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## A novel process towards the industrial realization of large-scale oxymethylene dimethyl ether production – COMET†

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Oxymethylene dimethyl ethers (OME) show promising solubility and combustion properties for applications in various chemical processes and sectors. OME enable clean and quasi soot-free combustion, which can consequently lead to considerable NO<sub>x</sub> emissions reduction. Besides reducing local emissions, OME can significantly reduce the global CO<sub>2</sub> emissions by substituting fossil diesel fuel if their production is based on sustainable methanol. Various process concepts for the OME production were proposed and investigated, but most of them have significant bottlenecks, which prevent their demonstration and scale-up in the near future. Only the production based on OME<sub>1</sub> and trioxane can already be demonstrated and scaled up, which, however, is complex and energy-intensive, considering a sustainable production based on H<sub>2</sub> and CO<sub>2</sub>. Therefore, the novel COMET (clean OME technology) process concept is introduced and experimentally demonstrated utilizing only state-of-the-art process units. The COMET process relies solely on methanol and formalin as feedstock and overcomes the challenging water management aspect in the OME value chain, using a reactive distillation column. The COMET process is evaluated at a scale of 100 kilotons per annum OME<sub>3-5</sub> product for the system boundary starting from H<sub>2</sub>O electrolysis and CO<sub>2</sub> capture. Key performance indicators are defined and compared with alternative processes from the literature. The COMET process shows a high carbon efficiency of 88% and overall energy efficiency of 54% in comparison to the alternative OME<sub>3-5</sub> production processes introduced in the literature. Moreover, the COMET process offers the forthwith large-scale production of OME in a relatively simple process chain and high technology readiness level.

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### 1. Introduction

The fluctuating availability of renewable energies can be compensated by various measures, such as storage technologies like batteries and heat reservoirs. However, the need for long-term storage and long-distance transportation are met more easily and flexibly using power-to-X (PtX) technologies and products.<sup>1</sup> Green H<sub>2</sub> will play a key role in meeting these needs with already rising demands, as is suggested by various roadmaps, strategies, and international agreements by various governments and companies.<sup>2,3</sup>

Currently, fossil energy carriers are essential to provide heat and power, and are important carbon sources for the chemical, petrochemical and plastic industries. However, at the end of the life cycle, they are usually combusted to CO<sub>2</sub>, which ends up in the atmosphere. Therefore, sustainable and circular carbon sources are required such as biogenic waste streams and CO<sub>2</sub> capture from air. One example of a circular carbon source is direct air capture (DAC) of CO<sub>2</sub>. Due to its locally independent availability, DAC is experiencing a growing governmental interest with increasing numbers and capacities of technological demonstrations.<sup>4</sup>

A combination of captured CO<sub>2</sub> and green H<sub>2</sub> enables the implementation of these sustainable solutions into various sectors and hard to defossilize processes. One suitable product of CO<sub>2</sub> and H<sub>2</sub> are oxymethylene dimethyl ethers (OME). OME show promising fuel and physical properties for a wide range of potential applications such as solvents or diesel fuel additives or substitutes. OME<sub>3-5</sub> have similar fuel properties to diesel fuel, a good solubility in diesel fuel and advantageous combustion behavior.<sup>5,6</sup> This makes OME

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attractive as a sustainable drop-in blending component for diesel fuel. Considering the high cetane number, OME can be an interesting dual fuel or ignition promoter in marine engines. Due to the absence of C–C bonds and the high amount of molecular bound oxygen, OME combust with distinct lower levels of emissions of particle matter than diesel fuel. Therefore, the NO<sub>x</sub> and soot emission trade-off of diesel fuel can be avoided.<sup>7,8</sup>

The basis for large-scale sustainable production of OME is MeOH, which can be produced by reacting captured CO<sub>2</sub> with H<sub>2</sub> from renewable-powered H<sub>2</sub>O electrolysis. Following this power-to-liquid (PtL) concept, the well-to-wheel (WtW) CO<sub>2</sub> emissions can be reduced by up to 93% compared to fossil fuels.<sup>9,10</sup> Furthermore, Voelker *et al.*<sup>10</sup> estimated an NO<sub>x</sub> reduction of 57% and an almost complete reduction of soot using OME instead of diesel fuel. Moreover, small blending rates of OME in diesel fuel already show a clearly positive impact on global CO<sub>2</sub> emissions, as well as local NO<sub>x</sub> and soot emissions.<sup>10,11</sup> With a worldwide demand of 26.5 million barrels diesel fuel per day,<sup>12</sup> small blending rates of OME showcase the need for large-scale production plants. Suitable compositions of the final OME<sub>3–5</sub> product are defined by key properties, such as density, viscosity, cetane number and flash point, which are standardized in a new fuel pre-standard for OME DIN/TS 51699.<sup>7,13</sup>

Various process concepts have been proposed to produce OME from MeOH which will be discussed and compared in detail in the following sections. One of the main challenges of these processes, in terms of technical feasibility and energy demand, is the separation of the by-product H<sub>2</sub>O from the target OME fraction. This H<sub>2</sub>O is formed in various synthesis steps from MeOH to OME. The OME production processes discussed in the literature use different techniques to separate H<sub>2</sub>O.<sup>14–25</sup> These techniques have different advantages and disadvantages, partly coupled with methods which are still in an early phase of investigation and demonstrations. In the present work, a new process concept, clean OME technology (COMET), is proposed which solves the challenging H<sub>2</sub>O separation utilizing the state-of-the-art reactive distillation technique. This makes the COMET process a technically feasible process for large OME production capacities.

### 1.1. Objective of this work

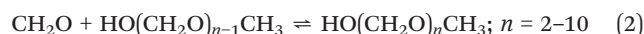
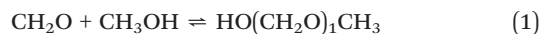
The main objective of this work is the introduction of the novel COMET process concept for the production of OME from MeOH and aqueous formaldehyde (FA(aq.)) solutions, which solves the challenging H<sub>2</sub>O management problem, using a reactive distillation column. Complementary, the experimental demonstration of the main COMET process units is introduced. The work further compares the COMET process concept major technical metrics to literature discussed OME production processes.

## 2. Theory and background

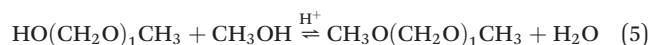
### 2.1. Synthesis of OME

OME are synthesized in an acidic environment from two methyl capping groups and *n* oxymethylene groups –CH<sub>2</sub>O–. Suppliers for the methyl capping groups are methanol (H<sub>3</sub>C–OH, MeOH), methylal (H<sub>3</sub>C–O–(CH<sub>2</sub>O)<sub>1</sub>–CH<sub>3</sub>, OME<sub>1</sub>) or dimethyl ether (H<sub>3</sub>C–O–CH<sub>3</sub>, DME). For the oxymethylene groups, formalin (FA(aq.)), paraformaldehyde (HO–(CH<sub>2</sub>O)<sub>*n*</sub>–H with *n* = 8–100, pFA) or trioxane ((CH<sub>2</sub>O)<sub>3</sub>, TRI) can be used.

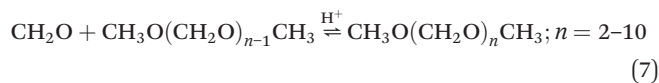
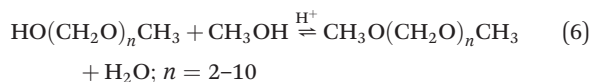
In a solution of MeOH, FA, and H<sub>2</sub>O, poly(oxymethylene) hemiformals (HO–(CH<sub>2</sub>O)<sub>*n*</sub>–CH<sub>3</sub> with *n* = 1–10, HF<sub>*n*</sub>) and poly(oxymethylene) glycols (HO–(CH<sub>2</sub>O)<sub>*n*</sub>–H with *n* = 1–10, MG<sub>*n*</sub>) bound most of the FA, as described by eqn (1)–(4). These are fast reactions even in absence of a catalyst. The amount of monomeric FA in solutions with MeOH and H<sub>2</sub>O is very small in chemical equilibrium.<sup>26</sup>



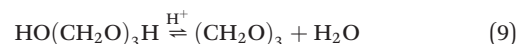
HF<sub>*n*</sub> are intermediate products for the production of OME. In an acidic environment HF<sub>1</sub> and MeOH react to OME<sub>1</sub> following the acetalization reaction described by eqn (5).



The formation of OME<sub>>1</sub> follows a similar acetalization mechanism, as described by eqn (6), and a sequential addition or chain propagation mechanism, as described by eqn (7).<sup>27</sup>



By using TRI as a feedstock for the OME synthesis, three CH<sub>2</sub>O units are formed in a first step, as described by eqn (8).



Furthermore, TRI can also be formed as a by-product following eqn (8), or by dehydration of MG<sub>3</sub>, as described by eqn (9).<sup>28</sup>

Using DME as a feedstock for the OME synthesis, OME<sub>1</sub> is formed by the incorporation of FA into DME, as described by eqn (10).





Furthermore, with DME and TRI as a feedstock for the OME synthesis, OME<sub>3</sub> is formed by direct incorporation of TRI into DME, as described by eqn (11).<sup>29–31</sup>

In addition to the main reaction network, side reactions lead to the formation of side products. Besides HF<sub>n</sub>, MG<sub>n</sub>, TRI and DME, the formation of tetroxane ((CH<sub>2</sub>O)<sub>4</sub>), methyl formate (HCOOCH<sub>3</sub>, MEFO) and formic acid (HCOOH, FOAC) was observed, which strongly depends on the catalyst systems and increases with increasing temperatures.<sup>32–34</sup>

## 2.2. OME<sub>3–5</sub> production processes

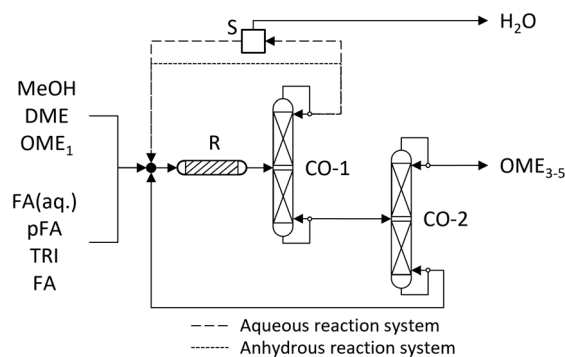
Different feedstock combinations can be used to produce OME<sub>3–5</sub> due to various potential suppliers of methyl groups and oxymethylene groups. Depending on the formation of H<sub>2</sub>O as a side product during the synthesis of OME<sub>n</sub>, the reaction systems are classified as anhydrous or aqueous. Methyl group suppliers for anhydrous reaction systems are mainly DME and OME<sub>1</sub>, because their conversion to OME<sub>n</sub> only requires a chain propagation with oxymethylene groups, as described by eqn (7) and presented in Table 1. Oxymethylene group suppliers for anhydrous reaction systems are mainly TRI and monomeric FA, which do not contain H<sub>2</sub>O or MeOH. Formalin, concentrated FA(aq.) or pFA contain H<sub>2</sub>O and are used as oxymethylene group suppliers for aqueous reaction systems.

Feedstocks containing MeOH generally lead to the formation of H<sub>2</sub>O as a side product in the aqueous OME synthesis, as described by eqn (5) and (6). This H<sub>2</sub>O needs to be separated and extracted from the process loop to prevent accumulation. Fig. 1 shows a simplified scheme for the production of OME<sub>3–5</sub> from various feedstocks. It consists mainly of a reactor for the OME synthesis R, two distillation columns CO-1 and CO-2 for product purification and a H<sub>2</sub>O separation unit S for aqueous reaction systems. For the H<sub>2</sub>O separation, various methods were proposed in the literature, such as extraction, adsorption or membrane, as discussed in the following section.

Various process concepts for the OME<sub>3–5</sub> production were proposed in patents and other publications and some of them are realized in large-scale production plants in China. However, details regarding their performance, the quality and composition of the final OME product and the long-term operation are scarce.<sup>35,36</sup> Table 2 lists the main OME<sub>3–5</sub> production processes discussed in the literature,

**Table 1** Methyl group suppliers and oxymethylene group suppliers for anhydrous and aqueous OME reaction systems

	Methyl group supplier	Oxymethylene group supplier
Anhydrous	DME, OME <sub>1</sub>	TRI, monomeric FA
Aqueous	MeOH, DME, OME <sub>1</sub>	FA(aq.), pFA, TRI, monomeric FA



**Fig. 1** OME<sub>3–5</sub> production process for various feedstocks, following aqueous and anhydrous reaction systems. CO, distillation column; R, reactor; S, H<sub>2</sub>O separator.

emphasizing the feedstock, main advantages, and hurdles. A detailed description is provided in the ESI.† A comparison with the COMET process based on their performances in terms of OME<sub>3–5</sub> yield, energy demand and technical feasibility is discussed in the results and discussion section. Further process concepts were proposed in the literature which show significant disadvantages in comparison to the process concepts presented in Table 2, as discussed in the ESI.†

## 2.3. H<sub>2</sub>O separation from the production of OME

One of the main challenging and energy-intensive process steps is the separation of the by-product H<sub>2</sub>O from the production chain towards OME<sub>3–5</sub>. Considering a sustainable production of OME<sub>3–5</sub> based on MeOH produced from H<sub>2</sub> and CO<sub>2</sub>, Fig. 2 shows how the synthesis of MeOH, intermediate products such as FA(aq.), DME and OME<sub>1</sub>, as well as the synthesis of OME<sub>n</sub> all result in the formation of H<sub>2</sub>O.

In the MeOH synthesis from H<sub>2</sub> and CO<sub>2</sub>, the OME<sub>1</sub> synthesis and the DME synthesis, H<sub>2</sub>O is a by-product and separated using distillation columns.<sup>19,37,38</sup> In the partial oxidation of MeOH towards FA, as described by eqn (15), H<sub>2</sub>O is formed as a by-product and used as a washing liquid in the absorber column. Downstream, H<sub>2</sub>O is partly separated from FA(aq.) in a concentration step using evaporation techniques.<sup>19,39</sup> Therefore, H<sub>2</sub>O is introduced into the TRI synthesis and separated in an energy-intensive cascade of distillation columns.<sup>28,40</sup> Only in the anhydrous FA synthesis, which is still in its very early stages, no H<sub>2</sub>O is present.<sup>41</sup> Regarding the synthesis of OME<sub>≥2</sub>, H<sub>2</sub>O is not formed as a by-product when the oxymethylene group suppliers TRI or monomeric FA are combined with the methyl group suppliers DME or OME<sub>1</sub>, as described by eqn (7) and (8).

This simplifies the final product purification. When starting from the cheaper and established reactant FA(aq.), H<sub>2</sub>O will always be present in the OME<sub>3–5</sub> sub-process and needs to be separated from the loop to circumvent accumulation. However, H<sub>2</sub>O cannot be separated



**Table 2** Advantages and main hurdles of various OME<sub>3-5</sub> production process concepts

Feedstock	Anhydrous synthesis			Aqueous synthesis		
	OME <sub>1</sub> and TRI <sup>14-16</sup>	DME and TRI <sup>17,18</sup>	OME <sub>1</sub> and monomeric FA <sup>19</sup>	MeOH and FA(aq.) <sup>20,21</sup>	MeOH and monomeric FA <sup>19</sup>	OME <sub>1</sub> and FA(aq.) or pFA <sup>19</sup>
(+) Main advantages and (-) main hurdles	+ High OME <sub>3-5</sub> yield after the synthesis  + Simple product purification  - Complex and energy-intensive preparation of TRI <sup>28,39,40,42</sup>	+ DME is cheaper than OME <sub>1</sub> <sup>43</sup>  - Complex and energy-intensive preparation of TRI <sup>28,39,40,42</sup> - High MEFO selectivity <sup>29-31,44,45</sup>	+ High OME <sub>3-5</sub> yield after the synthesis  + Potentially simpler and cheaper production of monomeric FA - Very low TRL of the monomeric FA production	+  Comparatively cheap feedstock - Formation of H <sub>2</sub> O as a side product - Low OME <sub>3-5</sub> yield after the synthesis - Low TRL of the H <sub>2</sub> O separation methods	+ Comparatively cheap feedstock  - Similar hurdles to MeOH and FA(aq.) - Very low TRL of the monomeric FA production	+ Fairly high OME <sub>3-5</sub> yield after the synthesis  - Similar hurdles to MeOH and FA(aq.)

individually simply *via* distillation due to a complex phase behavior of the synthesis product mixture containing mainly FA, H<sub>2</sub>O, MeOH, OME<sub>1-10</sub>, HF and MG with several azeotropes with similar boiling points. The separation of H<sub>2</sub>O from the loop is still a major challenge regarding the implementation of a potentially cheaper and scalable aqueous OME<sub>3-5</sub> production process.

**2.3.1. Extraction.** Using an extractant for the separation of H<sub>2</sub>O from the OME synthesis mixture separates the mixture into two phases, one organic phase mainly containing OME and the extractant and one aqueous phase mainly containing H<sub>2</sub>O, FA and MG. Downstream to the extraction the organic phase can be separated and purified using distillation columns. The extractant is also separated and recycled back to the extraction. Various extractants were investigated in the literature showing that toluene, *p*-xylene and *n*-heptane enable promising liquid-liquid separation behaviors between OME and H<sub>2</sub>O, FA and MG.<sup>46-52</sup>

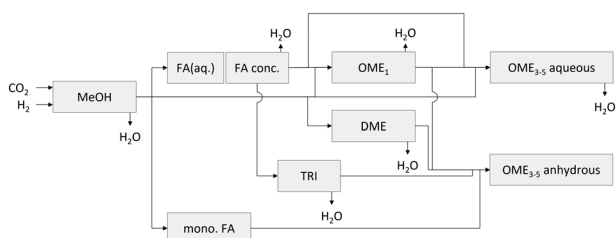
Results of Li *et al.*<sup>47</sup> show the separation of the OME synthesis product using toluene. About 70% of OME are separated in the organic phase and only 14% of FA and H<sub>2</sub>O migrate in the organic phase, as indicated by the split fraction. However, the organic phase mainly consists of toluene, which needs to be separated to be recycled. Furthermore, FA and H<sub>2</sub>O still represent a large proportion of

the organic phase and the aqueous phase still contains a large proportion of OME. A graphical illustration of the results of Li *et al.*<sup>47</sup> is presented in the ESI† in section 2.

An extraction method for the preparation of blends of OME in diesel fuel was proposed by Oestreich *et al.*<sup>53</sup> and discussed in the ESI.†

**2.3.2. Adsorption.** Schmitz *et al.*<sup>20</sup> investigated the adsorption of H<sub>2</sub>O from a mixture containing FA, H<sub>2</sub>O, MeOH and OME<sub>1-4</sub> using zeolite 3A. Their results show that zeolite 3A has a good selectivity for H<sub>2</sub>O with only small amounts of FA and MeOH being separated from the feed mixture. A graphical illustration of the results is presented in the ESI† in section 2. Ferre *et al.*<sup>54</sup> investigated the adsorption of H<sub>2</sub>O using zeolite 3A from binary and ternary mixtures with MeOH and FA. Their results show that an increasing amount of FA or MeOH in the feed mixture leads to an increased adsorption of these components. However, the selectivity for H<sub>2</sub>O is still far higher.

Regarding the separation of H<sub>2</sub>O from an OME<sub>3-5</sub> production process, the adsorption has the advantage of selectively separating H<sub>2</sub>O from the loop, which enables the recycle of all other components to the OME synthesis. Due to the reaction network between H<sub>2</sub>O and FA as described by eqn (3) and (4) not only the monomeric H<sub>2</sub>O is separated, but also H<sub>2</sub>O from MG<sub>*n*</sub>. Therefore, a significant reduction of the overall H<sub>2</sub>O content can be achieved. However, without H<sub>2</sub>O, FA from MG<sub>*n*</sub> remains in the mixture and either bounds with HF<sub>*n-1*</sub> to long chain HF<sub>*n*</sub>, or with MG<sub>*n-1*</sub> to long chain MG<sub>*n*</sub> or it remains in monomeric form. Either way, it increases the risk of local precipitations and, therefore, deactivation of the adsorbents. Therefore, a regeneration might be necessary. To reduce the risk of precipitation the temperature can be lifted, or the remaining H<sub>2</sub>O content can be increased. The latter would, however, decrease the yield of OME<sub>3-5</sub> in the OME synthesis and, therefore, increase the recycle streams and heat demand for separation. A suitable remaining H<sub>2</sub>O content should be experimentally investigated and confirmed

**Fig. 2** H<sub>2</sub>O separation from a sustainable production of OME<sub>3-5</sub> based on H<sub>2</sub> and CO<sub>2</sub>.

by long-term stability tests with alternating sequences of adsorption and regeneration. Furthermore, the scale-up potential should be investigated to ensure its feasibility for large-scale production plants.

Regarding the heat demand for the separation of H<sub>2</sub>O *via* adsorption, Schemme *et al.*<sup>43,55</sup> estimated that 2.1 kW h kg<sup>-1</sup> H<sub>2</sub>O are required. Their estimations are based on the results from Schmitz *et al.*<sup>20,56</sup> and assume that the adsorbents are heated up from 25 °C to 235 °C for the regeneration using high pressure steam. Furthermore, it was assumed that the heat demand is mainly based on the heat of adsorption and the heat capacity of the adsorbents.

**2.3.3. Membrane.** Schmitz *et al.*<sup>57</sup> tested two zeolite membranes type NaA and type T from Mitsui & Co. as well as three PVA-based polymer membranes PERVAP 4100, PERVAP 4101 and PERVAP 4102 from DeltaMem AG for a mixture containing FA, H<sub>2</sub>O, MeOH, OME<sub>1</sub> and OME<sub>2</sub>. Their results show that the zeolite membranes and the PERVAP 4102 were not suitable for the separation task, while the PVA-based polymer membranes PERVAP 4100 and PERVAP 4101 could separate H<sub>2</sub>O with a high selectivity, also for the repeated experiment with the tested zeolite membranes. A graphical illustration of the results by Schmitz *et al.*<sup>57</sup> using the PERVAP 4100 membrane is presented in the ESI† in section 2.

However, Ferre *et al.*<sup>58</sup> reported the application of a different membrane from DBI Gas und Umwelttechnik. The long-term stability of the membrane, selectivities in the reaction mixture and the scale-up potential should be further investigated to ensure its feasibility for large-scale production plants.

The advantages of the membrane for the separation of H<sub>2</sub>O are similar to the advantages of the adsorption with a high selectivity for H<sub>2</sub>O. However, likewise to the adsorption, this results in a higher risk for local precipitation. Therefore, a compromise might be necessary between the long-term stability and the H<sub>2</sub>O concentration of the retentate. A disadvantage of a higher H<sub>2</sub>O concentration in the retentate is an increase of the recycle streams which results in higher heat demands for the product purification and reduces the overall energy efficiency of the process.

Regarding the heat demand for the separation of H<sub>2</sub>O *via* membranes, Held *et al.*<sup>39</sup> estimated that 0.7 kW h kg<sup>-1</sup> H<sub>2</sub>O are required. This results from the evaporation of H<sub>2</sub>O after passing through the membrane to the reduced pressure of the permeate of 0.03 bar. Held *et al.*<sup>39</sup> assumed that no external heat is required but the temperature of the process

stream is reduced from 84 °C to 36 °C. In comparison to the separation of H<sub>2</sub>O *via* adsorption, the heat demand is significantly lower.

Table 3 summarizes the main advantages and main hurdles of the H<sub>2</sub>O separation methods extraction, adsorption, and membrane.

### 3. COMET process description

The COMET process<sup>59</sup> is based on the commercially available MeOH and FA(aq.) feedstock and produces mainly high purity OME<sub>3-5</sub>, as illustrated in Fig. 3. For the separation of H<sub>2</sub>O from the loop, a reactive distillation column is used.

The COMET process starts at the concentration of 50–55 wt% FA(aq.) (stream 1), which can be the product stream of a state-of-the-art FA production process.<sup>19,60</sup> The stream is mixed with the distillate of the second evaporator E-2 and the bottom of the third evaporator E-3. The FA(aq.) is then concentrated in a cascade of two evaporator stages E-1 and E-2 to provide a concentrated FA(aq.) of 85–88 wt% FA (stream 5) and an aqueous stream containing 10–25 wt% FA (stream 3). The concentrated FA(aq.) (stream 5) is used for the production of OME and mixed with the recycle streams, containing the azeotropic mixture of OME<sub>1</sub> and MeOH (stream 10) and OME<sub>≥6</sub> (stream 14). The mixture is converted in a fixed bed reactor R filled with an acidic heterogeneous catalyst. In contrast to the OME production process based on MeOH and FA(aq.),<sup>20</sup> the reactor inlet stream contains mainly OME<sub>1</sub> as a methyl capping source. This improves the selectivity towards OME<sub>3-5</sub>. The comparatively high selectivity further increases with decreasing H<sub>2</sub>O and MeOH concentrations in the concentrated FA feedstock (stream 5) and OME<sub>1</sub> recycle (stream 10). The synthesis product mixture, mainly containing FA, H<sub>2</sub>O, MeOH and OME<sub>1-10</sub> (stream 7), is separated in a cascade of three distillation columns. In the first distillation column CO-1, OME<sub>≥3</sub> are separated from the more volatile components FA, H<sub>2</sub>O, MeOH and OME<sub>1-2</sub>. Importantly, OME<sub>3</sub> cannot be completely separated to the bottom product, a small fraction remains in the distillate. In the third distillation column CO-3, OME<sub>≥6</sub> are separated and recycled to the reactor and a final product mixture (stream 13) of OME<sub>3-5</sub> is withdrawn at the top of CO-3. The distillate product of CO-1 is mixed with MeOH (stream 9) and sent to a reactive distillation column CO-2, to separate an azeotropic mixture of OME<sub>1</sub> and MeOH (stream 10) from FA and H<sub>2</sub>O (stream 11). On the catalytic trays, two main conversions take place. First, OME<sub>2</sub> and OME<sub>3</sub> are converted to OME<sub>1</sub> and FA over an acidic heterogeneous catalyst, as described by eqn (7). In addition, MeOH and FA are converted to OME<sub>1</sub> and H<sub>2</sub>O, following the acetalization reaction as described by eqn (1) and (5). The mechanism on the catalytic trays is illustrated in Fig. 4. Due to the evaporation and, therefore, the separation of the volatile product OME<sub>1</sub> from the liquid reaction mixture, the equilibrium of eqn (5) and (7) shifts, and the reactions proceed towards the production of OME<sub>1</sub>. Therefore, with sufficient retention time, OME<sub>2</sub> is converted

**Table 3** Main advantages and main hurdles of the H<sub>2</sub>O separation methods extraction, adsorption, and membrane

Method	Extraction	Adsorption	Membrane
H <sub>2</sub> O selectivity	Low	High	High
Energy demand	High	High	Comparatively low
Long-term operation	Likely	Challenging	Challenging
Scale-up potential	Likely	Challenging	Challenging



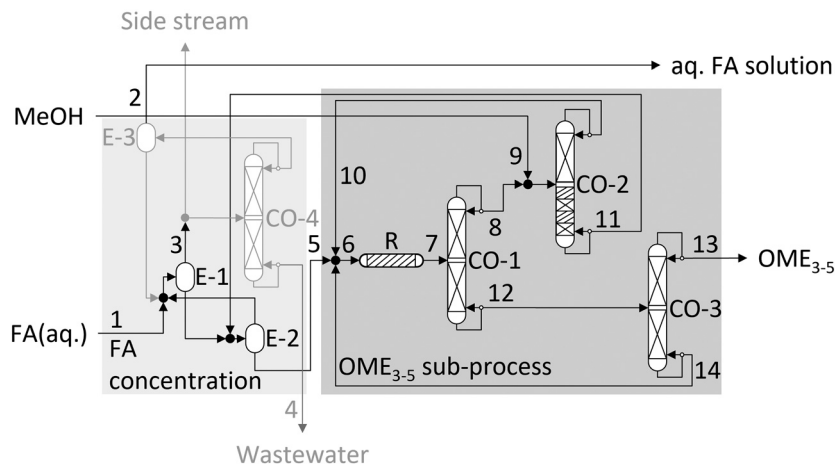


Fig. 3 COMET process concept for the production of OME<sub>3-5</sub> from MeOH and FA(aq.) feedstocks. The light grey arrows and process units were added in this work to the FA concentration sub-process to improve the recycle of FA. CO, distillation column; E, evaporator; R, reactor.

to a large extent to OME<sub>1</sub>, while the conversion of FA towards OME<sub>1</sub> is limited by the amount of MeOH. The mixture is separated into the azeotropic mixture of OME<sub>1</sub> and MeOH in the distillate (stream 10) and a mixture of FA and H<sub>2</sub>O in the bottom (stream 11). The distillate product of CO-2 is recycled back to the reactor and the bottom product is recycled to the evaporator E-2 for the FA concentration to separate H<sub>2</sub>O from the process and recycle concentrated FA back towards the OME reactor. Therefore, the reactive distillation column prevents the accumulation of H<sub>2</sub>O inside the loop and solves the challenging H<sub>2</sub>O management. In contrast to the H<sub>2</sub>O separation from the loop using adsorption or membranes, in the COMET process H<sub>2</sub>O is not separated selectively but together with the remaining FA. This significantly reduces the risk of precipitation, since enough H<sub>2</sub>O is left to convert the remaining FA to comparatively short-chain MG<sub>n</sub> which stay liquid at elevated temperature for sufficient retention time to downstream processing steps.

A similar concept for a reactive distillation column is applied in the OME<sub>1</sub> production process by Drunsel *et al.*<sup>61,62</sup> with the purpose to achieve a complete conversion of FA after the OME<sub>1</sub> reactor.

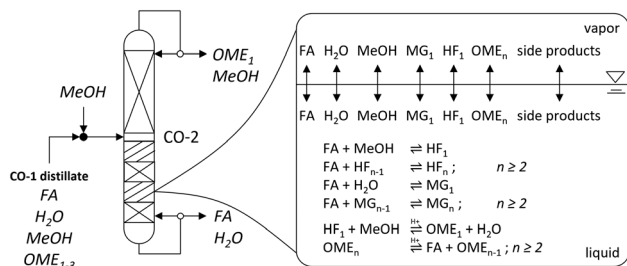


Fig. 4 H<sub>2</sub>O separation from the COMET process via reactive distillation. The left side shows the reactive distillation column with the main components of the feed and product streams. The illustration on the right side shows the interaction on a catalytic tray and was adopted from Schmitz *et al.*<sup>56</sup>

The amount of MeOH (stream 9) added to the feed of the reactive distillation column CO-2 defines the conversion of FA and oxymethylene groups with MeOH on the catalytic trays towards OME<sub>1</sub>, following eqn (1), (5) and (7). Therefore, a variation of the amount of MeOH (stream 9) varies the amount of OME<sub>1</sub> produced as the distillate product of the reactive distillation column CO-2. For the OME synthesis in the fixed bed reactor R a constant ratio of OME<sub>1</sub> to concentrated FA(aq.) is required before the reactor. Therefore, the amount of MeOH (stream 9) can be defined to exactly produce the amount of OME<sub>1</sub> required for the OME synthesis. Or the amount of MeOH (stream 9) can be increased to produce more OME<sub>1</sub> than required by the OME synthesis and the excess OME<sub>1</sub> can be extracted as a by-product. Another advantage of the COMET process is that the process offers a tunable product portfolio of OME. In the present work the amount of MeOH added to the reactive distillation column CO-2 was limited to only produce the required amount of distillate product (stream 10) for the OME synthesis and, therefore, achieve higher OME<sub>3-5</sub> selectivity. Considering the production of OME<sub>1</sub> as a target side product of the COMET process, another distillation column can be added to achieve high purities of the OME<sub>1</sub> side product, similar to the second distillation column of the production process for OME<sub>1</sub>.<sup>61</sup>

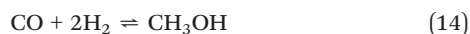
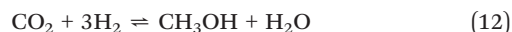
Besides the OME<sub>3-5</sub> product (stream 13), wastewater (stream 3) is produced with FA concentrations of about 10–25 wt%. This by-product stream is not limited to the COMET process but part of all OME<sub>3-5</sub> production processes using FA(aq.) as an intermediate product. Instead of its disposal and to increase the carbon yield of the process, several strategies are possible to handle this stream. In the present work, the stream was partly sent to the absorber column of the FA(aq.) production and concentrated in an additional distillation column CO-4. The concentrated FA stream was further concentrated in another evaporator E-3 to recycle the concentrated FA stream and to separate the stream with a low FA concentration (stream 2). This stream is also the purge stream for traces of MeOH and other volatile



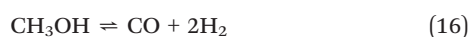
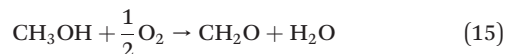
components to avoid accumulation in the loop. Instead of its disposal, this stream can be used to dilute an FA(aq.) product stream to prepare a stable formalin product.

### 3.1. Expanding the system boundary starting from H<sub>2</sub> and CO<sub>2</sub> feedstocks including the intermediate production of FA and MeOH

To enable a consistent basis of comparison with alternative OME<sub>3-5</sub> production processes, the system boundary is extended to account for a sustainable OME<sub>3-5</sub> production based on green H<sub>2</sub> and captured CO<sub>2</sub>. The intermediate production of MeOH and FA(aq.) is described in detail by Mantei *et al.*<sup>19</sup> A simplified process flow diagram of the extended COMET process concept starting from H<sub>2</sub> and CO<sub>2</sub> is illustrated in Fig. 5. The MeOH production starts from the compression of the feedstock H<sub>2</sub> and CO<sub>2</sub>, the mixing with the recycle stream, and the synthesis of MeOH, as described by eqn (12)–(14).



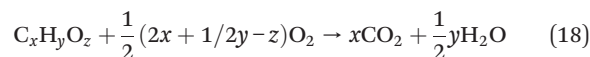
The synthesis product is purified in a cascade of two phase separators followed by a distillation column to prepare high purity MeOH. Part of the MeOH is used for the synthesis of FA. Therefore, MeOH and H<sub>2</sub>O are mixed with the recycle stream and air for the conversion to FA, as described by eqn (15)–(17).



The synthesis product is purified in an absorber column to separate FA and H<sub>2</sub>O from H<sub>2</sub>, CO, CO<sub>2</sub> and N<sub>2</sub> using part of

the wastewater from the FA concentration as a washing liquid. The product mixture of FA and H<sub>2</sub>O is concentrated in the FA concentration sub-process, as described in the previous section.

To use the heating value of the purge streams, a combustion sub-process was implemented. Following the assumptions from Mantei *et al.*,<sup>19</sup> excess air was added to achieve complete combustion and keep the adiabatic temperature rise below 800 °C. The stoichiometric amount of O<sub>2</sub> required for a complete combustion is described by eqn (18).



## 4. Materials and methods

### 4.1. Experiments

**4.1.1. Chemicals and materials.** The reactants OME<sub>1</sub> (purity > 99.8%) and MeOH (purity > 99.8%) were purchased from Brenntag GmbH and provided from ChemCom Industries B.V. The reactant FA(aq.) was provided as a stabilized aqueous FA (approx. 37 wt%) solution with low amounts of MeOH (≤0.5 wt%) by ChemCom Industries B.V.. The catalyst Amberlyst® 46 (A46) was provided by INAQUA Vertriebsgesellschaft mbH. Ion exchange resin (IER) III was purchased from Merck Chemicals GmbH.

**4.1.2. Analysis.** The samples were analyzed by ASG Analytik-Service AG. The FA content was determined by the sodium sulfite method for concentrations higher than 0.2 wt% and by voltametric analysis for smaller concentrations.<sup>63</sup> The H<sub>2</sub>O content was determined by Karl-Fischer titration. The content of OME<sub>1-10</sub>, MeOH, TRI, tetroxane and MEFO was determined by a gas chromatographic method using flame ionization detection (GC-FID). An online GC equipped with a thermal conductivity detector (GC-TCD) was used for online measurements. The applied methods for the GC analysis were ASG 2506 GC-FID for the organic compounds and ASG 2504 GC-FID for TRI and tetroxane.

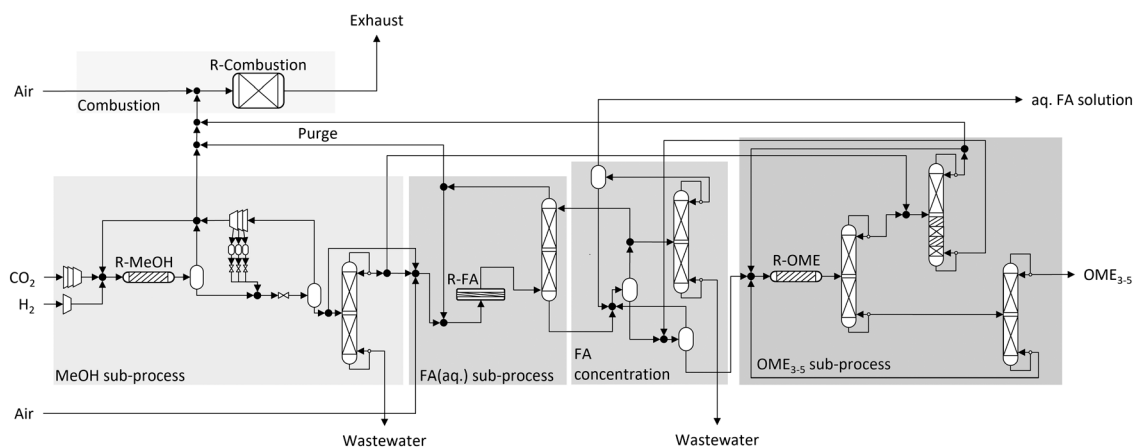


Fig. 5 COMET process concept for the production of OME<sub>3-5</sub> from H<sub>2</sub> and CO<sub>2</sub> feedstock with the intermediate production of MeOH and FA.



## 4.2. Experimental setups

**4.2.1. Concentrated FA(aq.) feed preparation.** A cascade of two thin film evaporators was used to provide the concentrated FA(aq.) solution. The setup was provided and operated by VTA Verfahrenstechnische Anlagen GmbH & Co. KG and directly connected to the OME synthesis setup. As a preparation, the stabilized FA(aq.) containing about 37 wt% FA was concentrated to about 55 wt% FA, which is the typical concentration of the product of a conventional FA(aq.) production plant. The product was stored in a heated tank and further concentrated prior further concentration. Two concentration steps followed. In a first step the FA(aq.) was concentrated to about 75 wt% FA and in a second step to about 85–89 wt% FA. The evaporators were operated under vacuum at about 200–600 mbar and temperatures of the heating fluid of 100–150 °C. A simplified process flow diagram is illustrated in Fig. S9 in the ESI.†

**4.2.2. OME synthesis.** For the synthesis of OME, a fixed bed reactor with a capacity of 1–5 L h<sup>-1</sup> was used. The setup contains a pump and heat exchanger to pressurise and heat up the reactant OME<sub>1</sub> to meet the synthesis conditions, as well as a pump and heated tubes for the concentrated FA(aq.) from the thin film evaporators. After mixing the reactants, the stream was converted in a fixed bed reactor filled with the IER catalyst A46 and heated to about 90 °C. The synthesis product was mixed with additional MeOH, cooled to ambient temperature, depressurized, and directly analyzed using the GC-TCD before it was stored at room temperature. A simplified process flow diagram is illustrated in Fig. S10 in the ESI.†

Before using the catalyst A46, it was stored in a mixture of FA, H<sub>2</sub>O, MeOH and OME<sub>1</sub> to prevent further swelling inside the reactor unit.

The addition of MeOH to the OME synthesis product allowed for a stable storage and transport. The stabilization was a preventive measure to ensure a homogeneous liquid solution without precipitation even at low temperatures and long storage periods. The amount of additional MeOH was determined to meet the demand for the reactive distillation column in an integrated process, see Fig. 3 and the description in section 3. The targets of the reactive distillation column were a bottom product with a concentration of about 60 wt% FA and H<sub>2</sub>O, as well as an almost complete conversion of MeOH. The latter target assumes that OME<sub>2-3</sub> are converted to FA and OME<sub>1</sub> as described by eqn (7), MeOH and FA are converted to OME<sub>1</sub> and H<sub>2</sub>O as described by eqn (1) and (5), and the distillate product is the azeotropic mixture of OME<sub>1</sub> and MeOH at ambient pressure.

**4.2.3. OME synthesis product neutralization.** The OME synthesis product was pumped through a neutralization bed of IER III at ambient temperature and stored prior to the thermal separation in the distillation column.

**4.2.4. Thermal separation in CO-1.** A DN50 glass distillation column with one upper and one lower section of

70 cm height each was used for all thermal separation steps.<sup>64</sup> For the investigation of the first separation step in CO-1, the two sections of the column below and above the feed were filled with Montz 750 structured packings. A horizontal reboiler was used which prevented the flooding of the column in the start-up phase due to strong foaming of the mixture at boiling conditions. The column was continuously operated at ambient pressure with a feed rate of 1–2 L h<sup>-1</sup> for about 250 kg OME synthesis product for about 200 h. After achieving steady state, distillate and bottom products were withdrawn continuously from the column. A simplified process flow diagram is illustrated in Fig. S11 the ESI.†

**4.2.5. Reactive distillation in CO-2.** The core step of the COMET process concept takes place in CO-2. The distillate product of CO-1 was used as the feedstock for CO-2. The same distillation setup of CO-1 was used for the experimental investigation of the reactive distillation CO-2. The lower section of the column was filled with a fixed bed of the catalyst A46 and Montz 750 structured packing on top. The upper section above the feed was filled with Montz 750 structured packing. The column was continuously operated at ambient pressure with a feed rate of 0.5–1 L h<sup>-1</sup> and for about 150 kg feedstock under continuous withdrawal of distillate and bottom product at steady state conditions.

**4.2.6. Thermal separation in CO-3.** The bottom product of CO-1 containing mainly OME<sub>≥3</sub> was fractionated in CO-3. The same distillation setup of CO-1 was used for the experimental investigation of the thermal separation in CO-3 with Montz 750 packings in the upper and lower sections. The setup was continuously operated at about 100 mbar with a feed rate of 2–3 L h<sup>-1</sup> and for about 50 kg feedstock under continuous withdrawal of distillate and bottom product at steady-state conditions.

## 4.3. Process simulation and evaluation

**4.3.1. Process modelling and simulation.** The process simulation software Aspen Plus® V12 was used for the steady-state simulation of the COMET process based on H<sub>2</sub> and CO<sub>2</sub>. Aspen Energy Analyzer V12 was used for heat integration.

The components H<sub>2</sub>, CO<sub>2</sub>, CO, N<sub>2</sub>, O<sub>2</sub>, FA, MeOH, H<sub>2</sub>O, OME<sub>1-10</sub>, HF<sub>1-10</sub> and MG<sub>1-10</sub> were considered. The properties for OME<sub>2-10</sub>, HF<sub>1-10</sub> and MG<sub>1-10</sub> were not included in the Aspen database. Their properties were introduced as new components according to a previous work of our group.<sup>19</sup> The rest of the components properties were adopted from Aspen database. Details are described by Mantei *et al.*,<sup>19</sup> who also describe the UNIFAC-based model which was used as a basis for the process simulation.

Pressure values presented in this work describe the absolute pressure.

The methodology for the process simulation including the individual simulation of all sub-processes, the material integration and interconnection, convergences, the



adjustment of the production capacity and finally the heat integration followed the procedure from Mantei *et al.*<sup>19</sup>

For the simulation of the reactive distillation column, the kinetic model from Schmitz *et al.*<sup>27</sup> for the OME synthesis over A46 for feedstocks comprising MeOH, FA, H<sub>2</sub>O and OME<sub>1</sub> was used. The model was implemented in a Fortran subroutine and activated on the catalyzed trays inside the RadFrac column. The implementation of the kinetic model was validated with the experimental results from Drunsel,<sup>61</sup> who investigated a reactive distillation column in the OME<sub>1</sub> production process for a similar reactive separation task, however, using the catalyst A15 instead of A46. A good agreement was obtained between experimental and simulation results. Furthermore, the subroutine was slightly adjusted to be used in a reactor unit and was validated with the experimental results from Schmitz *et al.*<sup>27</sup> with a good agreement. In contrast to the kinetics of the OME formation as described by eqn (5)–(7), the model assumes the formation of HF<sub>*n*</sub> and MG<sub>*n*</sub> as described by eqn (1)–(4) to be in chemical equilibrium at all retention times.

Regarding the side products HF<sub>*n*</sub> and MG<sub>*n*</sub> for mixtures containing FA, H<sub>2</sub>O and MeOH, the true composition was used for the process simulation, which considers the presence of HF<sub>*n*</sub> and MG<sub>*n*</sub>. The overall composition, considering the stoichiometric decomposition of HF<sub>*n*</sub> and MG<sub>*n*</sub> to their reactants MeOH, H<sub>2</sub>O and FA, was used for the evaluation and the presentation of the results.

The formation of the side products TRI, DME, MEFO, FOAC and tetroxane was not considered in the process simulation, due to very small concentrations in the synthesis product when A46 is used as a catalyst.

**4.3.2. Process evaluation and comparison criteria.** The COMET process was evaluated using various key performance indicators (KPIs). This allows a consistent basis of comparison with other OME<sub>3–5</sub> production processes which are described by Mantei *et al.*<sup>19</sup> Therefore, the same equations were used to determine the process energy efficiency, carbon efficiency and the material selectivity, as described by eqn (19)–(21).

$$\eta_{\text{energy}} = \frac{\dot{m}_{\text{OME}_{3-5}} \cdot \text{LHV}_{\text{OME}_{3-5}}}{\sum_k \dot{Q}_k + \sum_l W_l + \sum_i \dot{m}_i \cdot \text{LHV}_i} \quad (19)$$

$$\eta_C = \frac{C_{\text{OME}_{3-5}}}{\sum_i C_i} \quad (20)$$

$$\eta_{\text{mass}} = \frac{\dot{m}_{\text{OME}_{3-5}}}{\sum_i \dot{m}_i} \quad (21)$$

where  $\dot{m}_i$  denotes the mass flow rate of the reactants  $i$ , LHV denotes the lower heating value at 298 K,  $\dot{Q}_k$  and  $W_l$  represent the externally supplied heat flows and electric power and  $C$  denotes the number of carbon atoms.

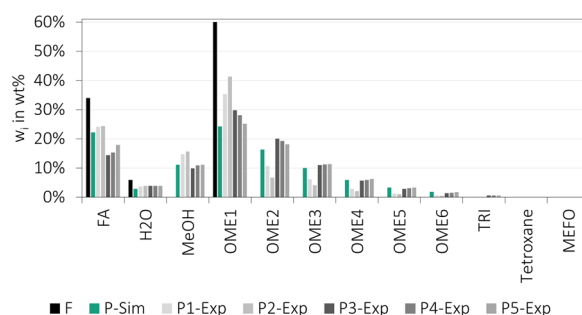
## 5. Results and discussion

### 5.1. Experimental demonstration

In the following sections the experimental results of the demonstration of the main COMET process units are presented and discussed. All investigations were carried using state-of-the-art experimental setups. The FA concentration units and the OME synthesis unit were interconnected. The products were collected and subsequently further processed in the distillation units.

**5.1.1. OME synthesis.** The continuous OME synthesis was investigated experimentally by feeding pure OME<sub>1</sub> and a concentrated FA(aq.) solution in a fixed bed reactor with 2.7–3.5 L h<sup>-1</sup> at about 90 °C and 10 bar over the catalyst A46. Fig. 6 illustrates the composition of the feed mixture (F), the simulated equilibrium product composition (P-Sim), two preliminary experimental products (P1–2-Exp) from the starting phase and the three product barrels (P3–5-Exp) which contained about 250 kg of the OME synthesis product.

The compositions of the three product barrels (P3–5-Exp) show a good agreement with the simulated equilibrium composition (P-Sim). Comparing the three product compositions among each other, a small shift towards longer chain OME<sub>≥3</sub> and FA with increasing time on stream was observed. This is mainly a result of the slightly fluctuating FA concentration in the concentrated FA(aq.) feed stream (85–88 wt% FA) from the evaporation units and the feed stream flowrates. The OME<sub>1</sub> flowrate was regulated to meet a constant ratio between OME<sub>1</sub> and FA, while the FA flowrate was regulated to stabilize the level of the small storage between the second thin film evaporator and the OME synthesis sub-process. Although there was good agreement of the chemical equilibrium between simulation and experiment results, the kinetics of the OME synthesis were predicted to be much faster than those found experimentally. The simulation predicted the chemical equilibrium at a weight hourly space velocity (WHSV, feed mass flow in



**Fig. 6** OME synthesis from OME<sub>1</sub> and concentrated FA(aq.) over A46 (conditions: concentrated FA(aq.) with 85–89 wt% FA, (concentrated FA(aq.))/OME<sub>1</sub> = 0.6 g g<sup>-1</sup>, A46/(OME<sub>1</sub> + concentrated FA(aq.)) = 0.34 g g<sup>-1</sup> h<sup>-1</sup>, approx. 3 L h<sup>-1</sup>, 90 °C, 10 bar, fixed bed reactor). F represents the feed composition. P-Sim, P1–5-Exp represent the product composition of the simulated equilibrium, two experimental preliminary products P1–2 from the starting phase and the three product barrels P3–5, respectively.



relation to the amount of catalyst) of about  $70 \text{ h}^{-1}$ . However, the experiments were carried out at a WHSV of approximately  $16 \text{ h}^{-1}$  and  $3 \text{ h}^{-1}$ , whereby only the lower WHSV was sufficient to obtain chemical equilibrium, as presented by P3–5-Exp in Fig. 6. The WHSV of  $16 \text{ h}^{-1}$  led to high amounts of unreacted FA, low concentrations of  $\text{OME}_{\geq 3}$  and, therefore, solidification of the synthesis product after cooling and without adding MeOH for stabilization, as presented by P1–2-Exp in Fig. 6. The kinetic model from Schmitz *et al.*<sup>27</sup> was used for the simulation, which was initially regressed on experimental results of the OME synthesis from MeOH and FA with partly higher concentrations of  $\text{H}_2\text{O}$  or OME. Therefore, the feed mixture already contained high concentrations of HF which can directly react to OME, as described by eqn (5) and (6). Furthermore, the model was based on the assumption that the reactions towards HF and MG, as described by eqn (1)–(4), are in equilibrium at all retention times since their kinetics are much faster than the kinetics of the formation of OME. In the COMET process this assumption is not met. The concentration of MeOH in the feed is very low because  $\text{OME}_1$  was used as methyl group supplier instead. Therefore, FA is bound mainly in MG which need to depolymerize to be converted to OME, as described in the ESI† in the section about the  $\text{OME}_{3-5}$  production process based on  $\text{OME}_1$  and FA(aq.) or pFA. This is the limiting step for the reaction kinetics of the COMET process but not significant for the OME synthesis based on MeOH and FA. Therefore, the kinetic model<sup>27</sup> is a suitable basis but needs to be further extended to realistically describe the reaction progress of other feed mixtures, which is required to correctly design the reactor unit.

Besides the main components, small fractions of the side products MEFO, TRI and tetroxane were detected in the product barrels P3–5 of about 0.1 wt%, 0.6 wt% and 0.1 wt%, respectively. For P1–2 (during the starting phase) concentrations of about 0.1 wt%, 0.1 wt% and 0.03 wt% were obtained. Therefore, these concentrations strongly depend on the WHSV, which is a matter of investigation for high yields of  $\text{OME}_{3-5}$  at low concentrations of the side product.

**5.1.2. Synthesis product neutralization.** Before the separation of the OME synthesis products P3–5 in the continuous distillation setup, the thermal stability was tested with a similar procedure to the investigations from Mantei *et al.*<sup>33</sup> The pre-tests were conducted in a micro distillation setup using about 50 mL of P3–5 with a stepwise increase of the reboiler temperature. The distillation led to a solidification of the bottom products for all three samples. However, while P3 solidified at about  $130 \text{ }^\circ\text{C}$ , P4 only solidified at about  $170 \text{ }^\circ\text{C}$  and P5 solidified only after heating up to  $170 \text{ }^\circ\text{C}$  and cooling down the bottom product. Additionally, the electric conductivity was measured for the three samples with a reduction from  $6.3 \mu\text{S cm}^{-1}$  for P3,  $3.2 \mu\text{S cm}^{-1}$  for P4 and  $2.1 \mu\text{S cm}^{-1}$  for P5. This indicates that the reduced thermal stability of the OME synthesis product is an initial phenomenon. Nevertheless, for the thermal separation of P3–5 a neutralization was necessary. Further pre-tests were

conducted with different retention times of the IER III in the OME product mixtures for neutralisation. The results indicated that a WHSV of about  $12 \text{ h}^{-1}$  was sufficient to neutralize the OME synthesis product at ambient temperature. This WHSV includes a high safety margin and can probably be increased, especially for the OME synthesis product after longer times on stream of the catalyst. Furthermore, the results showed a connection between the thermal stability of the synthesis product as verified by the micro distillation and the electric conductivity. Below an electric conductivity of  $0.2 \mu\text{S cm}^{-1}$  the synthesis product was thermally stable, and the distillation did not change the product composition. Above  $1.0 \mu\text{S cm}^{-1}$  changes in the composition were detected. Finally, the OME synthesis product of the three product barrels P3–5 was neutralized at ambient temperature in a fixed bed of IER III at a WHSV of about  $12 \text{ h}^{-1}$  using about 500 g IER III. A deactivation of the IER III with increasing time on stream was not observed for the OME synthesis product, of about 200 kg which was continuously neutralized in the fixed bed.

During the investigation of the OME synthesis for about 80 h on stream, stable catalytic activity was observed. Also under reactive distillation conditions, the catalyst performance did not show an obvious deactivation for about 600 h on stream. However, further investigations are required to verify if the changing thermal stability is an initial phenomenon of the catalysts time on stream. In addition to the impact on the process design, the cause of this behavior should be further investigated. It might only be the leaching of the catalyst as emphasized by Fink *et al.*<sup>65,66</sup> and Baranowski *et al.*,<sup>67</sup> but it could also be influenced by the side product formation, especially formic acid, which was not analyzed in this work but reported in the literature.<sup>32</sup>

**5.1.3. Synthesis product separation in CO-1.** After the neutralization of the OME synthesis products P3–5, the distillation of the volatile components FA,  $\text{H}_2\text{O}$ , MeOH and  $\text{OME}_{1-2}$  from  $\text{OME}_{\geq 3}$  was investigated in the distillation setup. The results illustrated in Fig. 7 are an example but representative result of the continuous distillation

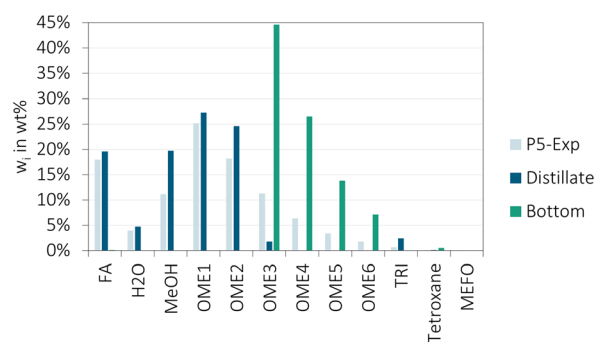


Fig. 7 CO-1, OME synthesis product separation (conditions:  $2 \text{ L h}^{-1}$ , reflux/distillate =  $0.5\text{--}2 \text{ s}^{-1}$ , distillate/feed = 81 wt%, Montz 750 structured packing,  $85\text{--}175 \text{ }^\circ\text{C}$ , ambient pressure). The values describe the mass fractions of the feed mixture, here P5-Exp as presented in Fig. 6, the distillate product and bottom product.



experiment and show that the targeted separation between OME<sub>2</sub> and OME<sub>3</sub> was successfully realized. It also shows that FA, H<sub>2</sub>O and MeOH can be separated from OME<sub>≥3</sub>. The side products MEFO and TRI are also separated from the bottom product, but the side product tetroxan has a higher boiling point than OME<sub>3</sub> and stays in the bottom product.

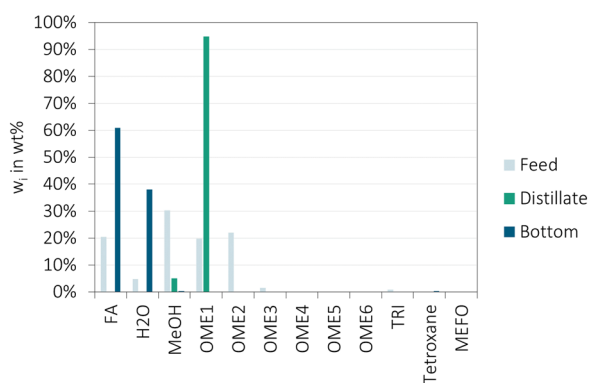
The distillation setup was operated at a feed temperature of 87 °C, a condensation temperature of 85 °C and a reboiler temperature of 175 °C. The distillate to feed ratio was about 0.81 and the time-based reflux ratio was varied as a controlled variable between 0.5–2 s s<sup>-1</sup> (time controlled) to achieve a constant condensation temperature.

Fig. 7 shows that OME<sub>2</sub> was completely separated to the distillate product. However, also a small fraction of OME<sub>3</sub> went to the distillate product, which was about 14% of the feed amount of OME<sub>3</sub>. Besides OME<sub>≥3</sub>, traces of FA, H<sub>2</sub>O and MeOH were detected in the bottom product which were mainly below 0.6 wt%. MEFO was not detected in the bottom product.

Regarding the continuous operation of the distillation setup, an increasing precipitation of FA inside the condenser was challenging in the initial phase but could be prevented by increasing the temperature of the cooling fluid to above 25 °C. However, as a result the temperature difference decreased between the cooling fluid and the boiling points of the most volatile components MEFO, the azeotropic mixture of OME<sub>1</sub> and MeOH, as well as OME<sub>1</sub>. Thus, the area of the condenser was relatively small to obtain a complete condensation and a small fraction of the most volatile components accumulated in a cool trap. As a result, the ratio of OME<sub>1</sub> to OME<sub>2</sub> in the feed mixture P5-Exp differs from the ratio of OME<sub>1</sub> to OME<sub>2</sub> in the distillate product.

**5.1.4. Reactive distillation in CO-2.** After the separation of the OME synthesis product in CO-1, the distillate product of CO-1 was separated and converted in a reactive distillation column CO-2.

A representative result of the continuous reactive distillation experiment is illustrated in Fig. 8. The distillate



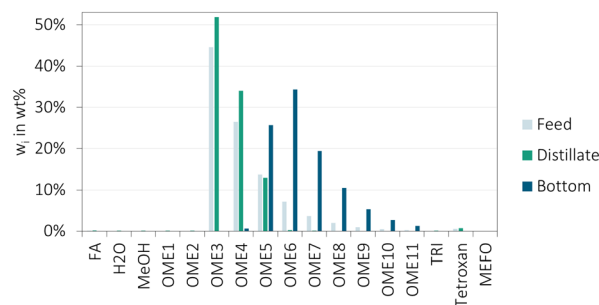
**Fig. 8** CO-2, reactive distillation of the distillate product of CO-1 over A46 (conditions: A46/(feed stream) = 0.35 g g<sup>-1</sup> h<sup>-1</sup>, 1 L h<sup>-1</sup>, distillate/feed = 63 wt%, Montz 750 structured packing, 45–104 °C, ambient pressure). The values describe the mass fractions of the feed mixture, the distillate product and bottom product.

and bottom product compositions show that the targets of the reactive distillation column were obtained. OME<sub>≥2</sub> were converted to OME<sub>1</sub> and FA, the composition of the distillate product is the azeotropic mixture of OME<sub>1</sub> and MeOH and the bottom product contains mainly FA and H<sub>2</sub>O. Regarding the bottom product composition, small concentrations of MeOH of about 0.3 wt% were detected besides the desired range of FA and H<sub>2</sub>O. Furthermore, traces of OME<sub>1–6</sub> were detected with concentrations far below 0.1 wt%. However, as a result of the high H<sub>2</sub>O and FA content in the bottom product, the quantification of traces is complex due to the fast precipitation of the bottom product solidifies fast if not heated or diluted.

CO-2 was operated at a condensation temperature of 45 °C and reboiler temperature of 104 °C with a distillate to feed ratio of about 0.63.

The results confirm that the reactive distillation column is a feasible instrument for the separation of H<sub>2</sub>O from the OME<sub>3–5</sub> production loop of the OME<sub>3–5</sub> production and that an almost complete conversion of MeOH can be achieved. Furthermore, the results indicate, that the variation of the amount of MeOH in the feed mixture to the reactive distillation column can be used to tune the amount of OME<sub>1</sub> produced as the distillate product.

**5.1.5. Product separation in CO-3.** The representative results of the continuous distillation experiment for the separation of the final product mixture OME<sub>3–5</sub> from the bottom product of the distillation column CO-1 are illustrated in Fig. 9. The target was a cut between OME<sub>5</sub> and OME<sub>6</sub>. A significant amount of OME<sub>5</sub> stayed in the bottom product, which, however, can be separated to the distillate product by increasing the reboiler temperature or reducing the operational pressure. Regarding the distillate product, besides OME<sub>3–5</sub>, small fractions of OME<sub>6</sub> of 0.2 wt%, tetroxan of 0.7 wt% and traces of FA and H<sub>2</sub>O were detected. The concentration of tetroxan is mainly a result of the retention time, temperature, and selection of catalyst in the reactor, which can be improved to reduce the side product formation. However, the pre-standard DIN/TS 51699 (ref. 13) does not limit the concentration of tetroxan. The concentration of TRI is limited to 0.1 wt% and was detected smaller than 0.01



**Fig. 9** CO-3, product separation (conditions: 5.5 L h<sup>-1</sup>, distillate/feed = 82 wt%, Montz 750 structured packing, 100–210 °C, 200 mbar). The values describe the mass fractions of the feed mixture, here the CO-1 bottom product, the distillate product and bottom product.



wt% in the distillate product of CO-3. The final OME<sub>3-5</sub> product from the COMET process was compatible with the pre-standard DIN/TS 51699.

Due to the solidification of the bottom product OME<sub>≥5</sub> at room temperature, it was diluted in THF with a ratio of 1:10 g/g to enable the GC analysis. However, this also impairs the detection limits and accuracy of the analysis. Alternatively, to liquify the bottom product it can also be heated up. At 80 °C, the bottom product is completely liquid, which enables its recycling to the reactor, as illustrated in Fig. 3.

The distillation setup was operated at a condensation temperature of 140 °C and a reboiler temperature of 210 °C. The high distillate temperature was a result of the high feed flow rate and the limited area for condensation. As a result, a complete condensation was not obtained and small fraction of OME<sub>3</sub> accumulated in a cool trap. In contrast to the other distillation experiments the operation pressure was reduced to 200 mbar to reduce the reboiler temperature for the separation between OME<sub>5</sub> and OME<sub>6</sub>. The distillate to feed ratio was about 0.82.

## 5.2. Process simulation and evaluation

**5.2.1. Mass balance.** Stream compositions and conditions of the COMET process simulation in Aspen Plus are listed in Table 4, following the stream numbering of Fig. 3. Stream compositions and conditions for all sub-processes are presented in the ESI.†

The feedstock (stream 1) containing about 50 wt% FA and 49 wt% H<sub>2</sub>O is mixed with the distillate of the second evaporator E-2 and the bottom of the third evaporator E-3. The mixture is concentrated in a cascade of two evaporators E-1 and E-2 operated at 400 and 500 mbar respectively and low retention times. The pressure levels were selected to obtain similar evaporation and condensation temperatures as experimentally verified. However, in practice, the pressure level might be lower to achieve the desired concentrations.

This is a result of the simplified modelling of the evaporators which require more detailed considerations of the reaction kinetics of eqn (1)–(4) as recently introduced by Tönges and Burger.<sup>68</sup> The FA concentration is similar to the conventional production of pFA and generates a concentrated FA solution containing about 88 wt% FA (stream 5) and a solution containing about 18 wt% FA (stream 3). Stream 3 is split to be used as a washing liquid for the FA absorber column and to be purified in the distillation column CO-4 operated at 5.5 bar to pure H<sub>2</sub>O (stream 4) (<200 ppm FA) and a concentrated FA solution with 44 wt% FA. To prevent the accumulation of MeOH and other impurities in the loop, the concentrated FA solution is sent to another evaporator E-3 operated at ambient pressure. This prepares a by-product of the COMET process with a higher MeOH concentration (stream 2) and a FA solution with a similar composition to the FA feedstock, which is recycled to the evaporator cascade. The by-product (stream 2) has a low FA concentration of about 14 wt%. Furthermore, its mass flow is about 17.6% of the mass flow of the target OME<sub>3-5</sub> product. This high mass flow is similar to alternative OME<sub>3-5</sub> production processes using FA(aq.) solution as an intermediate product.<sup>19</sup>

The concentrated FA product (stream 5) is pressurized to about 10 bar, then mixed with the recycle streams containing mainly OME<sub>1</sub> and OME<sub>>5</sub> (stream 10 and stream 14) and converted to OME in a fixed bed reactor at about 10 bar and 90 °C, over A46 catalyst, as used for the experimental demonstration. The reactor product contains about 20 wt% OME<sub>3-5</sub>, which is relatively high in comparison to the process based on MeOH and FA(aq.), which has 0 to 15 wt% OME<sub>3-5</sub> in the reactor product,<sup>39</sup> as presented in Table 8. The reactor product is purified in a first distillation column CO-1 operated at a slight overpressure of 1.8 bar, where OME<sub>≥3</sub> are separated from FA, MeOH, H<sub>2</sub>O, OME<sub>1-2</sub> and a small fraction of OME<sub>3</sub>. The slight overpressure improves the separation efficiency and reduces the losses of OME<sub>3</sub> to the distillate product (stream 8) to about 12%. The FA concentration of the

**Table 4** Stream compositions and conditions of the COMET process presented in Fig. 3

Stream	Overall mass fractions													
	1	2	3	4	5	6	7	8	9	10	11	12	13	14
<i>T</i> in °C	64.9	30.0	90.0	30.0	90.4	90.0	90.0	81.5	81.0	41.5	117.4	200.5	86.6	194.9
<i>p</i> in bar	1.0	1.0	0.3	1.0	10.3	10.0	10.1	1.8	1.8	1.0	1.0	1.8	0.07	0.07
<i>m</i> in kg h <sup>-1</sup>	18 509	2203	14 753	7870	22 957	66 666	66 666	51 796	5288	41 330	15 753	14 871	12 490	2380
FA	0.502	0.142	0.184	0.000	0.880	0.303	0.186	0.239	0.000	0.000	0.727	0.000	0.000	0.000
H <sub>2</sub> O	0.491	0.778	0.796	1.000	0.120	0.042	0.022	0.028	0.000	0.002	0.268	0.000	0.000	0.000
MeOH	0.007	0.080	0.020	0.000	0.000	0.028	0.100	0.129	1.000	0.045	0.005	0.000	0.000	0.000
OME <sub>1</sub>	0.000	0.000	0.000	0.000	0.000	0.591	0.276	0.356	0.000	0.953	0.000	0.000	0.000	0.000
OME <sub>2</sub>	0.000	0.000	0.000	0.000	0.000	0.000	0.179	0.230	0.000	0.000	0.000	0.000	0.000	0.000
OME <sub>3</sub>	0.000	0.000	0.000	0.000	0.000	0.000	0.107	0.017	0.000	0.000	0.000	0.419	0.499	0.000
OME <sub>4</sub>	0.000	0.000	0.000	0.000	0.000	0.000	0.061	0.000	0.000	0.000	0.000	0.271	0.323	0.000
OME <sub>5</sub>	0.000	0.000	0.000	0.000	0.000	0.000	0.033	0.000	0.000	0.000	0.000	0.149	0.177	0.000
OME <sub>6</sub>	0.000	0.000	0.000	0.000	0.000	0.018	0.018	0.000	0.000	0.000	0.000	0.080	0.000	0.496
OME <sub>7</sub>	0.000	0.000	0.000	0.000	0.000	0.009	0.009	0.000	0.000	0.000	0.000	0.042	0.000	0.262
OME <sub>8</sub>	0.000	0.000	0.000	0.000	0.000	0.005	0.005	0.000	0.000	0.000	0.000	0.022	0.000	0.136
OME <sub>9</sub>	0.000	0.000	0.000	0.000	0.000	0.002	0.002	0.000	0.000	0.000	0.000	0.011	0.000	0.070
OME <sub>10</sub>	0.000	0.000	0.000	0.000	0.000	0.001	0.001	0.000	0.000	0.000	0.000	0.006	0.000	0.036



**Table 5** Overall mass balance for the production of OME<sub>3-5</sub> following the COMET process and the processes P1 and P4 by Mantei *et al.*<sup>19</sup> The processes were simulated with a capacity of 100 kilotons per annum OME<sub>3-5</sub>

	COMET	P1	P4
Total input in kg kg <sub>OME<sub>3-5</sub></sub> <sup>-1</sup>	6.60	7.54	8.53
H <sub>2</sub>	0.25	0.27	0.21
CO <sub>2</sub>	1.82	1.96	2.20
Air	4.53	5.32	5.92
Total output in kg kg <sub>OME<sub>3-5</sub></sub> <sup>-1</sup>	6.60	7.54	8.53
OME <sub>3-5</sub>	1.00	1.00	1.00
OME <sub>3</sub>	0.50	0.43	0.43
OME <sub>4</sub>	0.32	0.38	0.36
OME <sub>5</sub>	0.18	0.19	0.21
Wastewater	1.03	1.30	1.00
Aq. FA solution	0.18	—	—
Exhaust gas	4.39	5.24	6.54

bottom product (stream 12) is reduced to about 100 ppm. In the third distillation column CO-3 operated at 70 mbar, the main product OME<sub>3-5</sub> (stream 12) is extracted from the process with about 50 wt% OME<sub>3</sub>, 32 wt% OME<sub>4</sub>, 18 wt% OME<sub>5</sub> and traces of FA and H<sub>2</sub>O in compliance with the pre-standard DIN/TS 51699 specifications. The distillate product (stream 8) of the first distillation column CO-1 is mixed with MeOH (stream 9) and introduced to the reactive distillation column CO-2. This column is operated at ambient pressure. The selection of the pressure level is a compromise between the condensation temperature of the distillate, the reaction kinetics on the catalytic trays and the composition of the azeotropic mixture of OME<sub>1</sub> and MeOH in the distillate. A pressure reduction would favorably improve the azeotropic composition to higher OME<sub>1</sub> concentrations. However, it would also lead to a reduction of the condenser temperature below 41 °C which can lead to more expensive cooling utilities and decelerate the reaction kinetics on the catalytic trays. Increased pressure levels would benefit from higher reaction kinetics due to the higher temperature level on the catalytic trays, but lower OME<sub>1</sub> concentrations in the distillate product. This would decrease the OME<sub>3-5</sub> selectivity in the OME synthesis reactor and necessarily increase the recycle streams and, therefore, the specific heat demand for product purification. The mixture is separated into the azeotropic mixture of 95 wt% OME<sub>1</sub> and 4.5 wt% MeOH in

**Table 6** Overall energy demand for the production of OME<sub>3-5</sub> following the COMET process and the processes P1 and P4 by Mantei *et al.*<sup>19</sup> The processes were simulated with a capacity of 100 kilotons per annum OME<sub>3-5</sub>

	COMET	P1	P4
Total input kW h kW <sup>-1</sup> h <sub>OME<sub>3-5</sub>,LHV</sub> <sup>-1</sup>	1.60	1.70	1.34
H <sub>2</sub>	—	—	—
Total output kW h kW <sup>-1</sup> h <sub>OME<sub>3-5</sub>,LHV</sub> <sup>-1</sup>	1.00	1.00	1.00
OME <sub>3-5</sub>	—	—	—
Energy demand kW h kW <sup>-1</sup> h <sub>OME<sub>3-5</sub>,LHV</sub> <sup>-1</sup>	—	—	—
Electricity	0.11	0.09	0.14
LPS, 4 bar	0.09	-0.10	0.24
MPS, 23 bar	0.05	0.30	-0.07
Cooling water	-1.02	-1.05	-0.79
Heat. T > 250 °C	—	—	0.19

**Table 7** Overall process efficiencies for the production of OME<sub>3-5</sub> following the COMET process and the processes P1 and P4 by Mantei *et al.*<sup>19</sup> The processes were simulated with a capacity of 100 kilotons per annum OME<sub>3-5</sub>

	COMET	P1	P4
$\eta_{\text{energy}}$ in %	54.1	50.3	54.4
$\eta_{\text{C}}$ in %	88.1	81.6	72.5
$\eta_{\text{mass}}$ in %	41.1	38.1	41.4

the distillate (stream 10) and a mixture of 73 wt% FA and 27 wt% H<sub>2</sub>O in the bottom (stream 11). A summary of the overall mass balance of the COMET process is listed in Table 5.

For the comparison with alternative OME<sub>3-5</sub> production processes the results from Mantei *et al.*<sup>19</sup> are presented for some selected processes, namely: P1, which produced OME<sub>3-5</sub> *via* MeOH and a concentrated FA solution (aqueous OME synthesis) feedstock, and P4 which produces OME<sub>3-5</sub> *via* OME<sub>1</sub> and monomeric FA (anhydrous OME synthesis) feedstock. The processes are described in detail in the ESI† in section 1.4 and 1.5.

The results show that the overall COMET process requires less H<sub>2</sub> than P1 but more H<sub>2</sub> than P4. The difference to P1 is mainly based on the FA concentration sub-process, in which the simulation of this work contains a modified separation of FA from H<sub>2</sub>O due to the addition of a distillation column and a third evaporator. This results in a smaller amount of FA, which exits the process in the form of an aqueous FA solution by-product stream (see stream 2 in Fig. 3).

The difference to P4 is mainly based on the advantages of the anhydrous FA synthesis from MeOH which produces H<sub>2</sub> as a by-product which can be separated and recycled to the MeOH sub-process. The lower CO<sub>2</sub> demand of the COMET process in comparison to P1 and P4 is also based on the FA concentration sub-process and the anhydrous FA synthesis. P1 requires more CO<sub>2</sub> due to the higher amount of FA in the by-product stream (see stream 2 in Fig. 3). The lower demand of air of the COMET process is mainly a result of the consideration of smaller purge streams which are oxidized in the combustion sub-process. The oxygen demand for the partial oxidation of MeOH towards FA(aq.) is only slightly lower for the COMET process than for P1.

The composition of the final OME<sub>3-5</sub> product mixture also shows significant differences. While Mantei *et al.*<sup>19</sup> chose a composition close to the highest yield of OME<sub>3-5</sub> after the synthesis, the composition in this work was selected to meet the requirements for the pre-standard DIN/TS 51699.

Regarding the wastewater production, the COMET process produces less wastewater than P1 but more wastewater than P4. The difference to P1 is mainly based on the composition of the wastewater. While the simulation of the COMET process produces high-purity wastewater and an aqueous FA solution by-product, Mantei *et al.*<sup>19</sup> considered the aqueous FA solution to be part of the wastewater. The difference to P4 is also explained by the anhydrous FA synthesis.<sup>19</sup>



**Table 8** Comparison of various OME<sub>3-5</sub> production processes based on the results of Held *et al.*,<sup>39</sup> Mantei *et al.*<sup>19</sup> and Schemme *et al.*<sup>43</sup>

Feedstock	Anhydrous synthesis			Aqueous synthesis			COMET
	OME <sub>1</sub> and TRI	DME and TRI	OME <sub>1</sub> and monomeric FA	MeOH and FA(aq.)	OME <sub>1</sub> and FA(aq.) or pFA	MeOH and monomeric FA	
w <sub>OME<sub>3-5</sub></sub> in wt%	5, 34 <sup>39</sup> 0, 35 <sup>43</sup>	— <sup>a</sup>	5, 29 <sup>19</sup>	0, 15 <sup>39</sup> 4, 16 <sup>19</sup> 0, 14 <sup>43</sup>	4, 19 <sup>19</sup>	3, 19 <sup>19</sup>	0, 20
$Q_{\text{Reboiler}}/H_{\text{OME}_{3-5}}$ in $kW_{\text{Heat}}/kW_{\text{OME}_{3-5}}$ in	7.6% <sup>39</sup> 5.5% <sup>43</sup>	— <sup>a</sup>	15% <sup>19</sup>	47% <sup>39</sup> 39% <sup>19</sup> 78% <sup>43</sup>	26% <sup>19</sup>	48% <sup>19</sup>	35%
$\eta_{\text{energy,overall}}$ in %	29–37 <sup>39</sup> 22–26 <sup>43</sup>	— <sup>a</sup>	27–36 <sup>19</sup>	30–36 <sup>39</sup> 25–31 <sup>19</sup> 24–29 <sup>43</sup>	26–32 <sup>19</sup>	28–37 <sup>19</sup>	28–34
Scale-up potential in the near future	Likely	Unlikely	Unlikely	Less likely	Less likely	Unlikely	Likely

<sup>a</sup> Further investigations and an adjusted process concept are required to estimate the process performance.

The exhaust gas flow is lower for the COMET process than for P1 and P4 which is the result of the smaller purge streams and, therefore, the lower air demand for the combustion.

**5.2.2. Energy demand.** The specific energy demand and operation conditions for the main process units evaluated by the COMET process simulation are listed in the ESI.†

The summary of the overall energy demand of the COMET process after the heat integration in comparison to P1 and P4 is listed in Table 6.

The different H<sub>2</sub> demands between the COMET process, P1 and P4 directly reflect on the total process energy demand. Furthermore, the electricity demand of the COMET process is higher than for P1 but lower than for P4. Compared to P1, the operation conditions of the phase separators in the MeOH sub-process were adjusted resulting in higher recycling rates and, therefore, higher compression demand. Furthermore, P1 and P4 did not consider the compression demand for the combustion sub-process.<sup>19</sup> The higher electricity demand of P4 is a result of the anhydrous FA synthesis, which requires higher recycle streams in comparison to the partial oxidation of MeOH in P1 and the COMET process.<sup>19</sup>

The demand for low pressure steam (LPS) of the COMET process is higher than the demand of P1 but lower than the demand of P4, which is mainly a result of the heat integration strategies. P1 generates more LPS than it consumes, while P4 and the COMET process show higher demands than generated. However, the demand for medium pressure steam (MPS) is lower for the COMET process. P1 shows a very high demand of MPS, while P4 generates more than it consumes. The MPS demand for the COMET process is significantly lower than for P1. The main consumers of MPS are CO-1 and CO-3 in the OME<sub>3-5</sub> sub-process. However, MPS is also generated in the MeOH synthesis reactor and the combustion sub-process. While the combustion sub-process generates a similar amount of MPS of about  $-0.15 \text{ kW h kW}^{-1} h_{\text{OME}_{3-5},\text{LHV}}^{-1}$  comparing P1 and the COMET process, the amount differs for the MeOH

reactor with  $-0.04$  and  $-0.11 \text{ kW h kW}^{-1} h_{\text{OME}_{3-5},\text{LHV}}^{-1}$ , respectively. The lower MPS generation of P1 is mainly a result of the inlet temperature to the MeOH reactor. The inlet temperature of P1 is about 185 °C and, therefore, needs to be heated up to the operation temperature of 250 °C using generated MPS and the exothermic heat of the methanol synthesis reactor. The inlet temperature of the COMET process simulation is about 240 °C, which requires a larger heat transfer area but improves the energy efficiency. Furthermore, the demand for MPS of the distillation columns in the OME<sub>3-5</sub> sub-process differ significantly. P1 requires about  $0.32 \text{ kW h kW}^{-1} h_{\text{OME}_{3-5},\text{LHV}}^{-1}$  of MPS for the purification of the OME<sub>3-5</sub> product stream, while the COMET process requires only  $0.20 \text{ kW h kW}^{-1} h_{\text{OME}_{3-5},\text{LHV}}^{-1}$ . This is mainly a result of the higher OME<sub>3-5</sub> yield of the COMET process after the OME synthesis reactor as discussed in the previous section.

The demand for cooling water is similar between P1 and the COMET process but significantly lower for P4.<sup>19</sup> The cooling is required mainly for the temperature level between 90 °C and 30 °C and is, therefore, hardly utilizable for the heat integration of the COMET process. Mantei *et al.*<sup>69</sup> proposed the utilization of heat pumps instead of cooling water and evaluated a significant enhancement potential for the overall energy efficiency.

Only P4 has a demand for heat above 250 °C, due to the endothermic anhydrous FA synthesis.

Regarding the heat demand for the separation of H<sub>2</sub>O *via* reactive distillation, the COMET process requires about  $1.1 \text{ kWh kg}^{-1} \text{ H}_2\text{O}$  at 117 °C. This is based on the assumption that the main target of the reactive distillation column is the separation of H<sub>2</sub>O from the loop. Therefore, the heat demand of the reboiler and the feed preheater can be allocated to the amount of H<sub>2</sub>O separated from the loop.

**5.2.3. Process efficiencies.** The summary of the overall process efficiencies of the COMET process after the heat integration is listed in Table 7 and compared with P1 and P4.

The overall energy efficiency of the COMET process is higher than P1 and similar to P4. Mantei *et al.*<sup>19</sup> reported an



efficiency of 49–50% for processes considering the FA(aq.) sub-process. Furthermore, the carbon efficiency is considerably higher than P1 or P4. The low carbon efficiency of P4 is mainly a result of the high CO side-product formation during the anhydrous FA synthesis.<sup>19</sup> While the lower carbon efficiency of P1 is mainly a result of the more efficient H<sub>2</sub>O separation of the FA concentration sub-process considered for the COMET process simulation. As a result, the carbon efficiency of P1 could also be increased by adjusting the FA concentration sub-process that can be considered in a future work. The OME<sub>3–5</sub> yield based on the feedstock H<sub>2</sub> and CO<sub>2</sub> is also higher for the COMET process than for P1 which is also a result of the more efficient H<sub>2</sub>O separation of the FA concentration sub-process. The OME<sub>3–5</sub> yield is similar to P4 since H<sub>2</sub>O, formed in the FA(aq.) synthesis, is separated from the process loop, compared to the formation, separation and recycling of H<sub>2</sub> in the anhydrous FA synthesis of P4.

**5.2.4. Comparison to alternative OME<sub>3–5</sub> production processes.** To compare alternative OME<sub>3–5</sub> production process concepts, important performance parameters are listed in Table 8. These performance parameters include the mass fraction of OME<sub>3–5</sub> before and after the OME reactor  $w_{\text{OME}_{3–5}}$ . Furthermore, the heat demand of the reboiler of the distillation columns and feed preheaters of the OME sub-process  $Q_{\text{Reboiler}}$  in relation to the OME<sub>3–5</sub> product mass flow times its LHV  $H_{\text{OME}_{3–5}}$  is considered. Another key performance parameter is the overall energy efficiency  $\eta_{\text{energy,overall}}$ . This considers the entire process chain starting from H<sub>2</sub>O electrolysis and CO<sub>2</sub> via the production of the intermediate products towards the target product mixture OME<sub>3–5</sub>.

Regarding the electricity and heat demand for the H<sub>2</sub>O electrolysis and CO<sub>2</sub> preparation, the assumptions from Held *et al.*<sup>39</sup> were considered. For the CO<sub>2</sub> preparation all three scenarios from Held *et al.*<sup>39</sup> were considered, comprising CO<sub>2</sub> from point sources (CPS), post combustion capture (PCC) using mono-ethanol amine scrubbing and direct air capture (DAC). The key assumptions for the expanded system boundary evaluation are summarized in ESI† in Table S28. In addition, the scale-up potential in the near future is qualitatively evaluated. The key performance parameters are based on the results from Held *et al.*,<sup>39</sup> Mantei *et al.*<sup>19</sup> and Schemme *et al.*<sup>43</sup>

Regarding the yield of OME<sub>3–5</sub> after the reactor as illustrated in Fig. 1, the anhydrous process concepts show far higher OME<sub>3–5</sub> concentrations than the aqueous process concepts, as indicated by  $w_{\text{OME}_{3–5}}$ . This also reflects on the heat demand for the OME<sub>3–5</sub> product purification, which is compared based on  $Q_{\text{Reboiler}}/H_{\text{OME}_{3–5}}$ . The anhydrous process concepts show significantly lower heat demands for the product purification. Exceptions are the production based on OME<sub>1</sub> and FA(aq.) or pFA and the COMET process, which despite comparatively low yields of OME<sub>3–5</sub> in the reactor require less heat for the separation of the target product than the other aqueous process concepts. For a consistent basis of comparison, the production of the intermediate products for

different OME production processes was considered in the evaluation, indicated by  $\eta_{\text{energy,overall}}$ . The result is a low overall energy efficiency of <40% for all processes, with minor differences between anhydrous and aqueous process concepts. Greater differences were reported between different literature sources, which is especially significant comparing the results for the OME<sub>3–5</sub> production based on OME<sub>1</sub> and TRI as well as MeOH and FA(aq.) from Held *et al.*,<sup>39</sup> Mantei *et al.*<sup>19</sup> and Schemme *et al.*<sup>43</sup> Those differences are discussed in detail by Mantei *et al.*<sup>19</sup> and mainly result from different heat integration strategies and the simulation procedure. Schemme *et al.*<sup>43</sup> only integrated the heat between individual sub-processes, while Mantei *et al.*<sup>19</sup> and Held *et al.*<sup>39</sup> considered the heat integration between all sub-processes. However, Held *et al.*<sup>39</sup> did use stoichiometric material balances and literature data, while Mantei *et al.*<sup>19</sup> and Schemme *et al.*<sup>43</sup> used the process simulation software Aspen Plus.

Regarding the low overall energy efficiency of all OME<sub>3–5</sub> production processes, Mantei *et al.*<sup>69</sup> showed the potential of including high temperature heat pumps (HTHP) to lift the temperature of the excess heat streams and, therefore, supply internal heat demands and, in addition, external heat demands. This strategy has the potential to lift the overall energy efficiency above 61% considering heat as a valuable by-product of the process. Besides only small differences in the energy efficiency, the production costs of the OME<sub>3–5</sub> product also show no significant differences between different production processes.<sup>19,43</sup>

Regarding a sustainable large-scale production of OME<sub>3–5</sub> in the near future the COMET process and a production based on OME<sub>1</sub> and TRI feedstock show great potential to be scaled up today. However, the latter is comparatively complex, comprising five sub-processes for the production of MeOH, FA(aq.), TRI, OME<sub>1</sub> and OME<sub>3–5</sub>. A sustainable OME<sub>3–5</sub> production based on the COMET process on the other hand comprises three sub-processes for the production of MeOH, FA(aq.) and OME<sub>3–5</sub>. The OME production process based on DME and TRI requires further investigations, mainly due to the high MEFO formation during the synthesis and the low activities for the conversion of DME to OME. A fast scale-up of the processes based on monomeric FA is mainly prevented by the low TRL of the monomeric FA production. Finally, the aqueous process concepts require the separation of H<sub>2</sub>O from the loop of the OME<sub>3–5</sub> sub-process, which is the main bottleneck for a fast scale-up. Various concepts for separating H<sub>2</sub>O from the loop were already proposed, and some show promising results, as discussed before and demonstrated in this work for the reactive distillation column. This enables a scale-up for the processes based on MeOH and FA(aq.) and OME<sub>1</sub> and FA(aq.) or pFA. In comparison to the OME<sub>3–5</sub> production based on OME<sub>1</sub> and TRI, the aqueous process concepts enable a considerable simplification, which typically improves the robustness and therefore feasibility for large-scale application.



## 6. Conclusion

The COMET process, which solves the challenging H<sub>2</sub>O management of aqueous OME processes, was introduced in this work. The process benefits from a simple feedstock preparation, a short process chain from H<sub>2</sub> and CO<sub>2</sub> to OME<sub>3-5</sub>, a comparatively high OME<sub>3-5</sub> yield after the reactor, and the possibility of extracting the by-product H<sub>2</sub>O from the loop using a state-of-the-art reactive distillation unit.

Other H<sub>2</sub>O separation methods, as discussed in the literature, were presented, and their main advantages and hurdles were evaluated quantitatively. The main advantage of the H<sub>2</sub>O separation in the COMET process *via* reactive distillation is the scale-up potential and the feasible application in large-scale production plants.

Starting solely from MeOH and FA(aq.) commercial feedstocks, the main COMET process units, comprising all evaporation, reaction and separation process steps, were experimentally demonstrated on a pilot scale. Importantly, the technical feasibility of the reactive distillation column – the heart of the COMET process concept – was demonstrated for a long duration of around 600 h on stream. In addition, the purification of the final OME<sub>3-5</sub> product was successfully realized with a product compliant with the pre-standard DIN/TS 51699.

The COMET process was simulated and evaluated using Aspen Plus and compared with relevant alternative OME<sub>3-5</sub> production processes. Therefore, the system boundary was expanded, including H<sub>2</sub> production *via* H<sub>2</sub>O electrolysis, CO<sub>2</sub> capture and all intermediate production sub-processes. With an overall energy efficiency of 28–34%, depending on the CO<sub>2</sub> source, the energy demand of the COMET process is similar to the alternative OME<sub>3-5</sub> production processes, in which overall energy efficiencies were evaluated in the range of 25–36%. Moreover, the COMET process shows a higher carbon efficiency of 88%.

The OME market is limited by the lack of technologically feasible large-scale processes. However, compared to relevant alternative OME<sub>3-5</sub> production processes, the novel COMET process shows the smallest technological hurdles and can already be demonstrated and scaled up.

## Symbols

$C$ [—]	Number of carbon atoms
$H$ [kW]	Energy content based on the LHV
$m, \dot{m}_i$ [kg h <sup>-1</sup> ]	Mass flow rate
$p$ [bar]	Pressure
$\dot{Q}_k$ [kW]	Heat flow
$T$ [K]	Temperature
$w_i$ [wt%]	Mass fraction
$W_i$ [kW]	Electric power
$\eta$ [%]	Efficiency

## Sub- and superscripts

C	Carbon
$i$	Reactant, component

## Abbreviations

A15	Amberlyst® 15
A46	Amberlyst® 46
CO	Distillation column
COMET	Clean OME technology
DAC	Direct air capture
DME	Dimethyl ether
EGR	Exhaust gas recirculation
EPDM	Ethylene propylene diene monomer
FA	Formaldehyde
FA(aq.)	Aqueous FA solution, formalin
FFKM	Perfluoroelastomer
FOAC	Formic acid
GC-FID	Gas chromatograph equipped with a flame ionization detector
GC-TCD	Gas chromatograph equipped with a thermal conductivity detectors
HF	Poly(oxyethylene) hemiformals
HTHP	High temperature heat pump
HVO	Hydrogenated vegetable oil
IER	Ion exchange resin
LHV	Lower heating value
LPS	Low pressure steam
MEFO	Methyl formate
MeOH	Methanol
MG	Poly(oxyethylene) glycols
MPS	Medium pressure steam
OME	Oxymethylene dimethyl ethers
OME <sub>1</sub>	Methylal
pFA	Paraformaldehyde
PTFE	Polytetrafluoroethylene
PtL	Power-to-liquid
PtX	Power-to-X
R	Reactor
S	Separator
THF	Tetrahydrofuran
TRI	Trioxane
TRL	Technology readiness level
WtW	Well-to-wheel

## Author contributions

Conceptualization, F. M. and O. S.; methodology, F. M. and C. S.; experimental investigation, F. M., C. S. and O. S.; analytics, F. M., C. S. and H. S.; simulation and evaluation, F. M., A. E., F. F. and A. P.; writing – original draft preparation, F. M.; writing – review and editing, O. S. and M. K.; scientific supervision, O. S.; project administration, O. S. All authors have read and agreed to the published version of the manuscript.

## Conflicts of interest

There are no conflicts to declare.



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## References

- C. Hank, A. Sternberg, N. Köppel, M. Holst, T. Smolinka, A. Schaadt, C. Hebling and H.-M. Henning, *Sustainable Energy Fuels*, 2020, **4**, 2256.
- IEA, *Hydrogen*, Paris, 2022.
- Hydrogen*, <https://www.iea.org/fuels-and-technologies/hydrogen>, (last accessed December 2022).
- Direct Air Capture*, <https://www.iea.org/reports/direct-air-capture>, (last accessed June 2022).
- M. Härtl, K. Gaukel, D. Pélerin and G. Wachtmeister, *MTZ worldwide*, 2017, vol. 78, p. 52.
- A. Omari, B. Heuser, S. Pischinger and C. Rüdinger, *Appl. Energy*, 2019, 1242.
- P. Dworschak, V. Berger, M. Härtl and G. Wachtmeister, *SAE Int. J. Fuels Lubr.*, 2022, **15**, DOI: [10.4271/04-15-02-0008](https://doi.org/10.4271/04-15-02-0008).
- A. D. Gelner, D. Rothe, C. Kykal, M. Irwin, A. Sommer, C. Pastoetter, M. Härtl, M. Jaensch and G. Wachtmeister, *Environ. Sci.: Atmos.*, 2022, **2**, 291.
- C. Hank, L. Lazar, F. Mantei, M. Ouda, R. J. White, T. Smolinka, A. Schaadt, C. Hebling and H.-M. Henning, *Sustainable Energy Fuels*, 2019, **3**, 3219.
- S. Voelker, S. Deutz, J. Burre, D. Bongartz, A. Omari, B. Lehrheuer, A. Mitsos, S. Pischinger, A. Bardow and N. von der Assen, *Sustainable Energy Fuels*, 2022, **6**, 1959.
- S. Deutz, D. Bongartz, B. Heuser, A. Kätelhön, L. Schulze Langenhorst, A. Omari, M. Walters, J. Klankermayer, W. Leitner, A. Mitsos, S. Pischinger and A. Bardow, *Energy Environ. Sci.*, 2018, **11**, 331.
- IEA, *World Energy Outlook 2022*, Paris, 2022.
- Kraft- und Brennstoffe - Polyoxymethylendimethylether (OME) - Anforderungen und Prüfverfahren (DIN/TS 51699:2023-04 - Entwurf)*, Beuth Verlag GmbH, Berlin, 2023, <https://www.beuth.de/de/vornorm-entwurf/din-ts-51699/364871868>, (last accessed March 2023).
- H. Schelling, E. Ströfer, R. Pinkos, A. Haunert, G.-D. Tebben, H. Hasse and S. Blagov, Method for producing Polyoxymethylene Dimethyl Ethers, *US 20070260094A1*, BASF Aktiengesellschaft, 2007.
- J. Burger, A novel process for the production of diesel fuel additives by hierarchical design, *Ph.D. Thesis*, University of Kaiserslautern, Kaiserslautern, 2012.
- J. Burger, M. Siegert, E. Ströfer and H. Hasse, *Fuel*, 2010, **89**, 3315.
- E. Ströfer, H. Schelling, H. Hasse and S. Blagov, Method for the Production of Polyoxymethylene Dialkylethers From Troxan and Dialkylethers, *US007999140B2*, 2011.
- C. F. Breitzkreuz, N. Schmitz, E. Ströfer, J. Burger and H. Hasse, *Chem. Ing. Tech.*, 2018, **90**, 1489.
- F. Mantei, R. E. Ali, C. Baensch, S. Voelker, P. Haltenort, J. Burger, R.-U. Dietrich, N. von der Assen, A. Schaadt, J. Sauer and O. Salem, *Sustainable Energy Fuels*, 2022, **6**, 528.
- N. Schmitz, E. Ströfer, J. Burger and H. Hasse, *Ind. Eng. Chem. Res.*, 2017, **56**, 11519.
- J. Burger, N. Schmitz, H. Hasse and E. Ströfer, Process for preparing Polyoxymethylene Dimethyl Ethers from Formaldehyde and Methanol in aqueous solutions, *US 2018/0134642 A1*, 2018.
- R. Palkovits, C. H. Gierlich and I. Delidovich, Verfahren zur Trennung von Oxymethylenethern, *EP 3 514 134 A1*, 2019.
- C. H. Gierlich, Herstellung von Oxymethylenethern anhand von alternativen Reaktionskonzepten, *Ph.D. Thesis*, Aachen, 2021.
- G. P. Hagen and M. J. Spangler, Preparation of Polyoxymethylene Dimethyl Ethers by Catalytic Conversion of Formaldehyde Formed by Oxidation of Dimethyl Ether, *US 6392102 B1*, 2002.
- T. Qiang, W. Jinfu, W. Tiefeng and Z. Yanyan, Method for Preparing DMM3–5 from Hypercoagulable Polyoxymethylene Dimethyl Ether Component DMM6+ and Dimethoxymethane DMM, *CN105152882 A*, 2017.
- N. Schmitz, F. Homberg, J. Berje, J. Burger and H. Hasse, *Ind. Eng. Chem. Res.*, 2015, **54**, 6409.
- N. Schmitz, J. Burger and H. Hasse, *Ind. Eng. Chem. Res.*, 2015, **54**, 12553.
- T. Grützner, H. Hasse, N. Lang, M. Siegert and E. Ströfer, *Chem. Eng. Sci.*, 2007, **62**, 5613.
- M. Drexler, P. Haltenort, U. Arnold and J. Sauer, *Chem. Ing. Tech.*, 2022, **94**, 256.
- C. F. Breitzkreuz, N. Hevert, N. Schmitz, J. Burger and H. Hasse, *Ind. Eng. Chem. Res.*, 2022, **61**, 7810.
- P. Haltenort, K. Hackbarth, D. Oestreich, L. Lautenschütz, U. Arnold and J. Sauer, *Catal. Commun.*, 2018, **109**, 80.
- J. Voggenreiter and J. Burger, *Ind. Eng. Chem. Res.*, 2021, **60**, 2418.
- F. Mantei, S. Kopp, A. Holfelder, E. Flad, D. Kloeters, M. Kraume and O. Salem, *React. Chem. Eng.*, 2023, **8**, 917.
- I. Bogatykh, T. Osterland, H. Stein and T. Wilharm, *Energy Fuels*, 2020, **34**, 3357.
- K. Hackbarth, P. Haltenort, U. Arnold and J. Sauer, *Chem. Ing. Tech.*, 2018, **90**, 1.
- H. Liu, Z. Wang, Y. Li, Y. Zheng, T. He and J. Wang, *Appl. Energy*, 2019, **233-234**, 599.
- D. Bongartz, L. Doré, K. Eichler, T. Grube, B. Heuser, L. E. Hombach, M. Robinius, S. Pischinger, D. Stolten, G. Walther and A. Mitsos, *Appl. Energy*, 2018, 757.
- D. Bongartz, J. Burre and A. Mitsos, *Ind. Eng. Chem. Res.*, 2019, **58**, 4881.
- M. Held, Y. Tönges, D. Pélerin, M. Härtl, G. Wachtmeister and J. Burger, *Energy Environ. Sci.*, 2019, **12**, 1019.
- C. F. Breitzkreuz, M. Dyga, E. Forte, F. Jirasek, J. de Bont, J. Wery, T. Grützner, J. Burger and H. Hasse, *Chem. Eng. Process.*, 2021, 108710.
- J. Sauer and G. Emig, *Chem. Eng. Technol.*, 1995, **18**, 284.



- 42 J. Burre, D. Bongartz and A. Mitsos, *Ind. Eng. Chem. Res.*, 2019, **58**, 5567.
- 43 S. Schemme, J. L. Breuer, M. Köller, S. Meschede, F. Walman, R. C. Samsun, R. Peters and D. Stolten, *Int. J. Hydrogen Energy*, 2020, **45**, 5395.
- 44 P. Haltenort, L. Lautenschütz, U. Arnold and J. Sauer, *Top. Catal.*, 2019, **62**, 551.
- 45 M. Drexler, P. Haltenort, U. Arnold, J. Sauer, S. A. Karakoulia and K. S. Triantafyllidis, *Catal. Today*, 2022, 113847.
- 46 X. Li, J. Cao, M. A. Nawaz, Y. Hu and D. Liu, *J. Chem. Eng. Data*, 2019, **64**, 5548.
- 47 X. Li, H. Tian, W. Zhang and D. Liu, *International Journal of Chemical and Molecular Engineering*, 2018, 536.
- 48 L. Wang, S. Zhou, P. Li, P. Li, Q. Li and Y. Yu, *J. Chem. Eng. Data*, 2018, **63**, 3074.
- 49 M. Shi, X. Yu, G. He and Q. Li, *Can. J. Chem. Eng.*, 2018, **96**, 968.
- 50 M. Shi, G. He, F. Gan, X. Yu and Q. Li, *J. Chem. Eng. Data*, 2017, **62**, 2183.
- 51 Z. Zhuang, J. Zhang and D. Liu, *Huagong Xuebao*, 2016, **67**, DOI: [10.11949/j.issn.0438-1157.20160391](https://doi.org/10.11949/j.issn.0438-1157.20160391).
- 52 Z. Zhuang, J. Zhang, X. Liu and D. Liu, *J. Chem. Thermodyn.*, 2016, 190.
- 53 D. Oestreich, L. Lautenschütz, U. Arnold and J. Sauer, *Fuel*, 2018, **214**, 39.
- 54 A. Ferre and J. Burger, *Ind. Eng. Chem. Res.*, 2021, **60**, 15256.
- 55 S. Schemme, Techno-ökonomische Bewertung von Verfahren zur Herstellung von Kraftstoffen aus H<sub>2</sub> und CO<sub>2</sub>, *Ph.D. Thesis*, Aachen, 2020.
- 56 N. Schmitz, Production of poly(oxymethylene) dimethyl ethers from formaldehyde and methanol, *Ph.D. Thesis*, Scientific Report Series, Kaiserslautern, 2018.
- 57 N. Schmitz, C. F. Breitzkreuz, E. Ströfer, J. Burger and H. Hasse, *J. Membr. Sci.*, 2018, **564**, 806.
- 58 A. Ferre, J. Voggenreiter, Y. Tönges and J. Burger, *Motortech. Z.*, 2021, **82**, 28.
- 59 F. Mantei, M. Ouda and A. Schaadt, Method for Producing Polyoxymethylene Dimethyl Ethers, WO22013132 A1, 2022, <https://worldwide.espacenet.com/patent/search/family/077168217/publication/WO2022013132A1?q=pn%3DWO2022013132>, (last accessed July 2022).
- 60 *Ullmann's Encyclopedia of Industrial Chemistry: Formaldehyde*, ed. A. W. Franz, H. Kronemayer, D. Pfeiffer, R. D. Pilz, G. Reuss, W. Disteldorf, A. O. Gamer and A. Hilt, Wiley-VCH Verlag GmbH & Co. KG, Weinheim, 2016.
- 61 J.-O. Drunsel, Entwicklung von Verfahren zur Herstellung von Methylal und Ethylal, *Ph.D. Thesis*, Scientific Report Series, Kaiserslautern, 2012.
- 62 H. Hasse, J.-O. Drunsel, J. Burger and U. Schmidt, Process for the production of pure Methylal, WO 2012/062822 A1, 2012.
- 63 I. Bogatykh, T. Osterland, H. Stein and T. Wilharm, *Energy Fuels*, 2019, **33**, 11078.
- 64 Kontinuierliche Destillation PD250 AD, <https://asg-analytik.de/technikum/z1-sp2/glaskolonnen>, (last accessed February 2023).
- 65 A. Fink, C. H. Gierlich, I. Delidovich and R. Palkovits, *ChemCatChem*, 2020, **12**, 5710.
- 66 A. Fink, Struktur-Aktivitätsbeziehungen von Zeolithen als feste Säurekatalysatoren in der Synthese von Oxymethylendimethylethern, *Ph.D. Thesis*, Aachen, 2022.
- 67 C. J. Baranowski, A. M. Bahmanpour and O. Kröcher, *Appl. Catal., B*, 2017, **217**, 407.
- 68 Y. Tönges and J. Burger, *Chem. Eng. Res. Des.*, 2022, **189**, 572.
- 69 F. Mantei, M. Kraume and O. Salem, *Chem. Ing. Tech.*, 2022, **25**, 912.

