



**Enhanced catalytic performance of  $Mn_xO_y$ - $Na_2WO_4/SiO_2$   
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## COMMUNICATION

## Enhanced catalytic performance of $Mn_xO_y$ - $Na_2WO_4/SiO_2$ for the oxidative coupling of methane using an ordered mesoporous silica support

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**Abstract** The oxidative coupling of methane is a highly promising reaction for the direct conversion of methane. Silica supported  $Mn_xO_y$ - $Na_2WO_4$  is a suitable catalyst for this reaction. In this study, a variety of different  $SiO_2$  materials have been tested as support. Surprisingly, the application of ordered mesoporous silicas, here exemplarily shown for SBA-15 as support material, greatly enhances the catalytic performance. The  $CH_4$  conversion increased two fold and also the  $C_2$  selectivity is strongly increased.

The proven reserves of natural gas have an enormous potential as an alternative to the decreasing reserves of crude oil.<sup>1</sup> The main component of natural gas is  $CH_4$ , the most stable hydrocarbon. Its conversion into value added products, particularly its direct conversion, remains a difficult challenge.<sup>2,3</sup> One possible direct conversion is the oxidative coupling of methane (OCM), as shown in Equation 1.



Although, reams of catalysts have been studied,<sup>4</sup> a breakthrough has not been achieved yet, especially because many catalysts deactivate due to the harsh reaction conditions.<sup>5,6</sup>  $Mn_xO_y$ - $Na_2WO_4/SiO_2$  is a very active, selective and stable catalyst, a fact which has been confirmed by several research groups.<sup>7-9</sup> The current knowledge on this catalyst has recently been reviewed.<sup>7</sup> Moreover, a fluidized bed processing procedure was developed for the large scale preparation of this material<sup>10</sup> allowing its application in the OCM mini-plant at the Technische Universität Berlin.<sup>11</sup>

To optimize the catalytic performance and to understand the structure-activity relationship, a variation of different support materials for  $Mn_xO_y$ - $Na_2WO_4$  are under investigation in our research

groups.<sup>12</sup> However, it can be concluded that most support materials show inferior or just comparable performance to  $SiO_2$  as support. On the other hand we observed a remarkable influence on the catalytic performance when different types of silica supports were used.

In this communication, we want to report on the observation that the application of SBA-15 leads to a greatly enhanced catalytic performance in comparison with any other studied  $SiO_2$  supports. SBA-15 is an ordered mesoporous silica<sup>13</sup> which is widely used as catalyst support in fundamental research.<sup>14</sup> Indeed as very narrow pore size distributions and highly ordered cylindrical mesopores can be prepared in this material, SBA-15 has significant advantages to study the dispersion of the supported active phase.<sup>15</sup> On the other hand, so far SBA-15 has not found its way to industrial applications, probably because much cheaper porous silicas can be prepared by other approaches. Furthermore, the ordered cylindrical pore structures has been described to be detrimental in terms of transport of substrates in the material, thus might cause severe diffusion limitations.<sup>16,17</sup> In here, we show that the application of SBA-15 can largely enhance the catalytic performance of an OCM catalyst, compared to commercially available porous silica supports.

This is most surprising as the silica support completely loses its ordered mesoporous structure during preparation of the catalyst.

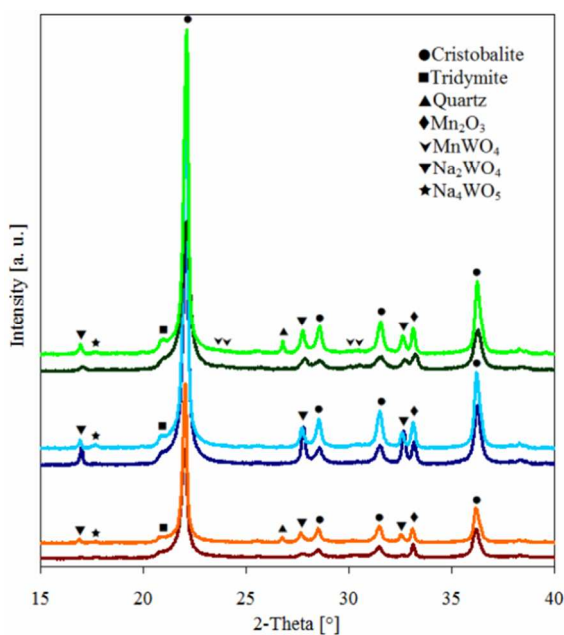
The prepared catalysts are shown in Table 1. For the preparation of the  $Mn_xO_y$ - $Na_2WO_4/SiO_2$  catalysts, a standard wet impregnation procedure was used, as described in the literature.<sup>7</sup> The detailed synthetic protocols can be found in the supporting information. Catalytic tests were performed with a 6-fold parallel reactor set-up and 50 mg catalyst. Details about the experimental setup can be found in the supporting information.

**Table 1** The codes of  $Mn_xO_y-Na_2WO_4/SiO_2$  catalysts, origin of support materials and surface areas of the applied silica support materials and catalysts.

Catalyst Code	Silica support material		Surface area of catalyst ( $m^2/g$ )		
	Origin	Comment	Surface area ( $m^2/g$ )	Before reaction	After reaction
Cat-1	Synthesized	SBA-15	616.9	6.8	4.2
Cat-2	BASF	D 11-10	105.4	6.7	3.1
Cat-3	Sigma	Silica gel, grade 923	492.3	2.8	3.9

The measured surface areas of the prepared catalysts showed a drastic reduction compared to the pure support material cf. Table 1, which is caused by the phase transformation from amorphous  $SiO_2$  to  $\alpha$ -cristobalite yielding complete collapse of the ordered mesoporous structure of SBA-15.<sup>7</sup> This transformation is also observed by XRD measurements, as shown in Figure 1. The loss of surface area during catalysis (before and after) is very small.

In Figure 1, XRD patterns for the fresh and tested catalysts are represented in darker and lighter colors, respectively. In the fresh samples  $\alpha$ -cristobalite, tridymite,  $MnWO_4$ ,  $Na_4WO_5$  and  $Na_2WO_4$  were detected. Moreover,  $Mn_2O_3$  or braunite is found, but an unambiguous assignment was not possible due to the small number of signals and their low intensity. For all catalysts,  $\alpha$ -cristobalite is formed as the main  $SiO_2$  phase.



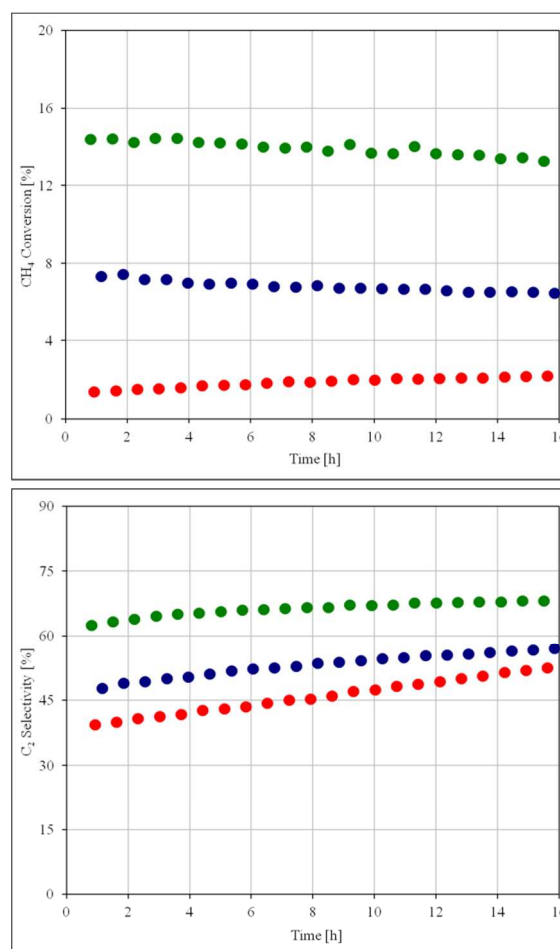
**Fig. 1** XRD patterns of Cat-1 (green), Cat-2 (blue) and Cat-3 (red).

For the used samples of Cat-1 and Cat-3, quartz was additionally found. It is important to note that substantial differences in the phase compositions were not observed for all catalysts before and after OCM reaction.

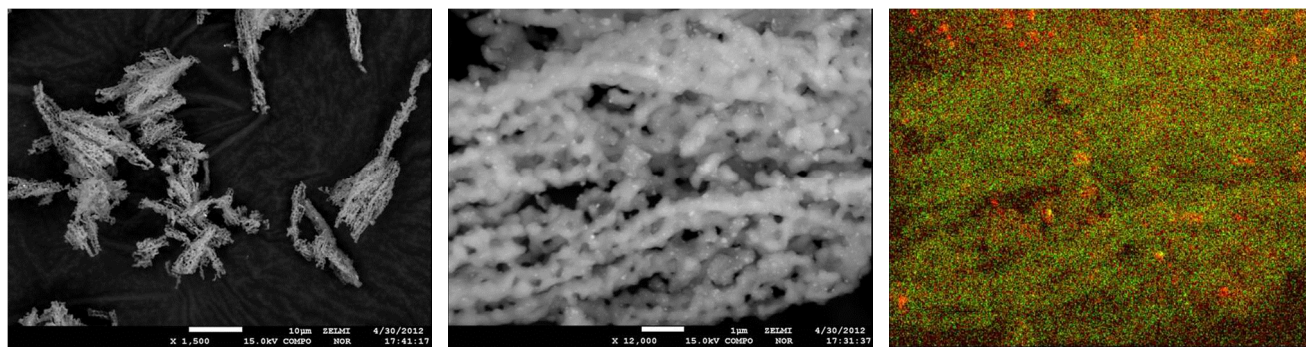
Elemental analysis furthermore revealed that the Mn, W and Na content on all silica supports is very similar (Table S2). The three catalysts thus show very similar results regarding surface area and

composition and therefore no significant difference in their catalytic activity would be expected at this point.

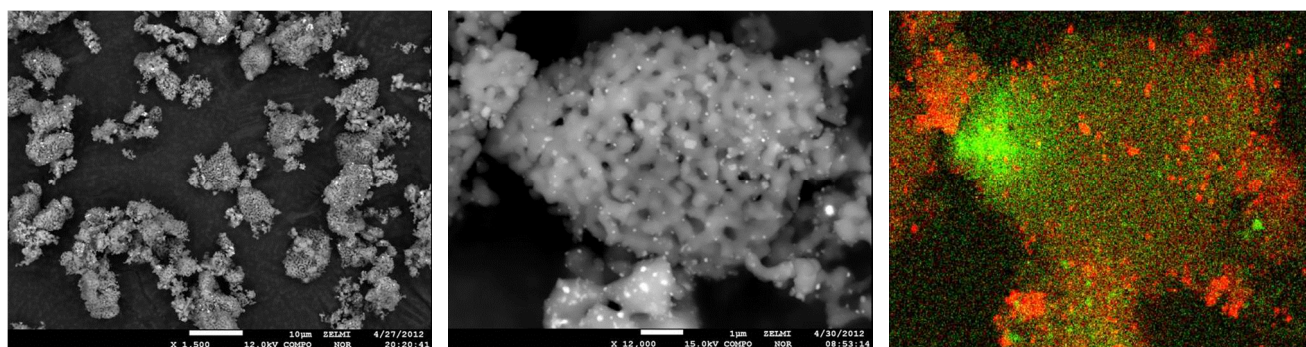
The results of the catalytic tests are shown in Figure 2. All tested catalysts exhibited a stable catalytic performance, with slightly increasing selectivities.



**Fig. 2**  $CH_4$  conversion as a function of time on stream (above) and the  $C_2$  selectivity as a function of time on stream (below) for Cat-1 (green), Cat-2 (blue) and Cat-3 (red) catalysts.



**Fig. 3** SEM images of fresh Cat-1 ( $\text{Mn}_x\text{O}_y\text{-Na}_2\text{WO}_4/\text{SBA-15}$ ) and EDX-mapping of W L-edge (green) and Mn K-edge (red).



**Fig. 4** SEM images of fresh Cat-2 ( $\text{Mn}_x\text{O}_y\text{-Na}_2\text{WO}_4/\text{SiO}_2$ ) and EDX-mapping of W L-edge (green) and Mn K-edge (red)

Cat-3 was almost inactive for the oxidative coupling of methane under the applied reaction conditions, which is surprising, because its catalytic performance is even worse than most catalysts reported in the literature.<sup>7</sup> Cat-2 showed a comparable catalytic performance when compared to the other commercial SiO<sub>2</sub> support materials investigated in our laboratory (not shown). On the other hand, the performance of Cat-1 (SBA-15 supported) was outstanding with approximately 14% CH<sub>4</sub> conversion, i.e. it showed two fold increase in conversion compared to Cat-2 with an even higher C<sub>2</sub> selectivity. As stated above, this significant increase in catalytic performance can be hardly explained by surface area or composition of the three catalysts. However, BET, XRD and elemental analysis give no information on the distribution of active components on the catalysts.

In Figure 3 and 4, SEM images and EDX mapping measurements are shown for the fresh catalyst Cat-1 and Cat-2. In Figure 3, the rod shaped morphology of the SBA-15 support can still be seen in the SEM images of fresh Cat-1. The EDX mapping shows the homogeneous distribution of elements, especially tungsten. In contrast to this, for fresh Cat-2 irregular spherical silica particles can be seen with a more inhomogeneous distribution of elements, cf. Figure 4. In addition, EDX-mapping of fresh catalyst Cat-1 showed even distribution of Mn with smaller particle sizes, while in Cat-2, Mn rich phases are observed as larger agglomerates on the silica support material.

For the first time in the research on the oxidative coupling of methane, one catalyst exists which we can reproducibly prepare with a good, a medium and a low catalytic performance and which possess a stable catalytic performance. The ongoing detailed studies will enable a first real view on the structure-activity relationship of this catalyst soon.

## Conclusions

The catalytic performance of the Mn<sub>x</sub>O<sub>y</sub>-Na<sub>2</sub>WO<sub>4</sub>/SiO<sub>2</sub> catalyst is greatly enhanced by the application of SBA-15 as silica precursor, approaching a level which might allow an industrial application. The explanation for this enhancement might be that the Mn<sub>x</sub>O<sub>y</sub>-Na<sub>2</sub>WO<sub>4</sub> precursors are better dispersed in the small pores and the high surface area of SBA-15, which is reflected in the enhanced dispersion of the final catalyst even when the mesostructure of the SBA-15 has collapsed after the thermal treatment. An optimization of this silica precursor, e.g. via adjusting the pore size and volume, could result in further improvements of the catalytic performance. Moreover, a detailed structural characterization and comparison of these three catalysts, currently in progress, might give a first insight into the structure-activity relationship of this catalyst, a missing feature hindering the understanding of many catalytic systems, particularly for metal oxides. Unraveling such kind of relationship could give room for further, perhaps even concerted, improvements.

## Notes and references

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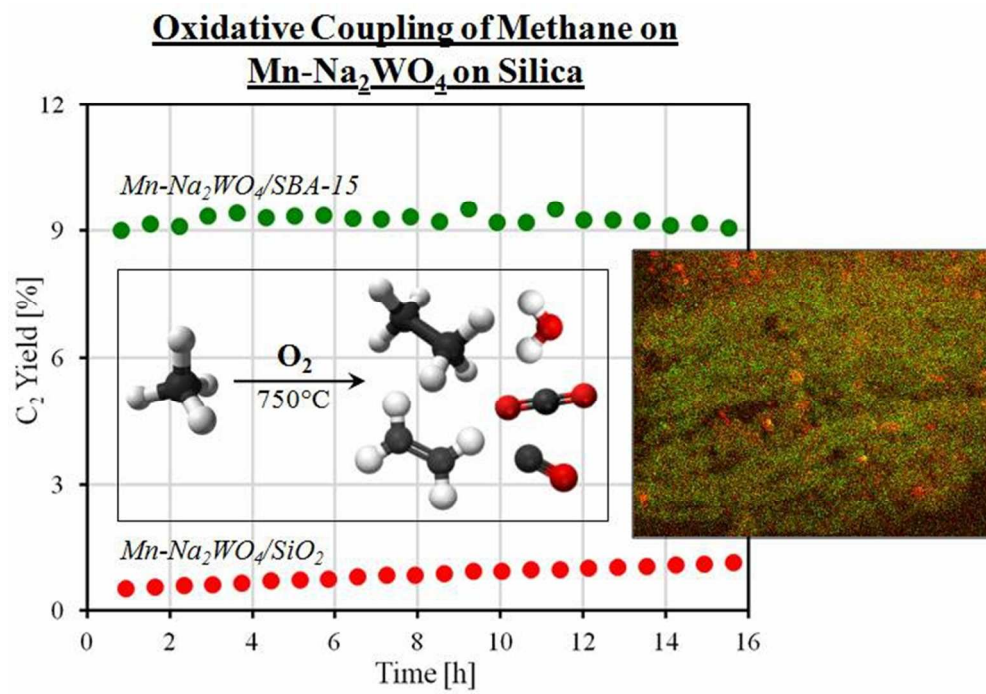
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