
  comprehensive review. *Chemical Engineering and Processing Process Intensification*,
  147, 107771. https://doi.org/10.1016/j.cep.2019.107771
- 433 Blümich Bernhard. (2000). *NMR imaging of materials*. Clarendon Press Oxford University Press.
- Bothe, D., Stemich, C., & Warnecke, H.-J. (2006). Fluid mixing in a T-shaped micro-mixer. *Chemical Engineering Science*, 61(9), 2950–2958.
  https://doi.org/10.1016/j.ces.2005.10.060
- 437 Callaghan, P. T., & Xia, Y. (1991). Velocity and diffusion imaging in dynamic NMR microscopy.
  438 *Journal of Magnetic Resonance (1969)*, *91*(2), 326–352. https://doi.org/10.1016/0022439 2364(91)90196-Z
- 440 Callaghan P. T. (1991). *Principles of nuclear magnetic resonance microscopy*. Clarendon press.
- 441 Caprihan, A., & Fukushima, E. (1990). Flow measurements by NMR. *Physics Reports*, *198*(4),
  442 195–235. https://doi.org/10.1016/0370-1573(90)90046-5
- Ciobanu, L., Jayawickrama, D. A., Zhang, X., Webb, A. G., & Sweedler, J. V. (2003). Measuring
  Reaction Kinetics by Using Multiple Microcoil NMR Spectroscopy. *Angewandte Chemie*

445 International Edition, 42(38), 4669–4672. https://doi.org/10.1002/anie.200351901

- Commenge, J.-M., & Falk, L. (2011). Villermaux–Dushman protocol for experimental
  characterization of micromixers. *Chemical Engineering and Processing: Process Intensification*, 50(10), 979–990. https://doi.org/10.1016/j.cep.2011.06.006
- 449 Chung, K. Y. (1993). Dean Vortices in Curved Tube Flow: 5. 3-D MRI and Numerical Analysis
- 450 of the Velocity Field. *AIChE Journal*, 39(10), 1592-1602.
- 451 https://doi.org/10.1002/aic.690391003

- 452 Dechow, J., Forchel, A., Lanz, T., & Haase, A. (2000). Fabrication of NMR Microsensors for
  453 nanoliter sample volumes. *Microelectronic Engineering*, 53(1–4), 517–519.
  454 https://doi.org/10.1016/S0167-9317(00)00368-3
- 455 Dreher, S., Kockmann, N., & Woias, P. (2009). Characterization of Laminar Transient Flow
- 456 Regimes and Mixing in T-shaped Micromixers. *Heat Transfer Engineering*, 30(1–2), 91–
  457 100. https://doi.org/10.1080/01457630802293480
- Engler, M., Kockmann, N., Kiefer, T., & Woias, P. (2004). Numerical and experimental
  investigations on liquid mixing in static micromixers. *Chemical Engineering Journal*, *101*(1–3), 315–322. https://doi.org/10.1016/j.cej.2003.10.017
- Eroglu, S., Gimi, B., Roman, B., Friedman, G., & Magin, R. L. (2003). NMR spiral surface
  microcoils: Design, fabrication, and imaging. *Concepts in Magnetic Resonance*, *17B*(1), 1–
  10. https://doi.org/10.1002/cmr.b.10068
- 464 Falk, L., & Commenge, J.-M. (2010). Performance comparison of micromixers. *Chemical*465 *Engineering Science*, 65(1), 405–411. https://doi.org/10.1016/j.ces.2009.05.045
- 466 Ferrari, M., Moyne, C., & Stemmelen, D. (2018). Study of Dispersion in Porous Media by Pulsed
- 467 Field Gradient NMR: Influence of the Fluid Rheology. *Transport in Porous Media*, 123(1),
- 468 101–124. https://doi.org/10.1007/s11242-018-1027-0
- 469 Fu, R., Brey, W. W., Shetty, K., Gor'kov, P., Saha, S., Long, J. R., Grant, S. C., Chekmenev, E.
- 470 Y., Hu, J., Gan, Z., Sharma, M., Zhang, F., Logan, T. M., Brüschweller, R., Edison, A.,
- 471 Blue, A., Dixon, I. R., Markiewicz, W. D., & Cross, T. A. (2005). Ultra-wide bore 900MHz
- 472 high-resolution NMR at the National High Magnetic Field Laboratory. *Journal of Magnetic*
- 473 *Resonance*, *177*(1), 1–8. https://doi.org/10.1016/j.jmr.2005.07.013
- 474 Galletti, C., Mariotti, A., Siconolfi, L., Mauri, R., & Brunazzi, E. (2019). Numerical investigation
- 475 of flow regimes in T-shaped micromixers: Benchmark between finite volume and spectral

- 476 element methods. *The Canadian Journal of Chemical Engineering*, 97(2), 528–541.
  477 https://doi.org/10.1002/cjce.23321
- Gladden, L. F., & Alexander, P. (1996). Applications of nuclear magnetic resonance imaging in
  process engineering. *Measurement Science and Technology*, 7(3), 423–435.
  https://doi.org/10.1088/0957-0233/7/3/026
- Harel, E., & Pines, A. (2008). Spectrally resolved flow imaging of fluids inside a microfluidic chip
  with ultrahigh time resolution. *Journal of Magnetic Resonance*, *193*(2), 199–206.
  https://doi.org/10.1016/j.jmr.2008.04.037
- 484 Hessel, V., Löwe, H., & Schönfeld, F. (2005). Micromixers—A review on passive and active
- 485 mixing principles. *Chemical Engineering Science*, 60(8–9), 2479–2501.
   486 https://doi.org/10.1016/j.ces.2004.11.033
- Hoult, D. I., & Richards, R. E. (2011). The signal-to-noise ratio of the nuclear magnetic resonance
  experiment. *Journal of Magnetic Resonance*, 213(2), 329–343.
  https://doi.org/10.1016/j.jmr.2011.09.018
- Hsieh, S.-S. (2013). Mixing efficiency of Y-type micromixers with different angles. *International Journal of Heat and Fluid Flow*, 10.
- 492 Ionin, B. I., & Ershov, B. A. (1995). NMR Spectroscopy in Organic Chemistry. Springer US.
  493 https://doi.org/10.1007/978-1-4684-1785-2
- 494 Klein, M., Perrin, J. C., Leclerc, S., Guendouz, L., Dillet, J. & Lottin, O. (2013). Spatially and
- 495 Temporally Resolved Measurement of Water Distribution in Nafion Using NMR Imaging.
- 496 *ECS Transactions*, 58 (1), 283-289. https://doi.org/10.1149/05801.0283ecst

Klein, M., Perrin, J.-C., Leclerc, S., Guendouz, L., Dillet, J., & Lottin, O. (2013). Anisotropy of
Water Self-Diffusion in a Nafion Membrane under Traction. *Macromolecules*, 46(23),

499 9259–9269. https://doi.org/10.1021/ma401511t

- 500 Kovacs, H., Moskau, D., & Spraul, M. (2005). Cryogenically cooled probes—A leap in NMR
- 501 technology. Progress in Nuclear Magnetic Resonance Spectroscopy, 46(2–3), 131–155.
- 502 https://doi.org/10.1016/j.pnmrs.2005.03.001
- Lacey, M. E., Subramanian, R., Olson, D. L., Webb, A. G., & Sweedler, J. V. (1999). High Resolution NMR Spectroscopy of Sample Volumes from 1 nL to 10 μL. *Chemical Reviews*,

505 99(10), 3133–3152. https://doi.org/10.1021/cr980140f

- Lee, C.-Y., & Fu, L.-M. (2018). Recent advances and applications of micromixers. *Sensors and Actuators B: Chemical*, 259, 677–702. https://doi.org/10.1016/j.snb.2017.12.034
- 508 Lobasov, A. S., Minakov, A. V., Kuznetsov, V. V., Rudyak, V. Ya., & Shebeleva, A. A. (2018).

509 Investigation of mixing efficiency and pressure drop in T-shaped micromixers. *Chemical* 

- 510 Engineering and Processing Process Intensification, 134, 105–114. 511 https://doi.org/10.1016/j.cep.2018.10.012
- Massin, C., Boero, G., Vincent, F., Abenhaim, J., Besse, P.-A., & Popovic, R. S. (2002). High-Q
  factor RF planar microcoils for micro-scale NMR spectroscopy. *Sensors and Actuators A: Physical*, 97–98, 280–288. https://doi.org/10.1016/S0924-4247(01)00847-0
- Meier, R. C., Höfflin, J., Badilita, V., Wallrabe, U., & Korvink, J. G. (2014). Microfluidic
  integration of wirebonded microcoils for on-chip applications in nuclear magnetic
  resonance. *Journal of Micromechanics and Microengineering*, 24(4), 045021.
  https://doi.org/10.1088/0960-1317/24/4/045021
- Mendez, D. L. M., Lemaitre, C., Castel C., Ferrari, M., Simonaire, H. & Favre E. (2017).
  Membrane contactors for process intensification of gas absorption into physical solvents:

- 521 Impact of dean vortices. Journal of Membrane Science, 530, 20-32,
  522 http://dx.doi.org/10.1016/j.memsci.2017.02.016
- 523 Milc, K. W., Serial, M. R., Philippi, J., Dijksman, J. A., van Duynhoven, J. P. M., & Terenzi, C.
- 524 (2022). Validation of temperature-controlled rheo-MRI measurements in a submillimeter-
- 525 gap Couette geometry. *Magnetic Resonance in Chemistry*, 60(7), 606–614. 526 https://doi.org/10.1002/mrc.5157
- Murphree, D., Cahn, S. B., Rahmlow, D., & DeMille, D. (2007). An easily constructed, tuning free,
  ultra-broadband probe for NMR. *Journal of Magnetic Resonance*, 188(1), 160–167.
  https://doi.org/10.1016/j.jmr.2007.05.025
- 530 Oliveira, A. V. S., Stemmelen, D., Leclerc, S., Glantz, T., Labergue, A., Repetto, G., & Gradeck,
- 531 M. (2020). Velocity field and flow redistribution in a ballooned 7×7 fuel bundle measured
- 532 by magnetic resonance velocimetry. *Nuclear Engineering and Design*, 369, 110828.

533 https://doi.org/10.1016/j.nucengdes.2020.110828

- 534 Olson, D. L., Peck, T. L., Webb, A. G., Magin, R. L., & Sweedler, J. V. (1995). High-Resolution
- 535 Microcoil <sup>1</sup> H-NMR for Mass-Limited, Nanoliter-Volume Samples. *Science*, 270(5244),
  536 1967–1970. https://doi.org/10.1126/science.270.5244.1967
- 537 Peck, T. L., Magin, R. L., & Lauterbur, P. C. (1995). Design and Analysis of Microcoils for NMR
  538 Microscopy. *Journal of Magnetic Resonance, Series B*, 108(2), 114–124.
  539 https://doi.org/10.1006/jmrb.1995.1112
- 540 Peng, K., Xu, F., Yang, L., Yao, C., & Chen, G. (2022). Dean instability and vortex-induced mixing
- 541 for two miscible fluids in T-micromixers. *Chemical Engineering and Processing Process*
- 542 Intensification, 176, 108975. https://doi.org/10.1016/j.cep.2022.108975

543	Perrin, JC., Waldner, C., Bossu, J., Chatterjee, A. & Hirn, U. (2022). Real time monitoring of the					
544	through thickness moisture profile of thin sheets using NMR. Chemical Engineering					
545	Science, 251, 117464. https://doi.org/10.1016/j.ces.2022.117464					
546	Pettegrew, J. W. (2012). NMR: Principles and Applications to Biomedical Research. Springer					
547	Science & Business Media.					
548	Robert M., Kaddouri A. E., Perrin JC., Leclerc S. & Lottin O. (2018). Towards a NMR-Based					
549	Method for Characterizing the Degradation of Nafion XL Membranes for PEMFC. Journal					
550	of The Electrochemical Society, 165, n°6F3209-F3216.					
551	https://doi.org/10.1149/2.0231806jes					
552	Seymour, J. D., & Callaghan, P. T. (1997). Generalized approach to NMR analysis of flow and					
553	dispersion in porous media. AIChE Journal, 43(8), 2096–2111.					
554	https://doi.org/10.1002/aic.690430817					
555	Stapf, S., & Han, SI. (2006). NMR Imaging in Chemical Engineering. John Wiley & Sons,					
556	Incorporated. http://ebookcentral.proquest.com/lib/ulorraine/detail.action?docID=481636					
557	Trumbull, J. D., Glasgow, I. K., Beebe, D. J., & Magin, R. L. (2000). Integrating microfabricated					
558	fluidic systems and NMR spectroscopy. IEEE Transactions on Biomedical Engineering,					
559	47(1), 3–7. https://doi.org/10.1109/10.817611					
560	Webb, A. G. (2008). Microcoil nuclear magnetic resonance spectroscopy. In NMR Spectroscopy					
561	in Pharmaceutical Analysis (p. 48).					
562	Webb, A. G. (2012). Increasing the Sensitivity of Magnetic Resonance Spectroscopy and Imaging.					
563	Analytical Chemistry, 84(1), 9-16. https://doi.org/10.1021/ac201500v					
564	Webb, A. G. (2013). Radiofrequency microcoils for magnetic resonance imaging and spectroscopy.					
565	Journal of Magnetic Resonance, 229, 55-66. https://doi.org/10.1016/j.jmr.2012.10.004					
	37					

566	Wensink, H., Benito-Lopez, F., Hermes, D. C., Verboom, W., Gardeniers, H. J. G. E., Reinhoudt,			
567	D. N., & van den Berg, A. (2005). Measuring reaction kinetics in a lab-on-a-chip by			
568	microcoil NMR. Lab on a Chip, 5(3), 280. https://doi.org/10.1039/b414832k			
569	Woytasik, M., Grandchamp, JP., Dufour-Gergam, E., Gilles, JP., Megherbi, S., Martincic, E.,			
570	Mathias, H., & Crozat, P. (2006). Two- and three-dimensional microcoil fabrication process			
571	for three-axis magnetic sensors on flexible substrates. Sensors and Actuators A: Physical,			
572	132(1), 2–7. https://doi.org/10.1016/j.sna.2006.06.062			
573	Wüthrich, K. (1990). Protein structure determination in solution by NMR spectroscopy. Journal of			
574	Biological Chemistry, 265(36), 22059–22062. https://doi.org/10.1016/S0021-			
575	9258(18)45665-7			
576	Zhang, J., & Balcom, B. J. (2010). Parallel-plate RF resonator imaging of chemical shift resolved			
577	capillary flow. <i>Magnetic Resonance Imaging</i> , 28(6), 826–833.			
578	https://doi.org/10.1016/j.mri.2010.03.033			
579	Zhang, X., & Webb, A. G. (2005). Magnetic Resonance Microimaging and Numerical Simulations			
580	of Velocity Fields Inside Enlarged Flow Cells Used for Coupled NMR Microseparations.			
581	Analytical Chemistry, 77(5), 1338–1344. https://doi.org/10.1021/ac048532b			
582				
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# **3D** Magnetic resonance velocimetry for hydrodynamics characterization in microdevices: application to micromixers and comparison to CFD simulations

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Fig.S1: visualization of the 3D NMR signal intensity image on the Paraview software. Front view (a) and side view (b) of the machined micromixer filled with the copper sulfate solution



Fig.S2 : analysis and quantification of the misalignment of the three axes of the micromixer channels. (a) Views in 3 cross-sectional planes of the projections of the points in each channel and of the centerlines which constitute the axes of the microchannels. (b) Zooms on the three views near the mixing point, showing misalignment of the three microchannels: the two inlet centerlines and the outlet centerline did not meet exactly at the mixing point. The measured distances between the axes were about 90  $\mu$ m and 55  $\mu$ m

7.5

× 10<sup>-3</sup>

6.5

1 L 6

6.5

8 × 10<sup>-3</sup>

7.5

6.5

7.5

× 10<sup>-3</sup>



Fig.S3: Analysis of the images near the intersection of the 3 microchannels. Views of the sections of the 3 channels at different heights from the bottom of the micromixer, showing the presence of PMMA residues.



Fig.S4: interpolation of CFD data over coarser grid to correspond to MRI measurement conditions. (a) for axial images and (b) for images in planes orthogonal to the axial images. Construction of a uniform mesh with the "PointVolume Interpolator" function of Paraview. Reconstruction of the resampled data with Python as 3D matrices for each velocity component to perform averaging and display the results.

(a)



Fig.S5: 3D velocity maps obtained with CFD simulation on ideal geometry of the micromixer. The flow rate on each inlet is 1.9ml/min corresponding to  $Re_{outlet}$  of 80. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig.S6: 3D velocity maps obtained with CFD simulation on ideal geometry of the micromixer. The flow rate on each inlet is 2.4ml/min corresponding to  $Re_{outlet}$  of 102. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig.S7: 3D velocity maps obtained with CFD simulation on ideal geometry of the micromixer. The flow rate on each inlet is 2.9ml/min corresponding to  $Re_{outlet}$  of 123. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig.S8: 3D velocity maps obtained with CFD simulation on ideal geometry of the micromixer. The flow rate on each inlet is 3.2ml/min corresponding to  $Re_{outlet}$  of 136. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig.S9: 3D velocity maps obtained with CFD simulation on ideal geometry of the micromixer. The flow rate on each inlet is 3.4ml/min corresponding to  $Re_{outlet}$  of 144. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig.S10: 3D velocity maps obtained with CFD simulation on ideal geometry of the micromixer. The flow rate on each inlet is 3.9ml/min corresponding to  $Re_{outlet}$  of 165. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig.S11: 3D velocity maps obtained with CFD simulation on ideal geometry of the micromixer. The flow rate on each inlet is 4.2ml/min corresponding to  $Re_{outlet}$  of 178. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig.S12: 3D velocity maps obtained with the developed milli-coil. The flow rate on each inlet is 1.9ml/min corresponding to  $Re_{outlet}$  of 80. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2 and P3 corresponding respectively to 2mm, 2.5mm and 3mm from the mixing point.



Fig. S13: 3D velocity maps obtained with the developed milli-coil. The flow rate on each inlet is 2.4ml/min corresponding to  $Re_{outlet}$  of 102. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2 and P3 corresponding respectively to 2mm, 2.5mm and 3mm from the mixing point.



Fig.S14: 3D velocity maps obtained with the developed milli-coil. The flow rate on each inlet is 2.9ml/min corresponding to  $Re_{outlet}$  of 123. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2 and P3 corresponding respectively to 2mm, 2.5mm and 3mm from the mixing point.



Fig. S15: 3D velocity maps obtained with the developed milli-coil. The flow rate on each inlet is 3.2ml/min corresponding to Re<sub>outlet</sub> of 136. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2 and P3 corresponding respectively to 2mm, 2.5mm and 3mm from the mixing point.



Fig. S16: 3D velocity maps obtained with CFD simulation on real geometry of the micromixer. The flow rate on each inlet is 1.9 ml/min corresponding to  $Re_{outlet}$  of 80. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig. S17: 3D velocity maps obtained with CFD simulation on real geometry of the micromixer. The flow rate on each inlet is 2.4 ml/min corresponding to  $Re_{outlet}$  of 102. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig. S18: 3D velocity maps obtained with CFD simulation on real geometry of the micromixer. The flow rate on each inlet is 2.9 ml/min corresponding to  $Re_{outlet}$  of 123. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig. S19: 3D velocity maps obtained with CFD simulation on real geometry of the micromixer. The flow rate on each inlet is 3.2 ml/min corresponding to  $Re_{outlet}$  of 136. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig. S20: 3D velocity maps obtained with CFD simulation on real geometry of the micromixer. The flow rate on each inlet is 3.4 ml/min corresponding to  $Re_{outlet}$  of 144. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig. S21: 3D velocity maps obtained with CFD simulation on real geometry of the micromixer. The flow rate on each inlet is 3.9 ml/min corresponding to  $Re_{outlet}$  of 165. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig. S22: 3D velocity maps obtained with CFD simulation on real geometry of the micromixer. The flow rate on each inlet is 4.2 ml/min corresponding to  $Re_{outlet}$  of 178. The velocity maps in the orthogonal plane to the main flow direction are obtained at slice positions P1, P2, P3 and P4 corresponding respectively to 2mm, 3mm, 4mm and 7mm from the mixing point.



Fig. S23: 3D velocity maps obtained with the developed milli-coil (3.9 ml/min per inlet corresponding to  $Re_{outlet}$  of 165) illustrating the limit of the MRI measurement