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Aldose to ketose interconversion: galactose and arabinose isomerization over heterogeneous catalysts

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Isomerization of glucose, galactose and arabinose to corresponding keto-sugars was studied in the present work over a range of heterogeneous catalysts. Magnesium aluminates with different ratios between oxides resulting in materials with a Mg/Al ratio from 0.2 to 0.9 were prepared, characterized and evaluated in terms of their catalytic behavior. The catalyst with a Mg/Al ratio close to hydrotalcites was the most efficient considering activity, selectivity and stability. The sugar structure was shown to have a minor influence on catalytic activity and selectivity.

Introduction

Interconversion of aldose to ketose has attracted a lot of attention recently as research efforts have been focused on valorization of carbohydrates derived from lignocellulosic biomass within the framework of the so-called biorefinery concept.^{1–3}

Formation of fructose from glucose in fact is a well-known reaction practiced on a million-tons-per-year scale in the production of high-fructose corn syrup (HFCS) using an immobilized enzymatic catalyst – glucose isomerase.⁴ This enzyme can be also used for interconversion of a C5 sugar – xylose to xylulose.⁴ The final equilibrium mixture of fructose and glucose contains *ca.* 55% of the former.

Chemical conversion of glucose to fructose can be also performed under alkaline conditions and at a temperature higher than that typically used (55–60 °C) for the enzymatic process. Transformations in an alkaline environment following the so-called Lobry de Bruyn–Alberda–van Ekenstein rearrangement^{5–7} are nonspecific leading to formation of side products. Conversion is typically limited resulting in fructose concentration of 40%.⁴ It should be mentioned, that higher fructose concentrations, *i.e.* 55% fructose, 41% glucose, and 4% mannose, have been recently reported using NaAlO₂ in water.⁸ *In situ* complexation of fructose under basic conditions by applying the concept of anionic extraction can increase the yield of fructose and as mentioned in ref. 7 could

reach 72%. Aldose–ketose rearrangement is accompanied by epimerization which occurs in parallel to the main reaction. Other reactions, such as aldolization/retro-aldolization, β-elimination and benzylic rearrangement^{8–11} also contribute to formation of side products. The mechanism of isomerization reaction (Fig. 1) is thought¹² to occur in the case of glucose *via* formation of glucose anions followed by generation of an ene-diol intermediate, which is then converted to fructose anions and finally into fructose.

Besides homogeneous bases (alkali or alkaline earth hydroxides) heterogeneous base catalysts would be of high interest being devoid of the apparent drawbacks of their homogeneous counterparts, such as low selectivity, and challenges with separation.

In connection with heterogeneous catalysts, several types of them could be mentioned. Hydrotalcite (HT) with different Mg/Al molar ratios was studied in isomerization of glucose,^{13–18} while zeolites modified with basic metals were investigated for the same reaction. Commercial Mg–Al hydrotalcites prepared as hydroxide, carbonate and mixed hydroxide–carbonate were reported in ref. 14 to be prone to deactivation after 15%

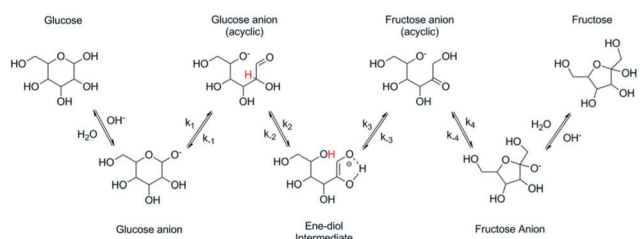


Fig. 1 Base catalyzed isomerization of glucose to fructose (reproduced with permission from ref. 12).

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conversion. Hydrotalcite with the atomic ratio Mg/Al equal to 3 afforded a 25% yield of fructose at 42% glucose conversion.¹³ The same ratio between Mg and Al was used in calcined and rehydrated Mg–Al hydrotalcite in ref. 15.

In contrast to ref. 13, when water was used as a solvent, in ref. 15, application of dimethylformamide instead of an aqueous solution, resulted in a *ca.* 35% yield of fructose at *ca.* 50% conversion for the rehydrated sample. The same solvent was used in ref. 19. Nevertheless water seems to be the preferred solvent. It was applied in the recent work of Delidovich and Palkovits¹⁷ where a fructose yield of up to 30% with 89% selectivity was reached in glucose isomerization to fructose and the optimal molar content of aluminum was reported to be $0.23 < \text{Al}/(\text{Mg} + \text{Al}) < 0.30$.¹⁷ Glucose conversion was seen to be linearly proportional to the basicity of hydrotalcite. The authors suggested that weak sites of hydrated hydrotalcites are more selective than the medium strong and strong sites. Moreover, HT in its carbonate or hydrated form was suggested to be catalytically active for glucose isomerization to fructose. As a consequence noncalcined HT was proposed to be more beneficial for the reaction than its calcined counterparts.¹⁷

Hydrotalcites were compared with other catalytic materials in a detailed review covering the applications of homogeneous and heterogeneous catalysts for isomerization of glucose.⁷ A viable alternative to hydrotalcites is magnesia. This implies that the presence of magnesium aluminate (MgAl_2O_4) or $(\text{M}^+\text{AlO}_2^-)$ formed during synthesis of hydrotalcites might not be needed to afford an active catalyst.

Commercially available MgO with increased basicity was reported as an efficient catalyst for glucose isomerization at 90 °C giving a yield of 33.3% fructose at a conversion of 44.1%.⁸ It should be noted that at the same time its selectivity was *ca.* 75% resulting not only in fructose but also in mannose, as well as oxidation products, such as glycolic, acetic, lactic and formic acids. Since the reaction conditions do not favor oxidation, formation of these products is most probably due to retro-aldol reactions.

The type of heterogeneous catalyst influences substantially selectivity. For instance non-acidic LiNbMoO_6 and strongly acidic HNbMoO_6 efficiently catalyze epimerization of glucose to mannose without any isomer formation.²⁰ In contrast, glucose isomerization into fructose *via* an intramolecular 1,2-hydride shift mechanism with Lewis acid catalysts such as Sn- β zeolites can be very efficient giving a 55% glucose conversion and a 32% yield of fructose in water at 110 °C.²¹ Besides fructose, an epimer – mannose – was also formed. The same type of catalyst was used for isomerization of the C5 sugar xylose resulting in an 18% yield of xylulose at 85% xylose conversion at 110 °C in water.²²

There is a limited amount of experimental data available for the isomerization of other sugars. Isomerization of lactose to lactulose using a range of homogeneous and heterogeneous catalysts was reported by Hajek *et al.*²³ Commercial hydrotalcite with a Mg/Al ratio equal to 3 in its noncalcined form was inactive, while the calcination at a temperature in the range 643–923 K had a minor influence on either its ac-

tivity or selectivity. For the same reaction Sn-MFI zeolites were reported in ref. 24.

Less is even known about isomerization of L-arabinose to L-ribulose (an intermediate for antiviral nucleosides) using heterogeneous catalysts, since the existing literature covers enzymatic and homogeneous catalysis. The same is valid for the isomerization of galactose (C4 epimer of glucose) to tagatose.

One of the very few reports on applications of hydrotalcites for isomerization of arabinose and galactose is related to one-pot synthesis of furanic compounds from arabinose and lactose (disaccharide of galactose and glucose) using a physical mixture of commercial HT with a Mg/Al ratio of 3 and acid catalysts of Amberlyst and Nafion types.²⁵ No information was provided in ref. 25 about the influence of the basicity, Mg to Al ratio or calcination temperature on the activity and selectivity.

Besides homogeneous and heterogeneous catalysts containing magnesia, there are also reports on utilization of cobalt compounds as catalysts²⁶ or co-factors in regard to homogeneous and enzymatic isomerization of glucose.²⁷ Application of cobalt and magnesium ions as co-factors in enzymatic transformations of glucose with glucose isomerase²⁷ is not necessarily related to their involvement in the binding of the reactant to the active site. However, according to ref. 26 Co(III) phthalocyanine species generated *in situ* from Co(II) phthalocyanines react with glucose giving a five membered ring chelate complex, which subsequently isomerizes to fructose. No reports are available on heterogeneous catalytic isomerization of carbohydrates with Co catalysts and it was therefore interesting to test if Co containing catalysts are active in this reaction.

The current work aims to explore isomerization of arabinose and galactose over heterogeneous catalysts, which contain MgO as such or as part of magnesium aluminates or CoMnAlMg mixed oxides. To this end besides commercial magnesia, several catalysts covering a broad range of magnesia to alumina ratios were synthesized, characterized, calcined at different temperatures and evaluated in the isomerization of carbohydrates. For the sake of comparison glucose isomerization was also performed with some catalysts.

A special emphasis was put on the evaluation of catalyst stability against leaching. The latter is especially important because catalyst deactivation due to its contact with aqueous reaction mixtures is often disregarded in the available literature.

Results and discussion

Catalyst characterization results

The elemental composition of monometallic cobalt catalysts was determined by ICP and corresponded well to the nominal loadings. The surface area (BET) decreased with increasing Co loading. For 10 wt% Co it decreased in the order $\text{Co}/\text{SBA-15} > \text{Co}/\text{Al}_2\text{O}_3 > \text{Co}/\text{TiO}_2$ and the BET values were 470, 215 and 98 $\text{m}^2 \text{g}^{-1}$.

The elemental composition of the CoMnAlMg catalyst is given in Table 1, while characterization data for magnesium aluminates are presented in Fig. 2 and 3 and Tables 2–4.



Fig. 2 displays the XRD data for the synthesized magnesium aluminate materials. As can be seen from Fig. 2 the samples with a higher content of alumina are X-ray amorphous, while an increase of MgO content affords crystalline materials. Already the $(\text{MgO})_{0.45}(\text{Al}_2\text{O}_3)_{0.55}$ material calcined at 680 °C contains reflections typical for MgAl_2O_4 spinels which is in agreement with the phase diagram. Further increase of MgO content in magnesium aluminates ($(\text{MgO})_{0.7}(\text{Al}_2\text{O}_3)_{0.3}$ and $(\text{MgO})_{0.9}(\text{Al}_2\text{O}_3)_{0.1}$ calcined at 680 °C) results in reflections, which can be attributed to the cubic structure of MgO, which somewhat shifted in terms of 2θ values compared to neat magnesia. The reason for the shift can be attributed to replacement of some Mg by Al. Reflections typical for the basal planes of double layer hydroxide (hydrotalcite) at 2θ values of 11.4°, 22.9° and 34.7° were not observed not only in the samples with a low Mg content, but also in materials with a high Mg content. According to XRD analysis there were no significant differences in the values of the coherent scattering region (CSR) for the studied samples. The CSR values obtained by the Rietveld method varied within 3–6 nm. The range of CSR values is within the error of the method (significant deviation of ~5 nm).

The MgO and Al_2O_3 composition in the synthesized materials was evaluated by XRF. The results presented in Table 2 clearly demonstrate that calcination did not influence the composition of magnesia–alumina materials.

The measured mass ratios displayed some deviations from the theoretical ones which can be at least partially explained by the presence of some organic leftovers in the catalysts even after calcination and inherent errors of the applied methods associated with for example high concentration of elements (ICP-AES).

The values of surface areas for the tested magnesium aluminates given also in Table 2 are in line with the characterization data typical for these types of materials.¹⁶

Basicity data presented in Table 3 and Fig. 3 clearly show that the basicity of magnesium aluminates increases by increasing the content of MgO. All catalysts displayed well-defined CO_2 TPD peaks with the peak maxima at ca. 450 K. This could be assigned to weak and medium basic sites, while some additional peaks present at the high temperature domain could be related to strong basic sites. In ref. 16 the activity for isomerization was related to the weak base sites; therefore for calculations of weak and medium basic sites, the cut-off temperatures mentioned in Table 3 were applied. The strong sites present in the catalyst might lead to the formation of degradation products as mentioned in ref. 16. A special case was 20 wt% MgO/MWNT displaying a more complicated TPD pattern, which could be related to the properties of the support only, in particular, the decomposition of

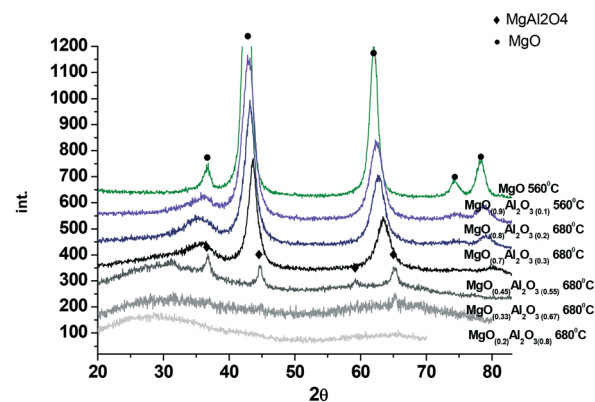


Fig. 2 XRD of the $(\text{MgO})_x(\text{Al}_2\text{O}_3)_y$ samples after calcination at 560 °C or 680 °C.

various functional groups present in MWNTs during TPD. In particular, decomposition of carboxylic ($-\text{COOH}$) and more stable keto groups ($-\text{C}=\text{O}$) occurs respectively at 250–300 °C and 680–720 °C.

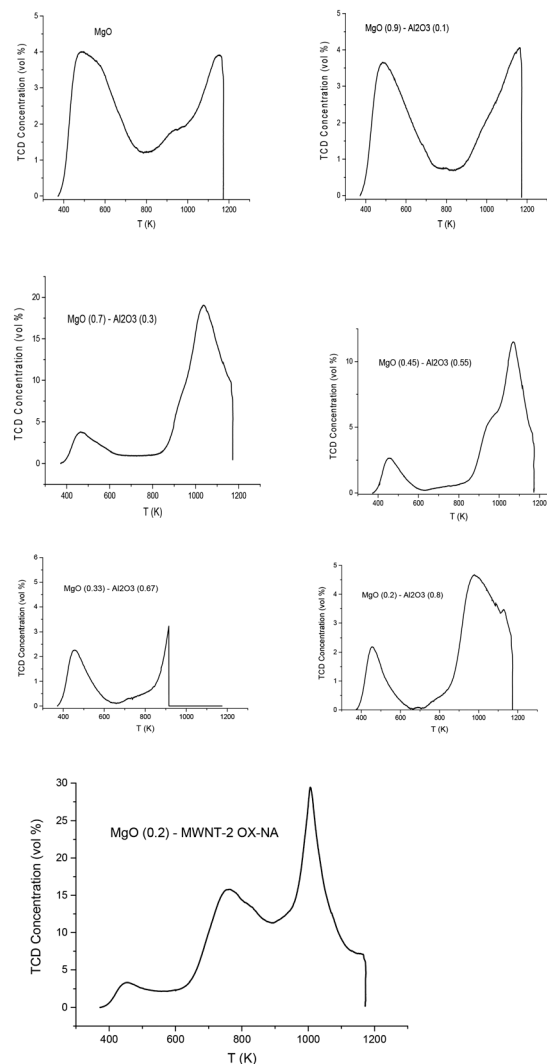


Fig. 3 CO_2 TPD for the catalysts in Table 3.

Table 1 Elemental composition of the CoMnAlMg catalyst in atom% according to XPS

Mg	Ni	Co	Mn	O	N	C	Al
15.6	0.3	4.2	5.0	59.5	0.8	4.7	9.9



Table 2 Analysis of the MgO–Al₂O₃ catalysts by XRF, ICP-AES and nitrogen physisorption

Sample		MgO, wt%	Al ₂ O ₃ , wt%	MgO/Al ₂ O ₃ mass ratio (XRF)	MgO/Al ₂ O ₃ mass ratio (XRF) ^a	MgO/Al ₂ O ₃ mass ratio ICP-AES	MgO/Al ₂ O ₃ mole ratio (XRF)	S _{BET} , m ² g ⁻¹
MgO	560 °C	99.2						176
(MgO) _{0.2} (Al ₂ O ₃) _{0.8}	T = 560 °C T = 680 °C	11.0	87.2	0.126			0.319	165
(MgO) _{0.33} (Al ₂ O ₃) _{0.67}	T = 560 °C T = 680 °C	19.7	78.5	0.251	0.116	0.091		97
(MgO) _{0.45} (Al ₂ O ₃) _{0.55}	T = 560 °C T = 680 °C	29.3	69.0	0.425	0.233		0.634	nm
(MgO) _{0.7} (Al ₂ O ₃) _{0.3}	T = 560 °C T = 680 °C	54.0	43.8	1.233	0.397	0.318	1.075	130
(MgO) _{0.8} (Al ₂ O ₃) _{0.2}	T = 560 °C T = 680 °C	69.7	29.6	2.354	1.215		3.118	132
(MgO) _{0.9} (Al ₂ O ₃) _{0.1}	T = 560 °C T = 680 °C	82.9	16.4	5.055		1.55	5.956	205
						1.55	5.607	272
						3.59	12.784	276
								nm

^a After hydrothermal stability tests at 110 °C; nm – not measured.

Catalytic results: screening

None of the monometallic cobalt catalysts displayed any meaningful activity in the isomerization of galactose at 110 °C and 5 bar of nitrogen with a catalyst loading of 0.2 g.

The results for the activity tests with commercial hydrotalcite and MgO are presented in Table 4. As follows from Table 4 besides the desired ketoses, some by-products, such as oligomers, epimers and degradation products were also observed with different selectivities depending on the catalyst.

It follows from Table 4 that commercial hydrotalcite with a Mg/Al ratio equal to 3 in its noncalcined form was rather inactive in isomerization independent of the type of substrate, while calcination at 500 °C was sufficient to boost the catalytic activity. Although the results in terms of conversion are like those for glucose to fructose isomerization at the same temperature reported in ref. 17, the influence of calcination seems to be different. In ref. 17 noncalcined HT was proposed to be more beneficial for the reaction than the calcined counterparts based on a suggestion that hydrotalcite in its carbonate or hydrated forms are the catalytically active species. In the current work as well as in the previous study on lactose isomerization²³ calcination was found to be beneficial for catalytic activity. Inactivity of noncalcined HT compared to that of the calcined counterpart in general could be related to basicity. As it is challenging to perform TPD–CO₂ for samples when carbonates are decomposed at ca. 200–300 °C, a comparison of the pH value of water suspensions with HT prior to and after calcination was performed. It was observed that the pH value of the water suspension with

the noncalcined hydrotalcite being equal to 8.95 was slightly elevated when hydrotalcite was calcined for 4 h at 680 °C reaching 9.55. Apart from these small differences which probably in itself cannot be responsible for the differences in the catalytic behaviour, it should be noted that the noncalcined sample was clearly hydrophobic in contrast to its calcined counterpart. This is probably the reason for their different performances in isomerization of sugars occurring in aqueous solutions.

Similar to ref. 8 where glucose to fructose isomerization with commercially available MgO was studied, in the current work isomerization of galactose was investigated. The catalyst is also active in isomerization allowing high conversion exceeding 50%. At the same time selectivity to ketose clearly decreased with conversion as the reaction resulted not only in tagatose, but also in oligomers and degradation products (Fig. 4). Another serious issue is catalyst deactivation through leaching. In a separate experiment on leaching of magnesia in galactose solution at 110 °C severe dissolution of MgO in the solution was observed.

Thus, even if magnesia is superficially active in the isomerization of sugars there is still a need for obtaining an active and at the same time stable and selective catalyst which, on the one hand, would be stable against leaching and on the other would avoid or at least minimize formation of oligomers or degradation products. It was thus interesting to test catalysts where MgO was deposited on an inert support: multiwall carbon nanotubes. Fig. 5 displays the results illustrating that application of supported MgO resulted in limited conversion, thereafter the reaction stopped completely. The hydrothermal stability was very moderate and will be discussed below.

Synthesis of carbon nanotubes is typically performed using a suitable growth catalyst. One such catalyst is the CoMnAlMg mixed oxide which was synthesized following ref. 28. The content of magnesium oxide in this catalyst was enough to expect non-negligible catalytic activity. Experiments were conducted with three sugars (arabinose, galactose and glucose) at three temperatures allowing a straightforward comparison between different substrates and calculations of activation energies. The results are presented in Fig. 6. Due to a narrow temperature

Table 3 Basicity data by CO₂ TPD for the samples calcined at 680 °C

Catalyst	Cut off T, K	Weak and medium basic sites, μmol g ⁻¹
20 wt% MgO/MWNT	600	218
(MgO) _{0.2} (Al ₂ O ₃) _{0.8}	700	118
(MgO) _{0.33} (Al ₂ O ₃) _{0.67}	650	125
(MgO) _{0.45} (Al ₂ O ₃) _{0.55}	650	140
(MgO) _{0.7} (Al ₂ O ₃) _{0.3}	650	268
(MgO) _{0.9} (Al ₂ O ₃) _{0.1}	750	350
MgO	750	507



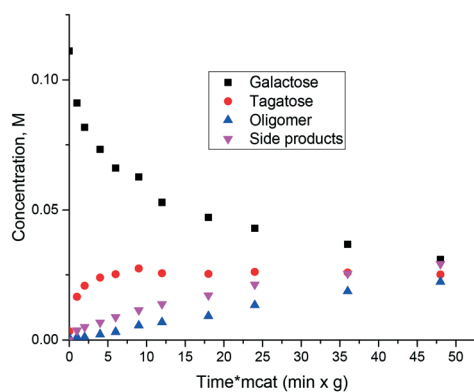
Table 4 Isomerization of sugars with MgO and hydrotalcite (HT)^a

Catalyst	Substrate	<i>T</i> , °C	<i>m</i> × <i>t</i> , (g × min)	<i>C</i> _{aldose}	<i>C</i> _{ketose} ^b	Conv.%	Selec%
MgO	Gal ^c	110	12.0	0.05	0.03	55	50
MgO	Gal	110	48	0.03	0.02	73	25
MgO	Gal	90	12.0	0.07	0.03	36	75
MgO	Gal	90	48.0	0.05	0.03	45	50
HT, noncalcined	Gal	110	48.0	0.10	0.00	9	0
HT, calcined at 500 °C	Gal	110	24.0	0.07	0.02	36	50
HT, calcined at 500 °C	Gal	110	48.0	0.07	0.02	36	50
HT, calcined at 500 °C	Glu ^c	120	48.0	0.07	0.03	36	75
HT, noncalcined	Ara ^d	110	48.0	0.12	0.00	8	0
HT, calcined at 500 °C	Ara	110	6.0	0.08	0.01	38	20
HT, calcined at 500 °C	Ara	110	48.0	0.06	0.01	54	14

^a *m*_{cat} = 0.2 g. ^b The rest of the oligomers, epimers and degradation products. ^c *C*_{initial} = 0.11 M. ^d *C*_{initial} = 0.13 M. Gal, Glu and Ara stand for galactose, glucose and arabinose, respectively. Concentrations are given in M.

interval it was challenging to establish very precisely the temperature dependence. For example, while some of the data points for glucose might coincide at 100 and 110 °C there was still a difference. For instance, a concentration of 0.09 M is reached within 50 min at a higher temperature, while at 100 °C the same concentration is obtained after 100 min. There is also a clear difference for arabinose. In order to have a more reliable evaluation of the temperature dependence the kinetic constants for first order reactions were calculated by numerical data fitting for respective kinetic curves across the whole experimental duration (Fig. 6d). The apparent activation energy for all sugars can be in fact considered the same being equal to 99 ± 8 kJ mol⁻¹. Information about the activation energy of sugar isomerization on magnesium aluminate, available in the literature, is limited. In ref. 16 data of the final conversion after 5 h in glucose to fructose isomerization are presented as a function of temperature. By using the conversion values in the temperature range 80–100 °C reported in ref. 16 the value of activation energy equal to 100 kJ mol⁻¹ can be calculated.

The conclusion that different sugars display almost the same reactivity was further confirmed by studying the mixtures of arabinose and galactose (Fig. 7a) and glucose and galactose (Fig. 7b). In both cases, there was a non-negligible formation of oligomers, which eventually deactivates the catalyst.

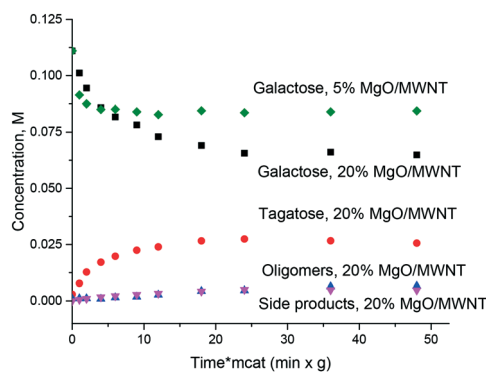
**Fig. 4** Isomerization of galactose with commercial MgO at 90 °C.

Catalytic results: Mg/Al ratio

The influence of the Mg/Al ratio on catalytic activity in isomerization of glucose to fructose was studied in ref. 17 for hydrotalcites. The latter materials being double layer hydroxides have a limited composition range. Namely in ref. 17, the content of Mg was varied between 0.78 and 0.89. In the current work a much broader range of Mg/Al was tested. In the prepared magnesium aluminates the hydrotalcite phases were not present, thus it can be suggested that the layered structure of hydrotalcite is not necessarily needed for catalytic activity.

The experimental data of galactose isomerization are displayed in Fig. 8 for materials calcined at either 560 or 680 °C. As can be seen from this figure an increase in the concentration of MgO brought along an increase of activity. Elevation of the calcination temperature did not influence substantially the catalytic activity.

A simple recycling test with magnesium aluminate having the composition 0.7 MgO 0.3 Al₂O₃ was performed by just filtering the catalyst after the reaction, washing with water and subsequent drying at 110 °C for 24 h and demonstrated a substantial decrease of catalytic activity (Fig. 8b). Therefore, catalyst regeneration should involve also burning of organic compounds by calcination similar to the procedure described in ref. 17. In particular previously performed recycling tests

**Fig. 5** Isomerization of galactose with MgO supported on MWNTs at 110 °C.

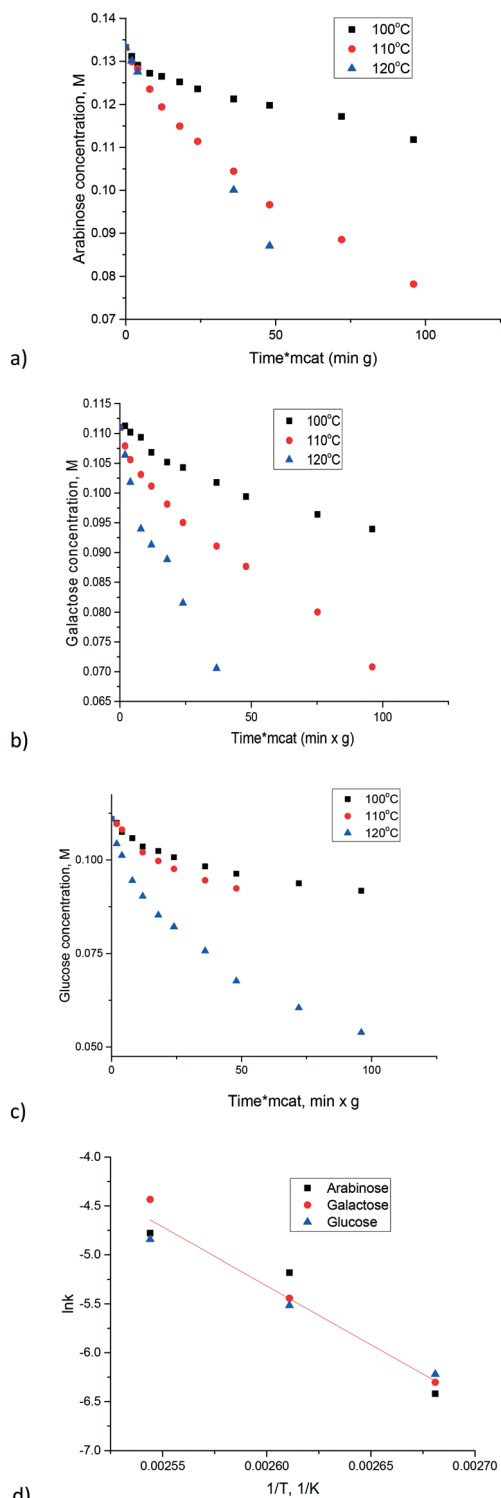


Fig. 6 Isomerization of sugars over the CoMnAlMg mixed oxide catalyst: a) arabinose, b) galactose, c) glucose, d) Arrhenius plot.

with hydrotalcites having a similar MgO content to that in the recycling experiment in the present study, were conducted by first regenerating the catalyst by calcination followed by rehydration in a sodium carbonate solution.¹⁷ While the molar ratio of Mg to Al changed from 3.34 to 2.03

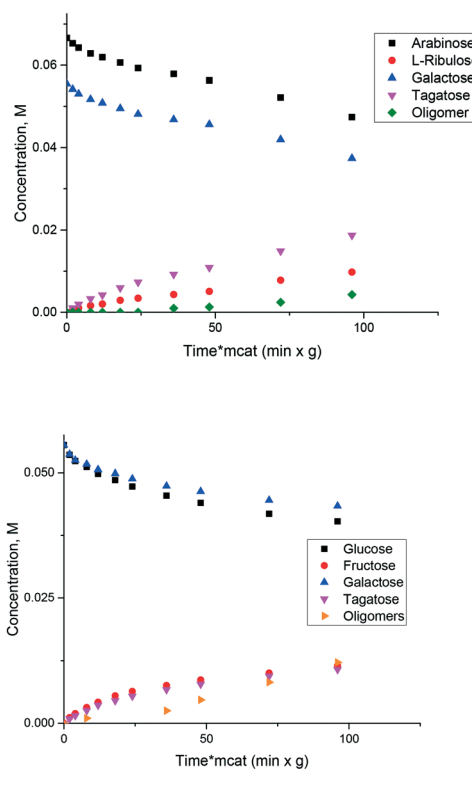


Fig. 7 Isomerization of the mixtures of sugars over the CoMnAlMg mixed oxide catalyst at 110 °C: a) arabinose–galactose, b) glucos–galactose.

reflecting a substantial loss of Mg during the recycling tests,¹⁷ the conversion did not change much in recycling experiments, being around 18–20%.

The TOF values reported in ref. 17 were in the range of 18–23 h⁻¹. Direct comparison of the TOF values is not straightforward as this parameter depends on conversion for first order reactions. Nevertheless, calculation of the TOF for the (MgO)_{0.7}(Al₂O₃)_{0.3} catalyst using XRF values from Table 3 and conversion after 30 min gave a TOF equal to 22 h⁻¹, which is the same as the values reported in ref. 17. To avoid uncertainties in the determination of TOF values and rationalize the influence of catalyst composition on the reactivity the first order kinetic constants have been plotted against magnesia content and overall basicity (Fig. 9). The errors in the determination of the rate constants were 15 ÷ 20% depending on the Mg/Al ratio.

As can be seen from Fig. 9 there is a clear linear correlation between the first order rate constant and the molar content of MgO or basicity as the latter parameters are related to each other. Previously glucose conversion in isomerization was related to the basicity of hydrotalcites with a rather narrow range of Mg/Al ratios.¹⁷ The same conclusion can be also applied for another sugar – galactose – and can be extended to a broader range of Mg/Al ratios.

The influence of the sugar structure for magnesium aluminates was tested by conducting experiments with galactose–glucose and galactose–arabinose mixtures using the (MgO)_{0.8}(Al₂O₃)_{0.2} catalyst.



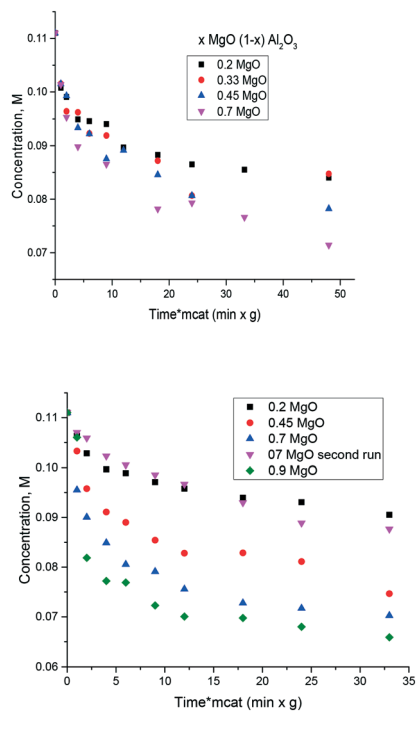


Fig. 8 Isomerization of galactose over the magnesium aluminates with different Mg/Al ratios at 110 °C and different calcination temperatures, a) 560 °C and b) 680 °C.

$\text{O}_3)_{0.2}$ catalyst calcined at 680 °C. Fig. 10 clearly illustrates that the kinetic behavior was independent of the sugar structure.

Hydrothermal stability tests

Literature data along with the catalytic results presented above clearly indicate that magnesium based materials (MgO, hydrotalcite and magnesium aluminate) could be promising catalysts for isomerization of sugars. A serious issue with this type of catalyst is leaching of the active phase into the solution. To address this issue hydrothermal activity tests were conducted at 110 °C for 2.5 h using 0.215 wt% catalyst loading in 2 wt% galactose solutions. Analysis of the liquid phase by AAS for several catalyst samples is shown in Table 5.

The sample containing only MgO was completely dissolved during hydrothermal treatment giving the brownish color of the resulting solution. 20 wt% MgO/MWNT was also prone to leaching resulting in a relatively high concentration of magnesium ions after hydrothermal stability tests in a galactose solution at 110 °C ($0.1967 \text{ mg ml}^{-1}$). The color of the solution after hydrothermal stability tests was yellowish. XRD of the hydrothermally treated 20 wt% MgO/MWNT (not shown) displayed only the MgO phase with lower intensity, which can be attributed to the leaching of magnesia from the catalyst.

Interestingly enough, leaching in the presence of galactose was much more severe compared to that of the treatment in neat water at the same temperature (Table 5). XRD data for

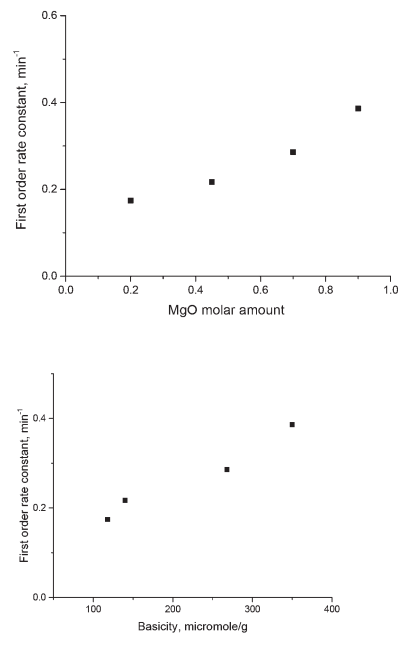


Fig. 9 Galactose isomerization first order rate constant vs. a) molar Mg content and b) basicity of magnesium aluminates. Reaction conditions: 110 °C, $m_{\text{cat}} = 9.2 \text{ g}$, 5 bar, 1000 rpm and calcination temperature = 680 °C.

the $(\text{MgO})_{0.7}(\text{Al}_2\text{O}_3)_{0.3}$ catalyst calcined at 680 °C and hydrothermally treated at 110 °C for 2.5 h are presented in Fig. 11.

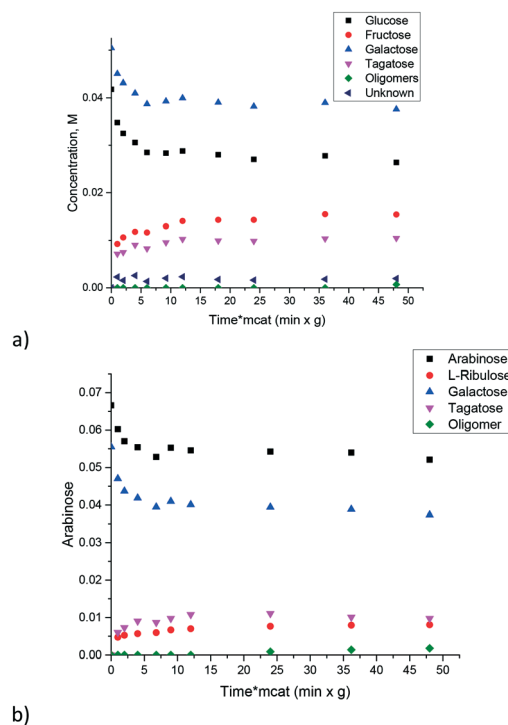


Fig. 10 Isomerization of sugar mixtures over the $(\text{MgO})_{0.8}(\text{Al}_2\text{O}_3)_{0.2}$ catalyst for a) galactose–glucose and b) galactose–arabinose mixtures. Reaction conditions: temperature = 110 °C, calcination temperature = 680 °C, $m_{\text{cat}} = 0.2 \text{ g}$, 5 bar and 1000 rpm.



Table 5 Concentration of leached ions after hydrothermal stability tests in 2 wt% galactose solution for 2.5 h at 110 °C for catalysts calcined at 680 °C

Catalyst	Mg content, mg ml ⁻¹	Al content, mg ml ⁻¹
(MgO) _{0.2} (Al ₂ O ₃) _{0.8} ^a	0.0015	0.00002
(MgO) _{0.33} (Al ₂ O ₃) _{0.67} ^a	0.0006	0.00002
(MgO) _{0.33} (Al ₂ O ₃) _{0.67}	0.008	0.00002
(MgO) _{0.45} (Al ₂ O ₃) _{0.55} ^a	0.0017	0.00002
(MgO) _{0.7} (Al ₂ O ₃) _{0.3} ^a	0.0007	0.0014
(MgO) _{0.8} (Al ₂ O ₃) _{0.2}	0.213	0.0013
(MgO) _{0.9} (Al ₂ O ₃) _{0.1}	0.318	0.0013
MgO	0.750	—
5% MgO/MWNT	0.0825	—
20% MgO/MWNT	0.1967	—

^a Hydrothermal treatment was performed in the absence of galactose.

Comparison of Fig. 11 with XRD data for non-treated catalyst (Fig. 2) clearly shows that the structure of the material changed significantly after hydrothermal treatment resulting in a structure close to hydrotalcite. This is an interesting observation showing that rehydration of magnesium aluminate results in double layer hydroxide. The activity of calcined hydrotalcite (Table 5) in galactose isomerization was similar to that of (MgO)_{0.7}(Al₂O₃)_{0.3} (Fig. 8b). This and the absence of any induction period or inflection points can indicate that if solid phase transformations are slow, different phases possess similar activities. Therefore catalytic activity can be attributed to the same MgO content in both samples, rather than to the presence of a HT phase as there was a linear increase of activity as a function of MgO content and even catalysts with low MgO loading not possessing a hydrotalcite structure were moderately active. Transformation of magnesium aluminate to hydrotalcite could have happened during the initial heating to the reaction temperature of the catalyst in a water solution, as well as during the reaction.

It was interesting to check if another magnesium aluminate with a much lower Mg content would undergo structural changes during hydrothermal treatment at 110 °C in galactose solution. For initially amorphous (MgO)_{0.33}(Al₂O₃)_{0.67} calcined

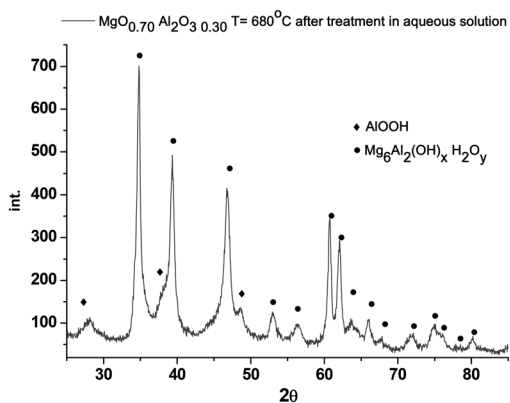


Fig. 11 XRD data for the (MgO)_{0.7}(Al₂O₃)_{0.3} catalyst calcined at 680 °C and hydrothermally treated at 110 °C for 2.5 h in galactose solution.

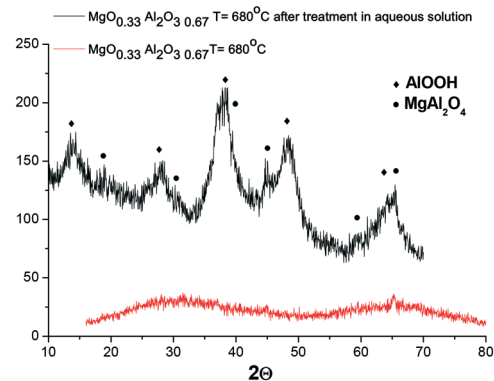


Fig. 12 XRD diffractogram of (MgO)_{0.33}(Al₂O₃)_{0.67} calcined at 680 °C prior to and after hydrothermal treatment at 110 °C for 2.5 h in galactose solution.

at 680 °C, hydrothermal treatment also resulted in the formation of a crystal structure, which is visible by the appearance of reflexes attributed to MgAl₂O₄ and AlO(OH) (Fig. 12).

Investigation of the Mg/Al ratio by XRF (Table 2) shows that the ratio between Mg and Al does not change substantially after hydrothermal treatment. At the same time the overall content of Mg and Al oxides decreased from 98 wt% to 87 wt%, which can be related to the appearance of water in the crystal structure as well as the existence of complex oxy- and hydroxy-mixed compounds containing Mg and Al.

For the (MgO)_{0.7}(Al₂O₃)_{0.3} catalyst the leached amount of MgO was very minor in contrast to the catalysts with higher MgO content, such as (MgO)_{0.9}(Al₂O₃)_{0.1} and (MgO)_{0.8}(Al₂O₃)_{0.2}, although the latter catalysts were more active in isomerization of aldoses that have resulted in a larger amount of side-products and displayed significant leaching with concentration of magnesium ions being 0.318 and 0.213 mg ml⁻¹, respectively. It can be thus concluded that the (MgO)_{0.7}(Al₂O₃)_{0.3} catalyst exhibited a Mg/Al ratio close to the hydrotalcites often reported as efficient base catalysts for isomerization of aldoses. The TEM images for this material are illustrated in Fig. 13.

As can be seen from the micrographs the catalyst is rather heterogeneous featuring particles ranging from 3 to 200 nm. Large (transparent) particles are spinel species with a large number of defects and in part are not well crystallized. No special phase of MgO is visible which can be also related to the poor leaching behavior of this catalyst as magnesia is present in the catalyst in a bound form. Smaller particles could tentatively be assigned to alumina even if the corresponding reflexes were not pronounced in the XRD patterns.

Conclusions

Isomerization of several aldoses (arabinose, galactose, glucose) was performed at 100–120 °C using a range of base catalysts including magnesia, hydrotalcites, magnesium aluminates with varying Mg/Al ratios and mixed oxides containing magnesia and alumina.



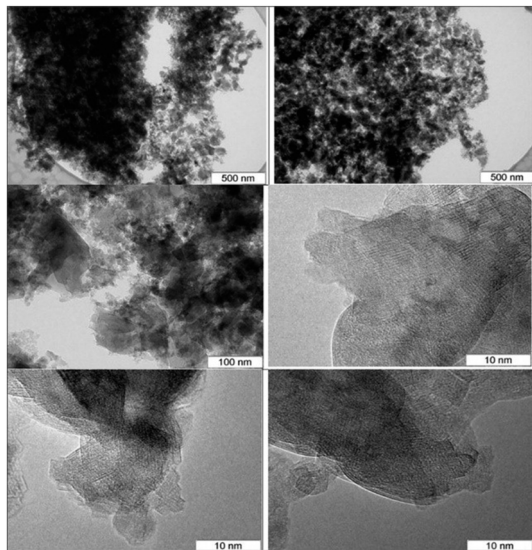


Fig. 13 TEM of the $(\text{MgO})_{0.7}(\text{Al}_2\text{O}_3)_{0.3}$ catalyst.

Experiments with different aldoses showed that the sugar structure did not influence the catalytic properties. The apparent activation energy for all sugars obtained for a mixed oxide catalyst containing, in addition to magnesia and alumina, cobalt and manganese oxide was close to 99 kJ mol^{-1} .

There was a linear correlation between the catalytic activity in galactose isomerization and the molar content of MgO in magnesium aluminates covering a broad range of Mg/Al ratios. The amount of MgO in the catalysts was closely related to their basicity.

Even if magnesium based materials (MgO, hydrotalcite and magnesium aluminate) could be applied for isomerization of sugars a serious issue is catalyst leaching. Hydrothermal stability tests with galactose solutions demonstrated that while a sample containing only MgO was dissolved completely, reasonable hydrothermal stability was displayed by the $(\text{MgO})_{0.7}(\text{Al}_2\text{O}_3)_{0.3}$ catalyst calcined at 680°C . The latter catalyst was active in isomerization of galactose exhibiting a TOF equal to 22 h^{-1} .

Experimental section

Catalysts

Commercial samples of MgO (Fluka, 63093) and hydrotalcite (Mg/Al ratio of 3, Aldrich, 652288) were used as-received and after calcination. The CoMnAlMg mixed oxide catalyst ($\text{Mn}:\text{Co}:\text{Al}_2\text{O}_3:\text{MgO}$) was synthesized from the corresponding nitrates following the procedure described in ref. 28. This catalyst, as well as several supported cobalt catalysts, was applied to assess a potential role of cobalt and in combination with magnesia and alumina. The metal loading was 10 wt% and as supports alumina, silica, titania and SBA-15 were applied.

Magnesium aluminate (Mg–Al–O) catalysts were prepared using the polymeric precursor or Pechini method.²⁹ Predetermined amounts of magnesium nitrate ($\text{Mg}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, 98%, Vekton) and aluminum nitrate ($\text{Al}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$,

98%, Vekton) were mixed and stirred at room temperature in distilled water. Thereafter, appropriate amounts of citric acid (99%) and ethylene glycol (>99%) (Sigma Aldrich) were added to this solution at room temperature. The polymerization reaction was conducted at 120°C until no gas evolution was visible. The dark solid was calcined at 560°C for 4 h under air with a heating rate of $10^\circ\text{C min}^{-1}$ to remove all organic components. Incomplete removal of organic compounds was observed at 560°C judging by the color of the obtained samples; therefore, some samples were calcined at 680°C for 24 h. It was decided to synthesize catalysts with the molar ratio $\text{MgO}/(\text{MgO} + \text{Al}_2\text{O}_3)$ equal 0.2; 0.33; 0.45; 0.7; 0.8 and 0.9 which corresponds to different phases in the phase diagram (Fig. 14).

Multiwall nanotubes (MWNTs) were synthesized by a CVD method of ethylene decomposition over bimetallic Fe–Co catalysts at 680°C (surface area of $305 \text{ m}^2 \text{ g}^{-1}$, average particle diameter of 9.4 nm).³⁰ The functionalized MWCNTs containing surface carboxylic groups (0.76 groups per 1 nm^2) were produced by treatment with boiling in concentrated nitric acid (denoted as MWNT–Ox–NA).³¹ Mg-containing samples were prepared by impregnation of functionalized MWNTs with the aqueous solutions of magnesium nitrate ($\text{Mg}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, 98%, Vekton).³² After impregnation for 6 h the samples were dried at 110°C for 4 hours followed by calcination at 350°C for 4 hours under argon. By varying the concentration of the magnesium nitrate solutions, samples of 5 and 20 wt% Mg loadings were obtained. The cobalt catalysts were prepared by an impregnation method using $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ (Lach-ner, s.r.o.) described previously.³³

Catalyst characterization

X-ray diffraction. Powder XRD measurements were carried out using an ARL X'TRA diffractometer (Thermo Electron Corporation, Switzerland), equipped with a vertical Theta-theta geometry (Bragg–Brentano), $\text{CuK}\alpha$ radiation ($\lambda = 0.15418 \text{ nm}$) and a Peltier cooled Si(Li) solid state detector. The range of 2θ values from 20° to 85° was scanned with a

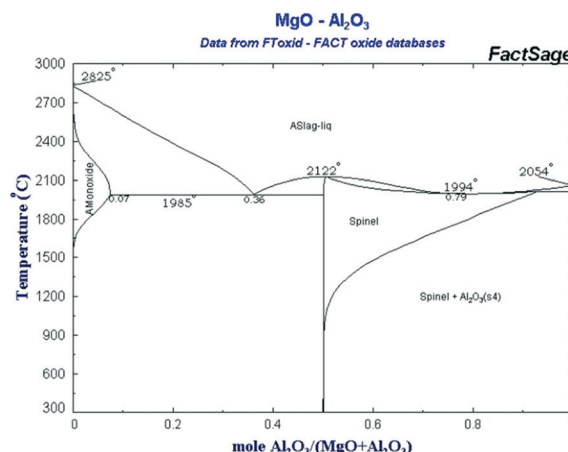


Fig. 14 Phase diagram of magnesium aluminates.



step of 0.05° and counting time of 3 s. The phase analysis was performed using the ICDD PDF-2 database. XRD analysis was performed using the Rietveld method in Topas software.

ICP-AES and XRF. The content of Al and Mg in the samples was determined by atomic emission spectroscopy with inductively coupled plasma on an Optima 4300 DV spectrometer (Perkin Elmer). The relative error in the determination of Al and Mg turned to be rather high, which could be associated with the high content of elements and the presence of carbon, which was released in the form of CO_2 . Therefore XRF was also performed.

The metal content in the catalysts was determined by X-ray fluorescence analysis (XRF) on an ARL Perform'X analyzer with a Rh-anode X-ray tube. The element content was assessed by the UniQuant program for standard-less analysis. Prior to analysis, the samples were ground in an agate mortar, and then mixed with cellulose in a ratio of 1:5 (wt) in order to obtain the required volume for filling the cuvette. The sample mixed with cellulose was pressed into a tablet and placed in a cassette for analysis. Thereafter, the cassette was loaded into the vacuum chamber of the spectrometer and the measurements were carried out. To cover a range of elements from F to U, 5 scans were conducted for each sample.

Energy dispersive X-ray spectroscopy. Local elemental analysis was performed with energy-dispersive X-ray spectroscopy using a Phoenix spectrometer (EDAX, USA) equipped with a Si(Li) detector giving energy resolution not worse than 130 eV. Quantitative EDX measurements were carried out *e.g.* using a Pd $L\alpha$ line, which is not overlapping with the other X-ray emission lines.

Nitrogen physisorption. The textural properties of the samples were measured with an automated flow type sorption unit Sorbi 4.2 (Meta) allowing the determination of the specific surface areas. All samples were subjected to thermal pre-training under argon at 200°C for 2 hours. The specific surface area (S , m^2g^{-1}) was calculated by the BET method. The error in determining the surface area values using this method is $\pm 10\%$.

Reactor set-up

Isomerization of D(+) galactose (Sigma-Aldrich, $\geq 99\%$ purity), L(+)-arabinose (Sigma, $\geq 99\%$ purity) and D(+) glucose (Fluka, $\geq 98\%$ purity) over different catalysts was investigated using a Parr 4561 autoclave. The liquid load was typically 100 ml of the overall volume of 300 ml. The sugar concentration in aqueous solution was equal to 2 wt% corresponding to 0.11 (galactose and glucose) and 0.13 M (arabinose). The autoclave was equipped with a gas entrainment impeller, baffles, heating jacket and cooling coil, sampling line, pressure, temperature and stirring rate controllers. The sugar solution was pre-heated under a nitrogen flow in a separate chamber. The catalyst (0.2–0.4 g) was put in the reactor which was flushed with nitrogen before heating. The reaction temperature was in the range 90 – 120°C with the major part of experiments conducted at 110°C . The stirring rate was 1000 rpm to over-

come potential influence of external mass transfer, while application of fine catalyst powder (below $63\ \mu\text{m}$) ensured negligible influence of internal mass transfer. The catalytic experiments were performed under nitrogen with the overpressure 5 bar. Samples (1–2 ml) were periodically withdrawn through a $0.5\ \mu\text{m}$ sinter during the semi-batch experiments. The experiments were carried out typically for 240 minutes.

The pH of sugar and the catalyst slurry was measured for some cases after the reaction. As mentioned in the Introduction section, isomerization of glucose to fructose can occur at high alkaline pH (*ca.* 12–13). Much milder conditions in the present work rule out contribution of the homogeneous base catalysts to the experimental data.

HPLC analysis

The concentrations of reactants and products were determined by high-performance liquid chromatography (HPLC) (HITACHI Chromaster HPLC) equipped with an RI detector. A Biorad HPX-87C carbohydrate column was used and the mobile phase was 1.2 mM CaSO_4 . The temperature of the column was 70°C , the flow rate of the mobile phase was $0.5\ \text{ml}\ \text{min}^{-1}$ and the detector was at 40°C . Calibrations were made for all reacting components as well as potential side products.

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Notes and references

- 1 J. J. Bozell and G. R. Petersen, *Green Chem.*, 2010, **12**, 539.
- 2 D. M. Alonso, J. Q. Bond and J. A. Dumesic, *Green Chem.*, 2010, **12**, 1493.
- 3 D. Yu. Murzin and I. L. Simakova, in *Comprehensive Inorganic Chemistry II, vol. 7, From Elements to Applications*, ed. R. Schlögl and J. W. Niemantsverdriet, 2013, vol. 7, p. 559.
- 4 S. H. Bhosale, M. B. Rao and V. V. Deshpande, *Microbiol. Rev.*, 1996, **60**, 280.
- 5 C. A. Lobry de Bruyn and W. Alberda van Ekenstein, *Recl. Trav. Chim. Pays-Bas*, 1895, **14**, 201–206.
- 6 S. J. Angyal, *Carbohydr. Res.*, 1997, **300**, 279–281.
- 7 I. Delidovich and R. Palkovits, *ChemSusChem*, 2016, **9**, 547.
- 8 A. A. Marianou, C. M. Michailof, A. Pineda, E. F. Iliopoulou, K. S. Triantafyllidis and A. A. Lappas, *ChemCatChem*, 2016, **8**, 1100.
- 9 C. Kooyman, K. Vellenga and H. de Wilt, *Carbohydr. Res.*, 1977, **54**, 33.
- 10 B. Y. Yang and R. Montgomery, *Carbohydr. Res.*, 1996, **280**, 27.



- 11 C. J. Knill and J. F. Kennedy, *Carbohydr. Polym.*, 2003, **51**, 281.
- 12 J. M. Carraher, C. N. Fleitman and J.-P. Tessonnier, *ACS Catal.*, 2015, **5**, 3162.
- 13 C. Moreau, R. Durand, A. Roux and D. Tichit, *Appl. Catal., A*, 2000, **193**, 257.
- 14 J. Lecomte, A. Finiels and C. Moreau, *Starch/Staerke*, 2002, **54**, 75.
- 15 C. Moreau, J. Lecomte and A. Roux, *Catal. Commun.*, 2006, **7**, 941.
- 16 S. Yu, E. Kim, S. Park, I. K. Song and J. C. Jung, *Catal. Commun.*, 2012, **29**, 63.
- 17 I. Delidovich and R. Palkovits, *J. Catal.*, 2015, **327**, 1.
- 18 I. Delidovich and R. Palkovits, *Catal. Sci. Technol.*, 2014, **4**, 4322.
- 19 A. Takagaki, M. Ohara, S. Nishimura and K. Ebitani, *Chem. Commun.*, 2009, 6276.
- 20 A. Takagaki, S. Furusato, R. Kikuchi and S. T. Oyama, *ChemSusChem*, 2015, **8**, 3769.
- 21 M. Moliner, Y. Roman-Leshkov and M. E. Davis, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 6164.
- 22 V. Choudhary, A. B. Pinar, S. I. Sandler, D. G. Vlachos and R. F. Lobo, *ACS Catal.*, 2011, **1**, 1724.
- 23 J. Hajek, D. Yu. Murzin, T. Salmi and J.-P. Mikkola, *Top. Catal.*, 2013, **56**, 839.
- 24 L. Ren, Q. Guo, P. Kumar, M. Orazov, D. Xu, S. M. Alhassan, K. A. Mkhoyan, M. E. Davis and M. Tsapatsis, *Angew. Chem., Int. Ed.*, 2015, **54**, 10848.
- 25 J. Tuteja, S. Nishimura and K. Ebitani, *Bull. Chem. Soc. Jpn.*, 2012, **85**, 275.
- 26 K. K. Yadav, S. Ahmad and S. M. S. Chauhan, *J. Mol. Catal. A: Chem.*, 2014, **394**, 170.
- 27 J. A. Moulijn, M. Makee and A. E. van Diepen, *Chemical Process Technology*, Wiley, Chichester, 2013.
- 28 S. Buchholz, V. Michele, L. Mleczko, C. Muennich, R. Rudolf and A. Wolf, Patent WO2007093337 to Bayer Material Science AG, 2007.
- 29 J. Rufner, D. Anderson, K. van Benthem and H. R. Castro, *J. Am. Ceram. Soc.*, 2013, **96**, 2077.
- 30 V. L. Kuznetsov, S. N. Bokova-Sirosh, S. I. Moseenkov, A. V. Ishchenko, D. V. Krasnikov, M. A. Kazakova, A. I. Romanenko, E. N. Tkachev and E. D. Obratsova, *Phys. Status Solidi B*, 2014, **251**, 2444.
- 31 I. Mazov, V. Kuznetsov, I. Simonova, A. Stadnichenko, A. Ishchenko, A. Romanenko, E. Tkachev and O. Anikeeva, *Appl. Surf. Sci.*, 2012, **258**, 6272.
- 32 A. S. Andreev, M. A. Kazakova, A. V. Ishchenko, A. G. Selyutin, O. B. Lapina, V. L. Kuznetsov and J.-B. d'Espinose de Lacaillerie, *Carbon*, 2017, **114**, 39.
- 33 D. Kubička and R. Černý, *Ind. Eng. Chem. Res.*, 2011, **51**, 8849.

