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In situ X-ray diffraction computed tomography studies examining the thermal and chemical stabilities of working $\text{Ba}_{0.5}\text{Sr}_{0.5}\text{Co}_{0.8}\text{Fe}_{0.2}\text{O}_{3-\delta}$ membranes during oxidative coupling of methane[†]

Dorota Matras, ^{ab} Antonis Vamvakarios, ^{bcd} Simon D. M. Jacques, ^c Vesna Middelkoop, ^e Gavin Vaughan, ^f Miren Agote Aran, ^{bcd} Robert J. Cernik^a and Andrew M. Beale ^{bcd}

In this study we present the results from two *in situ* X-ray diffraction computed tomography experiments of catalytic membrane reactors (CMRs) using $\text{Ba}_{0.5}\text{Sr}_{0.5}\text{Co}_{0.8}\text{Fe}_{0.2}\text{O}_{3-\delta}$ (BSCF) hollow fibre membranes and $\text{Na}-\text{Mn}-\text{W}/\text{SiO}_2$ catalyst during the oxidative coupling of methane (OCM) reaction. The negative impact of CO_2 , when added to the inlet gas stream, is seen to be mainly related to the C_{2+} yield, while no evidence of carbonate phase(s) formation is found during the OCM experiments. The main degradation mechanism of the CMR is suggested to be primarily associated with the solid-state evolution of the BSCF phase rather than the presence of CO_2 . Specifically, *in situ* XRD-CT and post-mortem SEM/EDX measurements revealed a collapse of the cubic BSCF phase and subsequent formation of secondary phases, which include needle-like structures and hexagonal $\text{Ba}_6\text{Co}_4\text{O}_{12}$ and formation of a BaWO_4 layer, the latter being a result of chemical interaction between the membrane and catalyst materials at high temperatures.

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1. Introduction

Over the past decades, the potential of membrane reactors towards a cleaner and more sustainable future has become more apparent.^{1–3} Catalytic membrane reactors (CMRs) are able to simultaneously perform separation and reaction processes offering a unique opportunity to increase the overall system efficiency and decrease the energy consumption in a single unit. More specifically, the application of oxygen permeable membranes with mixed ionic and electronic conductivity (MIEC) properties is an alternative technology for oxygen

separation from air.^{4–6} It offers both reduced costs and high oxygen permeation flux, and more importantly it can be directly integrated with catalytic reactor systems.^{7–10} The use of perovskite membranes in the oxidative coupling of methane (OCM) process^{11–15} is believed to provide a more uniform distribution of oxygen along the axial direction when compared to conventional fixed bed reactors. This is expected to directly influence the C_{2+} yield as the methane can be activated by the surface oxygen ionic species and the contribution of gas phase homogenous combustion reactions can be decreased.¹⁶

Wang *et al.*¹⁷ studied the OCM reaction using a catalytic membrane reactor with a $\text{Ba}_{0.5}\text{Sr}_{0.5}\text{Co}_{0.8}\text{Fe}_{0.2}\text{O}_{3-\delta}$ tubular membrane and a $\text{La}-\text{Sr}/\text{CaO}$ catalyst packed-bed. Both, the packed-bed catalytic membrane reactor and the catalytic membrane reactor (*i.e.* membrane only) configurations were tested using similar OCM conditions. Although both configurations showed similar performance for CH_4 conversion and C_{2+} yield, the packed-bed catalytic membrane reactor led to significantly higher $\text{C}_2\text{H}_4/\text{C}_2\text{H}_6$ ratio (12.5 as opposed to 1 for the catalytic membrane reactor). Czuprat *et al.*¹⁸ investigated the $\text{Na}-\text{Mn}-\text{W}/\text{SiO}_2$ catalyst packed inside the hollow fibre $\text{BaCo}_x\text{Fe}_y\text{Zr}_z\text{O}_{3-\delta}$ ($x + y + z = 1$) membrane. A C_{2+} yield of 17% and $\text{C}_2\text{H}_4/\text{C}_2\text{H}_6$ ratio of 4 was obtained for a CH_4 conversion of

^a School of Materials, University of Manchester, Manchester, Lancashire M13 9PL, UK. E-mail: matras.dorota@gmail.com

^b Research Complex at Harwell, Harwell Science and Innovation Campus, Rutherford Appleton Laboratory, Didcot, Oxon, OX11 0FA, UK

^c Finden Limited, Merchant House, 5 East St Helen Street, Abingdon, OX14 5EG, UK. E-mail: simon@finden.ac.uk

^d Department of Chemistry, University College London, 20 Gordon Street, London, WC1H 0AJ, UK. E-mail: andrew.beale@ucl.ac.uk

^e Sustainable Materials Management, Flemish Institute for Technological Research, VITO NV, Boeretang 200, Mol, Belgium

^f ESRF – The European Synchrotron, Grenoble, 38000, France

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35% (using a pure stream of air on the core/feed side). In the study of Bhatia *et al.*¹⁹ three configurations were investigated for the OCM reaction: packed-bed reactor, packed-bed catalytic membrane reactor and catalytic membrane reactor with coated Na–Mn–W/SiO₂ catalyst. The latter configuration was found to perform the best among the three configurations, with a C₂₊ yield of 35% and C₂H₄/C₂H₆ ratio of 1.4. The changes in the catalytic membrane material were analysed postreaction with XRD and SEM analysis; catalyst components were found to migrate towards the reaction surface, whereas cation species from the membrane migrated towards the oxygen-enriched surface.

In this work we investigate a Ba_{0.5}Sr_{0.5}Co_{0.8}Fe_{0.2}O_{3- δ} (BSCF) hollow fibre membrane during the oxidative coupling of methane; this perovskite is known to have excellent oxygen permeability^{7,20-22} at operating temperatures required for this reaction (~ 2 S mL min⁻¹ cm⁻²²³). However, there are still several challenges related to the long-term phase stability and chemical poisoning of this material that prevent its application under real industrial conditions. More precisely, at intermediate temperatures (*i.e.* below 850 °C) the cubic BSCF phase has been observed to decompose to secondary Ba and Co hexagonal phases with a simultaneous decrease in oxygen permeation.²⁴⁻²⁹ The reason behind the instability of the cubic phase has been attributed to a change in the oxidation state of Co upon exposure to air below 850 °C. According to the Goldschmidt tolerance factor,³⁰ the cubic structure is not able to accommodate Co with higher oxidation states³¹ and thus the more stable hexagonal phase is expected to form. The formation of the secondary hexagonal phase was identified to take place at the grain boundaries from the feed side (air exposed side), but over a long duration experiment (*i.e.* 1000 s of hours of operation) this phase can also be identified in the grains.³²⁻³⁴ The high internal stress resulting from this phenomenon was suggested to be responsible for the formation of cracks in the membrane reactor.³⁵

The second major challenge is related to the interaction of the BSCF material with CO₂. In many studies, the formation of BaCO₃ (or Ba_xSr_{1-x}CO₃) has been reported to have a detrimental effect on the oxygen permeation.^{36,37} A thin layer of this carbonate phase forming at the surface of the membrane is considered to act as a barrier for the oxygen ion transport. In a study by Engels *et al.*,³⁸ 15% of CO₂ in the permeated side gradually decreased the initial oxygen permeation flux by 80% after 200 min of operation. Although the effect of CO₂ was found to be reversible, as the initial oxygen permeation flux could be restored when changing the composition of the sweep side to pure He, minor structural changes in the membrane were induced during this process; these may be considered important for the long term stability of BSCF membranes. Finally, as recently reported by Vamvakarios *et al.*,^{39,40} Ba-containing membranes may also be susceptible to chemical poisoning by WO₄²⁻ species. During the OCM reaction, a thin layer of BaWO₄ was found to form at the interface between a Na–Mn–W/SiO₂ catalyst and a hollow fibre BaCo_xFe_yZr_zO_{3- δ} (BCFZ) membrane. Although there was no direct change in the

reactor performance, the loss of catalyst components coupled with the formation of the stable BaWO₄ layer was proposed to have a significant impact on the long-term performance of this integrated reactor system.

The aim of this work was to assess the aforementioned issues and their implication on the performance of working catalytic membrane reactors for the oxidative coupling of methane reaction. We employed the X-ray diffraction computed tomography (XRD-CT) technique^{41,42} to investigate the evolution of the solid-state chemistry in the BSCF membrane in two experiments. The first experiment involved exposing the BSCF membrane to 20% CO₂/CH₄ gas stream (permeate side) for 11 h. In the second experiment, the Na–Mn–W/SiO₂ catalyst was placed inside the BSCF membrane and the packed-bed catalytic membrane reactor was tested for the OCM reaction with different concentrations of CO₂ in the CH₄ stream (permeate side). The solid-state changes observed from the XRD-CT data are discussed and compared with laboratory mass spectrometry and post-reaction SEM/EDX measurements.

2. Materials and methods

2.1. Preparation of hollow fibre BSCF membrane

The BSCF perovskite hollow membranes were supplied by the Flemish Institute for Technological Research (VITO), prepared from the BSCF mixed ionic-electronic conducting perovskite powder obtained from CerPoTech (Norway). Membranes were prepared using the spinning and phase inversion technique previously reported in the work of Di Felice *et al.*²³ The dimensions of the BSCF membranes used for both experiments were: 3.6–3.7 mm outside diameter and 2.8–2.9 mm inside diameter.

2.2. *In situ* XRD-CT measurements at ID15A, ESRF

XRD-CT measurements were performed at ID15A beamline at the ESRF⁴³ using a monochromatic beam of 95 keV with a size of 40 μm × 90 μm (H × V). For both experiments BSCF membranes were placed inside a quartz reactor (6 mm outside diameter and 4 mm inside diameter). A schematic representation of the experimental setup can be found in Fig. S1 (ESI†). Two different experiments were performed: (1) with catalytic membrane reactor and (2) with packed-bed catalytic membrane reactor using 60 mg of Na–Mn–W/SiO₂ catalyst placed inside the BSCF membrane and supported between glass wool. During the first experiment (with pure membrane reactor) the sample was scanned using 150 translation steps (with 40 μm step) and 100 angular steps (with 1.8° step), whereas during the experiment with packed-bed catalytic membrane reactor sample was scanned using 140 translation steps (with 40 μm step) and 120 angular steps (1.5° step). Diffraction patterns were collected using a PILATUS3 X CdTe (Dectris) area detector. The calibration of the detector was performed using a CeO₂ NIST standard. Every collected 2D diffraction image was converted to 1D powder diffraction patterns using PyFAI software⁴⁴ using the nDTomo software⁴⁵ with implemented trimmed mean filter (20%) to remove the artefacts due to hot spots of (single)



crystalline materials.⁴⁶ The reconstructed images were obtained using the filtered back projection algorithm. For both experiments the XRD-CT data were corrected for the self-absorption using a zero-order approximation. The detailed description of the self-absorption correction strategy can be found elsewhere.⁴⁷

2.3. *In situ* OCM experiments at ID15A, ESRF

Fig. S2 (ESI[†]) shows the photograph of the experimental setup used during the *in situ* measurements. In both experiments two separate gas streams were delivered to the reactor cell. During the temperature ramp up to 820 °C, the sample was heated under atmosphere of 50 sccm Ar (inner side gas stream) and 50 sccm air (outer side gas stream) with a temperature ramp of 20 °C min⁻¹. Pure gases of air, CH₄, CO₂ and Ar were delivered to the reactor by mass flow controllers (Brooks). The OCM reaction was performed at atmospheric pressure and the outlet gases were analysed by mass spectrometry using an Ecosys portable mass spectrometer. The sample was heated using an in-house developed box furnace. In the case of the first experiment, we investigated the long-term stability (11 h) of the BSCF membrane during the OCM reaction, using a CH₄ stream co-fed with 20% of CO₂. The outer side gas stream delivered 50 sccm of air and the inner side gas stream delivered 80 sccm of CH₄ and 20 sccm of CO₂. In the case of the second experiment, the solid-state interaction between the membrane and catalyst materials was investigated, testing different compositions of OCM reaction mixtures (CH₄ stream also co-fed with CO₂). The outer side gas stream delivered 50 sccm of air and the inner side gas stream delivered OCM mixture co-feed with 0, 10, 20, 50 and 100% of CO₂ in CH₄/CO₂ mixture, keeping the 100 sccm as total flow of inner side gas stream. The experimental protocol details for both experiments can be found in the ESI[†] (Fig. S3 and S4).

2.4. Analysis of reconstructed diffraction patterns

The summed diffraction patterns from each XRD-CT dataset were analysed with the X'Pert HighScore Plus software to identify which phases were present and create the model necessary for the Rietveld refinement. The identified phases together with crystallographic information are presented in Table S1 (ESI[†]). The analysed diffraction patterns were extracted only from pixels where the membrane or catalyst particles were present (the regions of air, reactor and voids between particles were masked). The batch refinement analysis using the reconstructed diffraction patterns was performed using the software Topas v5.⁴⁸ For both measurements, individual reflections of the cubic perovskite (space group $Pm\bar{3}m$) were fitted, in order to follow the relative changes in peak intensities (referred as hkl _phase in Topas). In addition, the application of a 20% trimmed mean filter for data correction,⁴⁶ especially in the case of second experiment, affected the relative intensity of each peak (*i.e.* spotty data). Therefore, Pawley instead of Rietveld refinement analysis was used as a means to overcome the intensity mismatch problem. In addition, due to the significant thermal expansion of the ceramic material during the temperature ramp, the relative changes in peak intensities were compared only using the measurements performed

at high temperatures. The scattering signal originating from catalyst particles (low absorbing and weakly scattering material) was highly altered by the membrane signal (highly absorbing and strongly scattering material) and therefore it was not possible to perform refinement analysis using these diffraction patterns. The distribution of each component is based on the summed intensity of a corresponding diffraction peak in the XRD-CT data.

2.5. Laboratory mass spectrometry and gas chromatography measurements

Complementary OCM experiments to the *in situ* experiments performed at the ID15A beamline were also performed in the laboratory. In contrast to the *in situ* measurements, the reactor used in the laboratory measurements was closed from both ends; two inlets streams and two outlets streams were connected to the reactor using nonporous alumina rods. The sample was placed inside a quartz reactor, itself placed inside a horizontal tubular furnace (Carbolite). In the case of the packed-bed catalytic membrane reactor, 68 mg of Na–Mn–W/SiO₂ catalyst was placed inside the BSCF membrane supported by quartz wool.

A schematic representation of the experimental setup used in the laboratory measurements is presented in Fig. S5 (ESI[†]). Gases: 20% of O₂/He, 50% of CH₄/Ar, CO₂ and N₂ were delivered to the reactor by mass flow controllers (Brooks). The OCM reaction was performed at atmospheric pressure and the outflow gases were analysed by mass spectrometry, using an Ecosys portable mass spectrometer, and by gas chromatography, using the SRI 8610C Gas Chromatograph (Molecular Sieve 13X and Haysep 6D columns) equipped with FID and TCD detectors. The inner side gas stream was cooled with an ice bath to collect the produced water (in the case of the experiment with packed-bed catalytic membrane reactor). The experimental setup allowed, by adjusting accordingly the valves, switching between the two streams and choosing which one was probed by mass spectrometry. Prior to the experiments, the membrane sealing was tested by measuring the gas flow from the sweep outlet 2 (outer side gas stream) while using only an inert gas to the reactor through inlet 1 (inner side gas stream). This way the sealing of both membrane sections was tested. During the temperature ramp, only the gas stream flow outlet 2 was probed in order to ensure the sealing did not fail during the high temperature treatment. Before introducing the OCM reaction mixture the valves were adjusted to probe the outlet 1 (inner side gas stream). The experimental protocol for both experiments can be found in Tables S2 and S3 (ESI[†]).

3. Results

3.1. Catalytic membrane reactor

The initial distribution of the membrane components is presented in Fig. 1. The membrane composition was found to be non-uniform, as two distinct regions can be observed in the acquired cross section. On the left side, the majority of the material consisted of cubic BSCF perovskite, whereas the right



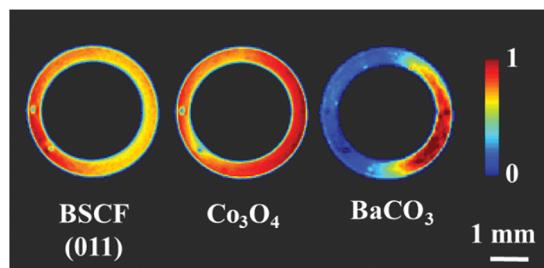


Fig. 1 Distribution of the BSCF, Co_3O_4 and BaCO_3 phases in the as-prepared membrane at room temperature.

side contained Co_3O_4 , BaCO_3 and an additional phase (Fig. S6 and S7, ESI[†]). This unusual distribution of membrane materials was not a measurement artefact and it was most likely formed during the membrane preparation. However, it is important to note that the membrane primarily consists of the BSCF phase while the other phases are present in minor quantities (Fig. S7, ESI[†]).

Fig. 2 shows the distribution of the membrane components (Co_3O_4 and $\text{Ba}_6\text{Co}_4\text{O}_{12}$) as well as the distribution of the cubic perovskite (the main (011) reflection) during the high temperature scans under Ar and under OCM reaction conditions. It is important to note that the initially present BaCO_3 decomposed during the temperature ramp and no other carbonate phase formed during the experiment. The results from the refinement analysis showed that the BSCF phase was not stable under OCM operating conditions (Fig. S7, ESI[†]). More precisely, after around 3 h of operation, a secondary hexagonal phase, identified as $\text{Ba}_6\text{Co}_4\text{O}_{12}$, started to form at the region where the cubic perovskite was initially present in the fresh material. As mentioned previously, the instability of the cubic perovskite is expected to occur during operation of the membrane at intermediate temperatures. Different compositions and morphologies of this secondary phase have been previously reported^{24,33,35} however the hexagonal phase is expected to contain mainly Ba and Co. During the OCM reaction the intensities of all BSCF reflections were seen to gradually decrease, whereas the Co_3O_4 remained stable for the duration of the experiment.

Although all BSCF reflections are seen to gradually decrease with time (Fig. 3), an unusual behaviour can be observed between the 1st and 2nd XRD-CT scan; the intensity of the (011), (111), (002), (211) and (022) reflections increased whereas the intensity of the (001) and (021) reflections decreased. In addition, after the introduction of the OCM reaction mixture, the lattice parameter of the cubic perovskite decreased from 4.015 to 4.013 Å (Fig. S8, ESI[†]). This observation can be explained by the contraction of the lattice parameter, through the oxidation of the transition metals occupying the perovskite B site. This would also suggest a decrease in oxygen vacancies in order to maintain the charge neutrality of the structure.⁴⁹ The gradual decrease in lattice parameter during the OCM reaction, from 4.013 to 4.011 Å is most likely caused by the change in the structure of the cubic perovskite; the formation of hexagonal $\text{Ba}_6\text{Co}_4\text{O}_{12}$ phase resulted in the decrease of Ba and Co in the resulting cubic perovskite, which is in agreement with previously reported studies.^{34,50}

In order to rationalise the subtle changes in the peak intensities, further analysis was performed focusing on how the structural changes in the cubic BSCF (*i.e.* occupancy of each element at A and B sites, oxygen occupancy) affect the intensity of certain reflections. The simulation of BSCF diffraction patterns with different chemical composition can be found in the ESI[†] (Section S3, Simulation of BSCF structure). Considering the relative changes in the simulated data, we conclude that the changes in the BSCF structure between HT/Ar and OCM XRD-CT scan were due to the increase in oxygen occupancy, which corresponds to a decrease in oxygen vacancies (Table S4, ESI[†]). Although the changes in the intensity of the (011) and (211) reflections are contradictory to the results of the simulated data, the relative changes in the simulated data are minor and therefore it is not possible to rule out that the changes in intensity may also be due to changes in the amounts of Co and Ba. The decrease in oxygen permeability is expected to occur during the introduction of the OCM reaction mixture (changing gas environment from He to CH_4), as the CO_2 (produced from combustion) reacts with the membrane forming a layer of BaCO_3 . However, the XRD-CT measurements (X-ray beam size of 40 $\mu\text{m} \times 90 \mu\text{m}$) did not detect the presence of any carbonate phases. The relative changes in the BSCF structure though is a

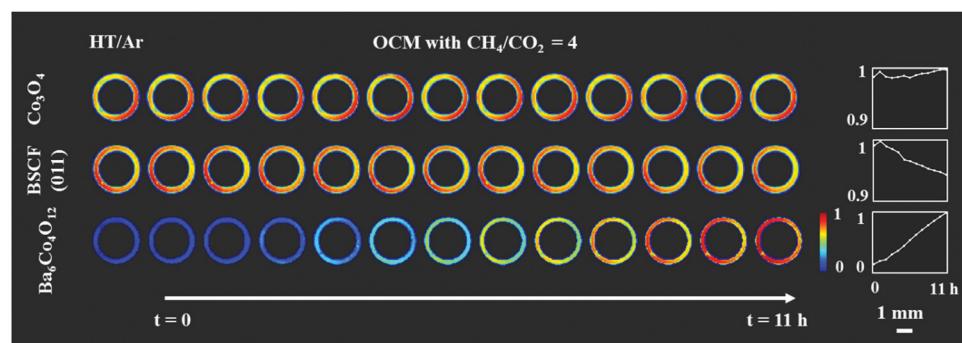


Fig. 2 Distribution of membrane components during the high temperature XRD-CT scan and OCM reaction. Plots on the right correspond to changes in the normalised scale factors of Co_3O_4 and $\text{Ba}_6\text{Co}_4\text{O}_{12}$ and the peak intensity of the BSCF (011) reflection.

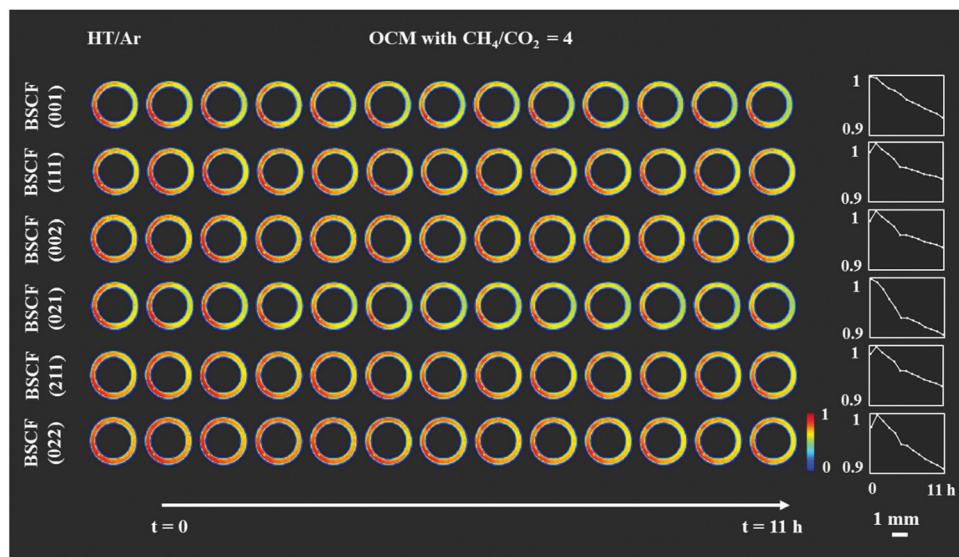


Fig. 3 Distribution of membrane components – reflections of cubic BSCF phase during the high temperature XRD-CT scan and OCM reaction. Plots on the right correspond to changes in the normalised peak intensities for the various BSCF reflections.

clear indication of the oxygen permeation changing as a function of the imposed operating conditions. As the perovskite was found to be unstable during the OCM reaction conditions, the decrease in intensity of all BSCF reflections during the experiment is attributed to the ongoing decomposition of this phase. The decomposition of BSCF during operation as well as the formation of Co_3O_4 , BaCO_3 and the unidentified phase(s) during the preparation indicate inherent problems associated with the chemical stability of the perovskite phase.

3.2. Catalytic membrane reactor with Na–Mn–W/SiO₂ packed-bed

The distribution of the fresh membrane components at ambient conditions is presented in Fig. 4. Initially, the membrane consisted primarily of the BSCF perovskite while minor quantities of CoO and Co_3O_4 were also present. It is important to note that the BaCO_3 phase seen in the first experiment was not observed here (Fig. S14, ESI[†]). During the temperature ramp performed under air/Ar, the quantity of CoO significantly decreased which could be explained by the oxidation of Co^{2+} to Co^{3+} in the presence of air at high temperatures.

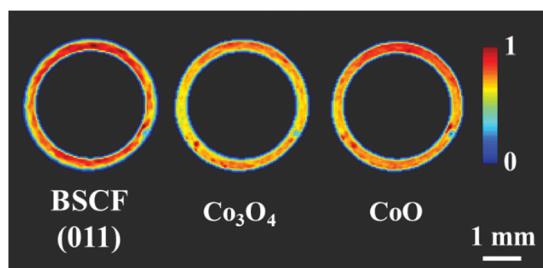


Fig. 4 Distribution of the BSCF, Co_3O_4 and CoO phases in the as-prepared membrane at ambient conditions.

The disappearance of CoO and simultaneous increase in Co_3O_4 is clearly visible by the relative changes in the diffraction patterns collected at room temperature and operating temperature (Fig. S14, ESI[†]).

The distribution of the membrane components as well as the relative changes during the OCM reaction are shown in Fig. 5. After the introduction of the OCM reaction mixture, the CoO was found to completely disappear from the membrane material, whereas the distribution and quantity of Co_3O_4 , after the initial formation/growth from the CoO, was found to remain stable until the end of the experiment. Regarding the BSCF phase, the relative changes in its primary reflection ((011) at $2\text{Th} = 2.6^\circ$ in Fig. S15, ESI[†]) showed that the quantity of the perovskite gradually decreased with the greatest reduction observed after the introduction of the OCM reaction mixture. A similar trend was observed for the other cubic BSCF reflections (Fig. S15, ESI[†]). However, in contrast to the previous experiment, no changes in the lattice parameter of the cubic structure occurred with time; its value remained constant and was determined to be 4.045 Å (Fig. S16, ESI[†]).

The results of the refinements clearly show that the BSCF material was not stable during the OCM experiment. More importantly, a new phase, identified as BaWO_4 was seen to form at the interface between membrane and catalyst material at the beginning of the second XRD-CT scan under OCM conditions with pure CH_4 . The distribution of all catalyst components and the BaWO_4 is presented in Fig. 6.

The BaWO_4 was formed as a result of an interaction between the catalyst and membrane material. As observed previously,³⁹ the Na^+ and WO_4^{2-} species are volatile and they can interact with the reactor vessel at high temperatures (*i.e.* above 780 °C).⁵¹ In this experiment the WO_4^{2-} species were found to migrate towards the membrane material where they interacted with Ba^{2+} present in the cubic perovskite, forming a continuous layer of



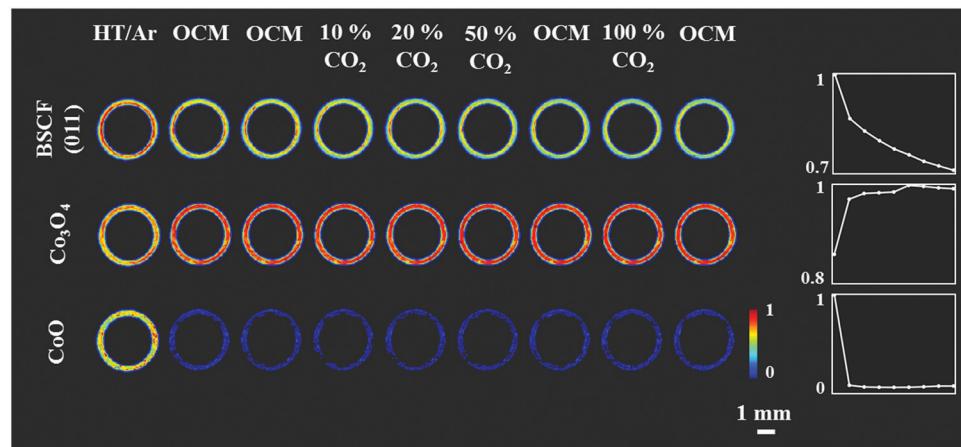


Fig. 5 Distribution of crystalline components present in the membrane material during the OCM experiment. Plots on the right correspond to the relative changes in the normalised mean scale factor or peak intensity obtained through refinement.

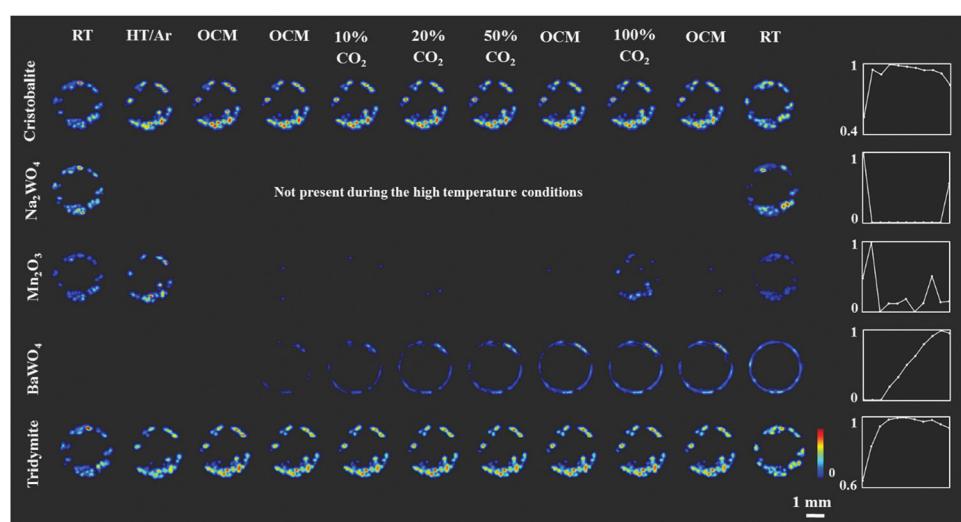


Fig. 6 Distribution of crystalline components present in the catalyst material during the OCM experiment. Plots on the right correspond to relative changes in the normalised mean summed peak intensity.

BaWO_4 at the membrane-catalyst interface. A similar observation was made in the case of *operando* studies with a $\text{BaCo}_{0.4}\text{Fe}_{0.4}\text{Zr}_{0.2}\text{O}_{3-\delta}$ perovskite membrane and a $\text{Na}-\text{Mn}-\text{W}/\text{SiO}_2$ catalyst.^{39,40} It is possible that this phenomenon can have a detrimental effect on the overall long-term performance of the CMR reactor as the newly formed layer is expected to have a negative impact on the oxygen permeability while the loss of catalyst active species is also bound to affect its performance. Similarly to the first experiment, no significant changes in the membrane were detected when CO_2 was co-fed with CH_4 ; there was no evidence for carbonate phases formation in this experiment.

Regarding the solid-state chemistry of the other catalyst components, the temperature induced a gradual transition between cristobalite and tridymite during the OCM reaction.^{51,52} In addition, the Mn_2O_3 phase was seen to disappear after the introduction of the OCM reaction mixture. However, it temporarily reappeared after the introduction of the reaction mixture containing pure CO_2 (possibly due to the CO_2 reduction by

$\text{Mn}^{2+}/\text{Mn}^{3+}$). This observation is in good agreement with previous work on a $\text{Na}-\text{Mn}-\text{W}/\text{SiO}_2$ catalyst-membrane system;^{51,53} the active catalyst does not need to contain the crystalline Mn_2O_3 phase. It is also clear that in the presence of CH_4 , the crystalline Mn_2O_3 loses its long-range order, most likely due to bulk reduction to Mn^{2+} . However, no crystalline phase containing Mn^{2+} (*i.e.* MnWO_4) was seen to form during the OCM reaction. It should finally be noted that minor changes in the distribution of catalyst components between room temperature and high temperature XRD-CT scans were attributed to minor movement of catalyst particles during the introduction of gas to the reactor (*i.e.* loose packing).

3.3. Laboratory performance testing and *ex situ* SEM/EDX characterisation

Complementary laboratory catalytic performance experiments were designed and performed in order to further support and rationalise the solid-state chemistry observations from the

in situ XRD-CT measurements. However, small changes to the experimental protocol had to be made in order to advance our understanding of the working system (*i.e.* duration of experiment, additional steps related to co-feeding CH₄/CO₂). The reactor outlet gases were analysed with mass spectrometry (MS) and gas chromatography (GC); however, the analysis is only qualitative due to the dead volume of the experimental setup introduced by the large condenser/catch pot after the reactor and before the analytical instruments which led to diluted streams and a corresponding delay in observing the changes in gas phase composition. The recovered samples after the laboratory experiments were analysed with scanning electron microscopy (SEM) coupled with energy dispersive X-ray analysis (EDX).

The results of the MS data collected during the OCM reaction with the catalytic membrane reactor are presented in the Fig. 7. With the addition of CO₂ in the reactor inlet stream

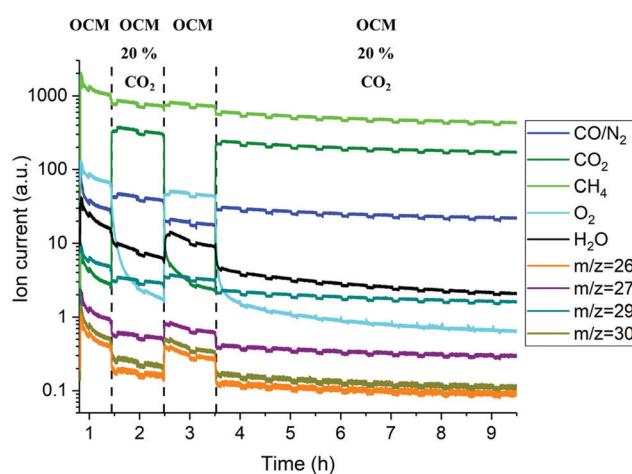


Fig. 7 Results of mass spectrometry data collected during the OCM experiment with catalytic membrane reactor. The y axis is in logarithmic scale and the ion current values were normalized with respect to the He signal (equivalent to instrument response with time). Note: oscillations in the mass spectrometry are due to the GC injection and corresponding valve movements.

(inner side stream) the most apparent changes are related to the O₂ signal, as it was found to gradually decrease after the introduction of 20% CO₂ in the CH₄ stream. Changing the gas composition to CH₄ (50% of CH₄ in Ar) resulted in the recovery of the oxygen permeability which was found to decrease again in the second cycle with 20% of CO₂. However, during this second cycle, the oxygen MS signal was seen to decrease faster than during the first cycle. The O₂ signal continued to gradually decrease until the end of experiment. The GC measurements showed that the BSCF membrane exhibits poor activity for the OCM reaction and the additional CO₂ in the reactor inlet stream further decreases it (Fig. S19, ESI†).

The elemental distribution of the membrane components at the inner side of the membrane is presented in Fig. 8. The results from the EDX mapping measurements showed that there is a segregation of CoO_x species in the membrane material (Fig. 9). At the same time, the SEM images revealed the presence of a thin layer of ~5 µm on the membrane surface which possesses a different structure (needle-like) when compared to membrane bulk (dense structure). The formation of the needle-like structure in the BSCF membrane has been previously reported in the work of Engels *et al.*³⁸ however no exact information regarding its composition and structure was presented. The EDX mapping measurements showed that there was no significant difference in the elemental composition between the surface layer and the dense bulk of the membrane. The morphology of this new structure resembles the morphology of the secondary phases formed in the BSCF during operation under temperatures below 900 °C.^{34,54,55}

The mass spectrometry results collected with the packed-bed catalytic membrane reactor are shown in Fig. 10. As mentioned previously, the changes in the MS signal were slower due to the large dead volume after the reactor, however it is still possible to observe the general trend in the gas phase composition. The gradual introduction of CO₂ into the OCM reaction mixture (from 10 to 50%) resulted in a decrease of oxygen permeation as well as lower production of C₂₊ molecules (Fig. S20, ESI†). It is important to note that the difference in the C₂₊ production between the 1st

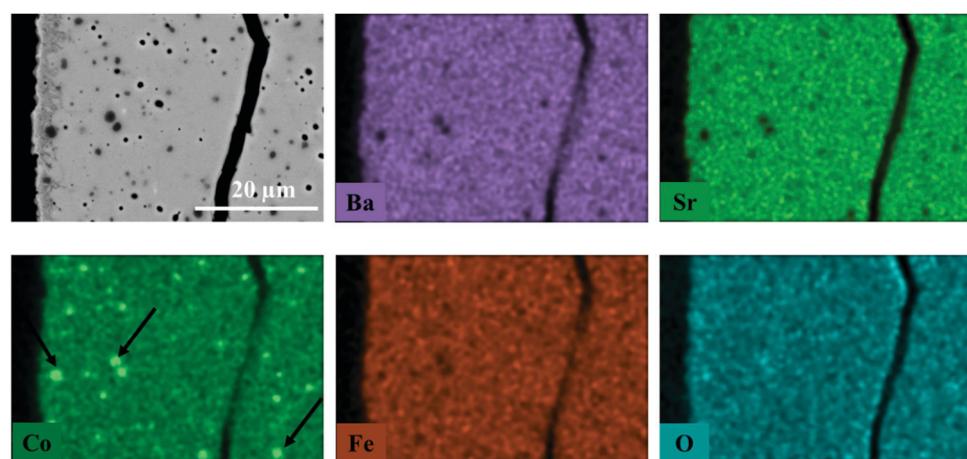


Fig. 8 Elemental distribution of membrane components at the inner side of membrane obtained by EDX mapping. This fragment of membrane cross section was obtained from the reactor inlet.



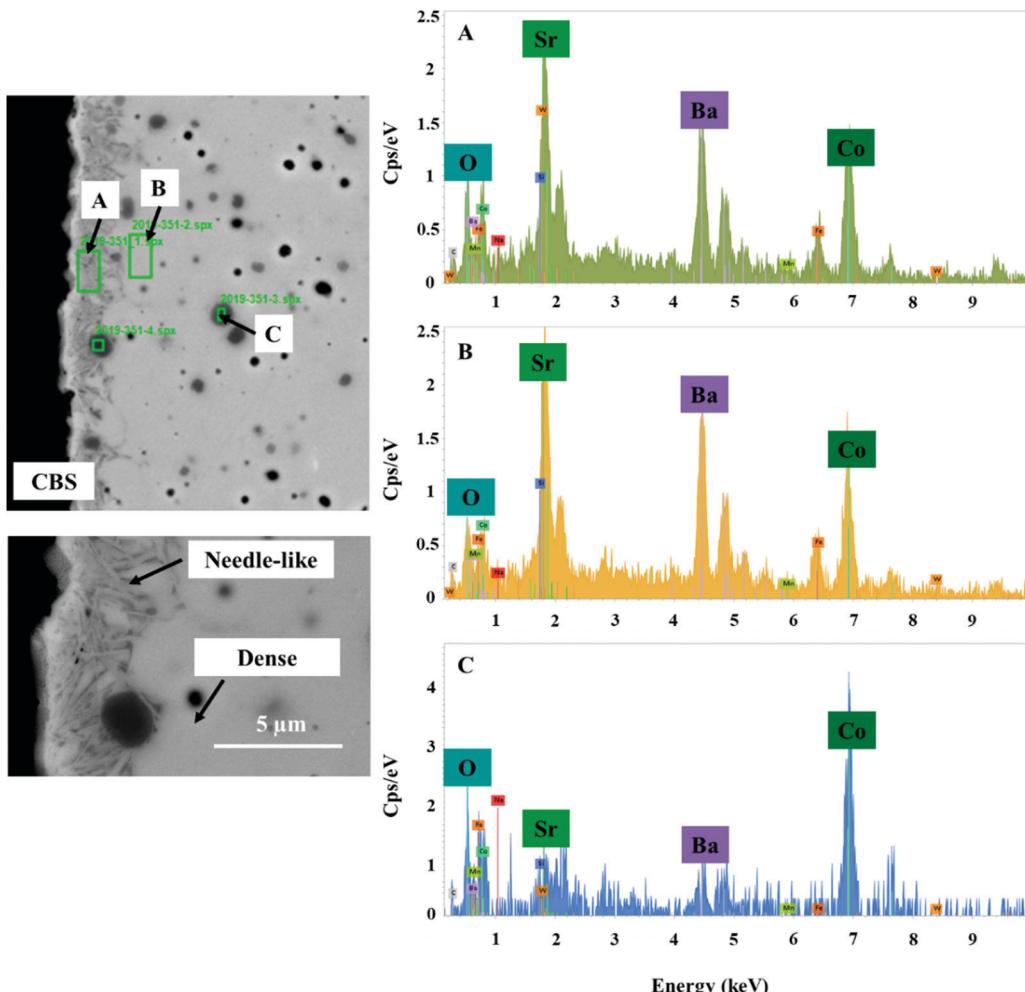


Fig. 9 SEM image of the inner side of the membrane and the corresponding energy spectrum from chosen regions. This fragment of the membrane cross section was obtained from the reactor inlet.

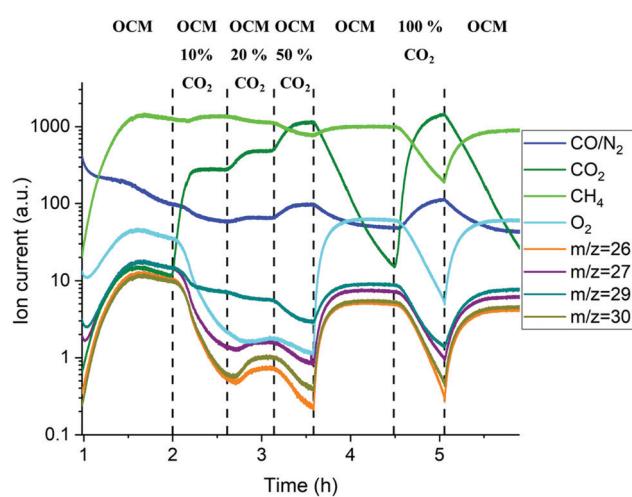


Fig. 10 Mass spectrometry results collected during the OCM experiment with packed-bed catalytic membrane reactor. The y axis is in logarithmic scale and the ion current values were normalized with respect to the He signal (equivalent to instrument response with time). Note: the 1st OCM measurements were performed with $\text{CH}_4:\text{O}_2$ of 3.5 whereas the 2nd and 3rd OCM measurements were performed with $\text{CH}_4:\text{O}_2$ of 2.5.

and 2nd stage of the OCM reaction (1–2 h and 3.5–4.5 h respectively) was mainly caused by the introduction of another $\text{CH}_4:\text{O}_2$ ratio (3.5 vs. 2.5). As reported previously,^{56,57} at higher $\text{CH}_4:\text{O}_2$ ratios the production of C_2+ is seen to increase. However, a lower production of C_2+ molecules, when compared to the 2nd and 3rd OCM measurement performed at the same $\text{CH}_4:\text{O}_2$ ratio, can be explained by the solid-state changes in the membrane primarily associated with the formation of the BaWO_4 layer.

The elemental distribution of the catalyst and membrane components at the inner side of the membrane near the reactor inlet is presented in Fig. 11. The results from the EDX mapping revealed that during the OCM reaction there was an interaction between volatile W-containing species and the BSCF membrane which led to the formation of thin deposits on the membrane surface, at the interface between membrane and catalyst particles. This observation further supports the results obtained from the *in situ* XRD-CT data where it was shown that a BaWO_4 formed during the OCM experiment. In addition, these deposits were seen to contain Na and Mn species. Similarly to the previous experiment with the BSCF membrane, Co species were present in

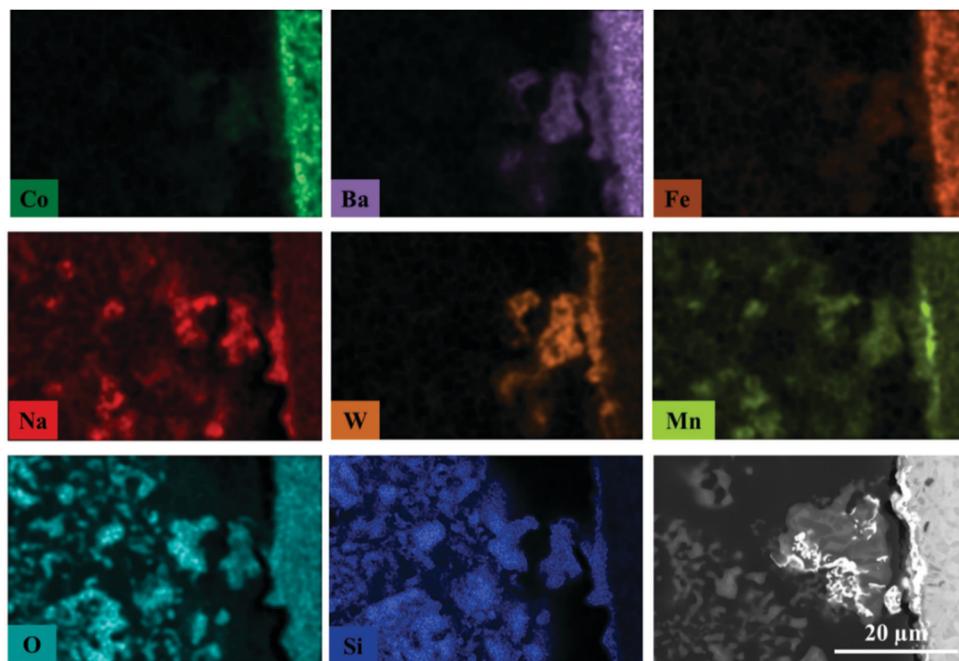


Fig. 11 Elemental distribution at the inner side of membrane obtained by EDX mapping. This fragment of the membrane cross section was obtained from the reactor inlet.

the form of large agglomerates containing CoO_x species but in addition they were also seen to agglomerate at the membrane surface, forming a uniform layer.

The EDX mapping measurements were also performed for inter and intra particle investigation of the distribution of the catalyst active components (Fig. S22, ESI[†]). The Na and Mn containing species are seen to be present in the catalyst particles as well as in the deposits on the membrane surface, whereas the W-containing species are seen to be present in the deposits and in the particles interacting with the membrane. These observations suggest that all active catalyst components are seen to migrate towards the membrane surface during the OCM reaction, with the W-containing species being the most prone to migration/volatilisation. In addition, the remaining Na and Mn containing species in catalyst particles are seen to have similar and rather uniform distribution; no formation of agglomerates with high concentration of material.

4. Summary and conclusions

In this work, we presented *in situ* XRD-CT studies of two catalytic membrane reactors consisting of BSCF hollow fibre membranes and a Na–Mn–W/SiO₂ catalyst. The spatially-resolved diffraction patterns obtained under operating conditions coupled with laboratory MS measurements and SEM/EDX post-reaction measurements revealed that there are several challenges that need to be addressed before the BSCF membrane can be used for the industrial application of the OCM process.

More specifically, the long-term operation of the BSCF membrane at intermediate temperatures (*i.e.* 825 °C) resulted

in the gradual decomposition of the BSCF to secondary phases; the XRD-CT measurements identified the formation of a hexagonal phase containing in majority Ba and Co species ($\text{Ba}_6\text{Co}_4\text{O}_{12}$) while the SEM measurements indicated the formation of a new structure (needle-like) around the edge of membrane. However, the EDX mapping measurements did not reveal any chemical difference between the needle-like structure and the membrane dense structure. In addition, the formation of the secondary phases during operation of the BSCF membrane was shown to cause a gradual decrease in oxygen permeation.

In the case of the packed-bed catalytic membrane reactor, a chemical interaction between the membrane and catalyst was observed; at high temperature the W species were seen to migrate towards the membrane surface forming a layer of BaWO_4 . The additional SEM/EDX measurements on the recovered samples revealed that also Na and Mn species migrated towards the membrane but to a lesser extent. With the MS measurements, the formation of the BaWO_4 layer was directly seen to have a detrimental effect on the membrane performance, as the production of C_{2+} molecules was seen to decrease during the OCM measurements performed in between the CO_2 cycling. The BaWO_4 formed at the interface between the catalyst particles and membrane is believed to suppress the oxygen permeation and may have indirectly led to a reactor type change, specifically to fixed-bed type with co-fed CH_4 and O_2 . The behavior of the W species results from the intrinsic instability of Na–Mn–W/SiO₂ catalyst at high temperatures and under operating conditions; this problem could potentially be solved through the chemical promotion and/or thermal treatment.^{51,52}



Regarding the poisoning effect of CO₂, the laboratory measurements showed that the addition of CO₂ (from 10% to 100%) to the reaction mixture significantly affected the oxygen permeation and thus the production of C₂₊ molecules. In all measurements, the CO₂ effect was reversible and the initial performance could be recovered when removing the CO₂ from the reaction mixture. It should be noted that both *in situ* XRD-CT and *ex situ* SEM/EDX mapping of the recovered membrane samples did not provide any evidence for carbonate species formation or elemental segregation. This implies that if any carbonate species formed at the membrane surface, these are in not in large quantities and indeed below the detection limit of the techniques applied in this work (*i.e.* instruments and experimental conditions). The structural changes and formation of BaWO₄ were seen to be the main mechanisms behind the long-term degradation of the investigated CMRs. It is concluded that the CO₂ produced from CH₄ combustion/oxidation processes during the OCM reaction does not significantly affect the oxygen permeation. However, as shown in this work, using the CO₂ as the reactor diluent has a negative effect on the membrane reactor performance and should be avoided.

Author contributions

Experimental plan, D. M., A. V., S. D. M. J. and A. M. B.; Material preparation, V. M. Synchrotron measurements, D. M., A. V., M. A. A. and A. M. B., Beamline support, G. V.; Data analysis, D. M. and A. V.; SEM/EDX measurements, V. M.; Original draft preparation, D. M. and A. V.; Writing—review and editing, all authors.; supervision, S. D. M. J., R. J. C. and A. M. B.; funding acquisition, S. D. M. J. and A. M. B.

Data availability

Copies of raw radially integrated XRD-CT data can be found at <https://rcahdrive.rc-harwell.ac.uk/index.php/s/75ZywcMDqmzi6Hn>. All data are available from the corresponding authors on reasonable request.

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Conflicts of interest

The authors declare no conflict of interest.

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