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# Co-electrolysis SOEC and internal reforming SOFC for achieving a carbon-neutral society

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Solid oxide electrolysis cells (SOECs) can produce carbon monoxide (CO) from carbon dioxide (CO<sub>2</sub>) in addition to hydrogen (H<sub>2</sub>) from steam using renewable energy sources such as solar and wind power. Methanol (MeOH) and dimethyl ether (DME) are industrially synthesized from syngas (H<sub>2</sub> + CO mixture) with an H<sub>2</sub>/CO ratio of 2. Solid oxide fuel cells (SOFCs) can generate power using MeOH and DME in addition to H<sub>2</sub>. To achieve a carbon-neutral society, H<sub>2</sub>O/CO<sub>2</sub> co-electrolysis SOECs and internal reforming SOFCs using MeOH and DME were investigated using the same negatrode-supported microtubular cells. As a result of quantitative analysis of the gas composition, the H<sub>2</sub>/CO ratios of the product gases were identical to the H<sub>2</sub>O/CO<sub>2</sub> ratios of the input gases during the co-electrolysis. The gas diffusion resistance increased with decreasing H<sub>2</sub>O/CO<sub>2</sub> ratios for co-electrolysis SOECs, and increased with time during co-electrolysis at H<sub>2</sub>O/CO<sub>2</sub> = 2 and internal reforming of MeOH and DME. H<sub>2</sub>O/CO<sub>2</sub> co-electrolysis SOECs and internal reforming SOFCs using MeOH and DME were successfully operated for 100 h.

## Sustainability spotlight

To achieve a carbon-neutral society, converting carbon dioxide (CO<sub>2</sub>) into raw materials and synthetic fuels (e-fuel) is very important. Solid oxide electrolysis cells (SOECs) can produce syngas (H<sub>2</sub> + CO mixture) from steam and CO<sub>2</sub> using renewable energy sources such as solar and wind power. Syngas can be converted into methanol (MeOH) and dimethyl ether (DME) in addition to e-fuel such as gasoline, kerosene, and diesel *via* Fischer–Tropsch synthesis. MeOH and DME are fuels for solid oxide fuel cells (SOFCs) in addition to raw materials for commodity plastics (e-chemical). The reversible SOECs and SOFCs using carbon-containing fuels contribute to the following UN sustainable development goals: affordable and clean energy (SDG 7), responsible consumption and production (SDC 12), and climate action (SDG 13).

## 1. Introduction

Solid oxide fuel cells (SOFCs) are expected to be power generation systems with high energy conversion efficiencies. SOFCs can directly use hydrocarbon and alcohol fuels in addition to hydrogen (H<sub>2</sub>), because oxide ions (O<sup>2-</sup>) conduct through an electrolyte and the electrochemical oxidation occurs at the negatrode (fuel electrode) at high temperatures above 600 °C.<sup>1–3</sup> Power generation with internal partial oxidation reforming of butane and steam reforming of ethanol has been investigated using microtubular SOFCs supported on a Ni-Gd-doped ceria (GDC) negatrode.<sup>4</sup> A long-term flight of a drone (UAV; unmanned aerial vehicle) was demonstrated using a portable SOFC system powered by liquefied petroleum gas (LPG) cartridges.<sup>5</sup> Because LPG is a fossil fuel, replacing LPG with carbon-neutral fuels is desirable. Although bio-ethanol helps to reduce carbon dioxide (CO<sub>2</sub>) emissions, sulfur-containing bio-

fuels deteriorate Ni-based SOFC negatrodes by promoting carbon deposition.<sup>6</sup>

Solid oxide cells can be used as electrolyzers as shown in Fig. 1. Solid oxide electrolysis cells (SOECs) can produce carbon monoxide (CO) from CO<sub>2</sub> in addition to H<sub>2</sub> from steam *via* the reverse reaction of SOFCs with O<sup>2-</sup> conduction through the electrolyte. Alkaline and polymer electrolyte membrane (PEM) electrolysis are widely demonstrated for H<sub>2</sub> production.<sup>7</sup> Some researchers have investigated low-temperature CO<sub>2</sub> electrolysis using alkaline aqueous electrolytes and PEMs. However, these require a high cell voltage above 2.5 V, and these exhibit a low faradaic efficiency.<sup>8</sup> On the contrary, SOECs can produce CO at 1.4 V *via* high-temperature CO<sub>2</sub> electrolysis, resulting in faradaic and energy efficiencies of 100% in principle. If external thermal energy can compensate for the latent heat of water vaporization and endothermic electrolysis, a further decrease in cell voltage and an increase in efficiency are possible.

CO<sub>2</sub> capture is one of the most important technologies for achieving a carbon-neutral society. Chemical adsorption method involving monoethanolamine and gas separation membranes have been demonstrated for CO<sub>2</sub> capture.<sup>9</sup> The initial target is treating the exhaust gases from plants, which

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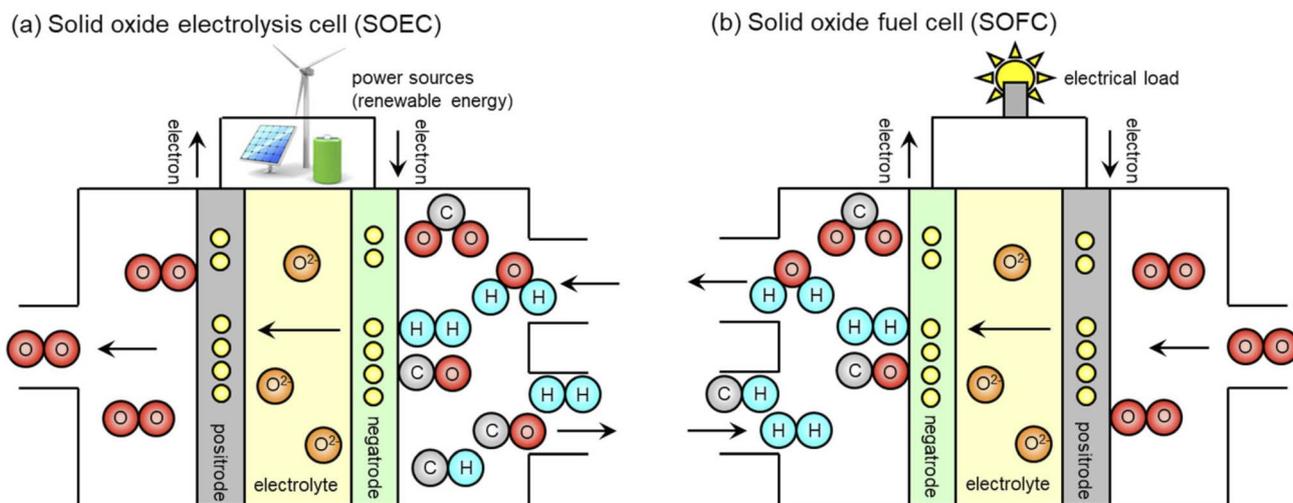
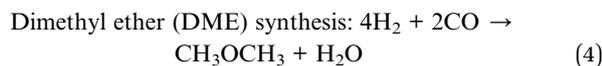
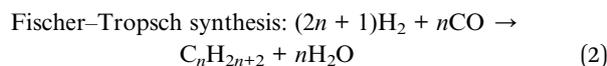


Fig. 1 Schematics of (a) co-electrolysis SOEC and (b) internal reforming SOFC.

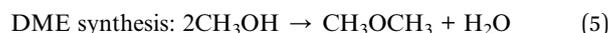
contain 4–14 vol% CO<sub>2</sub>. Recently, direct air capture (DAC), which extracts CO<sub>2</sub> directly from the atmosphere at any location, has been investigated.<sup>10</sup> Mixed ionic liquid membranes exhibit a high CO<sub>2</sub> permeability of  $2.5983 \times 10^{-6} \text{ cm}^3(\text{STP}) \cdot \text{cm cm}^{-2} \text{ s}^{-1} \text{ cmHg}^{-1}$  and a high CO<sub>2</sub>/N<sub>2</sub> selectivity of  $1.0059 \times 10^4$  at 400 ppm CO<sub>2</sub>.<sup>11</sup>

SOECs can simultaneously perform steam and CO<sub>2</sub> electrolysis, known as co-electrolysis.<sup>12–16</sup> Syngas, a mixture of H<sub>2</sub> and CO, is produced *via* the co-electrolysis SOECs. Syngas can be converted into many useful sources and fuels as follows:



Methanation is used to produce synthetic natural gas,<sup>17</sup> with the advantage of utilizing existing pipelines and infrastructure. However, the SOFCs using synthetic natural gas require a high temperature of 790 °C for methane reforming. Fischer–Tropsch (FT) synthesis is used to produce synthetic fuels (e-fuel) such as gasoline, kerosene, and diesel for airplanes, vessels, and large vehicles.<sup>18</sup> A Co/SiO<sub>2</sub> catalyst yields 1500 and 815 g kg<sub>cat</sub><sup>−1</sup> h<sup>−1</sup> of C<sub>5+</sub> and C<sub>10–20</sub> hydrocarbons, respectively, *via* FT synthesis at 776 °C and 1.1 MPa.<sup>19</sup> However, SOFCs easily deteriorate because of carbon deposition, when hydrocarbons with high carbon number are used as fuels. Therefore, the author focuses on methanol (MeOH) and dimethyl ether (DME). MeOH is industrially produced from syngas generated by natural gas reforming using Cu/ZnO-based catalysts.<sup>20–22</sup> Direct methanol fuel cells (DMFCs) using PEM electrolyte have been investigated owing to their low reforming temperatures below 150 °C.<sup>23</sup> However, DMFCs suffer from CO poisoning on a negatrocathode and

MeOH crossover through an electrolyte. For SOFCs, MeOH is a promising fuel, because it does not cause CO poisoning and MeOH crossover.<sup>24–27</sup> In addition, DME can be easily produced by the dehydration of methanol.<sup>28</sup>



The reforming temperature of DME is less than 180 °C, and its characteristics are similar to those of LPG, because it is easily liquefied at 0.6 MPa. Thus, DME is a possible alternative to LPG as a fuel for portable SOFC systems.<sup>29,30</sup>

In the present work, co-electrolysis SOECs and internal reforming SOFCs were investigated using microtubular negatrocathode-supported cells with the same cell configuration. Numerous researchers have demonstrated reversible SOFC/SOEC operation using H<sub>2</sub> and steam.<sup>31–33</sup> However, there have been few demonstrations of reversible operation using carbon-containing fuels. The H<sub>2</sub>/CO ratio of the syngas should be 2 for MeOH, DME, and FT syntheses. To investigate the effect of CO<sub>2</sub> concentration on electrochemical characteristics, co-electrolysis was performed with varying H<sub>2</sub>O/CO<sub>2</sub> ratios from 0 to infinity (∞). The compositions of the product gases after co-electrolysis were analyzed using gas chromatography. Assuming that MeOH and DME are synthesized from syngas, internal reforming SOFCs were demonstrated using MeOH and DME. Steam reforming was investigated for MeOH, because it is a liquid with water solubility. Partial oxidation reforming was investigated for DME in addition to steam reforming, since it easily mixes with ambient air at ordinary temperatures and pressures. Short-term durability tests of the co-electrolysis SOECs using an input gas at H<sub>2</sub>O/CO<sub>2</sub> = 2 and internal reforming SOFCs using MeOH and DME were conducted for 100 h.

## 2. Experimental

### 2.1 Cell preparation

Similar to previously reported internal reforming of butane and ethanol,<sup>4</sup> negatrocathode-supported microtubular cells with the same



cell configuration were used for co-electrolysis SOECs and internal reforming SOFCs using MeOH and DME. Negatode microtube substrates were constructed using 60 wt% NiO (Sumitomo Metal Mining)-40 wt%  $\text{Ce}_{0.9}\text{Gd}_{0.1}\text{O}_{1.95}$  (GDC; Shin-Etsu Chemical) adding pore former (acrylic resin; Sekisui Plastic) and binder (cellulose; Yuken Kogyo) powders. These powders were mixed using a kneading machine with adding an appropriate amount of water, and the negatode microtubes were extruded using a piston cylinder. After extrusion, the microtubes were dried overnight in air at room temperature. A slurry was prepared by mixing  $(\text{Y}_2\text{O}_3)_{0.08}(\text{ZrO}_2)_{0.92}$  (YSZ; Tosoh), binder (polyvinyl butyral; Sekisui Chemical) dispersant (tallow propylene diamine, Kao Chemicals) and plasticizer (dioctyl adipate; Wako Pure Chemical Industries) with ethanol and toluene solvents. The YSZ electrolyte was formed by dip-coating. The YSZ electrolyte thin-film and the NiO-GDC negatode microtube were co-sintered in air for 3 h at 1400 °C. The GDC interlayer and the positrode (air electrode) of 70 wt%  $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.2}\text{Fe}_{0.8}\text{O}_{3-\delta}$  (LSCF; Kusaka Rare Metal)-30 wt% GDC were coated in a similar manner. The interlayer and positrode thin-film layers were sintered sequentially in air for 2 h at 1200 °C and for 1 h at 1050 °C, respectively. The microtube diameter and positrode area after sintering were 2.8 mm and 2.6 cm<sup>2</sup>, respectively. The thicknesses of the negatode, electrolyte, interlayer, and positrode were 640, 10, 1, and 20 μm, respectively.

## 2.2 Evaluation of co-electrolysis SOECs and internal reforming SOFCs

Table 1 and Fig. 2 show the experimental conditions and setup for evaluating the co-electrolysis SOECs and internal reforming SOFCs. For internal steam reforming of MeOH, an aqueous MeOH solution (82 vol% at S/C = 0.5 and 69 vol% at S/C = 1.0) was supplied to the negatode using a pump and vaporizer heated at 250 °C. For co-electrolysis and internal steam reforming of DME, steam was fed to the negatode using a temperature-controlled bubbler. Before evaluation, a mixture of 20% H<sub>2</sub>-3% H<sub>2</sub>O-77% N<sub>2</sub> was supplied to the negatode for 2 h at 700 °C for Ni catalyst reduction. Subsequently, the input gases were adjusted to the flow rates listed in Table 1, and air was supplied to the positrode at a rate of 100 mL min<sup>-1</sup>. Hydrogen was supplied during co-electrolysis SOEC to prevent Ni catalyst oxidation in the negatode under open circuit voltage (OCV). Current-voltage (*I*-*V*) characteristics were evaluated using a potentiostat/galvanostat (Solartron Analytical 1470E) by varying the voltage from OCV to 0.4 and 1.4 V for SOECs and SOFCs, respectively, at a sweep rate of 5 mV s<sup>-1</sup>. Using a frequency response analyzer (Solartron Analytical 1455), the electrochemical impedance between the negatode and positrode was measured in the frequency range of 100 kHz-0.1 Hz with 20 steps per logarithmic decade. The polarization impedance can be deconvoluted using the distribution of relaxation times (DRT) analysis.<sup>34,35</sup> The DRT analysis was performed using

Table 1 Flow rates (mL min<sup>-1</sup>) of the input gases for co-electrolysis SOECs and internal steam (SR) and partial oxidation (POx) reforming SOFCs

	H <sub>2</sub> O/CO <sub>2</sub>	H <sub>2</sub>	H <sub>2</sub> O	CO <sub>2</sub>	O <sub>2</sub>	MeOH or DME	N <sub>2</sub>
Co-electrolysis	H <sub>2</sub> O/CO <sub>2</sub> = 0	3.0	0.0	20.0			77.0
	H <sub>2</sub> O/CO <sub>2</sub> = 1/3	3.0	5.0	15.0			77.0
	H <sub>2</sub> O/CO <sub>2</sub> = 1	3.0	10.0	10.0			77.0
	H <sub>2</sub> O/CO <sub>2</sub> = 2	3.0	13.3	6.7			77.0
	H <sub>2</sub> O/CO <sub>2</sub> = 3	3.0	15.0	5.0			77.0
	H <sub>2</sub> O/CO <sub>2</sub> = ∞	3.0	20.0	0.0			77.0
MeOH SR	S/C = 0.5		5.0			10.0	85.0
	S/C = 1.0		10.0			10.0	80.0
DME SR	S/C = 1.0		10.0			5.0	85.0
	S/C = 1.5		15.0			5.0	80.0
DME POx	O/C = 1.0				10.0	10.0	61.4
	O/C = 1.5				15.0	10.0	56.4

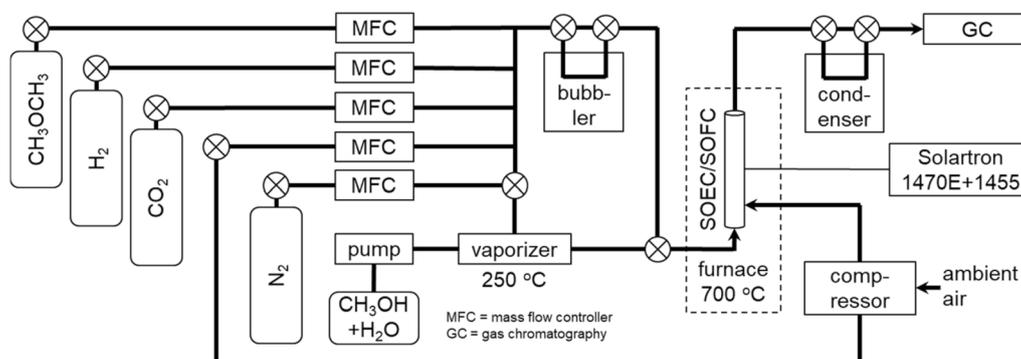


Fig. 2 Schematic of experimental setup for evaluation of co-electrolysis SOECs and internal reforming SOFCs.



Z-Assist software (Toyo Corp.).<sup>36</sup> It used FTIKREG software<sup>37</sup> to solve an ill-posed inverse problem in the DRT analysis *via* Tikhonov regularization. Real impedance was used for the DRT analysis because the impact of measurement errors and inductive components was lower than that with imaginary impedance.<sup>38</sup> Kramers–Kronig validation was performed using K–K test<sup>39</sup> and Lin-KK Tool<sup>40</sup> software. Residuals between measured and Kramers–Kronig transformed impedances were within 0.5%. The parameters obtained DRT analysis were refined by complex non-linear least squares (CNLS) fitting using ZView software (Scribner Associates), assuming an equivalent circuit model with a series connection of resistance ( $R_0$ ) and parallel resistance-capacitance ( $R_kC_k$ ) elements. The thermodynamic equilibrium compositions were calculated using HSC Chemistry 5.11 (Outokumpu). During co-electrolysis, the flow rates of the product gases were analyzed with a thermal conductivity detector (TCD) in a micro gas chromatograph (Agilent Technologies 490; Molsieve 5A column with Ar as the carrier gas for  $H_2$ ,  $N_2$ ,  $CH_4$ , and  $CO$ , and PorapLOT Q column with He as the carrier gas for  $CO_2$ ). Steam flow rates in the product gases were not measured by micro gas chromatography, because steam was removed using a condenser as shown in Fig. 2. The steam flow rates of the product gases were estimated by considering the mass balance of the hydrogen, oxygen and carbon species. Unfortunately, the flow rates of the reformat gases during internal reforming could not be analyzed, because MeOH and DME are soluble in water. Short-term durability tests of co-electrolysis SOECs and internal reforming SOFCs were conducted for 100 h at a constant voltage of 1.28 V and a constant current density of  $0.3 \text{ A cm}^{-2}$ , respectively.

### 3. Results and discussion

#### 3.1 Co-electrolysis SOECs

Fig. 3 shows the equilibrium amounts of the product gases as a function of  $H_2O + CO_2$  utilization at (a)  $H_2O/CO_2 = 0$  and (b)  $H_2O/CO_2 = 2$  at  $700^\circ\text{C}$ . The input gases are 0.2 mol of  $H_2O/CO_2$  and 0.03 mol of  $H_2$ . The product amounts of the  $H_2/CO$  mixtures

are assumed to obey faradaic law. At  $H_2O/CO_2 = 0$ , carbon is deposited, when  $H_2O + CO_2$  utilization exceeds 0.8. At low temperatures, carbon is easily deposited, even at lower  $H_2O/CO_2$  ratios. Moreover, methane is slightly produced, when  $H_2O + CO_2$  utilization is above 0.6 at  $H_2O/CO_2 = 2$ . When carbon and/or methane are not produced, the amounts of  $H_2$  and  $CO$  produced increase linearly with the  $H_2O + CO_2$  utilization, and the  $H_2/CO$  ratios of the product gases are identical to the  $H_2O/CO_2$  ratios of the input gases. Co-electrolysis is desirable to be operated at high temperatures to prevent carbon and methane production.

Fig. 4 shows the  $I$ - $V$  characteristics for co-electrolysis SOEC at various  $H_2O/CO_2$  ratios and  $700^\circ\text{C}$ . The flow rates of the input gases are listed in Table 1. The theoretical electromotive forces (EMFs) are 0.910, 0.906, 0.902, 0.899, 0.898, and 0.894 V at  $H_2O/CO_2 = 0, 1/3, 1, 2, 3,$  and  $\infty$ , respectively. The OCVs were 20–30 mV lower than the theoretical EMFs under all conditions, owing to slight gas leakage. The slopes of  $I$ - $V$  curves increased with decreasing  $H_2O/CO_2$  ratios. The  $H_2O + CO_2$  utilizations were 0.27–0.32 under 1.28 V at  $H_2O/CO_2 = 0 - \infty$ . The performances during  $H_2O/CO_2$  co-electrolysis and  $CO_2$  electrolysis

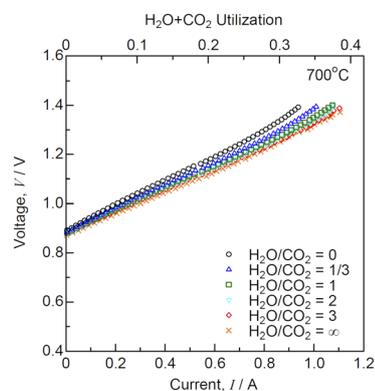


Fig. 4 Current–voltage characteristics for co-electrolysis SOEC at various  $H_2O/CO_2$  ratios and  $700^\circ\text{C}$ .

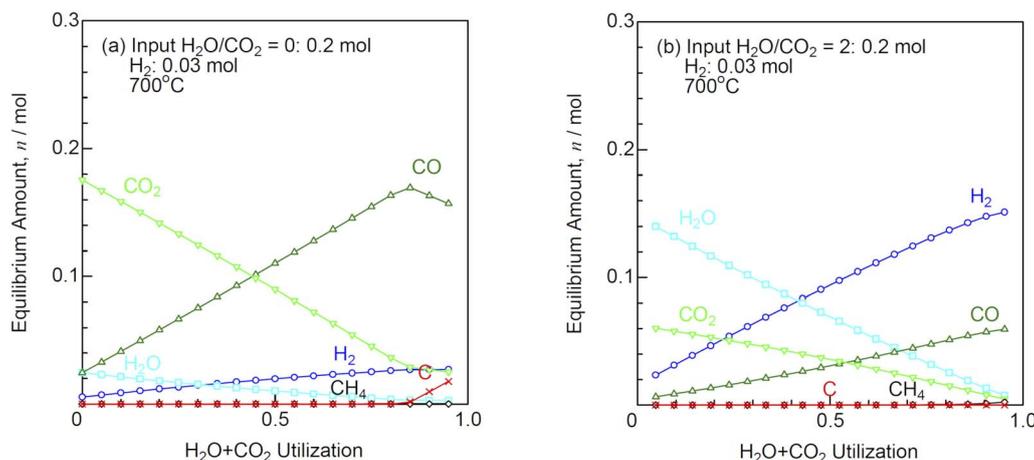


Fig. 3 Equilibrium amounts of the product gases as a function of  $H_2O + CO_2$  utilization at (a)  $H_2O/CO_2 = 0$  and (b)  $H_2O/CO_2 = 2$  (input  $H_2O/CO_2$ : 0.2 mol,  $H_2$ : 0.03 mol) at  $700^\circ\text{C}$ .



were lower than that during steam electrolysis.<sup>12,13,15,16</sup> The performance of the co-electrolysis SOEC decreased with increasing CO<sub>2</sub> concentrations in the input gases.

Fig. 5(a) and (b) show the Nyquist plots of impedance spectra for co-electrolysis SOEC under OCV and 1.28 V, respectively, at various H<sub>2</sub>O/CO<sub>2</sub> ratios and 700 °C. The ohmic losses ( $R_0$ ) derived from the intersections of the impedance arcs at high frequencies with the real axis remained unchanged with the H<sub>2</sub>O/CO<sub>2</sub> ratios. The ohmic losses under 1.28 V were slightly larger than those under OCV, because endothermic reaction of H<sub>2</sub>O/CO<sub>2</sub> co-electrolysis decreased the local temperature at the interface between electrolyte and electrode. The polarization resistances at low frequencies below 10 Hz increased with decreasing H<sub>2</sub>O/CO<sub>2</sub> ratios. DRT analysis was performed to deconvolute the impedance spectra as shown in Fig. 5(c) and (d). Six DRT peaks were observed under all the conditions, corresponding to the following physicochemical origins of the rate-determining elementary processes for negatrod-supported microtubular SOECs:<sup>36</sup>

$P_1$  (10–100 kHz): physical impedance originating from the grain boundaries, pores, and reaction products between the electrolyte and positrode

$P_2, P_3$  (20 Hz–10 kHz): charge transfer and ionic conduction processes at the negatrod

$P_4$  (2–200 Hz): oxygen surface exchange and diffusion processes at the positrode

$P_5, P_6$  (0.1–20 Hz): gas diffusion and nonstoichiometric oxygen variation processes at the Ni-GDC negatrod

Fig. 5(e) and (f) show the resistance against H<sub>2</sub>O/CO<sub>2</sub> ratios derived from the CNLS fitting using the equivalent circuit model with a series connection of  $R_0$  and six  $R_k C_k$  ( $k = 1-6$ ) elements.  $R_5$  and  $R_6$  increased with decreasing H<sub>2</sub>O/CO<sub>2</sub> ratios. Graves *et al.*<sup>12</sup> reported that the polarization resistance resulting from gas diffusion process at the negatrod increased at high CO<sub>2</sub> concentrations. This suggests that the CO/CO<sub>2</sub> mixture diffuses more difficultly than the H<sub>2</sub>/H<sub>2</sub>O mixture in the SOEC negatrod.

Syngas with an H<sub>2</sub>/CO ratio of 2 is desirable for methanol, DME, and FT syntheses. Therefore, a quantitative analysis of

**Table 2** Estimated flow rates (mL min<sup>-1</sup>) of the product gases for co-electrolysis SOEC under equilibrium, OCV, and 1.28 V using gas chromatography. The steam flow rates of the product gases were estimated in consideration for mass balance of hydrogen, oxygen and carbon species

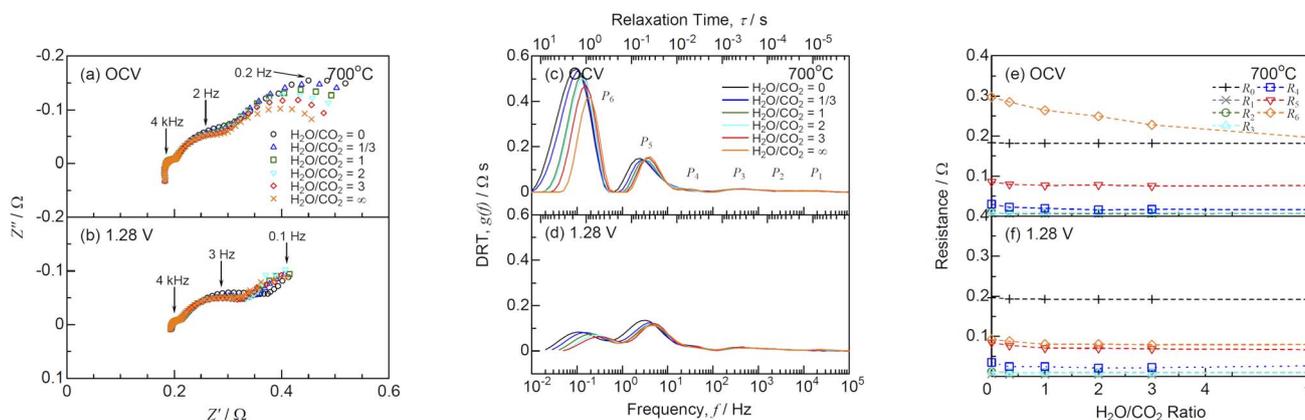
H <sub>2</sub> O/CO <sub>2</sub>	V	I	H <sub>2</sub>	H <sub>2</sub> O	CO	CO <sub>2</sub>	CH <sub>4</sub>	N <sub>2</sub>
0	Equilibrium		0.543	2.46	2.46	17.5	1 × 10 <sup>-6</sup>	77.0
	OCV	0 A	0.947	1.65	2.72	17.7	0	77.0
	1.28 V	0.770 A	1.12	2.43	7.70	11.7	0	77.0
1/3	Equilibrium		1.33	6.68	1.68	13.3	5 × 10 <sup>-6</sup>	77.0
	OCV	0 A	1.58	6.74	1.21	13.5	0	77.0
	1.28 V	0.830 A	3.13	5.33	5.51	9.03	0	77.0
1	Equilibrium		1.98	11.0	1.02	8.98	6 × 10 <sup>-6</sup>	77.0
	OCV	0 A	1.86	10.7	0.720	9.70	0	77.0
	1.28 V	0.871 A	4.96	7.61	3.80	6.63	0	77.0
2	Equilibrium		2.35	14.0	0.647	6.05	5 × 10 <sup>-6</sup>	77.0
	OCV	0 A	2.13	13.4	0.538	6.91	0	77.0
	1.28 V	0.886 A	6.22	10.0	2.44	4.31	0	77.0
3	Equilibrium		2.53	15.5	0.469	4.53	4 × 10 <sup>-6</sup>	77.0
	OCV	0 A	2.38	15.3	0.394	4.95	0	77.0
	1.28 V	0.902 A	7.22	10.6	1.84	3.39	0	77.0
∞	Equilibrium		3.00	20.0	0	0	0	77.0
	OCV	0 A	3.08	19.9	0	0	0	77.0
	1.28 V	0.909 A	9.46	13.5	0	0	0	77.0

product gas compositions is important for the co-electrolysis SOEC evaluation. Table 2 shows the estimated flow rates of the product gases for co-electrolysis SOEC under equilibrium, OCV and 1.28 V using gas chromatography. The steam flow rates were estimated in consideration for mass balance of hydrogen, oxygen and carbon species assuming no carbon deposition. The faradaic law for electrolysis is as follows:

$$I = \dot{n}zF \quad (6)$$

where  $I$  is the current (A),  $\dot{n}$  is the molar product rate (mol s<sup>-1</sup>),  $z$  is the valency of the ions (2), and  $F$  is the faradaic constant (96 485C mol<sup>-1</sup>). The ideal gas law is as follows:

$$p\dot{v} = 60\dot{n}R_g T \quad (7)$$



**Fig. 5** (a and b) Impedance, (c and d) DRT spectra, and (e and f) resistances for co-electrolysis SOEC under (a, c and e) OCV and (b, d and f) 1.28 V at various H<sub>2</sub>O/CO<sub>2</sub> ratios and 700 °C.



where  $p$  is the pressure (kPa),  $\dot{v}$  is the flow rate of the product gas ( $\text{L min}^{-1}$ ),  $R_g$  is the gas constant ( $8.3145 \text{ J mol}^{-1} \text{ K}^{-1}$ ), and  $T$  is the absolute temperature (K). In the standard state ( $p = 101.325 \text{ kPa}$  and  $T = 273.15 \text{ K}$ ),  $\dot{v}$  is derived from the following equation:

$$\dot{v} = 6.9652 \times 10^{-3} I \quad (8)$$

This indicates the production of approximately  $7 \text{ mL min}^{-1}$  of syngas per 1 A current. Under OCV, the equilibrium compositions differed from the input gas compositions owing to the reverse water gas shift reaction.



The differences between the equilibrium and measured gas flow rates were less than  $1 \text{ mL min}^{-1}$  under OCV. Under 1.28 V, the  $\text{H}_2 + \text{CO}$  flow rates in the range of  $5.15\text{--}6.38 \text{ mL min}^{-1}$  increased, when the current was  $0.770\text{--}0.909 \text{ A}$  at  $\text{H}_2\text{O}/\text{CO}_2 = 0 - \infty$ . The faradaic efficiencies were  $0.96\text{--}1$ , because carbon

and methane were not produced in the  $\text{H}_2\text{O} + \text{CO}_2$  utilizations ranges of  $0.27\text{--}0.32$ . Fig. 6 shows the evolution rates of the product gases for co-electrolysis SOEC at various  $\text{H}_2\text{O}/\text{CO}_2$  ratios and  $700^\circ\text{C}$ . The  $\text{H}_2/\text{CO}$  ratios of the product gases were almost the identical to the  $\text{H}_2\text{O}/\text{CO}_2$  ratios of the input gases.  $\text{CO}_2$  electrolysis was slightly more difficult than steam electrolysis, because the polarization resistances increased with decreasing  $\text{H}_2\text{O}/\text{CO}_2$  ratios as shown in Fig. 5. However, CO was produced *via* the reverse water gas shift reaction (eqn (9)) in addition to co-electrolysis. Thus, the product gas compositions easily reached equilibrium compositions.

The short-term durability test was performed under a constant voltage of 1.28 V at  $\text{H}_2\text{O}/\text{CO}_2 = 2$  and  $700^\circ\text{C}$  as shown in Fig. 7. Under 1.28 V, the initial current was 0.886 A, which decreased to 0.756 A after 100 h. The slope of the  $I\text{--}V$  curves increased, although the OCV remained unchanged during the durability test for 100 h. Fig. 8 shows the (a) impedance, (b) DRT spectra, and (c) resistances for co-electrolysis SOEC during the durability test under a constant voltage of 1.28 V at  $\text{H}_2\text{O}/\text{CO}_2 = 2$  and  $700^\circ\text{C}$ . The ohmic loss remained unchanged, whereas the polarization resistances at low frequencies below 10 Hz increased after 100 h. The DRT analysis suggested that  $R_5$  and  $R_6$  related to the gas diffusion process at the negatrod increased with time. The gas diffusion resistance should be decreased to improve durability in addition to the initial performance for co-electrolysis SOECs. Fig. 9 shows the evolution rates of the product gases during the durability test of the co-electrolysis SOEC under a constant voltage of 1.28 V at  $\text{H}_2\text{O}/\text{CO}_2 = 2$  and  $700^\circ\text{C}$ . The evolution rates of  $\text{H}_2$  and CO decreased with time, because the current decreased from 0.886 A to 0.756 A under 1.28 V after 100 h. The faradaic efficiency was remained above 0.95, confirming that the co-electrolysis SOEC produced syngas with a constant  $\text{H}_2/\text{CO}$  ratio of 2. MeOH and DME syntheses from syngas have been industrially established. Internal reforming SOFCs using MeOH and DME are discussed in the next section.

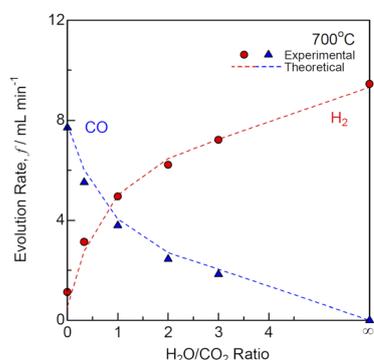


Fig. 6 Evolution rates of the product gases for co-electrolysis SOEC at various  $\text{H}_2\text{O}/\text{CO}_2$  ratios and  $700^\circ\text{C}$ .

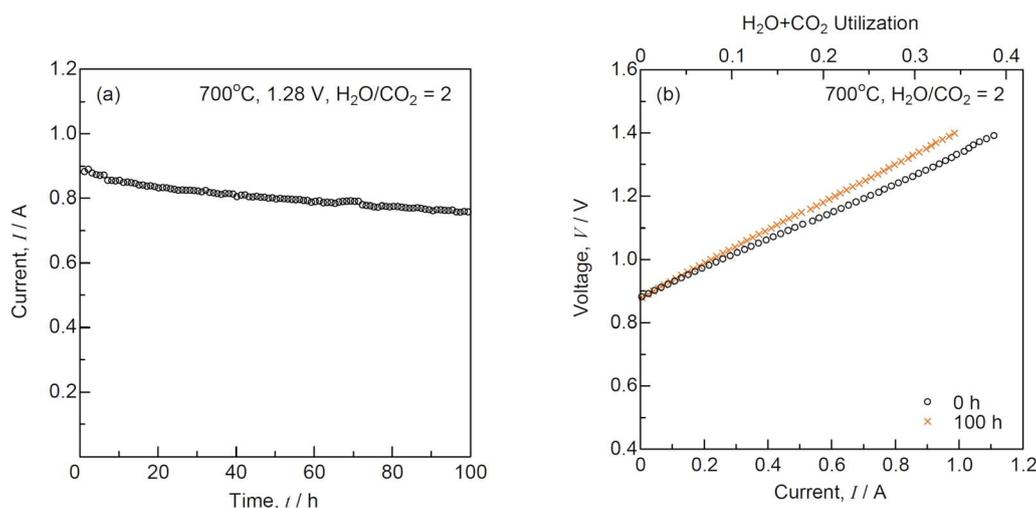


Fig. 7 (a) Time course of current under a constant voltage of 1.28 V and (b) current–voltage characteristics for co-electrolysis SOEC before and after the durability test at  $\text{H}_2\text{O}/\text{CO}_2 = 2$  and  $700^\circ\text{C}$ .



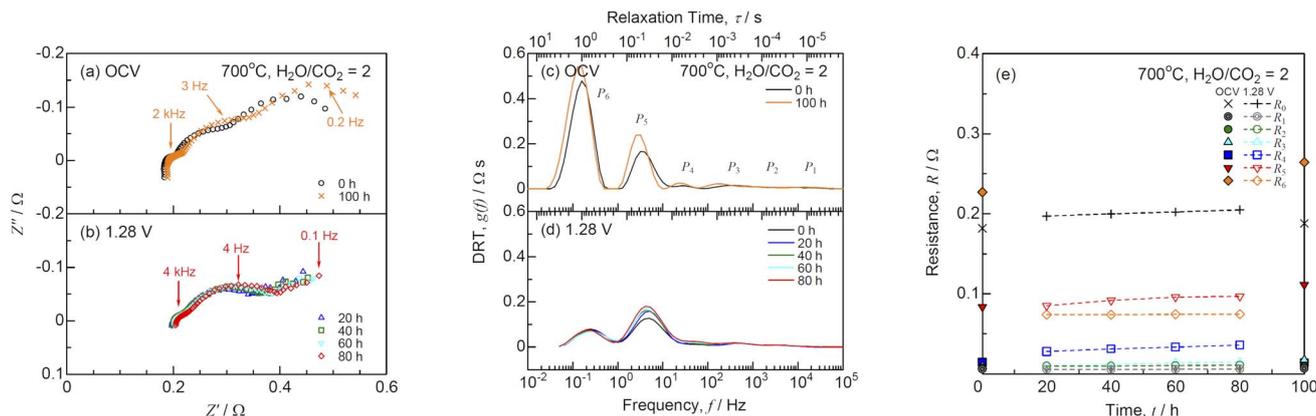


Fig. 8 (a and b) Impedance, (c and d) DRT spectra, and (e) resistances for co-electrolysis SOEC during the durability test under a constant voltage of 1.28 V at  $\text{H}_2\text{O}/\text{CO}_2 = 2$  and 700 °C.

### 3.2 Internal reforming SOFCs

MeOH and DME are easily reformed at low temperatures. The conversion of MeOH and DME reaches 99% at 150 and 180 °C, respectively, in thermodynamic equilibrium. The stoichiometric reaction for MeOH decomposition is as follows:



Fig. 10(a) shows the equilibrium amounts of the reformate gases as a function of S/C ratio during steam reforming of MeOH at 700 °C. Steam is not required stoichiometrically for MeOH decomposition as shown in eqn (10). However, steam reforming is required to prevent carbon deposition, because carbon is deposited at S/C ratios below 0.3 in equilibrium. Internal reforming SOFCs can be easily supplied with 82 vol% (S/C = 0.5) or 69 vol% (S/C = 1.0) MeOH water solutions. Meanwhile, DME can be reformed using partial oxidation (eqn (12)) in addition to steam (eqn (11)).



Fig. 10(b) and (c) show the equilibrium amounts of the reformate gases as a function of S/C and O/C ratios during steam reforming and partial oxidation reforming of DME at 700 °C. The stoichiometric S/C and O/C ratios are 0.5 for steam reforming and partial oxidation reforming of DME. However,

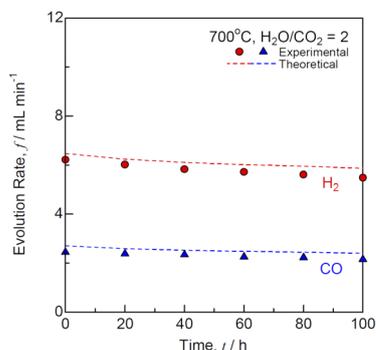


Fig. 9 Evolution rates of the product gases during the durability test of co-electrolysis SOEC under a constant voltage of 1.28 V at  $\text{H}_2\text{O}/\text{CO}_2 = 2$  and 700 °C.

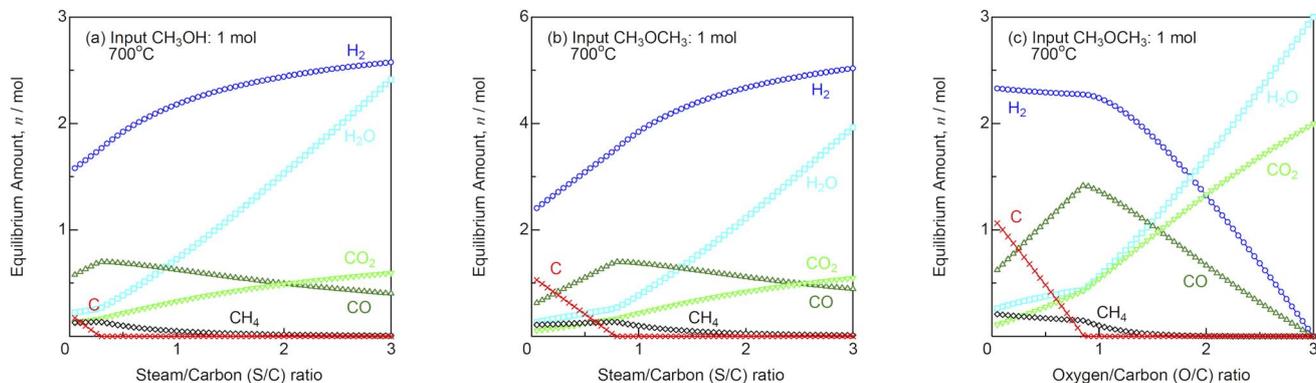


Fig. 10 Equilibrium amounts of the reformate gases as a function of S/C and O/C ratios for (a) steam reforming of MeOH, (b) steam reforming of DME, and (c) partial oxidation reforming of DME (input MeOH or DME: 1 mol) at 700 °C.



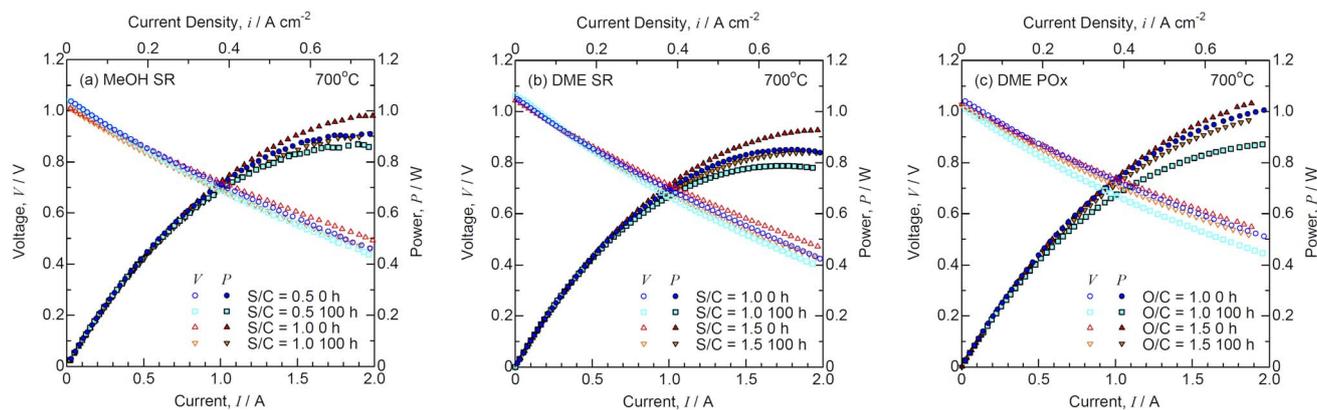


Fig. 11 Current–voltage characteristics for SOFCs during (a) steam reforming (SR) of MeOH, (b) SR of DME, and (c) partial oxidation reforming (POx) of DME before and after the durability tests at 700 °C.

carbon is deposited at S/C ratios <0.8 and O/C ratios <0.9 in equilibrium. The gas characteristics of DME are similar to those of LPG. Therefore, partial oxidation reforming can be easily performed by supplying DME/air mixture.

Fig. 11 shows the  $I$ - $V$  characteristics for SOFCs during (a) steam reforming of MeOH, (b) steam reforming of DME, and (c) partial oxidation of DME before and after the durability tests at 700 °C. The theoretical EMFs, OCVs, and voltages at  $0.3 \text{ A cm}^{-2}$  are summarized in Table 3. Unfortunately, the flow rate of the reformat gases during internal reforming could not be determined using the setup as shown in Fig. 2, because water-soluble MeOH and DME were removed from the condenser. The OCVs were almost identical to the theoretical EMFs under all the conditions, suggesting that MeOH and DME were reformed into  $\text{H}_2$  and  $\text{CO}$  by steam and partial oxidation. The OCVs remained unchanged during the durability tests. The  $I$ - $V$  characteristics were similar during internal reforming of MeOH and DME. Fig. 12 shows the time course of voltage for SOFCs during (a) steam reforming of MeOH, (b) steam reforming of DME, and (c) partial oxidation of DME during the durability test at a constant current density of  $0.3 \text{ A cm}^{-2}$  and 700 °C. Internal reforming SOFCs using MeOH and DME were continuously operated for 100 h. The degradation rates at different S/C ratios were almost the same during steam reforming of MeOH and DME, whereas the degradation rate at O/C = 1.0 was larger than that at O/C =

1.5 during partial oxidation reforming of DME. Previously, the voltage rapidly dropped during internal reforming of butane at  $\text{O/C} \leq 1.0$  and ethanol at  $\text{S/C} \leq 1.5$ ,<sup>4</sup> although Ni-ceria based negatrodes exhibit high tolerance against carbon deposition.<sup>3,24</sup> The voltage was stable during internal reforming of MeOH and DME, and no carbon deposition was detected at the Ni-GDC negatrode after the durability test. MeOH and DME contain C–O bonds but no C–C bonds, making carbon deposition difficult.<sup>25</sup>

The impedance spectra before and after the durability tests at 700 °C are shown in Fig. 13. Table 3 lists the ohmic losses ( $R_0$ ) and total polarization resistances ( $R_p$ ) before and after 100 h. The  $R_0$  values almost remained unchanged during the durability test, whereas those of  $R_p$  increased with time. The DRT analysis was performed to deconvolute the impedance spectra as shown in Fig. 14. Four DRT peaks corresponding to  $P_2$ – $P_5$  were detected under all the conditions. Fig. 15 shows the time course of the resistances derived from CNLS fitting using the equivalent circuit model with a series connection of  $R_0$  and four  $R_k C_k$  ( $k = 2$ – $5$ ) elements. The values of  $R_3$  and  $R_5$ , which were ascribed to charge transfer and ionic conduction processes, and gas diffusion processes in the negatrode, respectively, increased with time. The DRT peaks of  $P_3$  and  $P_5$  shifted to lower frequencies as shown in Fig. 14. The gas diffusion resistance of the negatrode-supported microtubular cells used in this work

Table 3 Theoretical EMFs, OCVs, voltages at  $0.3 \text{ A cm}^{-2}$ , ohmic losses and total polarization resistances under OCV for SOFCs during steam reforming (SR) of MeOH, SR of DME, and partial oxidation reforming (POx) of DME before and after the durability test at a constant current density of  $0.3 \text{ A cm}^{-2}$  and 700 °C

	MeOH SR		DME SR		DME POx	
Ratio	S/C = 0.5	S/C = 1.0	S/C = 1.0	S/C = 1.5	O/C = 1.0	O/C = 1.5
Theoretical EMF	1.04 V	1.02 V	1.04 V	1.02 V	1.03 V	0.997 V
OCV	1.05 V	1.02 V	1.05 V	1.04 V	1.05 V	1.03 V
$V_0 \text{ h}@0.3 \text{ A cm}^{-2}$	0.764 V	0.772 V	0.758 V	0.754 V	0.777 V	0.780 V
$V_{100 \text{ h}}@0.3 \text{ A cm}^{-2}$	0.749 V	0.748 V	0.734 V	0.733 V	0.730 V	0.761 V
$R_{0,0 \text{ h}}$	0.182 $\Omega$	0.160 $\Omega$	0.200 $\Omega$	0.180 $\Omega$	0.184 $\Omega$	0.165 $\Omega$
$R_{0,100 \text{ h}}$	0.182 $\Omega$	0.166 $\Omega$	0.202 $\Omega$	0.179 $\Omega$	0.186 $\Omega$	0.163 $\Omega$
$R_{p,0 \text{ h}}$	0.114 $\Omega$	0.115 $\Omega$	0.114 $\Omega$	0.121 $\Omega$	0.122 $\Omega$	0.115 $\Omega$
$R_{p,100 \text{ h}}$	0.137 $\Omega$	0.145 $\Omega$	0.151 $\Omega$	0.154 $\Omega$	0.162 $\Omega$	0.145 $\Omega$



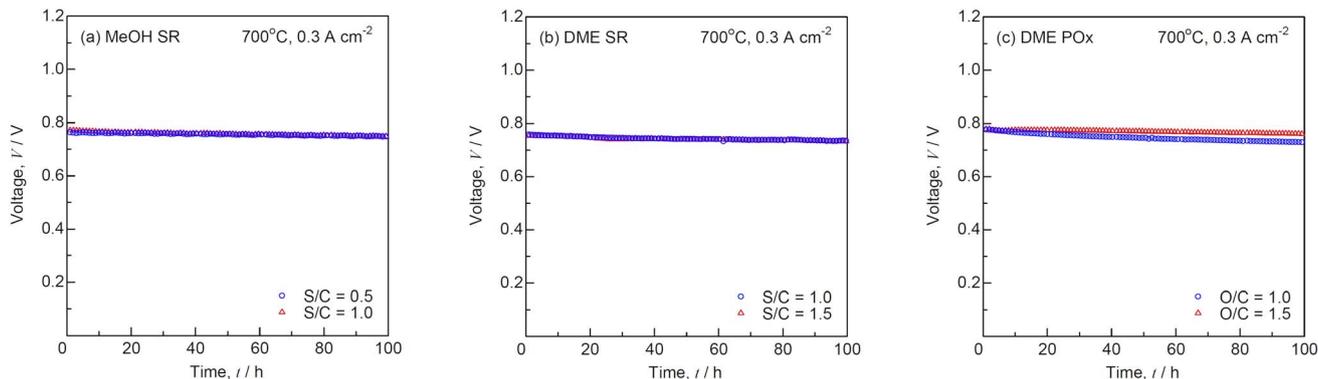


Fig. 12 Time courses of voltage for SOFCs during (a) steam reforming (SR) of MeOH, (b) SR of DME, and (c) partial oxidation reforming (POx) of DME during the durability test at a constant current density of 0.3 A cm<sup>-2</sup> and 700 °C.

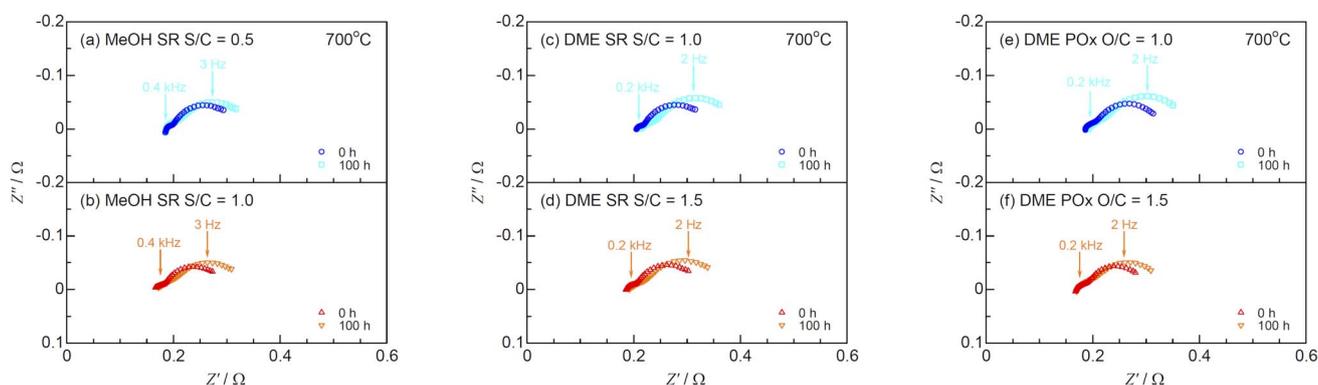


Fig. 13 Impedance spectra for SOFCs during (a and b) steam reforming (SR) of MeOH, (c and d) SR of DME, and (e and f) partial oxidation reforming (POx) of DME before and after the durability tests at 700 °C.

was large, degrading the negatode performance in charge transfer and gas diffusion processes during internal reforming of MeOH and DME. Thus, optimizing the negatode microstructure is important for improving the initial performance and durability of co-electrolysis SOECs and internal reforming SOFCs.

### 3.3 Carbon-neutral energy system using co-electrolysis SOECs and internal reforming SOFCs

Fig. 16 shows a schematic of the carbon-neutral energy system using co-electrolysis SOECs and internal reforming SOFCs. CO<sub>2</sub> is captured from the exhaust gas of plants and directly from the air (DAC).<sup>9–11</sup> Renewable energy sources such as solar and wind

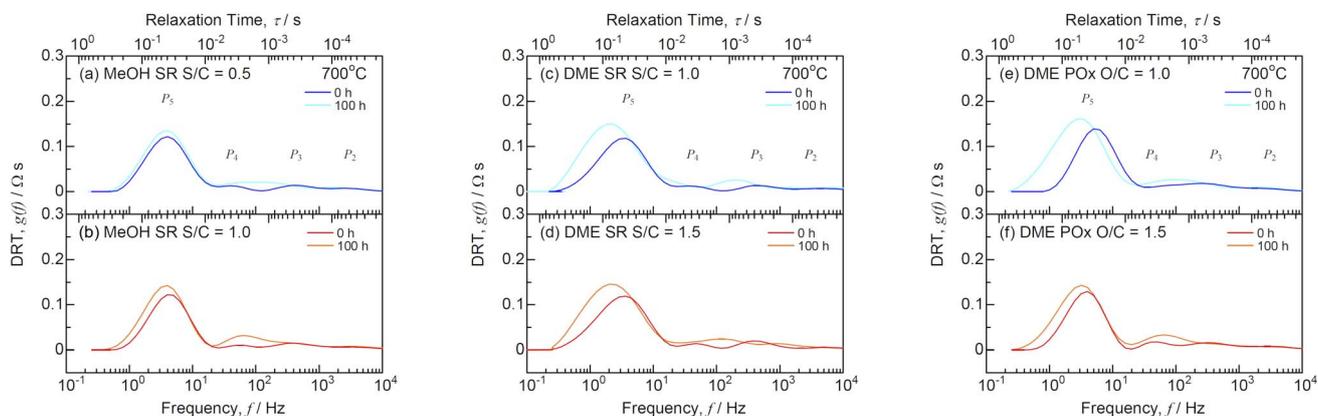


Fig. 14 DRT spectra for SOFCs with (a and b) steam reforming (SR) of MeOH, (c and d) SR of DME, and (e and f) partial oxidation reforming (POx) of DME before and after the durability tests at 700 °C.



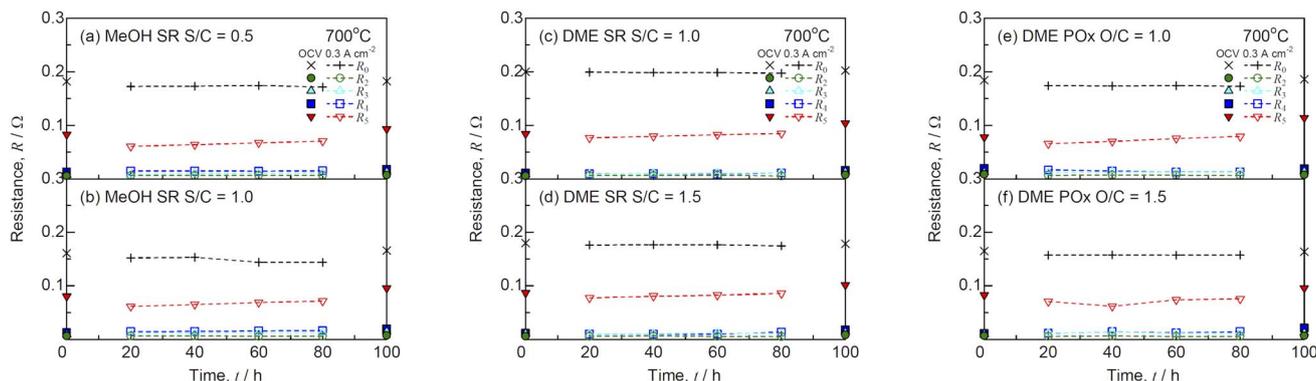


Fig. 15 Time courses of resistances for SOFCs with (a and b) steam reforming (SR) of MeOH, (c and d) SR of DME, and (e and f) partial oxidation reforming (POx) of DME during the durability test at a constant current density of  $0.3 \text{ A cm}^{-2}$  and  $700^\circ\text{C}$ .

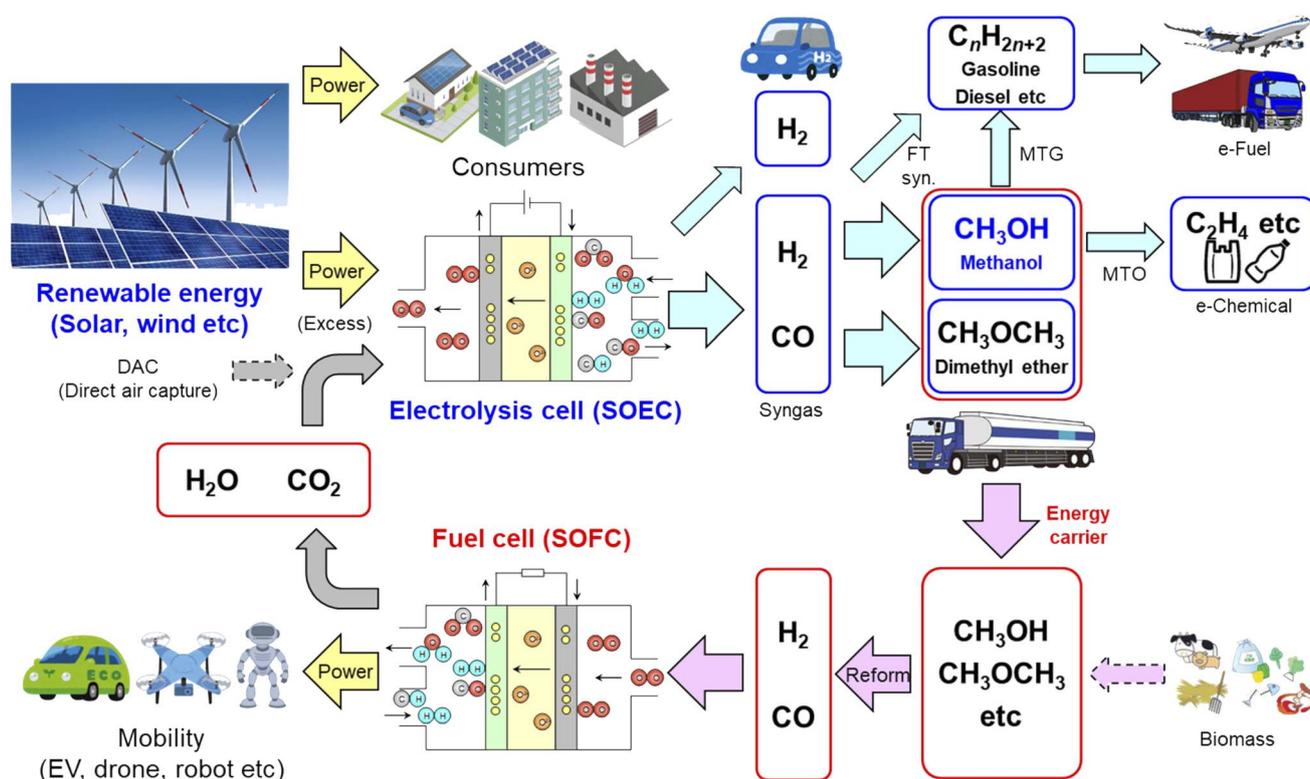


Fig. 16 Schematic of carbon-neutral energy system using co-electrolysis SOECs and internal reforming SOFCs.

power are supplied to consumers such as houses, buildings, and plants. The excess electric power is converted into chemical energy. Hydrogen is produced using SOECs in addition to alkaline and PEM electrolysis, and can be used in stationary fuel cells and fuel cell vehicles.<sup>7</sup> Syngas is also produced using SOECs, and converted into MeOH and DME.<sup>20–22,28</sup> Synthetic gasoline, kerosene, and diesel (e-fuel) are produced from syngas *via* FT synthesis, and can be used for airplanes, vessels, and large vehicles.<sup>18,19</sup> Methanol can be converted into gasoline (MTG; methanol to gasoline), olefin (MTO; methanol to olefins) such as ethylene and propylene, and aromatics (MTA; methanol to aromatics) such as benzene, toluene, and xylene. MTO widely

contributes to carbon-neutral manufacturing processes for commodity plastics (e-chemical), whereas the raw materials of engineering plastics such as polycarbonate and polyurethane are directly synthesized from  $\text{CO}_2$ .<sup>41</sup> MeOH is a liquid at ordinary temperatures and pressures, and DME is easily liquified at 0.6 MPa. Therefore, MeOH and DME are promising energy carriers and fuels for internal reforming SOFCs. Biomass is also used as a fuel in SOFCs after desulfurization and reforming. Although  $\text{CO}_2$  is exhausted from internal reforming SOFCs, it can be used for co-electrolysis SOECs. Mass and heat balance should be considered to achieve carbon neutrality. Thus, co-



electrolysis SOECs and internal reforming SOFCs significantly contribute to achieving a carbon-neutral society.

## 4. Conclusion

In the present work, H<sub>2</sub>O/CO<sub>2</sub> co-electrolysis SOECs and internal reforming SOFCs using MeOH and DME were investigated using negatrode-supported microtubular cells with the same cell configuration. The co-electrolysis performance decreased with increasing CO<sub>2</sub> concentration in the input gas owing to an increase in polarization resistance related to gas diffusion process at the negatrode. The diffusion of CO/CO<sub>2</sub> mixture was more difficult than that of H<sub>2</sub>/H<sub>2</sub>O mixture in the SOEC negatrode. The gas diffusion resistance increased with time during co-electrolysis SOECs at H<sub>2</sub>O/CO<sub>2</sub> = 2 and 700 °C. At various H<sub>2</sub>O/CO<sub>2</sub> ratios, the faradaic efficiencies exceeded 96%, and the H<sub>2</sub>/CO ratios of the product gases were almost identical to the H<sub>2</sub>O/CO<sub>2</sub> ratios of the input gases. During the durability test at H<sub>2</sub>O/CO<sub>2</sub> = 2, the faradaic efficiency and H<sub>2</sub>/CO ratio of the product gas were stable for 100 h.

MeOH and DME can be produced industrially from syngas with an H<sub>2</sub>/CO ratio of 2. The *I-V* characteristics were similar during internal reforming of MeOH and DME. The polarization resistances related to charge transfer and gas diffusion processes increased with time during internal reforming of MeOH and DME. Thus, the gas diffusion resistance should be decreased to improve the initial performance and durability of co-electrolysis SOECs and internal reforming SOFCs. In thermodynamic equilibrium, carbon is deposited at S/C < 0.3 for MeOH, and at S/C < 0.8 and O/C < 0.9 for DME during steam and partial oxidation reforming, respectively. The internal reforming SOFCs successfully operated for 100 h at S/C ≥ 0.5 for MeOH, S/C ≥ 1.0 and O/C ≥ 1.0 for DME. Therefore, MeOH and DME syntheses from syngas *via* co-electrolysis SOECs and power generation *via* internal reforming SOFCs using MeOH and DME will contribute to achieving a carbon-neutral society.

## Author contributions

Hirofumi Sumi: conceptualization, methodology, resources, data curation, investigation, writing – original draft preparation.

## Conflicts of interest

There are no conflicts to declare.

## References

- 1 K. Eguchi, H. Kojo, T. Takeguchi, R. Kikuchi and K. Sasaki, Fuel Flexibility in Power Generation by Solid Oxide Fuel Cells, *Solid State Ionics*, 2002, **152–153**, 411–416.
- 2 H. Sumi, K. Ukai, Y. Mizutani, H. Mori, C.-J. Wen, H. Takahashi and O. Yamamoto, Performance of Nickel-Scandia-Stabilized Zirconia Cermet Anodes for SOFCs in 3% H<sub>2</sub>O-CH<sub>4</sub>, *Solid State Ionics*, 2004, **174**, 151–156.
- 3 H. Sumi, T. Yamaguchi, K. Hamamoto, T. Suzuki and Y. Fujishiro, Impact of Direct Butane Microtubular Solid Oxide Fuel Cells, *J. Power Sources*, 2012, **220**, 74–78.
- 4 H. Sumi, T. Yamaguchi, H. Shimada, Y. Fujishiro and M. Awano, Internal Partial Oxidation Reforming of Butane and Steam Reforming of Ethanol for Anode-supported Microtubular Solid Oxide Fuel Cells, *Fuel Cells*, 2017, **17**, 875–881.
- 5 H. Sumi, S. Nakabayashi, T. Kawada, Y. Uchiyama, N. Uchiyama and K. Ichihara, Demonstration of SOFC Power Sources for Drones (UAVs; Unmanned Aerial Vehicles), *ECs Trans.*, 2019, **91**(1), 149–157.
- 6 Y. Shiratori, T. Ijichi, T. Oshima and K. Sasaki, Internal Reforming SOFC Running on Biogas, *Int. J. Hydrogen Energy*, 2010, **35**, 7905–7912.
- 7 S. A. Grigoriev, V. N. Fateev, D. G. Bessarabov and P. Millet, Current Status, Research Trends, and Challenges in Water Electrolysis Science and Technology, *Int. J. Hydrogen Energy*, 2020, **45**, 26036–26058.
- 8 R. Küngas, Review-Electrochemical CO<sub>2</sub> Reduction for CO Production: Comparison of Low- and High-temperature Electrolysis Technologies, *J. Electrochem. Soc.*, 2020, **167**, 044508.
- 9 A. A. Olajire, CO<sub>2</sub> Capture and Separation Technologies for End-of-pipe Applications - A Review, *Energy*, 2010, **35**, 2610–2628.
- 10 F. Bisotti, K. A. Hoff, A. Mathisen and J. Hovland, Direct Air capture (DAC) Deployment: A Review of the Industrial Deployment, *Chem. Eng. Sci.*, 2024, **283**, 119416.
- 11 Y. Kohno, M. Kanakubo, M. Iwaya, Y. Yamato and T. Makino, Ionic Liquid Mixtures for Direct Air Capture: High CO<sub>2</sub> Permeation Driven by Superior CO<sub>2</sub> Absorption with Lower Absolute Enthalpy, *ACS Omega*, 2022, **7**, 42155–42162.
- 12 C. Graves, S. D. Ebbesen and M. Mogensen, Co-electrolysis of CO<sub>2</sub> and H<sub>2</sub>O in Solid Oxide Cells: Performance and Durability, *Solid State Ionics*, 2011, **192**, 398–403.
- 13 S. D. Ebbesen, R. Knibbe and M. Mogensen, Co-electrolysis of Steam and Carbon Dioxide in Solid Oxide Cells, *J. Electrochem. Soc.*, 2012, **159**, F482–F489.
- 14 D. M. A. Duenas, M. Riedel, M. Riegraf, R. Costa and K. A. Friedrich, High Temperature Co-electrolysis for Power-to-X, *Chem. Ing. Tech.*, 2020, **92**, 45–52.
- 15 C. Xi, J. Sang, A. Wu, J. Yang, X. Qi, W. Guan, J. Wang and S. C. Singhal, Electrochemical Performance and Durability of Flat-tube Solid Oxide Electrolysis Cells for H<sub>2</sub>O/CO<sub>2</sub> Co-electrolysis, *J. Catal.*, 2021, **404**, 174–186.
- 16 E. Ioannidou, M. Chavani, S. G. Neophytides and D. K. Niakolas, Effect of the  $P_{H_2O}/P_{CO_2}$  and  $P_{H_2}$  on the Intrinsic Electro-catalytic Interactions and the CO Production Pathway on Ni/GDC during Solid Oxide H<sub>2</sub>O/CO<sub>2</sub> Co-electrolysis, *Int. J. Hydrogen Energy*, 2022, **47**, 10166–10174.
- 17 C. Mebrahtu, M. Nohl, L. Dittrich, S. R. Foit, L. G. J. de Haart, R.-A. Eichel and R. Palkovits, Integrated Co-Electrolysis and Syngas Methanation for the Direct Production of Synthetic Natural Gas from CO<sub>2</sub> and H<sub>2</sub>O, *ChemSusChem*, 2021, **14**, 2295–2302.



- 18 M. E. Dry, The Fischer-Tropsch Process: 1950-2000, *Catal. Today*, 2002, **71**, 227–241.
- 19 N. Koizumi, T. Mochizuki and M. Yamada, Preparation of Highly Active Catalysts for Ultra-clean Fuels, *Catal. Today*, 2009, **141**, 34–42.
- 20 G. Bozzano and F. Manenti, Efficient Methanol Synthesis: Perspectives, Technologies and Optimization Strategies, *Prog. Energy Combust. Sci.*, 2016, **56**, 71–105.
- 21 S. Mbatha, R. C. Everson, N. M. Musyoka, H. W. Langmi, A. Lanzini and W. Brilman, Power-to-methanol Process: A review of Electrolysis, Methanol Catalysts, Kinetics, Reactor Designs and Modelling, Process Integration, Optimisation, and Techno-economics, *Sustain. Energy Fuels*, 2021, **5**, 3490–3569.
- 22 Y. Choi, K. Futagami, T. Fujitani and J. Nakamura, The Difference in the Active Sites for CO<sub>2</sub> and CO Hydrogenations on Cu/ZnO-based Methanol Synthesis Catalysts, *Catal. Lett.*, 2001, **73**, 27–31.
- 23 A. S. Arico, S. Srinivasan and V. Antonucci, DMFCs: From Fundamental Aspects to Technology Development, *Fuel Cells*, 2001, **1**, 133–161.
- 24 T. Kim, K. Ahn, J. M. Vohs and R. J. Gorte, Deactivation of Ceria-based SOFC Anodes in Methanol, *J. Power Sources*, 2007, **164**, 42–48.
- 25 M. Liu, R. Peng, D. Dong, J. Gao, X. Liu and G. Meng, Direct Liquid Methanol-fueled Solid Oxide Fuel Cell, *J. Power Sources*, 2008, **185**, 188–192.
- 26 M. Lo Faro, A. Stassi, V. Antonucci, V. Modafferi, P. Frontera, P. Antonucci and A. S. Arico, Direct Utilization of Methanol in Solid Oxide Fuel Cells: An Electrochemical and Catalytic Study, *Int. J. Hydrogen Energy*, 2011, **36**, 9977–9986.
- 27 Z. Gao, R. Raza, B. Zhu and Z. Mao, Development of Methanol-fueled Low-temperature Solid Oxide Fuel Cells, *Int. J. Energy Res.*, 2011, **35**, 690–696.
- 28 K. Omata, Y. Watanabe, T. Umegaki, G. Ishiguro and M. Yamada, Low-pressure DME Synthesis with Cu-based Hybrid Catalysts using Temperature-gradient Reactor, *Fuel*, 2002, **81**, 1605–1609.
- 29 S. Wang, T. Ishihara and Y. Takita, Dimethyl Ether Fueled Intermediate Temperature SOFC Using LaGaO<sub>3</sub>-Based Perovskite Electrolytes, *Electrochem. Solid-State Lett.*, 2002, **5**, A177–A180.
- 30 C. Su, W. Wang, H. Shi, R. Ran, H. J. Park, C. Kwak and Z. Shao, Reducing the Operation Temperature of a Solid Oxide Fuel Cell Using a Conventional Nickel-based Cermet Anode on Dimethyl Ether Fuel through Internal Partial Oxidation, *J. Power Sources*, 2011, **196**, 7601–7608.
- 31 M. A. Laguna-Bercero, Recent Advances in High Temperature Electrolysis using Solid Oxide Fuel Cells: A Review, *J. Power Sources*, 2012, **203**, 4–16.
- 32 K. Hosoi, H. Hagiwara, S. Ida and T. Ishihara, La<sub>0.8</sub>Sr<sub>0.2</sub>FeO<sub>3-δ</sub> as Fuel Electrode for Solid Oxide Reversible Cells Using LaGaO<sub>3</sub>-Based Oxide Electrolyte, *J. Phys. Chem. C*, 2016, **120**, 16110–16117.
- 33 H. Uchida, H. Nishino, P. Puengjinda and K. Kakinuma, Remarkably Improved Durability of Ni-Co Dispersed Samaria-Doped Ceria Hydrogen Electrodes by Reversible Cycling Operation of Solid Oxide Cells, *J. Electrochem. Soc.*, 2020, **167**, 134516.
- 34 H. Schichlein, A. C. Muller, M. Voigts, A. Krugel and E. Ivers-Tiffée, Deconvolution of Electrochemical Impedance Spectra for the Identification of Electrode Reaction Mechanisms in Solid Oxide Fuel Cells, *J. Appl. Electrochem.*, 2002, **32**, 875–882.
- 35 A. Leonide, B. Rüger, A. Weber, W. A. Meulenber and E. Ivers-Tiffée, Evaluation and Modeling of the Cell Resistance in Anode-supported Solid Oxide Fuel Cells, *J. Electrochem. Soc.*, 2010, **157**, B234–B239.
- 36 H. Sumi, H. Shimada, Y. Yamaguchi, T. Yamaguchi and Y. Fujishiro, Degradation Evaluation by Distribution of Relaxation Times Analysis for Microtubular Solid Oxide Fuel Cells, *Electrochim. Acta*, 2020, **339**, 135913.
- 37 J. Weese, A Reliable and Fast Method for the Solution of Fredholm Integral Equations of the First Kind Based on Tikhonov Regularization, *Comput. Phys. Commun.*, 1992, **69**, 99–111.
- 38 E. Ivers-Tiffée and A. Weber, Evaluation of Electrochemical Impedance Spectra by the Distribution of Relaxation Times, *J. Ceram. Soc. Jpn.*, 2017, **125**, 193–201.
- 39 B. A. Boukamp, A Linear Kronig-Kramers Transform Test for Immittance Data Validation, *J. Electrochem. Soc.*, 1995, **142**, 1885–1894.
- 40 M. Schönleber, D. Klotz and E. Ivers-Tiffée, A Method for Improving the Robustness of Linear Kramers-Kronig Validity Tests, *Electrochim. Acta*, 2014, **131**, 20–27.
- 41 W. S. Putro, A. Ikeda, S. Shigeyasu, S. Hamura, S. Matsumoto, V. Y. Lee, J.-C. Choi and N. Fukaya, Sustainable Catalytic Synthesis of Diethyl Carbonate, *ChemSusChem*, 2021, **14**, 842–846.

