

This item is the archived peer-reviewed author-version of:

Dry reforming in a dielectric barrier discharge reactor with non-uniform discharge gap : effects of metal rings on the discharge behavior and performance

Reference:

Wang Jinxin, Zhang Kaimin, Meynen Vera, Bogaerts Annemie.- Dry reforming in a dielectric barrier discharge reactor with non-uniform discharge gap : effects of metal rings on the discharge behavior and performance
Chemical engineering journal - ISSN 1873-3212 - 465(2023), 142953
Full text (Publisher's DOI): <https://doi.org/10.1016/J.CEJ.2023.142953>
To cite this reference: <https://hdl.handle.net/10067/1956030151162165141>

Dry Reforming in a Dielectric Barrier Discharge Reactor with Non-uniform Discharge Gap: Effects of Metal Rings on the Discharge Behavior and Performance

Jinxin Wang^{a,b}, Kaimin Zhang^a, Vera Meynen^{a,*} and Annemie Bogaerts^{b,*}

^aLaboratory of Adsorption and Catalysis, Department of Chemistry, University of Antwerp, Universiteitsplein 1, 2610 Wilrijk, Antwerp, Belgium

^bPlasma Lab for Applications in Sustainability and Medicine - ANTwerp, Department of Chemistry, University of Antwerp, Universiteitsplein 1, 2610 Wilrijk, Antwerp, Belgium

*Email: vera.meynen@uantwerpen.be; annemie.bogaerts@uantwerpen.be

Abstract

The application of dielectric barrier discharge (DBD) plasma reactors is promising in various environmental and energy processes, but is limited by their low energy yield. In this study, we put a number of stainless steel rings over the inner electrode rod of the DBD reactor to change the local discharge gap and electric field, and we studied the dry reforming performance. At 50 W supplied power, the metal rings mostly have a negative impact on the performance, which we attribute to the non-uniform spatial distribution of the discharges caused by the rings. However, at 30 W supplied power, the energy yield is higher than at 50 W and the placement of the rings improves the performance of the reactor. More rings and with a larger cross-sectional diameter can further improve the performance. The reactor with 20 rings with a 3.2 mm cross-sectional diameter exhibits the best performance in this study. Compared to the reactor without rings, it increases the CO₂ conversion from 7% to 16 %, the CH₄ conversion from 12% to 23%, and the energy yield from 0.05 mmol/kJ supplied power to 0.1 mmol/kJ (0.19 mmol/kJ if calculated from the plasma power), respectively. The presence of the rings increases the local electric field, the displaced charge and the discharge fraction, and also makes the discharge more stable and with more uniform intensity. It also slightly improves the selectivity to syngas. The performance improvement observed by placing stainless steel rings in this study may also be applicable to other plasma-based processes.

Keywords: Dry reforming; Plasma-based process; Dielectric barrier discharge reactor; Non-uniform discharge gap; Energy yield improvement

1. Introduction

The application of plasma technology in chemical reactions is attractive and has been widely studied in a variety of processes, such as the decomposition of pollutants and synthesis of chemical products [1-3]. CO₂ reforming of CH₄ (dry reforming) is one of them. It can produce CO and H₂ (which could be used to produce clean fuels and chemicals) from renewable sources while reducing emissions of two greenhouse gases [4-7]. In the traditional thermal dry reforming, CH₄ and CO₂ need to be activated at a high temperature (at least 800 °C), resulting in problems of coking and energy losses [8]. Performing dry reforming in a plasma reactor can overcome these problems, because the high-energy electrons in non-equilibrium plasma are able to activate the molecules at relatively lower gas temperatures (lower than 250 °C) [9-14].

A common non-equilibrium plasma reactor is the dielectric barrier discharge (DBD) reactor, which is simple in design and can be operated at ambient temperature and pressure [5,15-17]. However, one of the main drawbacks of DBD reactors is their relatively low energy yield [8,18,19].

42 Khoja et al. studied the dry reforming of methane in DBD reactors with different dielectric materials
43 and configurations, and a better energy yield of 0.085 mmol CO₂ and CH₄ converted per kJ plasma
44 power was obtained for an alumina dielectric barrier [4]. Tu et al. [8] obtained an energy yield of
45 0.1 mmol/kJ in a DBD reactor combined with a catalyst, and this could be further improved to a
46 maximum of 0.19 mmol/kJ by increasing the gas flow rate and reducing the specific energy input.
47 Important to note that the energy yield in the above literature was calculated from the plasma power,
48 and it could be about half to two-thirds of the value if it was calculated from the supplied power.
49 The plasma discharge mode in a DBD reactor is mainly filamentary, and thus the plasma is not
50 uniform in the whole space [20,21]. High-energy electrons exist almost exclusively in the discharge
51 filaments [22,23]. The CH₄ and CO₂ molecules hit by the discharge filaments are dissociated upon
52 electron collisions into various radicals, which can recombine into the products within the filaments
53 or afterglow [23-25]. In addition to the dry reforming products, CO and H₂, some higher
54 hydrocarbons can also be formed by radical recombination reactions [24,25]. The strength of the
55 discharge filaments varies, which may result in poor product selectivity [26], and a too weak or too
56 strong discharge causes energy waste, which may be one of the reasons for the lower energy yield
57 of the DBD reactor [18,27].

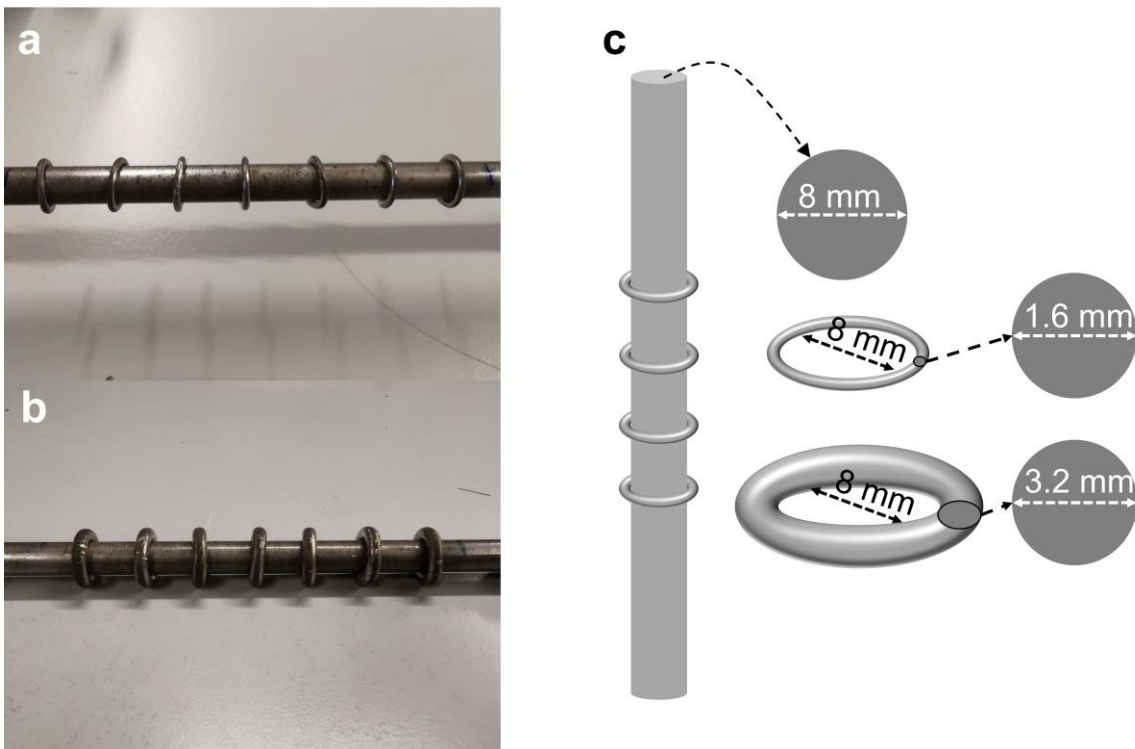
58 Changing the design of the reactor is one method to improve the performance. DBD reactors
59 with small discharge gaps (i.e., micro DBD reactors) may be able to increase the performance of
60 dry reforming, due to the enhancement of the electric field [28,29]. Previous studies found that a
61 reduced discharge gap did improve the conversion in DBD reactors at the same space time [30,31].
62 However, the same space time was maintained by reducing the gas flow rate, as the smaller gap
63 reduced the discharge volume, and thus the energy yield of the conversion process is actually
64 decreased.

65 In this study, we put a number of stainless steel rings with circular cross-sections over the
66 stainless steel inner electrode rod of the reactor (see Fig. 1). These rings reduce the local discharge
67 gap and change the electric field, which may promote the discharge with less effect on the discharge
68 volume. This may induce the discharge to take place on the rings, which might change the discharge
69 distribution and make the plasma more uniform, thereby changing the performance of the plasma-
70 based dry reforming. Since the material of the rings is similar to that of the stainless steel inner
71 electrode, and the rings cause a quite limited increase in the surface area of the inner electrode, the
72 possible catalytic effect of the stainless steel rings is not considered in this study. In addition, in a
73 plasma-based process with a catalyst, the metal active components on the catalyst surface will also
74 be in contact with the electrodes and might have a similar effect, but the performance caused by
75 the physical and chemical (surface) effects cannot be distinguished in a catalyst. Therefore, the role
76 of the metal rings in this study can also provide a reference for the study of catalysts with metal
77 components.

78 **2. Experimental**

79 **2.1. Set-up for plasma-based dry reforming**

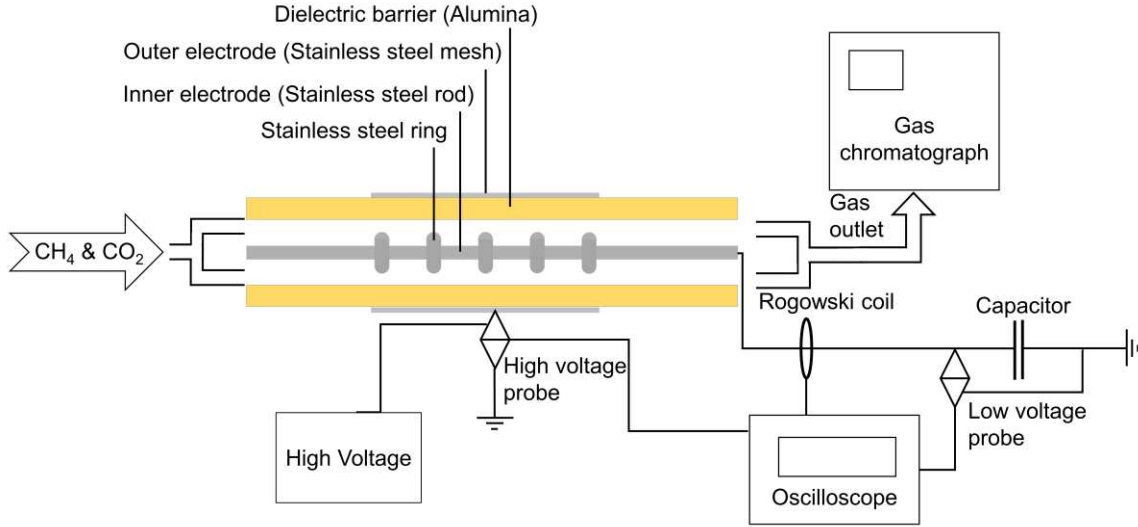
80 A coaxial cylindrical DBD reactor was applied for plasma-based dry reforming. The original
81 inner electrode was a smooth stainless steel rod with a diameter of 8 mm. A number of stainless
82 steel rings with an inner diameter of 8 mm were put over the inner electrode at even intervals, as
83 illustrated in Fig. 1. The cross-section of the rings is circular, and rings with two cross-sectional
84 diameters, i.e., 1.6 mm and 3.2 mm, were used in this study.



85

86 **Fig. 1.** Photographs of the stainless steel rod inner electrode, with stainless steel rings with cross-sectional diameters of
 87 (a) 1.6 mm and (b) 3.2 mm. (c) Schematic diagram of stainless steel rod, stainless steel rings and their dimensions.

88 Fig. 2 shows the whole set-up for the dry reforming experiments. The inner electrode was
 89 grounded. An alumina tube with an inner diameter of 17.4 mm and an outer diameter of 21.8 mm
 90 was coaxially placed over the inner electrode as a dielectric barrier, so the discharge gap of the
 91 ring-free reactor, which is the spacing between the 8 mm inner electrode and dielectric barrier, was
 92 about 4.7 mm. A stainless steel mesh tightly wound around the dielectric barrier tube was used as
 93 the external electrode. It was connected to high voltage supplied by a function generator (Tektronix,
 94 AFG 2021) and a high voltage amplifier (TREK, model 20/20C-HS). The length of the outer
 95 electrode was 100 mm, which defined the discharge length. According to the parameters of the
 96 reactor, the volume of the discharge zone was calculated to be 18.8 mL. A sinusoidal alternating
 97 input signal with a frequency of 3 kHz was provided by the function generator, and was then
 98 amplified by the amplifier. The voltage was measured by a high-voltage probe (Tektronix,
 99 P6015A), and the current was monitored by a Rogowski coil (Pearson 4100). A capacitor (10 nF)
 100 and a low-voltage probe (Picotech, TA150) connected in parallel with it were connected in series
 101 with the reactor to monitor the charge. The oscilloscope and connected PC collected and displayed
 102 all the electrical signals to calculate the power in real time. The power of the power supply was
 103 kept constant by adjusting the amplitude of the input signal from the amplifier according to the
 104 calculated power on the PC.



105

106 **Fig. 2.** DBD plasma set-up and analytical system for the dry reforming experiments.

107 **2.2. Performance of plasma-based dry reforming**

108 The feed gas into the reactor was composed of 10 mL/min of CH₄ and 10 mL/min of CO₂
 109 controlled by mass flow controllers (Bronkhorst EL-FLOW Select). Since the produced CO, H₂
 110 and unknown amounts of various (oxygenated) hydrocarbons by the dry reforming reaction causes
 111 an unknown expansion coefficient and thus a possible pressure increase, while the GC always
 112 samples at a constant ambient pressure, the outlet gas composition analyzed by the GC would have
 113 systematic errors. Therefore, an internal standard gas, 10 mL/min of N₂, was added into the outlet
 114 gas to exclude these errors [32]. An online gas chromatograph (Trace GC 1310, Interscience) with
 115 a thermal conductivity detector (TCD) and a flame ionization detector (FID) was applied to analyze
 116 the composition and concentration of the outlet gas. The composition of the gas after the gas circuit
 117 was flushed for 30 minutes and before turning on the plasma, was denoted as CO_{2,in} and CH_{4,in}.
 118 The power was then turned on to generate plasma and maintained at a constant supplied power for
 119 30 min. The outlet gases were analyzed and denoted with “out”, i.e., CO_{2,out}, CH_{4,out}, CO_{out}, H_{2,out}
 120 and C_xH_yO_{z,out}. The conversion of CO₂ and CH₄ were calculated by Eq. (1) and Eq. (2):

121
$$X_{\text{CO}_2}(\%) = \frac{\text{CO}_{2,\text{in}} - \text{CO}_{2,\text{out}}}{\text{CO}_{2,\text{in}}} \times 100\% \quad (1)$$

122
$$X_{\text{CH}_4}(\%) = \frac{\text{CH}_{4,\text{in}} - \text{CH}_{4,\text{out}}}{\text{CH}_{4,\text{in}}} \times 100\% \quad (2)$$

123 The (H-based) selectivity of H₂, and the C-based selectivity of CO and of the other chemicals
 124 were defined by Eq. (3) to Eq. (5):

125
$$S_{\text{H}_2}(\%) = \frac{\text{H}_{2,\text{out}}}{2 \times (\text{CH}_{4,\text{in}} - \text{CH}_{4,\text{out}})} \times 100\% \quad (3)$$

126
$$S_{\text{CO}}(\%) = \frac{\text{CO}_{\text{out}}}{(\text{CH}_{4,\text{in}} - \text{CH}_{4,\text{out}}) + (\text{CO}_{2,\text{in}} - \text{CO}_{2,\text{out}})} \times 100\% \quad (4)$$

127
$$S_{\text{C}_x\text{H}_y\text{O}_z}(\%) = \frac{x \times \text{C}_x\text{H}_y\text{O}_{z,\text{out}}}{(\text{CH}_{4,\text{in}} - \text{CH}_{4,\text{out}}) + (\text{CO}_{2,\text{in}} - \text{CO}_{2,\text{out}})} \times 100\% \quad (5)$$

128 In this study, the energy yield (EY) of dry reforming was defined as the mmol of CO₂ and CH₄
129 that can be converted per kJ of supplied energy, as shown in Eq. (6):

$$130 \quad EY \left(\frac{\text{mmol}}{\text{kJ}} \right) = \frac{V_{\text{CO}_2} X_{\text{CO}_2} + V_{\text{CH}_4} X_{\text{CH}_4}}{P_{\text{sply}} V_m} \times \frac{1000}{60} \left(\frac{\text{Wmin}}{\text{kJ}} \right) \quad (6)$$

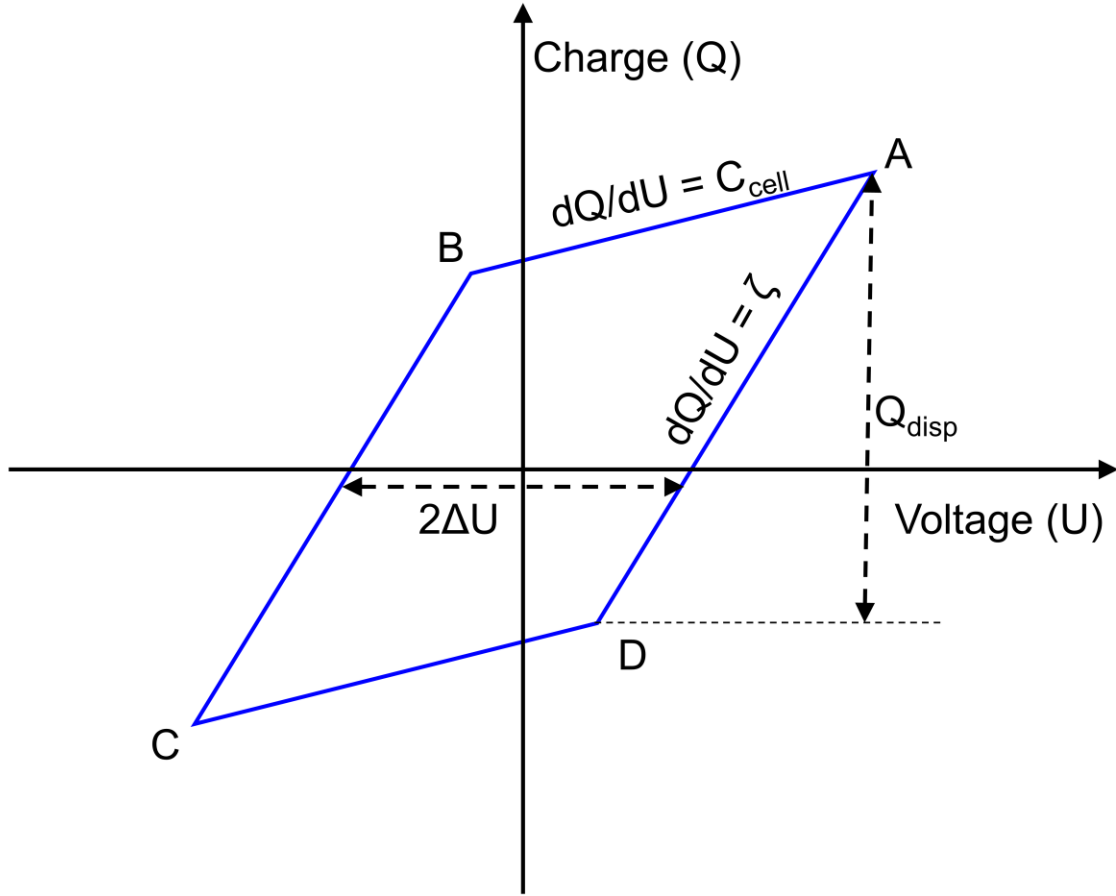
131 Where V_{CO₂} and V_{CH₄} were the volumetric flow rate of CO₂ and CH₄ in the feed gas, respectively
132 (in mL/min), and X_{CO₂} and X_{CH₄} are the conversion of CO₂ and CH₄, respectively. V_m is the molar
133 gas volume (24.4 mL/mmol). It should be noted that P_{sply} in Eq. 6 is the supplied power (in W), and
134 not the discharge power as used in most literature [8,33]. Indeed, we also want to include the effect
135 of the reactor on the discharge power in the calculation of the energy yield. If the discharge power
136 is used to calculate the EY, we will get a larger value, about 1.5–2 times larger, than the EY
137 presented in this paper.

138 2.3. Electrical characterization

139 To better understand the effect of adding the stainless steel rings on the reaction performance,
140 we performed an electrical characterization of the plasma. The above-mentioned high voltage probe,
141 low voltage probe, Rogowski coil and capacitor in the plasma set-up can collect the data of the
142 voltage, current and charge during the discharge. Errors were obtained from three repeat
143 experiments. The discharge power was calculated by voltage U and current I, via Eq. (7):

$$144 \quad P_{\text{sply}}(\text{W}) = \int_0^T UI \, dt \quad (7)$$

145 A Q-U graph, known as a Lissajous figure, was plotted using the voltage and charge data. Fig.
146 3 shows a typical Lissajous figure of the plasma in a DBD reactor. The slopes of the sides of the
147 Lissajous figure represent the equivalent capacitance of the DBD reactor at different phases, since
148 dQ/dU = C. The slopes of the Lissajous figures (dQ/dU) in this work were calculated by a Matlab
149 script [30]. In the AB and CD phases when the reactor is not discharged, the entire DBD reactor,
150 including the dielectric barrier and discharge gap, behaves as a capacitor. Therefore, the slope of
151 the AB and CD phases in the Lissajous figure is the capacitance of the capacitor formed by the
152 dielectric barrier and the discharge gap in series, denoted by C_{cell}. Plasma occurs in the BC and DA
153 phases. During the discharge, a part of the discharge gap behaves like a resistor due to the
154 breakdown of the reactive gas, while the dielectric barrier and another part of the discharge gap
155 (because the entire gap is not fully discharged in a DBD reactor) still behave as a capacitor.
156 Therefore, the slope of the BC and DA phases, denoted by ζ, represents the capacitance including
157 the dielectric barrier and the undischarged gap.



158

159 **Fig. 3.** Typical Lissajous figure of a discharge in a DBD reactor.

160 The capacitance C_{diel} of the dielectric barrier alumina tube, used in this work, is 0.266 nF, which
 161 was measured in a previous study [30]. Using C_{cell} , ζ and C_{diel} , the discharging area fraction,
 162 denoted as f , of the DBD reactor was calculated by Eq. 8 [34].

163
$$f = \frac{\zeta - C_{\text{cell}}}{C_{\text{diel}} - C_{\text{cell}}} \quad (8)$$

164 The burning voltage (U_{bur}) of the plasma in the reactor was calculated from C_{cell} , ζ , C_{diel} and the
 165 voltage at $Q = 0$ C in the Lissajous figures, i.e., ΔU , by Eq. 9 [34].

166
$$U_{\text{bur}} = \frac{1 - C_{\text{cell}}/C_{\text{diel}}}{1 - C_{\text{cell}}/\zeta} \Delta U \quad (9)$$

167 The mean electric field (E) in the reactors was calculated by the following equation [35]:

168
$$E = \frac{U_{\text{bur}}}{d_{\text{gap}}} \quad (10)$$

169 Where d_{gap} is the discharge gap, which is 4.7 mm in the reactor without rings. The electric field
 170 for the reactor with rings, as calculated in this study, was the mean electric field between the rings
 171 and the dielectric barrier, as we believe the discharge mainly takes place on the rings. The value of
 172 d_{gap} in the reactor with rings is 3.1 mm and 1.5 mm, respectively, for the rings with thickness of 1.6

173 mm and 3.2 mm. It should be noted that this mean electric field is only an approximation, because
174 in reality the potential will not drop linearly between the inner electrode (or the rings) and the outer
175 dielectric barrier, so the electric field will also not be constant, but it will be larger in the sheath
176 near the inner electrode (or rings). However, as we cannot measure the exact electric field
177 distribution inside the reactor, the above equation (10) is only an averaged estimate. Subsequently,
178 the mean electron energy and electron energy distribution function (EEDF) were calculated from
179 the mean electric field, by the Boltzmann equation using BOLSIG+ [36].

180 The charge difference between the points A and D in the Lissajous figure was the displaced
181 charge in the DA discharge phase. It was divided by the number of micro-discharges counted in
182 the current profile to get the average intensity of discharge filaments [30].

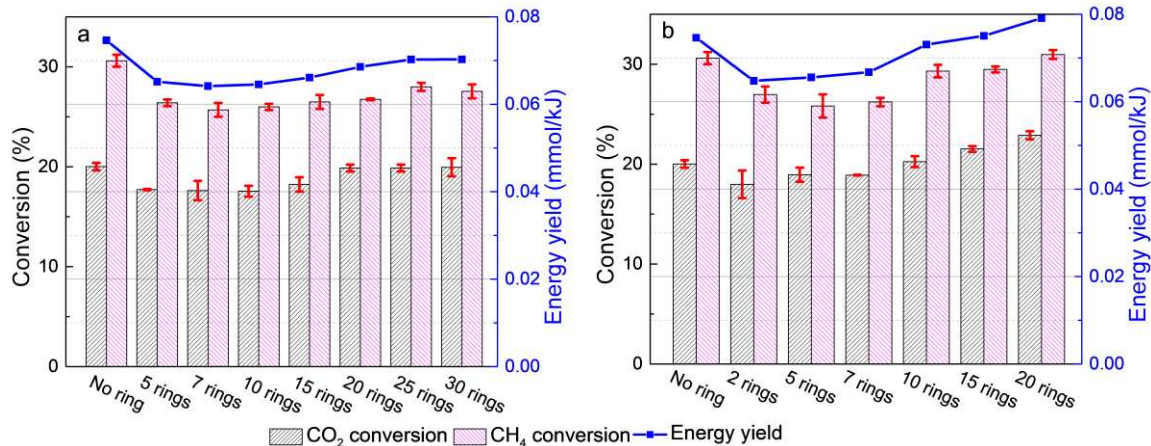
183 **3. Results and discussion**

184 The plasma-based dry reforming performance in a DBD reactor, with stainless steel rings over
185 the inner electrode, was tested at 50 W and 30 W supplied power.

186 **3.1. Plasma-based dry reforming at 50 W supplied power**

187 3.1.1. Conversion and energy yield

188 Fig. 4 illustrates the influence of placing stainless steel rings over the inner electrode rod on the
189 CH₄ and CO₂ conversions and on the energy yield of plasma-based dry reforming at 50 W supplied
190 power, for the rings with a cross-sectional diameter of 1.6 mm and 3.2 mm. A difference in
191 performance for the reactors with and without rings was observed, and also the number of rings
192 had an effect. Since the supplied power could not be controlled precisely at 50 W all the time (see
193 Table 1), sometimes changes in CH₄ and CO₂ conversions could not correctly reflect changes in
194 reactor performance. The energy yield can reflect a more accurate trend of the performance. Since
195 all experiments were performed at the same gas flow rate, and not the same space time, the specific
196 input energy (SEI) was only related to the power, and the energy yield follows a similar trend as
197 the conversion. As shown in Fig. 4a, for 1.6 mm rings, the energy yield and the reactant conversions
198 increase with the number of rings. However, they were always lower than in the reactor without
199 rings. The rings with a cross-sectional diameter of 3.2 mm (Fig. 4b) showed better performance
200 and a similar but more pronounced increasing trend upon placing more rings. The reactors with
201 only a few rings also had lower performance than the reactor without rings. However, the CO₂
202 conversion of the reactors with more than 10 rings exceeded that of the reactor with no rings, and
203 the reactor with more than 15 rings also showed a higher energy yield. The DBD reactor with 20
204 rings with 3.2 mm cross-sectional diameter exhibited the best performance, although the
205 improvement compared with the reactor without rings was limited.



206

207 **Fig. 4.** CH₄ and CO₂ conversion and energy yield of plasma-based dry reforming at 50 W supplied power, in a DBD
 208 reactor without rings, and with a varying number of stainless steel rings placed over the inner electrode, with cross-
 209 sectional diameters of (a) 1.6 mm and (b) 3.2 mm. The error bars were obtained from standard errors based on three
 210 repeat experiments with the rings kept in place.

211 Placing stainless steel rings over the inner electrode mainly has the following effects on the
 212 discharge in the DBD reactor. First of all, as introduced in the experimental section, the discharge
 213 gap between the inner electrode and the dielectric barrier was 4.7 mm (i.e., 17.4 mm – 8 mm,
 214 divided by 2). The metal rings reduced the discharge gaps at this location, to values of 3.1 mm and
 215 1.5 mm, for the 1.6 mm and 3.2 mm rings, respectively (i.e., 17.4 mm – 11.2 mm, divided by 2;
 216 and 17.4 mm – 14.4 mm, divided by 2). The reduction in discharge gap causes a higher reduced
 217 electric field strength and power density at the metal rings, leading to more displaced charge per
 218 period (see section 3.1.2 below) and a higher electron energy. Therefore, more gas molecules can
 219 be hit by the discharge filaments [37], and more successful electron impact excitation can occur in
 220 the plasma [38]. These are the positive effects of the metal rings on the plasma behavior, which
 221 should increase the performance of the reactor [39]. The 3.2 mm rings exhibit better performance
 222 than the 1.6 mm rings due to the smaller discharge gap. However, these positive effects are not well
 223 reflected between reactors with and without rings in Fig. 4, indicating the presence of negative
 224 effects, resulting in the majority of the ring-based configurations at 50 W supplied power being
 225 lower in performance than for the reactor without rings.

226 Second, the presence of rings in the discharge space of the DBD reactor occupies the available
 227 discharge volume. Since the same gas flow rate was applied in the experiments in Fig. 4, the
 228 discharge volume reduction reduced the space time of plasma-based dry reforming. This is
 229 unfavorable to the conversion of CH₄ and CO₂. In previous studies on a micro-gap DBD reactor
 230 [30], the reduction in discharge volume resulted in a lower energy yield than reactors with larger
 231 discharge gaps, despite the stronger electric field in the micro-gap reactor. However, the metal rings
 232 in this study only occupy part of the discharge volume at the location where the rings were placed,
 233 which is much smaller than the drop in volume brought by reducing the entire discharge gap in the
 234 micro-gap reactors. Moreover, as observed in Fig. 4, increasing the number or the cross-sectional
 235 diameter of the rings causes some improvement in conversion and energy yield (even when
 236 compared to the reactor without rings in some cases), although more rings means a larger discharge
 237 volume reduction. This suggests that although metal rings do reduce the discharge volume to some
 238 extent (~0.05 mL for each 1.6 mm ring and ~0.2 mL for each 3.2 mm ring), this is not the main
 239 negative factor that leads to the lower performance of the reactor with rings than the reactor without
 240 rings.

241 Finally, compared with the smooth and uniform surface of the inner electrode rod, discharges
242 are more likely to take place on the metal rings with stronger reduced electric fields. This makes
243 the discharge more stable and more uniform in intensity (see section 3.1.2 below), which may be
244 beneficial to reduce the formation of by-products and improve the selectivity of the desired
245 products [26]. However, the spatial distribution of the discharges becomes relatively non-uniform,
246 as it is more difficult for the discharge to take place where no metal rings are present. The place
247 where the discharge can take place on the inner electrode has changed from the 10 cm long
248 electrode rod to probably only the top of a few rings, so the volume of the discharge space is
249 expected to be greatly reduced. This reduces the chance of gas molecules being hit by the discharges,
250 resulting in an expected reduced conversion and energy yield [37]. Increasing the number of rings
251 will reduce the distance between the rings, and is thus expected to improve the spatial uniformity
252 of the plasma, in line with the trend in Fig. 4 that the performance increases with the number of
253 rings. This indicates that the non-uniform spatial distribution of the discharge is a more dominant
254 negative effect of the stainless steel rings on the reactor performance compared to the volume
255 occupied by the metal rings. The negative effects play a more dominant role at 50 W, making the
256 reactor with rings to perform worse in most cases than the original reactor without rings.

257 3.1.2. Electrical characterization

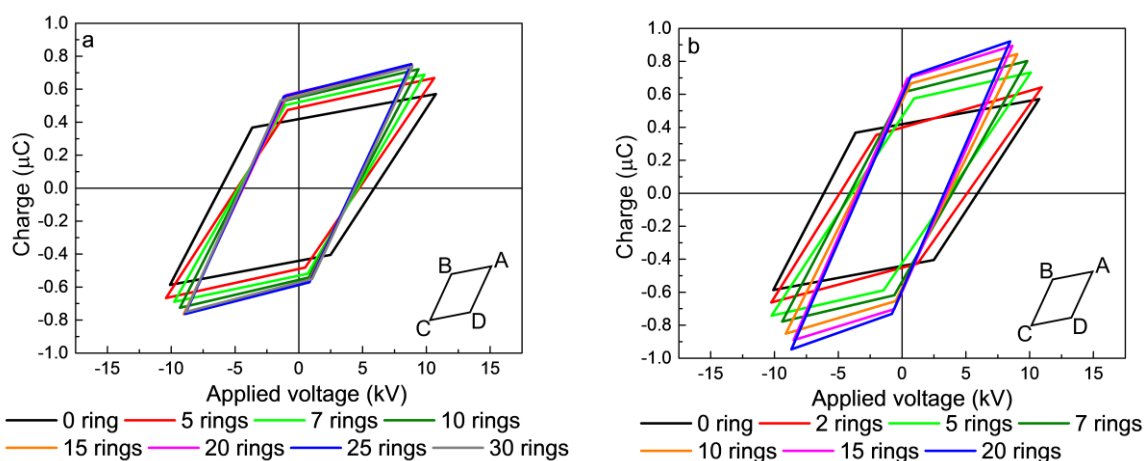
258 Electrical characterization data of the above experiments were collected and calculated to better
259 understand the effect of the stainless steel rings. The data that can be obtained directly from the raw
260 measurement data or through simple calculations like averaging, are listed in Table 1. In most cases,
261 the required applied voltage (presented in Table 1 as the root mean square voltage V_{RMS}) to achieve
262 a 50 W supplied power in the reactors with rings was smaller than in the reactor without rings.
263 Moreover, the required V_{RMS} decreased with the number of rings, while the root mean square
264 current (I_{RMS}) of the generated plasma increased. Stronger plasma currents (higher I_{RMS} values)
265 indicate stronger or more discharges induced by the rings in the reactor, possibly promoting the
266 production of active species, i.e., electrons, ions and radicals. In addition, the presence of the rings
267 enhanced the local electric field, making the discharge easier to take place in the reactor, so that a
268 certain power could be achieved at a lower applied voltage. Reactors with fewer than 15 rings with
269 1.6 mm diameter had a lower I_{RMS} than the ring-free reactor, due to the limited promotion induced
270 by the thinner rings on the discharge and because the discharge can only occur on fewer rings,
271 while reactors with more than 15 rings with 1.6 mm diameter and all reactors with 3.2 mm rings
272 had a higher I_{RMS} than the ring-free reactor, thus indicating stronger and more discharges, that can
273 promote the formation of active species, as mentioned above. The V_{RMS} trend of the reactors with
274 1.6 mm rings was not obvious when the number of rings was above 10. For the 3.2 mm rings, the
275 V_{RMS} was lower and the I_{RMS} was higher than for the same number of 1.6 mm rings, and the decrease
276 in V_{RMS} with the number of rings was more pronounced. The plasma power of the 1.6 mm rings
277 did not show a clear trend, while the 3.2 mm rings showed a trend of decreasing plasma power with
278 increasing number of rings, which is, in principle, negative to the performance of the reactor.

279
280
281

Table 1. Measured data from the input signals of the oscilloscope, including voltage, current, charge and power, for the DBD reactors without rings, and with stainless steel rings of varying number and two different cross-sectional diameters, at an almost constant supplied power of 50 W.

| Cross-sectional diameter of rings | Number of rings | V_{RMS} (kV) | I_{RMS} plasma (mA) | Plasma power (W) | Supplied power (W) | Displaced charge (nC/period) |
|-----------------------------------|-----------------|-----------------|-----------------------|------------------|--------------------|------------------------------|
| Without rings | 0 | 7.03 ± 0.04 | 19.5 ± 0.7 | 31.3 ± 0.8 | 51 ± 1 | 980 ± 35 |
| 1.6 mm | 5 | 6.86 ± 0.04 | 16.8 ± 0.4 | 32 ± 1 | 50.4 ± 0.8 | 1150 ± 29 |
| | 7 | 6.54 ± 0.04 | 18.3 ± 0.5 | 32.2 ± 0.5 | 50.2 ± 0.5 | 1210 ± 15 |
| | 10 | 6.26 ± 0.02 | 18.8 ± 0.2 | 32.7 ± 0.1 | 50.2 ± 0.1 | 1260 ± 19 |
| | 15 | 6.04 ± 0.04 | 19.7 ± 0.4 | 32.7 ± 0.6 | 50.3 ± 0.5 | 1320 ± 24 |
| | 20 | 6.09 ± 0.02 | 21.8 ± 0.9 | 33 ± 1 | 50.6 ± 0.4 | 1314 ± 8 |
| | 25 | 6.08 ± 0.04 | 22.1 ± 0.2 | 32.3 ± 0.6 | 50.7 ± 0.5 | 1320 ± 12 |
| 3.2 mm | 30 | 6.15 ± 0.07 | 22.4 ± 0.2 | 31.9 ± 0.3 | 50.3 ± 0.6 | 1290 ± 13 |
| | 2 | 7.14 ± 0.04 | 20.0 ± 0.3 | 30.8 ± 0.9 | 51.6 ± 0.9 | 1070 ± 16 |
| | 5 | 6.79 ± 0.04 | 21.4 ± 0.3 | 29.0 ± 0.2 | 51.1 ± 0.3 | 1320 ± 15 |
| | 7 | 6.44 ± 0.03 | 22.6 ± 0.3 | 28.5 ± 0.7 | 50.3 ± 0.7 | 1420 ± 6 |
| | 10 | 6.15 ± 0.04 | 24.5 ± 0.9 | 28.1 ± 0.5 | 50.5 ± 0.5 | 1498 ± 6 |
| | 15 | 5.90 ± 0.04 | 25.1 ± 0.4 | 27.9 ± 0.2 | 50.5 ± 0.6 | 1600 ± 11 |
| | 20 | 5.89 ± 0.01 | 24.9 ± 0.9 | 26.7 ± 0.6 | 50.6 ± 0.2 | 1652 ± 7 |

282 Fig. 5 shows the fitted Lissajous figures (consisting of the calculated slopes) of the plasma at 50
283 W supplied power, in reactors with varying number and two different cross-sectional diameter of
284 the stainless steel rings. The raw data of the Lissajous figures are shown in Fig. S1, in the supporting
285 information. Obvious differences can be observed from the Lissajous figures. First of all, the height
286 of the Lissajous figure of the reactors with rings is higher than that of the reactor without rings, and
287 it continues to increase as the number of rings increases. This indicates that the metal rings lead to
288 more charge displacement within the plasma, which supports the aforementioned discharge
289 promotion by the enhanced reduced electric field. The specific values of the displaced charges were
290 calculated by taking the charge difference between points A and D of the Lissajous figures, and are
291 listed in Table 1. For the 1.6 mm rings, only the reactors with less than 15 rings showed an obvious
292 trend of increasing the displaced charge with increasing number of metal rings, while the trend for
293 the 3.2 mm rings is still relatively obvious until the maximum of 20 rings. Furthermore, the
294 displaced charge in case of the 3.2 mm rings was always more than that of the 1.6 mm rings with
295 the same number of rings, which should be one of the reasons why the reactors with 3.2 mm rings
296 showed higher performance.



297

298 **Fig. 5.** The fitted Lissajous figures (calculated by the MATLAB script) of plasma-based dry reforming at 50 W supplied
299 power, in a DBD reactor without rings, and with varying number of stainless steel rings with cross-sectional diameters
300 of (a) 1.6 mm and (b) 3.2 mm.

301 Second, the voltage at $Q = 0$ C in the Lissajous figures, which is related to the burning voltage
302 (U_{bur}) of the plasma in the reactor, was changed by the rings. As shown in Table 2, after placing the
303 stainless steel rings on the inner electrode, U_{bur} was significantly reduced. Furthermore, as the
304 number of rings increases, U_{bur} shows a roughly decreasing trend. This suggests that the metal rings
305 made it easier to ignite and sustain the plasma at relatively lower voltages. Moreover, the 3.2 mm
306 rings lower the U_{bur} more than the 1.6 mm rings. The mean electric field can be calculated from
307 U_{bur} and the discharge gaps, and it is increased by the presence of the rings, due to the reduced
308 discharge gap. It should be noted that the electric field shown in Table 2 is the mean electric field
309 at the position of the rings (calculated from the gap between the rings and the dielectric barrier) in
310 the reactor with rings, as we believe that the discharge mainly take place on the rings. The electric
311 field decreases as the number of rings increases, due to the lower burning voltage of the plasma in
312 the reactor with more rings. The mean electron energy and electron energy distribution function
313 (EEDF) are also shown in Table 2 and Fig. S2, respectively. The metal rings enhance the mean
314 electron energy in the reactor, however, this electron energy is too high for efficient vibrational
315 excitation of CO_2 in plasma-based dry reforming [18,27]. As vibrational excitation is the most
316 energy-efficient pathway for dissociation, this might be one of the reasons why the performance of
317 the reactor with rings at 50 W is lower than the original reactor without rings. Similar with the trend
318 of the electric field, the mean electron energy also decreases with increasing number of rings, which
319 may be one of the reasons for the enhancement in reactor performance with increasing number of
320 rings. However, due to other physical changes that cause positive effects of the rings on the
321 discharge (as described above and below), the reactor with 3.2 mm rings performed better than the
322 reactor with 1.6 mm rings despite the higher mean electron energy. Even, some reactors (with more
323 than 15 rings) performed better than the reactor without rings at 50 W.

324 Finally, the slope of each side of the Lissajous figures also changes, indicating that the metal
325 rings change the capacitance of the DBD reactor. The slope (capacitance of the entire reactor) C_{cell}
326 of the undischarged phases and the slope ζ representative for the capacitance of the discharged
327 phases are listed in Table 2. The capacitance C_{cell} of all the reactors with rings is always higher than
328 that of the reactor without rings, because the discharge gaps are reduced by placing the metal rings.
329 There seems to be a slightly rising trend of C_{cell} with the number of rings, but some cases do not
330 follow it. On the other hand, ζ , has a clearly increasing trend with the number of rings. Since the
331 DBD reactor was not fully discharged, the capacitance represented by ζ includes the capacitance of
332 the undischarged gap and the dielectric barrier. The capacitance of the dielectric barrier is constant
333 because the same barrier was used in all experiments. Therefore, the rise of ζ indicates that the
334 undischarged fraction decreases upon increasing number of rings. The fractions of discharges
335 (denoted by f) in the reactors, calculated from C_{cell} and ζ , are also listed in Table 2. For the 1.6 mm
336 rings, the discharge fraction increases from 40.0% for 5 rings to 60.1% for 30 rings, while for the
337 3.2 mm rings, it increases from 38.7% for 2 rings to 62.9% for 20 rings, hence showing a larger
338 increase rate. Since the gas flow rate was approximately constant and the plasma power was even
339 reduced (at a constant supplied power) with the number of rings, the larger discharge fraction by
340 the stainless steel rings was achieved without increasing the specific energy input. It needs to be
341 noted that the plasma discharge fraction of the reactors with 5 rings or less, both for the 1.6 mm
342 and 3.2 mm rings, was smaller than that of the reactor without rings, but as the number of rings
343 increases, the discharge fraction becomes larger than in the reactor without rings. This is in line
344 with the effect mentioned above: the metal rings can promote the discharge but will induce the
345 discharge mainly taking place on the rings, resulting in a non-uniform spatial distribution of the
346 discharge. The spacing between the rings, where it is more difficult for the discharge to take place,
347 is large in reactors with fewer than five rings, and hence in this case the discharge fraction is even
348 smaller than that of a reactor without rings. As the number of rings increases, the negative effect
349 on the discharge fraction decreases and the positive effect increases, so the discharge fraction
350 increases substantially.

351
352
353

Table 2. Calculated data derived from the raw data of electrical characterization, obtained by a MATLAB script and BOLSIG+, for the DBD reactors without rings and with stainless steel rings of varying number and two different cross-sectional diameters, at an almost constant supplied power of 50 W.

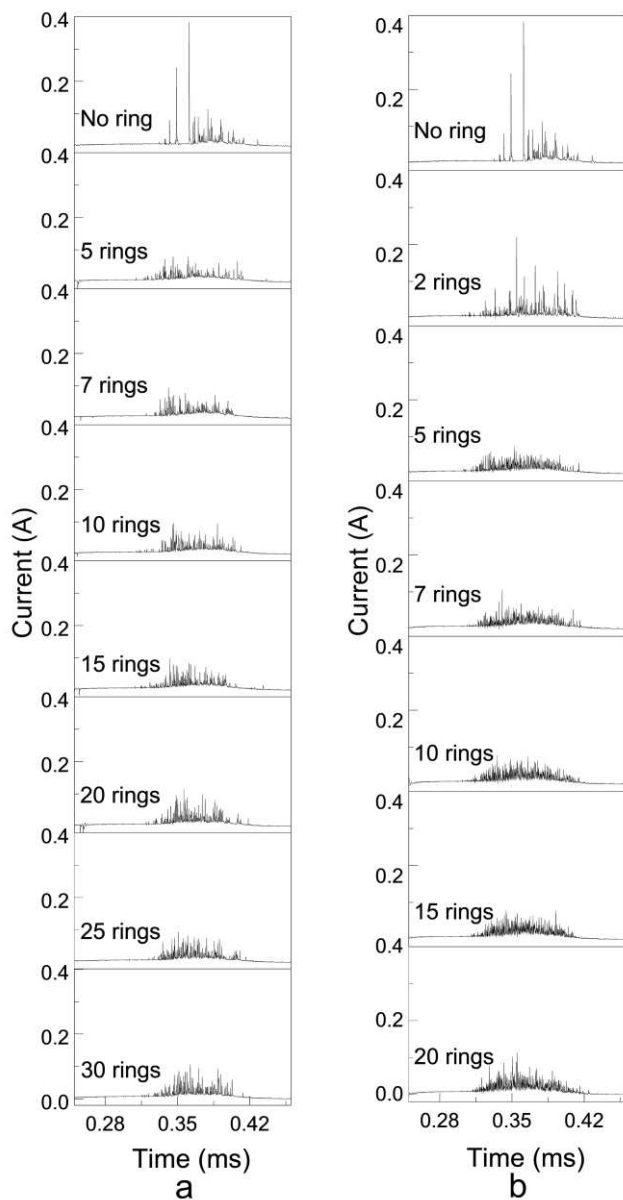
| Cross-sectional diameter of rings | Number of rings | U_{bur} (kV) | Electric field (kV/cm) | Mean electron energy (eV) | C_{cell} (pF) | ζ (pF) | f (%) | Number of micro-discharges (a.u./T) | Average filament charge (nC/disch.) |
|-----------------------------------|-----------------|-----------------|------------------------|---------------------------|-----------------|-----------------|----------------|-------------------------------------|-------------------------------------|
| Without rings | 0 | 6.3 ± 0.1 | 12.8 ± 0.2 | 3.31 ± 0.06 | 14.3 ± 0.5 | 133.1 ± 0.3 | 47.2 ± 0.2 | 90 ± 22 | 11 ± 3 |
| 1.6 mm | 5 | 5.2 ± 0.1 | 15.7 ± 0.3 | 3.75 ± 0.06 | 16.9 ± 0.4 | 116.5 ± 0.5 | 40.0 ± 0.3 | 75 ± 2 | 15 ± 1 |
| | 7 | 5.0 ± 0.1 | 15.1 ± 0.3 | 3.67 ± 0.06 | 16.6 ± 0.3 | 134.3 ± 0.7 | 47.2 ± 0.3 | 80 ± 5 | 15 ± 2 |
| | 10 | 4.7 ± 0.2 | 14.2 ± 0.6 | 3.53 ± 0.07 | 18.2 ± 0.3 | 149.7 ± 0.3 | 53.1 ± 0.1 | 81 ± 3 | 16 ± 1 |
| | 15 | 4.5 ± 0.2 | 13.6 ± 0.6 | 3.44 ± 0.07 | 19.5 ± 0.6 | 162.9 ± 0.2 | 58.2 ± 0.3 | 81 ± 7 | 16 ± 1 |
| | 20 | 4.5 ± 0.1 | 13.6 ± 0.3 | 3.44 ± 0.06 | 18.8 ± 0.3 | 165.7 ± 0.4 | 59.4 ± 0.2 | 96 ± 9 | 14 ± 1 |
| | 25 | 4.48 ± 0.03 | 13.5 ± 0.1 | 3.43 ± 0.06 | 19.7 ± 0.8 | 168.0 ± 0.7 | 60.2 ± 0.1 | 83 ± 7 | 16 ± 2 |
| 3.2 mm | 30 | 4.6 ± 0.1 | 13.9 ± 0.3 | 3.48 ± 0.06 | 20.1 ± 0.5 | 167.8 ± 0.6 | 60.1 ± 0.2 | 93 ± 9 | 14 ± 1 |
| | 2 | 5.7 ± 0.3 | 33.3 ± 1.8 | 5.4 ± 0.2 | 21.4 ± 0.7 | 116.0 ± 0.5 | 38.7 ± 0.2 | 140 ± 11 | 8 ± 2 |
| | 5 | 4.1 ± 0.1 | 23.9 ± 0.6 | 4.65 ± 0.07 | 17.3 ± 0.5 | 116.2 ± 0.9 | 39.8 ± 0.2 | 126 ± 5 | 10 ± 1 |
| | 7 | 4.2 ± 0.2 | 24.5 ± 1.2 | 4.7 ± 0.1 | 18.9 ± 0.4 | 141.6 ± 0.4 | 49.7 ± 0.3 | 163 ± 4 | 9 ± 2 |
| | 10 | 3.9 ± 0.1 | 22.8 ± 0.6 | 4.54 ± 0.07 | 22.2 ± 0.9 | 155.9 ± 0.7 | 54.8 ± 0.4 | 150 ± 6 | 10 ± 3 |
| | 15 | 3.7 ± 0.2 | 21.6 ± 1.2 | 4.4 ± 0.1 | 24 ± 1 | 175.2 ± 0.5 | 62.5 ± 0.4 | 165 ± 4 | 10 ± 1 |
| | 20 | 3.5 ± 0.1 | 20.4 ± 0.6 | 4.31 ± 0.07 | 27.0 ± 0.3 | 177.4 ± 0.7 | 62.9 ± 0.3 | 135 ± 4 | 12 ± 1 |

354
355
356
357
358
359
360
361
362
363
364
365
366

Fig. 6 shows the current profiles in the DBD reactors with varying number and two different cross-sectional diameters of the stainless steel rings. Comparing the reactors without and with fewer rings (≤ 5), the current intensity is more uniform in the reactor with rings, and the uniformity increases with the number of rings. The difference in current profiles between the reactors with relatively more rings is not significant, but it seems that the uniformity of current intensity slightly decreases with increasing number of rings, see for example the reactors with 5 and 20 rings with a diameter of 3.2 mm. This confirms that the rings can make the discharge more uniform in intensity, which may change the selectivity of some products. The discharge onset in the reactors with rings is earlier, which is consistent with one of the conclusions of Fig. 5 that the rings reduce the burning voltage of the plasma. The number of peaks in the current profiles is a measure for the number of micro-discharges in the plasma, this number is also listed in Table 2. The number of micro-discharges in the reactors with 1.6 mm rings is similar to that in the reactor without rings, while the number of micro-discharges in the reactor with the 3.2 mm rings is clearly higher (cf. Table 2).

367
368
369
370
371
372
373
374
375
376
377
378
379
380
381
382

Finally, the average filament charge of the micro-discharges can be obtained by dividing the displaced charge by the number of micro-discharges, and is a measure of the average discharge intensity. The average filament charge in case of the 1.6 mm rings is higher than that of the reactor without rings, while that of the 3.2 mm rings is similar to or even lower. Metal rings with both cross-sectional diameters improve the displaced charge within the plasma, as mentioned above, but the improvement for the 1.6 mm rings is on the average filament charge, or average discharge intensity, while the improvement for the 3.2 mm rings is on the number of micro-discharges. Hence, the change in discharge behavior caused by the stainless steel rings is different, possibly depending on the size of the remaining discharge gap between the metal and the dielectric barrier. Although the above electrical characterizations show a lot of positive effects of the rings, they are not sufficient to compensate for the negative effects of the non-uniform spatial distribution of the discharge and the high electron energy. The reactors with rings showed in most cases a worse performance than the reactor without rings. Only when having more rings (≥ 15) with a diameter of 3.2 mm, the reactor with rings performed better than the reactor without rings, due to its greater improvement on the discharge and less negative impact of the non-uniform discharge spatial distribution and the high electron energy.



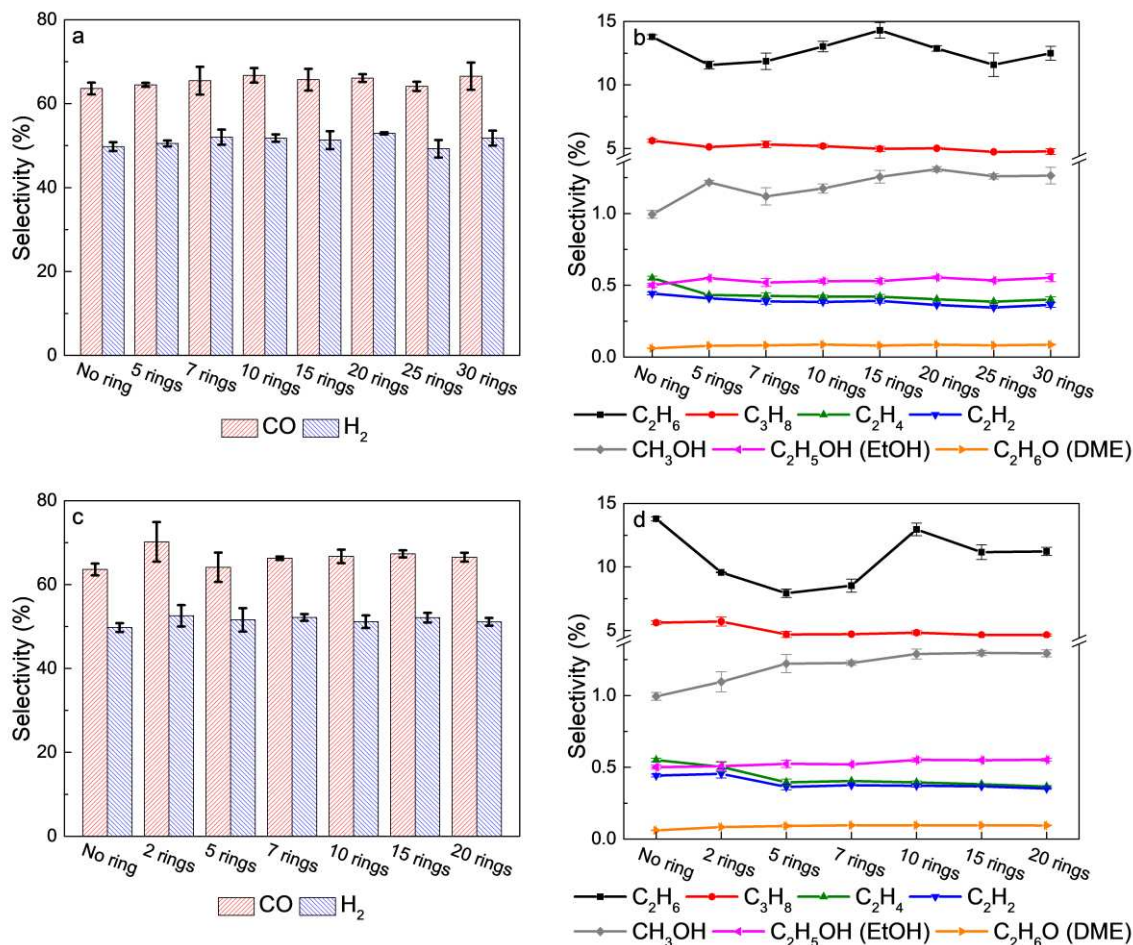
383

384 **Fig. 6.** Current profiles of a discharge phase in DBD reactors at 50 W supplied power, without rings, and with a varying
 385 number of stainless steel rings with cross-sectional diameters of (a) 1.6 mm and (b) 3.2 mm.

386 3.1.3. Selectivity

387 Due to the limited surface area of the rings and the fact that a similar material was used as for
 388 the stainless steel inner electrode, we believe that the conversion and selectivity changes are mainly
 389 caused by the physical effect of the rings on the discharge. Fig. 7 shows the selectivity of various
 390 products in the DBD reactor without rings and with varying numbers of stainless steel rings at 50
 391 W supplied power, for both 1.6 mm and 3.2 mm rings. Not only syngas is formed, but also other
 392 products, such as various hydrocarbons and oxygenations are formed in the DBD reactor, among
 393 others by radical recombination [24,25]. It needs to be noted that some possible liquid products
 394 (e.g., oily hydrocarbons, alcohols) and carbon deposits attached to the reactor, as well as other
 395 gaseous products not calibrated in the gas chromatograph, cannot be counted, and therefore, the
 396 carbon and hydrogen mass balance calculated from the product selectivity does not reach 100% (as

397 shown in Fig. S3 of the Supporting Information). As is clear from Fig. 7, at 50 W supplied power,
 398 the placement of the metal rings and the number of rings only had a small effect on the selectivity
 399 of most products. The exception is that there is a slightly increasing trend for the selectivity of
 400 methanol with the number of rings, and the selectivity of ethane is more affected, although without
 401 a clear trend.



402

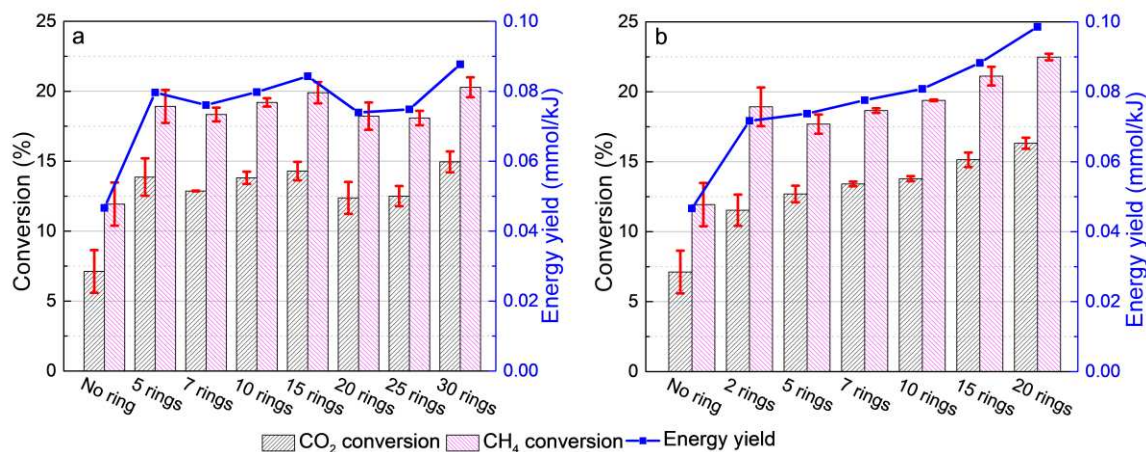
403 **Fig. 7.** Selectivity of various products of plasma-based dry reforming at 50 W supplied power, in a DBD reactor without
 404 rings, and with varying number of stainless steel rings. (a) Syngas and (b) ethane, propane, ethylene, acetylene, methanol,
 405 ethanol (EtOH) and dimethyl ether (DME) selectivity in reactors with rings with a diameter of 1.6 mm. (c) Syngas and
 406 (d) ethane, propane, ethylene, acetylene, methanol, ethanol and dimethyl ether selectivity in reactors with rings with a
 407 diameter of 3.2 mm. The error bars were obtained from standard errors based on three repeat experiments with the rings
 408 kept in place.

409 3.2. Plasma-based dry reforming at 30 W supplied power

410 3.2.1. Conversion and energy yield

411 Fig. 8 shows the influence of the same 1.6 mm and 3.2 mm rings on the CH₄ and CO₂
 412 conversions and the energy yield of plasma-based dry reforming, but at 30 W supplied power. The
 413 effect of the rings is clearly different from the case at 50 W. At 30 W, the energy yields and reactant
 414 conversions in all reactors with the rings are higher than in the reactor without rings. There is also
 415 a clear effect of the number of rings to the conversion and energy yield, and the performance of the
 416 reactor with 3.2 mm rings has a tendency to increase with the number of rings. Compared with the
 417 reactor without rings, the increase of reactor performance resulting from the rings at 30 W supplied

418 power is much greater than at 50 W. The DBD reactor with 20 rings with a 3.2 mm cross-sectional
 419 diameter has the best performance, with an energy yield double the value of the reactor without
 420 rings. It enhances the CO₂ conversion from 7.1% to 16.3 %, the CH₄ conversion from 11.9% to
 421 22.5%, and the energy yield from 0.05 mmol/kJ to 0.1 mmol/kJ. As already mentioned above, the
 422 energy yield in this work was calculated from the supplied power, and the energy yield of the
 423 reactor with the best performance is 0.19 mmol/kJ if calculated from the plasma power as in most
 424 of the literature. Furthermore, compared to the reactors with the same number and diameter of rings
 425 at a supplied power of 50 W, although the CO₂ and CH₄ conversions are reduced due to the lower
 426 power, the energy yield is actually improved by 17% on average at 30 W, except for the reactor
 427 without rings. The best performing reactor in this study, i.e., the one with 20 rings of 3.2 mm
 428 diameter, was investigated for its performance stability. As shown in Fig. S4, the conversion of
 429 CO₂ and CH₄ in this reactor did not change significantly during the 12-hour test.



430

431 **Fig. 8.** CH₄ and CO₂ conversion and energy yield of plasma-based dry reforming at 30 W supplied power, in a DBD
 432 reactor without rings, and with varying number of stainless steel rings with cross-sectional diameters of (a) 1.6 mm and
 433 (b) 3.2 mm. The error bars were obtained from standard errors based on three repeat experiments with the rings kept in
 434 place.

435 The effects of adding the stainless steel rings on the discharge has been described in section 3.1
 436 above, and also apply here, i.e., a higher displaced charge, a larger discharge fraction, a higher
 437 discharge intensity, etc. In addition, the discharge in the reactor without rings is unstable at 30 W
 438 supplied power (see section 3.2.2 below), which is one of the reasons that the improvement of the
 439 reactor performance by the metal rings is so large at 30 W. On the other hand, the positive effects
 440 of the metal rings on the discharge is indeed greater at 30 W supplied power, as can be deduced
 441 below from the larger changes in parameters such as the discharge fraction and burning voltage.

442 3.2.2. Electrical characterization

443 Table 3 shows the collected and calculated electrical characterization data of the experiments in
 444 Fig. 8, to better understand the effect of the stainless steel rings at 30 W supplied power. Similar to
 445 the case of 50 W, in most cases, the required applied V_{RMS} to achieve a 30 W supplied power in the
 446 reactors with rings is smaller than in the reactor without rings. The required V_{RMS} decreases with
 447 the number of rings, while the I_{RMS} of the generated plasma increase. The rings in the reactor
 448 promoted the generation of discharges and led to more or stronger discharges. The trend of V_{RMS}
 449 and I_{RMS} of the reactors with 1.6 mm rings is not obvious when the number of rings is more than
 450 10. For the 3.2 mm rings, the V_{RMS} is lower and the I_{RMS} was higher than for the same number of
 451 1.6 mm rings, and the trend of V_{RMS} with the number of rings was again more pronounced. Thicker
 452 rings also have a more pronounced improvement on the discharge. However, different from the

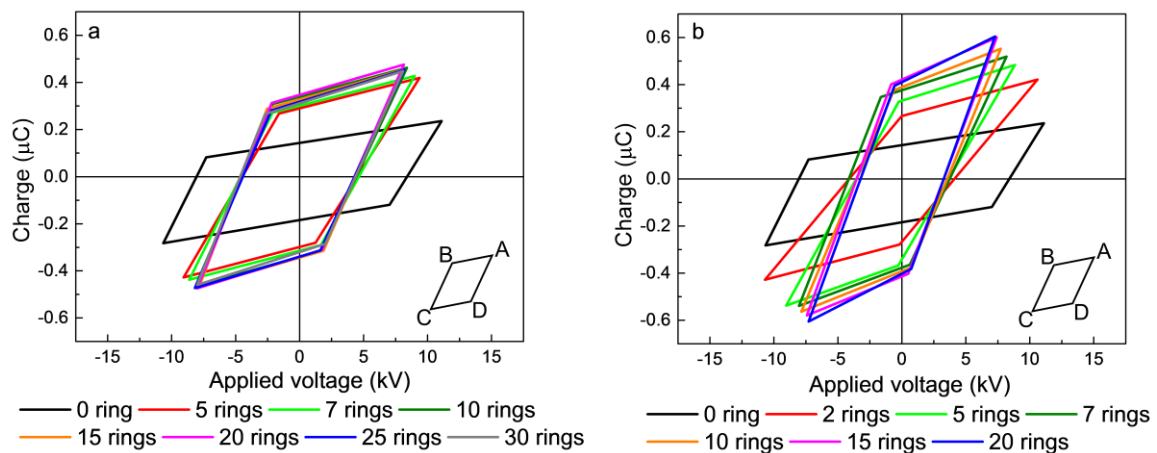
453 experiments at 50 W supplied power, the generated plasma power in all the reactors with rings is
 454 higher than in the reactor without rings. This is because the plasma in the reactor without rings is
 455 unstable, extinguishing or not discharging for some periods. As shown in Fig. S5 in the SI, in 6
 456 periods of 2 ms, there was no discharge in 4 half periods. This is also one of the reasons for the
 457 poor performance of the reactor without rings at 30 W supplied power. Since the plasma power is
 458 calculated as the average power over all periods, including the phases without discharge, the plasma
 459 power in the reactor without rings is lower than in the other reactors.

460 **Table 3.** Measured data from the input signals of the oscilloscope, including voltage, current, charge and power, for the
 461 DBD reactors without rings and with stainless steel rings of varying number and two different cross-sectional diameters,
 462 at an almost constant supplied power of 30 W.

| Cross-sectional diameter of rings | Number of rings | V_{RMS} (kV) | I_{RMS} plasma (mA) | Plasma power (W) | Supplied power (W) | Displaced charge (nC/period) |
|-----------------------------------|-----------------|-----------------|-----------------------|------------------|--------------------|------------------------------|
| Without rings | 0 | 7.42 ± 0.07 | 14.4 ± 0.6 | 10.2 ± 0.6 | 30.4 ± 0.9 | 360 ± 21 |
| 1.6 mm | 5 | 6.33 ± 0.04 | 13.9 ± 0.4 | 16.4 ± 0.6 | 30.6 ± 0.5 | 700 ± 7 |
| | 7 | 5.99 ± 0.04 | 14.6 ± 0.3 | 16.0 ± 0.4 | 30.5 ± 0.5 | 718 ± 6 |
| | 10 | 5.69 ± 0.04 | 17.2 ± 0.7 | 18.0 ± 0.3 | 30.8 ± 0.6 | 780 ± 13 |
| | 15 | 5.51 ± 0.07 | 16.6 ± 0.5 | 17.3 ± 0.6 | 30.2 ± 0.3 | 770 ± 11 |
| | 20 | 5.63 ± 0.04 | 17.7 ± 0.3 | 17.4 ± 0.7 | 30.8 ± 0.8 | 790 ± 11 |
| | 25 | 5.66 ± 0.02 | 17.5 ± 0.3 | 16.6 ± 0.5 | 30.4 ± 0.4 | 766 ± 5 |
| | 30 | 5.62 ± 0.04 | 17.8 ± 0.2 | 16.1 ± 0.4 | 29.9 ± 0.5 | 734 ± 7 |
| 3.2 mm | 2 | 6.89 ± 0.11 | 15.6 ± 0.3 | 14.2 ± 0.9 | 31.6 ± 0.7 | 700 ± 16 |
| | 5 | 5.87 ± 0.01 | 16.4 ± 0.7 | 15.4 ± 0.3 | 30.7 ± 0.8 | 850 ± 13 |
| | 7 | 5.55 ± 0.04 | 17.2 ± 0.5 | 15.4 ± 0.7 | 30.8 ± 0.5 | 884 ± 9 |
| | 10 | 5.30 ± 0.02 | 18.2 ± 0.8 | 15.7 ± 0.3 | 30.5 ± 0.4 | 930 ± 4 |
| | 15 | 5.06 ± 0.04 | 19.1 ± 0.4 | 15.9 ± 0.3 | 30.6 ± 0.4 | 1000 ± 12 |
| | 20 | 5.04 ± 0.02 | 19.4 ± 0.3 | 15.3 ± 0.5 | 30.2 ± 0.5 | 990 ± 11 |

463 Fig. 9 shows the fitted Lissajous figures of the plasma at 30 W supplied power, in the reactors
 464 without rings and with varying number and two different cross-sectional diameter of the stainless
 465 steel rings. The raw data of the Lissajous figures are shown in Fig. S6. It should be noted that the
 466 Lissajous figure of the reactor without rings is only in the discharge phases, because the Lissajous
 467 figure could not be formed in the undischarged period. The Lissajous figures at 30 W exhibit more
 468 obvious differences between the reactors with and without rings than at 50 W supplied power. First
 469 of all, the height of the Lissajous figures, reflecting the displaced charge (see Table 3), follows the
 470 same trend, which is higher for the reactors with rings than for the reactor without rings, and
 471 increases with the number of rings. After placing a certain number of rings (e.g. 15 of 1.6 mm
 472 rings), a larger number of rings does not enhance the displaced charge anymore. The enhancement
 473 of displaced charge by the rings is more than double or even nearly triple that of the reactor without
 474 rings, indicating that the promotion of stainless steel rings on the plasma is greater at 30 W.
 475 Furthermore, the displaced charge of the 3.2 mm rings was also more than that of the 1.6 mm rings
 476 with the same number of rings at 30 W, so the reactors with 3.2 mm rings also yield a higher
 477 performance. Second, the U_{bur} reflecting the voltage for igniting and sustaining the plasma,
 478 calculated from the Lissajous figures, was reduced by the metal rings, in a similar trend as at 50 W.
 479 At 30 W supplied power, the mean electric field and the mean electron energy also follow similar
 480 trends to those at 50 W, and even have similar values. They increase with the increase of the cross-
 481 sectional diameter of the rings, and decrease upon larger number of rings. The electron energy
 482 distribution function is shown in Fig. S7. Despite the similar values for electric field and mean
 483 electron energy for the reactor configurations with rings at 50 W and 30 W, the reactor without
 484 rings has higher values for the burning voltage, average electric field, and electron energy at 30 W
 485 than 50 W, probably due to the unstable discharge at low power, which could be one of the reasons
 486 for its poor performance at 30 W. At 30 W, the burning voltage in the reactors with rings is smaller
 487 than in the reactor without rings, and the mean electric field and mean electron energy as well, for
 488 the 1.6 mm rings, while it is higher for the 3.2 mm rings. In spite of this, the performance of the

489 reactors with rings is always better at 30 W than for the reactor without rings. Finally, the
 490 capacitance of different phases of the DBD reactor, C_{cell} and ζ , calculated from the slope of the
 491 Lissajous figures, are listed in Table 4, as well as the discharge fractions, which are also calculated
 492 from them. The tendency of capacitance and discharge fraction due to the presence of the rings is
 493 the same as at 50 W. Thicker and more rings increase the capacitance and the discharge fraction.
 494 The 1.6 mm rings increase the discharge fraction from 30.0% for 5 rings to 44.7% for 30 rings,
 495 while the 3.2 mm rings increase the discharge fraction from 20.1% for 2 rings to 51.4% for 20 rings,
 496 with a larger increase rate. The discharge fraction of the reactors with relatively few (≤ 5) rings
 497 were smaller than that of the reactor without rings, probably because the rings induce the discharge
 498 to mainly take place on them, as mentioned above in section 3.1.2. The discharge fraction of the
 499 same reactor at 30 W was less than at 50 W. However, compared with the ring-free reactor, some
 500 plasma parameters, including the discharge fraction, burning voltage and displaced charge, were
 501 improved more by the presence of rings at 30W than at 50W, which is another reason why the
 502 reactor with stainless steel rings showed better performance at 30W.



503

504 **Fig. 9.** The fitted Lissajous figures (calculated by the MATLAB script) of plasma-based dry reforming at 30 W supplied
 505 power, in a DBD reactor without rings and with varying number of stainless steel rings with cross-sectional diameters of
 506 (a) 1.6 mm and (b) 3.2 mm.

507 **Table 4.** Calculated data derived from the raw data of electrical characterization, obtained by a MATLAB script and
 508 BOLSIG+, for the DBD reactors without rings and with stainless steel rings of varying number and two different cross-
 509 sectional diameters, at an almost constant supplied power of 30 W.

| Cross-sectional diameter of rings | Number of rings | U_{bur} (kV) | Electric field (kV/cm) | Mean electron energy (eV) | C_{cell} (pF) | ζ (pF) | f (%) | Number of micro-discharges (a.u./T) | Average filament charge (nC/disch.) |
|-----------------------------------|-----------------|-----------------|------------------------|---------------------------|-----------------|-----------------|----------------|-------------------------------------|-------------------------------------|
| Without rings | 0 | 8.9 ± 0.3 | 18.1 ± 0.6 | 4.05 ± 0.07 | 8.8 ± 0.7 | 98.0 ± 0.5 | 34.7 ± 0.3 | 68 ± 9 | 5 ± 2 |
| | 5 | 5.1 ± 0.2 | 15.4 ± 0.6 | 3.71 ± 0.07 | 14.0 ± 0.5 | 89.7 ± 0.7 | 30.0 ± 0.3 | 97 ± 2 | 7 ± 1 |
| | 7 | 5.08 ± 0.07 | 15.3 ± 0.2 | 3.70 ± 0.06 | 14.2 ± 0.8 | 103.5 ± 0.4 | 35.5 ± 0.4 | 139 ± 4 | 5 ± 1 |
| | 10 | 4.8 ± 0.1 | 14.5 ± 0.3 | 3.58 ± 0.06 | 15.0 ± 0.8 | 122.6 ± 0.7 | 42.8 ± 0.2 | 110 ± 12 | 7 ± 3 |
| | 15 | 4.7 ± 0.1 | 14.2 ± 0.3 | 3.53 ± 0.06 | 16 ± 0.1 | 133.4 ± 0.9 | 47.0 ± 0.1 | 92 ± 6 | 8 ± 2 |
| | 20 | 4.6 ± 0.2 | 13.9 ± 0.6 | 3.48 ± 0.07 | 16.3 ± 0.4 | 129 ± 1 | 45.2 ± 0.4 | 99 ± 3 | 8 ± 2 |
| 1.6 mm | 25 | 4.67 ± 0.05 | 14.1 ± 0.2 | 3.52 ± 0.06 | 16.4 ± 0.9 | 121.5 ± 0.3 | 42.1 ± 0.4 | 101 ± 7 | 8 ± 3 |
| | 30 | 4.7 ± 0.2 | 14.2 ± 0.6 | 3.53 ± 0.07 | 17.2 ± 0.4 | 128.5 ± 0.9 | 44.7 ± 0.4 | 117 ± 2 | 6 ± 2 |
| | 2 | 5.0 ± 0.2 | 29.2 ± 1.2 | 5.09 ± 0.12 | 14.4 ± 0.8 | 65.0 ± 0.4 | 20.1 ± 0.3 | 71 ± 3 | 10 ± 2 |
| | 5 | 4.2 ± 0.1 | 24.5 ± 0.6 | 4.70 ± 0.07 | 18.3 ± 0.6 | 95.9 ± 0.2 | 31.3 ± 0.3 | 107 ± 6 | 8 ± 3 |
| | 7 | 4.08 ± 0.04 | 23.8 ± 0.2 | 4.64 ± 0.06 | 18.8 ± 0.5 | 127.8 ± 0.7 | 44.1 ± 0.2 | 161 ± 4 | 6 ± 2 |
| | 10 | 3.9 ± 0.1 | 22.8 ± 0.6 | 4.54 ± 0.07 | 21.5 ± 0.7 | 131.2 ± 0.6 | 44.9 ± 0.1 | 135 ± 5 | 7 ± 1 |
| | 15 | 3.6 ± 0.1 | 21.0 ± 0.6 | 4.37 ± 0.07 | 23.3 ± 0.4 | 147.1 ± 0.4 | 51.0 ± 0.1 | 101 ± 5 | 10 ± 1 |
| | 20 | 3.6 ± 0.1 | 21.0 ± 0.6 | 4.37 ± 0.07 | 27.2 ± 0.4 | 149.9 ± 0.8 | 51.4 ± 0.3 | 124 ± 3 | 8 ± 2 |

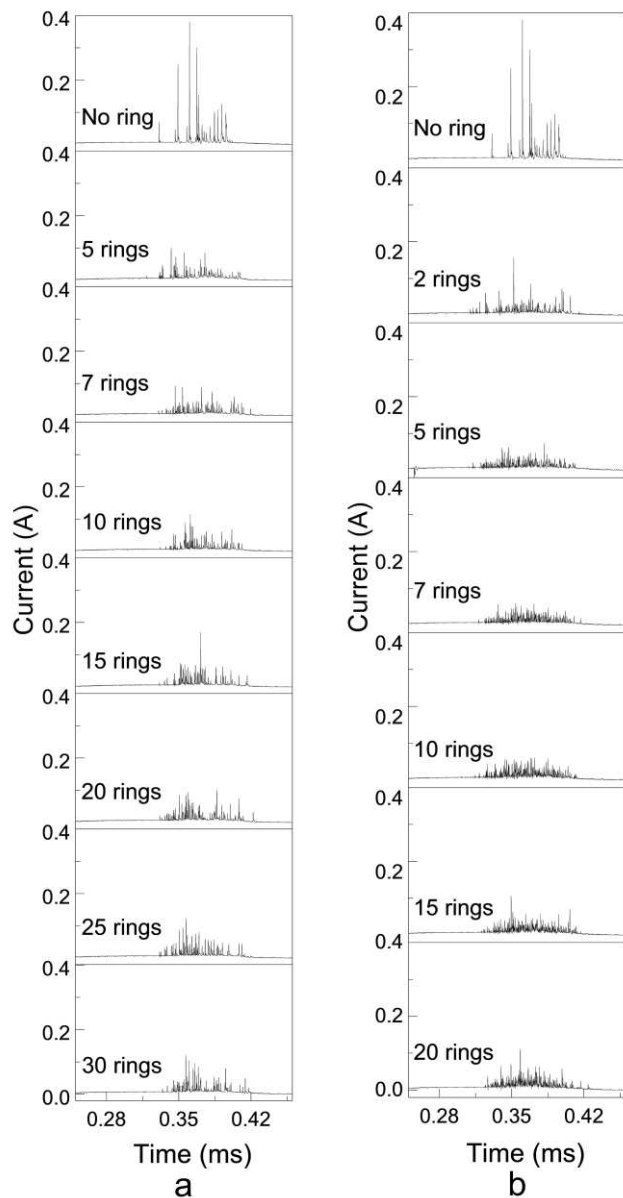
510

511

Fig. 10 shows the current profiles in the DBD reactors at 30 W supplied power. It should be noted that the current profile of the reactor without rings is selected from the stable discharge phases,

512 and actually it is not discharged in some phases, as shown in Fig. S5. At 30 W supplied power, the
513 stainless steel rings are again able to make the discharge more uniform in intensity, even with only
514 2 or 5 rings. As the number of rings increases, the uniformity of the current intensity of the plasma
515 decreases slightly, see for example the reactors with 5 and 20 rings with a diameter of 3.2 mm. The
516 onset of the discharge was earlier in the reactors with rings due to the reduction in U_{bur} . The number
517 of micro-discharges and average filament charge were calculated and listed in Table 4. Different
518 from the experiments at 50 W, where the 1.6 mm rings or 3.2 mm rings can only increase either the
519 average filament charge or the number of micro-discharges, at 30 W supplied power, both these
520 parameters are higher in the reactors with rings than in the reactor without rings. The reason for the
521 larger number of micro-discharges by the rings with both diameters is that the discharge in the
522 reactor without rings at 30 W is unstable and extinguishes in some periods. Moreover, with a larger
523 number of micro-discharges, the average displaced charge per discharge of the reactor with rings
524 is still higher than that of the reactor without rings. This indicates again that at 30 W supplied power,
525 the improvement of the discharge by the metal rings is greater than that at 50 W.

526



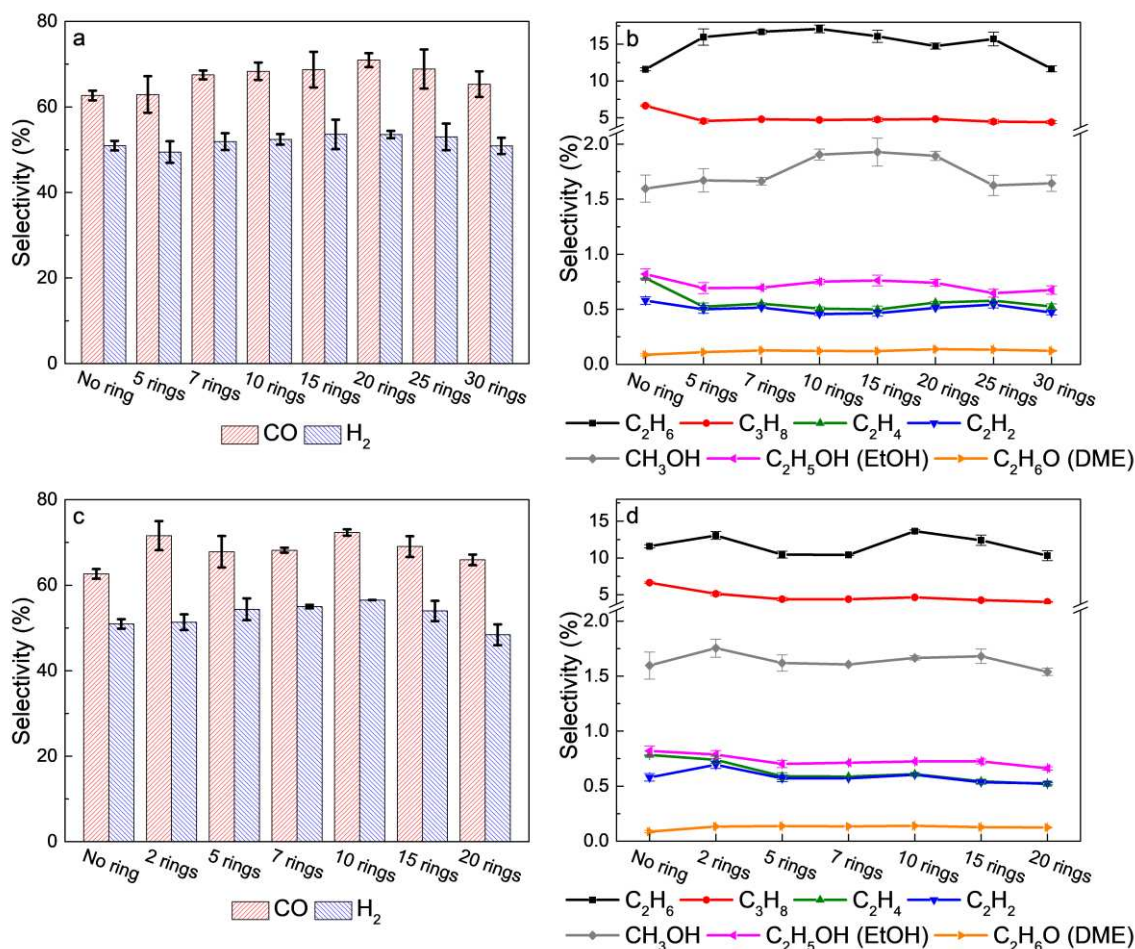
527

528 **Fig. 10.** Current profiles of a discharge phase in DBD reactors at 30 W supplied power, for the reactor without rings and
 529 with varying number of stainless steel rings with cross-sectional diameters of (a) 1.6 mm and (b) 3.2 mm.

530 3.2.3. Selectivity

531 Fig. 11 shows the selectivity of syngas and various by-products of the plasma-based dry
 532 reforming in the DBD reactors with stainless steel rings at 30 W supplied power. In most cases, the
 533 syngas selectivity in the reactors with rings is slightly higher than in the reactor without rings. In
 534 fact, the higher selectivity resulting from the rings seems to be present at 50 W as well, but is less
 535 obvious. Fig. S8 shows the carbon and hydrogen atomic balance, plotted from Fig. 11, and the
 536 atomic balance of the reactor with rings is higher than for the reactor without rings. This may be
 537 due to the fact that the products of plasma-based dry reforming were formed by the reaction of
 538 various radicals generated by the discharge, and the more uniform intensity of the discharge could
 539 reduce the types of formed by-products [26].

540 In addition, lower power seems to be favorable for improving the selectivity of some products.
 541 The selectivity of CO, H₂, ethylene, acetylene, methanol, and ethanol at 30 W was higher than at
 542 50 W, while the selectivity of other products was not much different. Close observation and
 543 comparison of the selectivity at 30 W and 50 W shows similarities in the trend of selectivity of
 544 some products at different powers. This also proves that although there are mainly physical effects,
 545 the stainless steel rings can have a certain effect on the product selectivity as well.



546

547 **Fig. 11.** Selectivity of various products of plasma-based dry reforming at 30 W supplied power, in a DBD reactor without
 548 rings and with varying number of stainless steel rings. (a) Syngas and (b) ethane, propane, ethylene, acetylene, methanol,
 549 ethane, ethanol (EtOH) and dimethyl ether (DME) selectivity in reactors with rings with a diameter of 1.6 mm. (c) Syngas and
 550 (d) ethane, propane, ethylene, acetylene, methanol, ethanol and dimethyl ether selectivity in reactors with rings with a
 551 diameter of 3.2 mm. The error bars were obtained from standard errors based on three repeat experiments with the rings
 552 kept in place.

553 3.3. Advantages of the reactor with rings

554 Due to different operating conditions (i.e., various gas composition and discharge parameters),
 555 it is difficult to accurately compare the reactor performance with literature. However, when we
 556 roughly compare the energy yields in literature (~0.1 mmol/kJ in empty reactors and ~0.2 mmol/kJ
 557 in reactors with catalyst) [4,7,23], the energy yield of 0.19 mmol/kJ achieved in our reactor
 558 configuration with rings but in absence of packing materials or catalyst, as obtained in this work,
 559 can be considered to be competitive. In addition, a lower power is generally favorable for higher

560 energy yield and higher liquefiable products selectivity, as demonstrated in this work and in the
561 literature [7,36]. However, it is difficult to stabilize the discharge at lower power in the original
562 reactor without rings. Therefore, another advantage of a reactor configuration with rings is to help
563 stabilize the discharge at lower power.

564 In addition to the above dry reforming processes, and in order to verify whether the DBD reactor
565 with rings can also be applied to other plasma-based reactions, we performed experiments for CO₂
566 decomposition at 30 W in the reactor with and without 3.2 mm rings. As shown in Fig. S9, the
567 conversion of CO₂ in the reactor with rings is again higher than in the reactor without rings,
568 indicating that this DBD reactor design has the potential to be extended to other plasma-based
569 processes. More in-depth research and analysis on CO₂ decomposition and other processes could
570 be the subject of future work, to get more insights in the underlying mechanisms.

571 **4 Conclusion**

572 In this study, we put stainless steel rings over the inner electrode rod of a cylindrical DBD
573 reactor, to change the local discharge gap and the electric field, and to study the dry reforming
574 performance. A varying number of rings with cross-sectional diameters of 1.6 mm and 3.2 mm
575 (leading to 3.1 mm and 1.5 mm discharge gaps) were used, and experiments were carried out at 50
576 W and 30 W supplied power. We found that at 50 W, the stainless steel rings mainly have negative
577 effects on the performance, while at 30 W, the rings greatly improve the performance, both in terms
578 of reactant conversion and energy yield. The reactor with 20 rings with 3.2 mm diameter showed
579 the best performance. Compared to the reactor without rings, it increases the CO₂ conversion from
580 7.1% to 16.3 %, the CH₄ conversion from 11.9% to 22.5%, and the energy yield from 0.05 mmol/kJ
581 to 0.10 mmol/kJ (0.19 mmol/kJ if it was calculated from the plasma power). All reactors with rings
582 have higher energy yield at 30 W than at 50 W.

583 Since the rings are made of similar material as the inner electrode and with small surface area,
584 we believe that the rings mainly cause physical effects on the discharge, and catalytic effects are
585 not considered. The difference in performance caused by the rings can be understood from studying
586 the electrical characteristics. The presence of the stainless steel rings changes the electric field
587 distribution in the reactor. The discharge gap is smaller where the rings are placed, increasing the
588 local electric field. Through electrical characterization, it was found that the displaced charge and
589 the discharge fraction, which can improve the conversion and energy yield of plasma-based dry
590 reforming, increase with the number and the cross-sectional diameter of rings at the same energy
591 input. This is the reason that the 3.2 mm rings always exhibit higher performance than the 1.6 mm
592 rings. The non-uniform discharge gap caused by the rings makes the discharge more stable and
593 more uniform in intensity, although its spatial distribution may be non-uniform. This spatial non-
594 uniformity was reduced by placing more rings, while the positive effects such as displaced charge
595 were enhanced, leading to better performance in reactors with more rings. At 30 W supplied power,
596 the metal rings can stabilize the discharge and have a greater improvement in the discharge than at
597 50 W, and thus yield better performance. Moreover, the effect of the stainless steel rings on the
598 discharge can also alter the selectivity of some products. The largest impact on selectivity is
599 however caused by lowering the supplied plasma power to 30W, increasing the selectivity of e.g.
600 methanol and ethane.

601 Besides dry reforming, the performance differences by placing metal rings in the DBD reactor
602 to alter the discharge gap distribution may be applicable to other plasma-based processes as well.
603 In addition, in plasma catalysis, the metal active components on the catalyst surface can also change
604 the discharge gap and electric field distribution in the reactor, and the effects mentioned above
605 might also play some role there. Catalyst studies to improve the conversion and selectivity of

606 plasma reactions need to consider not only its catalytic activity, but also these effects on the
607 discharge behavior.

608 **Declaration of competing interest**

609 The authors declare no known competing financial interest or personal relationships that could
610 inappropriately influence this work.

611

612 **Acknowledgements**

613 J.W acknowledges the financial support from the China Scholarship Council (No.
614 201806060123). K.Z acknowledges the EASiCHEM project funded by the Flemish Strategic Basic
615 Research Program of the Catalisti cluster and Flanders Innovation & Entrepreneurship
616 (HBC.2018.0484).

617 **Appendix A. Supporting information**

618 Supplementary data associated with this article can be found in the online version.

619 **References**

- 620 [1] A. Bogaerts, X. Tu, J.C. Whitehead, G. Centi, L. Lefferts, O. Guaitella, F. Azzolina-Jury, H.-
621 H. Kim, A.B. Murphy, W.F. Schneider, T. Nozaki, J.C. Hicks, A. Rousseau, F. Thevenet, A.
622 Khacef, M. Carreon, The 2020 plasma catalysis roadmap, *J. Phys. D Appl. Phys.* 53 (2020)
623 443001. <https://doi.org/10.1088/1361-6463/ab9048>.
- 624 [2] Y. Yi, J. Zhou, H. Guo, J. Zhao, J. Su, L. Wang, X. Wang, W. Gong, Safe Direct Synthesis of
625 High Purity H₂O₂ through a H₂/O₂ Plasma Reaction, *Angew. Chem. Int. Ed.* 52 (2013) 8446-
626 8449. <https://doi.org/https://doi.org/10.1002/anie.201304134>.
- 627 [3] Y. Yi, S. Li, Z. Cui, Y. Hao, Y. Zhang, L. Wang, P. Liu, X. Tu, X. Xu, H. Guo, A. Bogaerts,
628 Selective oxidation of CH₄ to CH₃OH through plasma catalysis: Insights from catalyst
629 characterization and chemical kinetics modelling, *Appl. Catal. B Environ.* 296 (2021) 120384.
630 <https://doi.org/10.1016/j.apcatb.2021.120384>.
- 631 [4] A.H. Khoja, M. Tahir, N.A.S. Amin, Dry reforming of methane using different dielectric
632 materials and DBD plasma reactor configurations, *Energy Convers. Manage.* 144 (2017) 262-
633 274. <https://doi.org/https://doi.org/10.1016/j.enconman.2017.04.057>.
- 634 [5] Y. Uytendhouwen, J. Hereijgers, T. Breugelmanns, P. Cool, A. Bogaerts, How gas flow design
635 can influence the performance of a DBD plasma reactor for dry reforming of methane, *Chem.*
636 *Eng. J.* 405 (2021) 126618. <https://doi.org/https://doi.org/10.1016/j.cej.2020.126618>.
- 637 [6] S. Zhang, Y. Gao, H. Sun, Z. Fan, T. Shao, Dry reforming of methane by microsecond pulsed
638 dielectric barrier discharge plasma: Optimizing the reactor structures, *High Voltage* 7 (2022) 718-
639 729. <https://doi.org/https://doi.org/10.1049/hve2.12201>.
- 640 [7] C.-j. Liu, G.-h. Xu, T. Wang, Non-thermal plasma approaches in CO₂ utilization, *Fuel*
641 *Process. Technol.* 58 (1999) 119-134. [https://doi.org/https://doi.org/10.1016/S0378-](https://doi.org/https://doi.org/10.1016/S0378-3820(98)00091-5)
642 [3820\(98\)00091-5](https://doi.org/https://doi.org/10.1016/S0378-3820(98)00091-5).
- 643 [8] X. Tu, J.C. Whitehead, Plasma-catalytic dry reforming of methane in an atmospheric
644 dielectric barrier discharge: Understanding the synergistic effect at low temperature, *Appl. Catal.*
645 *B Environ.* 125 (2012) 439-448. <https://doi.org/https://doi.org/10.1016/j.apcatb.2012.06.006>.
- 646 [9] R. Snoeckx, A. Bogaerts, Plasma technology – a novel solution for CO₂ conversion?, *Chem.*
647 *Soc. Rev.* 46 (2017) 5805-5863. <https://doi.org/10.1039/C6CS00066E>.

648 [10] B.S. Patil, N. Cherkasov, J. Lang, A.O. Ibhaddon, V. Hessel, Q. Wang, Low temperature
649 plasma-catalytic NO_x synthesis in a packed DBD reactor: Effect of support materials and
650 supported active metal oxides, *Appl. Catal. B Environ.* 194 (2016) 123-133.
651 <https://doi.org/10.1016/j.apcatb.2016.04.055>.

652 [11] L. Wang, Y. Yi, C. Wu, H. Guo, X. Tu, One-Step Reforming of CO₂ and CH₄ into High-
653 Value Liquid Chemicals and Fuels at Room Temperature by Plasma-Driven Catalysis, *Angew.*
654 *Chem. Int. Ed.* 56 (2017) 13679-13683. <https://doi.org/10.1002/anie.201707131>.

655 [12] Y. Uytendhouwen, K.M. Bal, E.C. Neyts, V. Meynen, P. Cool, A. Bogaerts, On the kinetics
656 and equilibria of plasma-based dry reforming of methane, *Chem. Eng. J.* 405 (2021) 126630.
657 <https://doi.org/10.1016/j.cej.2020.126630>.

658 [13] X. Tu, J.C. Whitehead, Plasma dry reforming of methane in an atmospheric pressure AC
659 gliding arc discharge: Co-generation of syngas and carbon nanomaterials, *Int. J. Hydrogen*
660 *Energy* 39 (2014) 9658-9669. <https://doi.org/10.1016/j.ijhydene.2014.04.073>.

661 [14] Y. Yi, C. Xu, L. Wang, J. Yu, Q. Zhu, S. Sun, X. Tu, C. Meng, J. Zhang, H. Guo, Selectivity
662 control of H₂/O₂ plasma reaction for direct synthesis of high purity H₂O₂ with desired
663 concentration, *Chem. Eng. J.* 313 (2017) 37-46.
664 <https://doi.org/10.1016/j.cej.2016.12.043>.

665 [15] Y. Yi, X. Wang, A. Jafarzadeh, L. Wang, P. Liu, B. He, J. Yan, R. Zhang, H. Zhang, X. Liu,
666 H. Guo, E.C. Neyts, A. Bogaerts, Plasma-Catalytic Ammonia Reforming of Methane over Cu-
667 Based Catalysts for the Production of HCN and H₂ at Reduced Temperature, *ACS Catalysis* 11
668 (2021) 1765-1773. <https://doi.org/10.1021/acscatal.0c04940>.

669 [16] X. Gao, Z. Lin, T. Li, L. Huang, J. Zhang, S. Askari, N. Dewangan, A. Jangam, S. Kawi,
670 Recent Developments in Dielectric Barrier Discharge Plasma-Assisted Catalytic Dry Reforming
671 of Methane over Ni-Based Catalysts, *Catalysts* 11 (2021). <https://doi.org/10.3390/catal11040455>.

672 [17] R. Vakili, R. Gholami, C.E. Stere, S. Chansai, H. Chen, S.M. Holmes, Y. Jiao, C. Hardacre,
673 X. Fan, Plasma-assisted catalytic dry reforming of methane (DRM) over metal-organic
674 frameworks (MOFs)-based catalysts, *Appl. Catal. B Environ.* 260 (2020) 118195.
675 <https://doi.org/10.1016/j.apcatb.2019.118195>.

676 [18] W.-C. Chung, M.-B. Chang, Review of catalysis and plasma performance on dry reforming
677 of CH₄ and possible synergistic effects, *Renewable and Sustainable Energy Reviews* 62 (2016)
678 13-31. <https://doi.org/10.1016/j.rser.2016.04.007>.

679 [19] A.H. Khoja, M. Tahir, N.A.S. Amin, Recent developments in non-thermal catalytic DBD
680 plasma reactor for dry reforming of methane, *Energy Convers. Manage.* 183 (2019) 529-560.
681 <https://doi.org/10.1016/j.enconman.2018.12.112>.

682 [20] A. Ozkan, T. Dufour, A. Bogaerts, F. Reniers, How do the barrier thickness and dielectric
683 material influence the filamentary mode and CO₂ conversion in a flowing DBD?, *Plasma Sources*
684 *Sci. Technol.* 25 (2016) 045016. <https://doi.org/10.1088/0963-0252/25/4/045016>.

685 [21] A. Ozkan, T. Dufour, T. Silva, N. Britun, R. Snyders, A. Bogaerts, F. Reniers, The influence
686 of power and frequency on the filamentary behavior of a flowing DBD—application to the
687 splitting of CO₂, *Plasma Sources Sci. Technol.* 25 (2016) 025013. <https://doi.org/10.1088/0963-0252/25/2/025013>.

688 [22] P. Vanraes, A. Nikiforov, A. Bogaerts, C. Leys, Study of an AC dielectric barrier single
689 micro-discharge filament over a water film, *Scientific Reports* 8 (2018) 10919.
690 <https://doi.org/10.1038/s41598-018-29189-w>.

691 [23] K. van 't Veer, Y. Engelmann, F. Reniers, A. Bogaerts, Plasma-Catalytic Ammonia
692 Synthesis in a DBD Plasma: Role of Microdischarges and Their Afterglows, *J. Phys. Chem. C*
693 124 (2020) 22871-22883. <https://doi.org/10.1021/acs.jpcc.0c05110>.

694 [24] C. De Bie, J. van Dijk, A. Bogaerts, The Dominant Pathways for the Conversion of Methane
695 into Oxygenates and Syngas in an Atmospheric Pressure Dielectric Barrier Discharge, *J. Phys.*
696 *Chem. C* 119 (2015) 22331-22350. <https://doi.org/10.1021/acs.jpcc.5b06515>.

697

698 [25] R. Snoeckx, R. Aerts, X. Tu, A. Bogaerts, Plasma-Based Dry Reforming: A Computational
699 Study Ranging from the Nanoseconds to Seconds Time Scale, *J. Phys. Chem. C* 117 (2013) 4957-
700 4970. <https://doi.org/10.1021/jp311912b>.

701 [26] D. Ray, P.M.K. Reddy, C. Subrahmanyam, Ni-Mn/ γ -Al₂O₃ assisted plasma dry reforming of
702 methane, *Catal. Today* 309 (2018) 212-218.
703 <https://doi.org/https://doi.org/10.1016/j.cattod.2017.07.003>.

704 [27] A. Bogaerts, G. Centi, Plasma technology for CO₂ conversion: a personal perspective on
705 prospects and gaps, *Frontiers in Energy Research* 8 (2020) 111.

706 [28] A. Añiral, T. Nozaki, M. Nakase, S. Yuzawa, K. Okazaki, J.G.E. Gardeniers, Gas-to-liquids
707 process using multi-phase flow, non-thermal plasma microreactor, *Chem. Eng. J.* 167 (2011) 560-
708 566. <https://doi.org/https://doi.org/10.1016/j.cej.2010.10.050>.

709 [29] H. Sekiguchi, M. Ando, H. Kojima, Study of hydroxylation of benzene and toluene using a
710 micro-DBD plasma reactor, *J. Phys. D Appl. Phys.* 38 (2005) 1722-1727.
711 <https://doi.org/10.1088/0022-3727/38/11/013>.

712 [30] Y. Uytendhouwen, S. Van Alphen, I. Michielsen, V. Meynen, P. Cool, A. Bogaerts, A
713 packed-bed DBD micro plasma reactor for CO₂ dissociation: Does size matter?, *Chem. Eng. J.*
714 348 (2018) 557-568. <https://doi.org/https://doi.org/10.1016/j.cej.2018.04.210>.

715 [31] Y. Uytendhouwen, K.M. Bal, I. Michielsen, E.C. Neyts, V. Meynen, P. Cool, A. Bogaerts,
716 How process parameters and packing materials tune chemical equilibrium and kinetics in plasma-
717 based CO₂ conversion, *Chem. Eng. J.* 372 (2019) 1253-1264.
718 <https://doi.org/https://doi.org/10.1016/j.cej.2019.05.008>.

719 [32] N. Pinhão, A. Moura, J.B. Branco, J. Neves, Influence of gas expansion on process
720 parameters in non-thermal plasma plug-flow reactors: A study applied to dry reforming of
721 methane, *Int. J. Hydrogen Energy* 41 (2016) 9245-9255.
722 <https://doi.org/https://doi.org/10.1016/j.ijhydene.2016.04.148>.

723 [33] W.-C. Chung, K.-L. Pan, H.-M. Lee, M.-B. Chang, Dry Reforming of Methane with
724 Dielectric Barrier Discharge and Ferroelectric Packed-Bed Reactors, *Energy & Fuels* 28 (2014)
725 7621-7631. <https://doi.org/10.1021/ef5020555>.

726 [34] F.J.J. Peeters, M.C.M. van de Sanden, The influence of partial surface discharging on the
727 electrical characterization of DBDs, *Plasma Sources Sci. Technol.* 24 (2014) 015016.
728 <https://doi.org/10.1088/0963-0252/24/1/015016>.

729 [35] Y. Wang, Y. Chen, J. Harding, H. He, A. Bogaerts, X. Tu, Catalyst-free single-step plasma
730 reforming of CH₄ and CO₂ to higher value oxygenates under ambient conditions, *Chem. Eng. J.*
731 450 (2022) 137860. <https://doi.org/https://doi.org/10.1016/j.cej.2022.137860>.

732 [36] G.J.M. Hagelaar, L.C. Pitchford, Solving the Boltzmann equation to obtain electron transport
733 coefficients and rate coefficients for fluid models, *Plasma Sources Sci. Technol.* 14 (2005) 722.
734 <https://doi.org/10.1088/0963-0252/14/4/011>.

735 [37] K. van 't Veer, S. van Alphen, A. Remy, Y. Gorbanev, N. De Geyter, R. Snyders, F. Reniers,
736 A. Bogaerts, Spatially and temporally non-uniform plasmas: microdischarges from the
737 perspective of molecules in a packed bed plasma reactor, *J. Phys. D Appl. Phys.* 54 (2021)
738 174002. <https://doi.org/10.1088/1361-6463/abe15b>.

739 [38] R. Aerts, W. Somers, A. Bogaerts, Carbon Dioxide Splitting in a Dielectric Barrier
740 Discharge Plasma: A Combined Experimental and Computational Study, *ChemSusChem* 8
741 (2015) 702-716. <https://doi.org/https://doi.org/10.1002/cssc.201402818>.

742 [39] B. Wang, W. Yan, W. Ge, X. Duan, Kinetic model of the methane conversion into higher
743 hydrocarbons with a dielectric barrier discharge microplasma reactor, *Chem. Eng. J.* 234 (2013)
744 354-360. <https://doi.org/https://doi.org/10.1016/j.cej.2013.08.052>.