


Cite this: *RSC Adv.*, 2017, 7, 26487

Direct conversion of carbohydrates to diol by the combination of niobic acid and a hydrophobic ruthenium catalyst†

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Tetrahydro-2,5-furandimethanol (THFDM) was obtained directly from a wide variety of carbohydrates by the combination of niobic acid and a hydrophobic ruthenium catalyst. Fructose, glucose, and polysaccharides consisting of fructose or glucose could be converted to THFDM in one-step. The selectivity to THFDM was kept around 60% while the glucose conversion varied from 9% to 49%. The as-synthesized niobic acid was characterized by TEM, N₂ adsorption/desorption, XRD, NH₃-TPD and FT-IR spectra of adsorbed pyridine. The niobic acid was proved to have medium and strong acid sites with a high Brønsted/Lewis ratio, which played a great role for keeping high THFDM selectivity using glucose as a substrate.

Received 6th April 2017
Accepted 11th May 2017

DOI: 10.1039/c7ra03939e

rsc.li/rsc-advances

Introduction

Production of chemicals and fuels from carbohydrates is an attractive way for alleviating the shortage of non-renewable fossil resources.^{1–5} One of the main strategies for the valorization of carbohydrates is the synthesis of platform compounds and their high value transformation. The conversion of carbohydrates to platform compounds such as 5-(hydroxymethyl) furfural (HMF),^{6–10} furfural,^{11–13} levulinic acid (LA)^{14,15} and the utilization of platform compounds have both been extensively researched. Furfuryl alcohol,^{16–20} tetrahydrofurfuryl alcohol,^{21,22} dimethylfuran,²³ 2,5-furandicarboxylic acid,²⁴ diformylfuran,^{25–31} cyclopentanone,^{32,33} γ -valerolactone,^{14,34–36} and tetrahydro-2,5-furandimethanol^{37,38} are the typical target products.

Considering that the platform compounds such as HMF are not stable and not easy to be separated, purified and stored, the conversion of carbohydrates to target products in one pot has attracted attention increasingly.^{39–47} A large number of works have been reported on the direct carbohydrate conversion. Sen and his co-workers gave reports on the direct conversion of fructose to 2,5-dimethyltetrahydrofuran or 5-methylfurfural by using RhCl₃ as the hydrogenation catalyst and HI as acid promoter.⁴⁵ Sels *et al.* obtained high yields of liquid straight-chain alkanes directly from cellulosic feedstock in an one-pot biphasic catalytic system.⁴⁶

Alcohols, including monobasic alcohols and diols, are important chemical products from fossil resources. The carbohydrates are ideal renewable raw materials for the production of monobasic alcohols and diols as they contain a large amount of alcohol hydroxyl.^{48–56} A lot of research had been reported on this topic. For example, hexanols could be produced in high yield by conversion of cellulose over Ir-ReO_x/SiO₂ catalyst in biphasic reaction system.⁵⁶ And 1,5-pentanediol could be obtained in high yield from biomass *via* furfural and tetrahydrofurfuryl alcohol.⁵⁵

Tetrahydro-2,5-furandimethanol (THFDM) is a valuable diol that can be used as solvent or monomer⁵⁷ and also could be converted to 1,6-hexanediol,⁵⁸ an important monomer in the plastics industry, in very high yield. Recently, we achieved the direct conversion of fructose to tetrahydro-2,5-furandimethanol (THFDM) by the combination of Amberlyst-15 and hydrophobic Ru/SiO₂-TM in a water/cyclohexane biphasic system.⁷ However, the reported procedure was only effective for the conversion of fructose, a monosaccharide not common in nature.

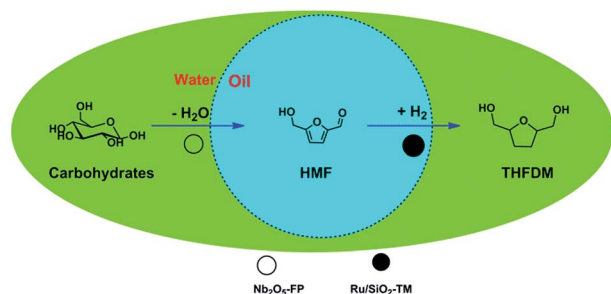
In this article, we reported a catalytic system which was able to provide THFDM directly from glucose, the monomer of most of the carbohydrates rich in nature. Based on this, we proved that this catalytic system was able to directly convert many kinds of carbohydrates including sucrose and starch to THFDM. The conversion of carbohydrates was illuminated in Scheme 1. The carbohydrates were first dehydrated to HMF in water phase on acids catalyst and then the HMF was hydrogenated to THFDM in oil phase on hydrophobic ruthenium catalyst. The hydrophobicity of ruthenium catalyst kept the ruthenium catalyst in oil phase. As a result the hydrogenation of carbohydrates under H₂ and the degradation of HMF were both avoided.

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ra03939e





Scheme 1 Conversion of carbohydrates into THFDM in the water/cyclohexane biphasic system.

Material and methods

Materials

All chemicals were of analytical grade and used as received unless otherwise stated. Inulin, sucrose, maltose, cellobiose, starch (soluble) and zirconium oxychloride were obtained from Aladdin Chemistry Co. Ltd. Fructose, glucose, and HF were purchased from Tianjin Kemiou Chemical Reagent Co. Ltd. Amberlyst-70 was obtained from Dow Chemical Shanghai Co. Ltd. RuCl_3 (Ru 37%) was purchased from Shenyang Research Institute of Nonferrous Metal. HY-zeolite (Si/Al = 5) was purchased from Nankai University Catalyst Co. Ltd. China. Niobium oxide was obtained from UCB Pharma. Pyridine and toluene were redistilled before used. All other reagents were commercially available.

Preparation of catalysts

The SiO_2 (4.00 g) was dispersed in the solution of RuCl_3 (RuCl_3 : 542.1 mg, 2.00 mmol, H_2O : 24.00 g). The mixture was stirred and kept at room temperature for 12 h and then dried at 393 K for 6 h. Before used, the catalyst was reduced at 673 K by H_2 for 2 h and then passivated in 1% O_2/N_2 at room temperature to afford Ru/SiO_2 . The Ru/SiO_2 -TM catalyst was prepared by CH_3 -functionalization of Ru/SiO_2 . Ru/SiO_2 (2.00 g) was transferred into a 100 mL round flask to which toluene (40 mL), then trimethylchlorosilane (10 mL) and pyridine (10 mL) were introduced. The mixture refluxed under N_2 for 24 h, and then filtered and washed with alcohol for five times. The resulted solid were dried at 323 K in vacuum overnight.

Nb_2O_5 (1.00 g) was added into H_3PO_4 aqueous solution (30 mL, 0.5 M) and stirred for 8 h at 343 K. Then the precipitation was washed with water (50 mL \times 3 times) and dried overnight at 393 K. The resulted white powder was calcined in air at 573 K for 3 h to afford the H_3PO_4 -treated niobic acid (Nb_2O_5 -P).

Nb_2O_5 (10.00 g) was added in HF aqueous solution (45 mL, 40 wt%). The mixture was stirred at 353 K for 24 h to afford a clear solution. Then concentrated ammonia water was added drop wise to adjust the pH 8–9 under drastic stirring. The precipitation was centrifuged and washed with ammonia water for 6 times. The solid was dried at 393 K to afford niobic acid. The H_3PO_4 -treated procedure was the same as the preparation of Nb_2O_5 -P to afford the Nb_2O_5 -FP.

$\text{ZrOCl}_2 \cdot 8\text{H}_2\text{O}$ (16.20 g) was dissolved in water (100 mL). Then concentrated ammonia water was added drop wise to adjust the pH 8–9 under drastic stirring. After kept stand for 24 h at room temperature, the precipitation was centrifuged and washed with water until no Cl^- was detected by AgNO_3 . The precipitation was dried at 373 K to afford zirconium hydroxide solid. Then the zirconium hydroxide was impregnated by H_2SO_4 (1 mol L^{-1}) under vigorous stirring at room temperature for 4 h. After centrifuged, the precipitation was dried at 393 K overnight and calcined at 773 K for 4 h to afford the $\text{SO}_4^{2-}/\text{ZrO}_2$.

Characterization of catalysts

Fourier transform infrared (FT-IR) spectra were collected on a Bruker Tensor 27 FT-IR spectrometer in KBr media. Samples were thoroughly dried before measured.

The X-ray powder diffraction (XRD) patterns were obtained using Rigaku D/Max 2500/PC powder diffractometer with $\text{Cu K}\alpha$ radiation ($\lambda = 0.15418$ nm) at 40 kV and 200 mA in a scanning rate of 5°min^{-1} .

The transmission electron microscopy (TEM) was conducted on a JEOLJEM-2000EX electron microscopy.

N_2 physical adsorption/desorption measurement was carried out at liquid nitrogen temperature on an Autosorb-1 Quantachrome instrument. Samples were pre-degassed at 300 $^\circ\text{C}$ for about 10 h to remove water and other physical adsorbed species.

Temperature-programmed desorption of ammonia (NH_3 -TPD) was performed on a Micromeritics AutoChem II 2920 Instrument with a thermal conductivity detector (TCD). The catalyst was degassed at 673 K in an atmosphere of Ar.

The FT-IR spectra for pyridine adsorption of the catalyst were conducted on a Bruker Tensor 27 FT-IR spectrometer. Prior to pyridine adsorption, the catalyst was evacuated in vacuum at 523 K. After that, the sample was used for the background reference collected. Subsequently, pulses of pyridine were introduced in an *in situ* cell for adsorption. Then the catalyst was heated to 523 K and evacuated for 30 min. The spectra of the catalysts were collected after the reaction cell was cooled to room temperature.

Catalytic reactions

The catalytic reactions were performed in a 60 mL stainless steel autoclave equipped with a magnetic stirrer, a pressure gauge, and automatic temperature control apparatus. The reactor was connected to a hydrogen cylinder for reaction pressure. In a typical experiment, an aqueous solution of carbohydrates (3 mL, 1.00 mmol by monosaccharide), cyclohexane (6 mL), and catalysts were put into the reactor. After sealing and purging with H_2 for 4 times to exclude air, the autoclave was heated to the desired temperature, and then H_2 (4 MPa) was charged into the reactor. The magnetic agitator was set to 1200 rpm to start the reaction. After reaction, the autoclave was cooled. The aqueous phase and oil phase were sampled and analyzed respectively. After sampling ethanol was added to make the acid and Ru/SiO_2 -TM in one phase. The catalysts were centrifuged



and washed with 50 mL ethanol for 3 times. After dried at 373 K under vacuum overnight, the catalyst was used for the next run.

Products analysis

The products were identified by Agilent 6890N GC/5973MS as well as by comparison with the retention times to corresponding standards in GC and HPLC traces. The amounts of products were determined based on GC and HPLC data. Gas chromatography measurements were conducted on Agilent 6890A GC. HP-5 capillary column (30 m × 530 μm × 1.5 μm) was used for separation of reaction mixtures. 1,2,4,5-Tetramethylbenzene (TMB) was used as the internal standard. HPLC analysis was carried out using a Waters e2695 HPLC system equipped with a 2414 refractive index detector maintained at 303 K. The aqueous sample was separated using a Waters Sugar-Pak 1 column (6.5 mm × 30 cm) at 358 K, using ethylenediaminetetraacetic acid calcium disodium salt hydrate (EDTACa) aqueous solution (50 mg L⁻¹) as the mobile phase at a flow rate of 0.55 mL min⁻¹. The conversion and selectivity of main products were evaluated as below:

$$\text{Conversion (\%)} = \left(1 - \frac{\text{moles of substrate}}{\text{moles of substrate loaded initially}}\right) \times 100\%$$

$$\text{Selectivity (\%)} = \frac{\text{moles of products}}{\text{moles of substrate converted}} \times 100\%$$

Results and discussion

We focused our attentions on the selection of different kinds of acid using glucose as the starting material. The HY, A-70, H₂SO₄, SO₄²⁻/ZrO₂ and Nb₂O₅ were selected as acids. The result was exhibited in Fig. 1. The acids were first used in the conversion of glucose to HMF (Fig. 1a). All the acids except Nb₂O₅-FP gave poor selectivity to HMF while the Nb₂O₅-FP had 78% selectivity to HMF after 1 h. And the selectivity of HMF was 56% after 4 h with 27% conversion of glucose. The results showed that the Nb₂O₅-PF was an effective solid acid for the conversion of glucose to HMF as reported in the literature.⁵⁹ When the acids were used in the conversion of glucose to THFDM, HY, A-70, H₂SO₄, SO₄²⁻/ZrO₂ resulted poor selectivity of THFDM (no more than 20% Fig. 1b) in the experiment range. This was attributed to the poor selectivity of HMF from glucose when those acids were used as the catalyst (Fig. 1a). The selectivity of THFDM is positively correlated to the selectivity of HMF on the specific acid. The selectivity of THFDM increased to 31% with 35% glucose conversion when H₃PO₄-treated niobic acid (Nb₂O₅-P) was employed. Surprisingly, the selectivity of THFDM increased significantly to 64% when H₃PO₄-treated fresh niobic acid (Nb₂O₅-PF) was used, in spite of the conversion of glucose was only 9% after 1 h. When the reaction time was extended to 4 h, the selectivity of THFDM was 63% at 32% glucose conversion. The fresh niobic acid was prepared by dissolution of

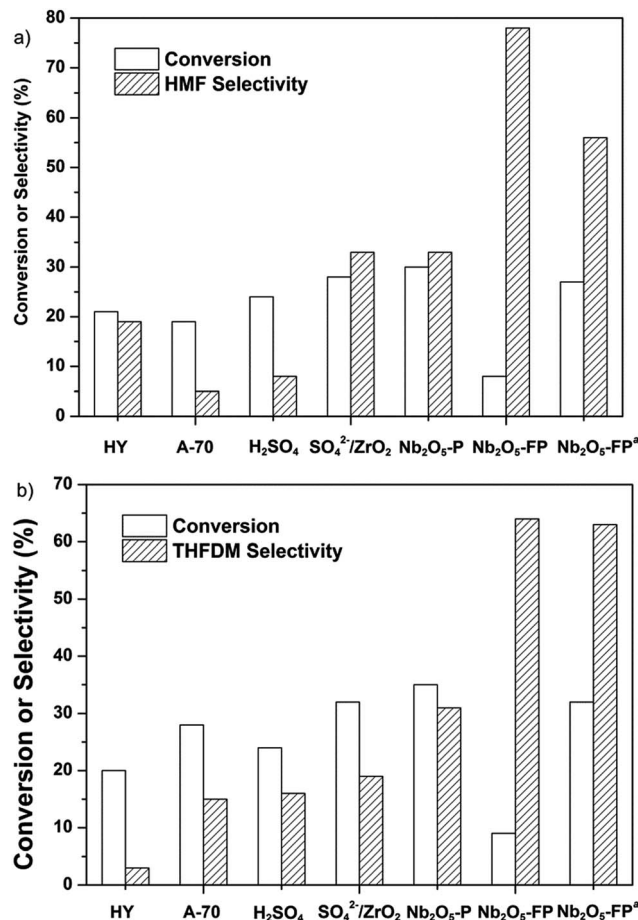


Fig. 1 The conversion of glucose by (a) acids (b) Ru/SiO₂-TM and acids. Reaction conditions: acids (40 mg), glucose aqueous solution (3 mL, glucose 1 mmol), cyclohexane (6 mL), H₂ (4 MPa), 433 K, 1 h. Ru/SiO₂-TM (30 mg) for (b). *Reaction time 4 h. HY: HY-zeolite (Si/Al = 5), A-70: Amberlyst-70.

niobium oxide and then precipitated by aqueous ammonia. And the Nb₂O₅-FP was used in subsequent reactions.

The performance of Nb₂O₅-FP was further studied by carrying out the reaction in two cases. The case I was the conversion of glucose to HMF under H₂ atmosphere by just using the Nb₂O₅-FP as the catalyst. And the combination of Nb₂O₅-FP and Ru/SiO₂-TM was used for the other case named case II. The conversion-time profiles of both cases were almost the same as shown in Fig. 2. This should be ascribed to the hydrophobicity of the Ru/SiO₂-TM. The Ru/SiO₂-TM was kept in the oil phase where there had no glucose. Thus, the conversion of glucose was catalyzed by Nb₂O₅-FP in both cases which led to the same conversion of glucose at the same reaction time.

The products selectivity *versus* glucose conversion was shown in Fig. 3. Very high selectivity of HMF was obtained at the initial stage for the case I (Fig. 3b). Meanwhile, 24% selectivity of fructose was gained in this case. It showed that the Nb₂O₅-PF was able to convert glucose to fructose. With the time prolonged the selectivity of fructose decreased to 3%. This was caused by the conversion of fructose to HMF. However, the selectivity of HMF decreased sharply from 76% to 27%, revealing that the



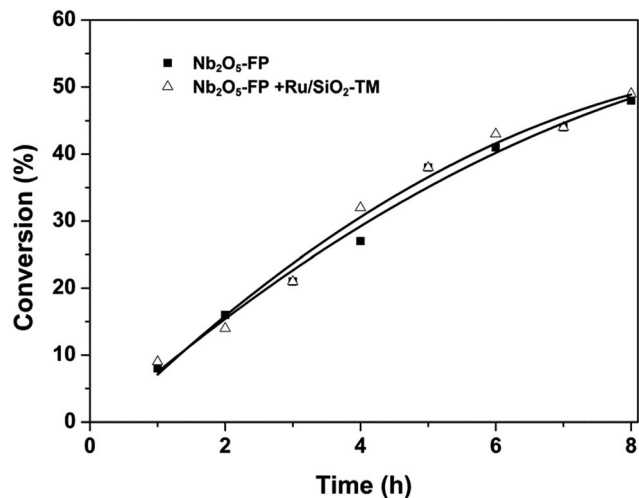


Fig. 2 Conversion–time profiles of glucose dehydration to HMF and glucose conversion to THFDM. Reaction conditions: Nb_2O_5 -FP (40 mg), glucose aqueous solution (3 mL, glucose 1 mmol), cyclohexane (6 mL), H_2 (4 MPa), 433 K. Δ : Ru/SiO₂-TM (30 mg).

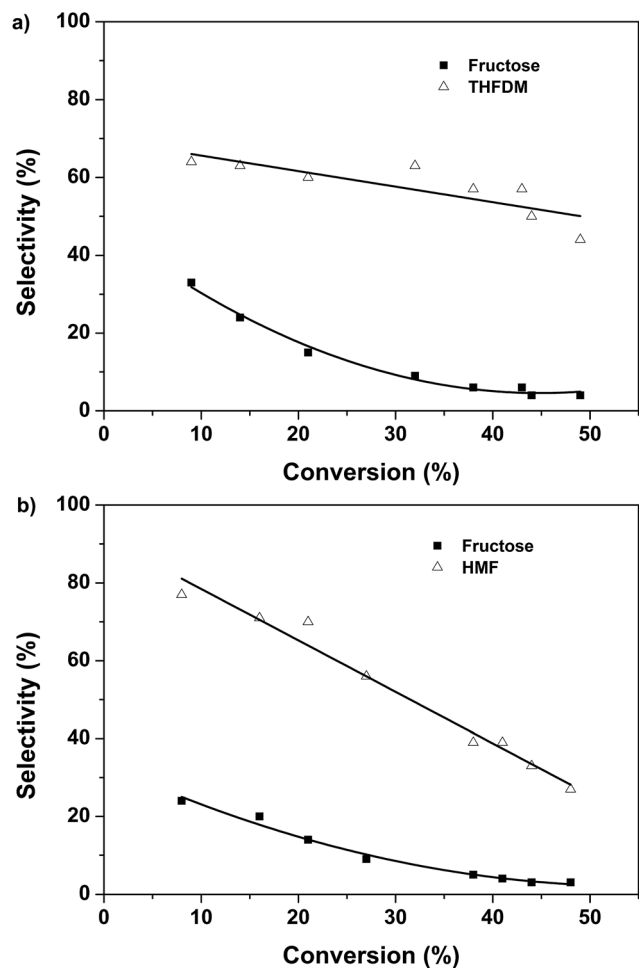


Fig. 3 Products selectivity versus glucose conversion over (a) Nb_2O_5 -FP and Ru/SiO₂-TM, (b) Nb_2O_5 -FP. Reaction conditions: Nb_2O_5 -FP (40 mg), glucose aqueous solution (3 mL, glucose 1 mmol), cyclohexane (6 mL), H_2 (4 MPa), 433 K. Ru/SiO₂-TM (30 mg) for (a).

HMF was not stable and was easy to degrade under this condition. This was in accordance with the results we obtained previously.⁷ The case II was the conversion of glucose to THFDM by the combination of Nb_2O_5 -FP and Ru/SiO₂-TM. The results at the initial stage were analogous to that of the case I (Fig. 3a). High selectivity of THFDM with quite a number of fructose was obtained. To our delight, the selectivity of THFDM kept around at 60% with no obviously decrease as the time prolonged (Fig. 3a). This phenomenon was quite different from the selectivity of HMF in the case I. The reason was that the THFDM was much more stable than the HMF. The HMF was converted to THFDM once generated in the case II. As a result, the degradation of HMF was inhibited and the selectivity of THFDM was well kept.

The effect of temperature played an important role in the conversion of glucose and the results were shown in Fig. 4. The conversion of glucose increased as the temperature increased from 413 K to 453 K as expected. The selectivity of THFDM kept around 60% when the temperature was not higher than 433 K. However, once the temperature exceeded 433 K the selectivity of THFDM decreased drastically. The best result was obtained at a temperature of 433 K. And the reaction was repeated at this temperature. Similar conversion of glucose and selectivity of THFDM was acquired showing the repeatability of the catalytic system.

Fig. 5 showed the recyclability of the catalytic system. After the analysis, ethanol was added into the system to obtain one liquid phase. The catalyst was then centrifuged and fully washed with ethanol after each cycle. After dried at 373 K under vacuum, the catalyst was used for the next run. It could be seen from Fig. 5 that there had a small gradual decrease in conversion value. This should be ascribed to the incomplete recovery of catalyst after each reaction. The selectivity of THFDM kept around 60% in all the four consecutive runs. This illustrated the catalyst was stable under the reaction conditions.

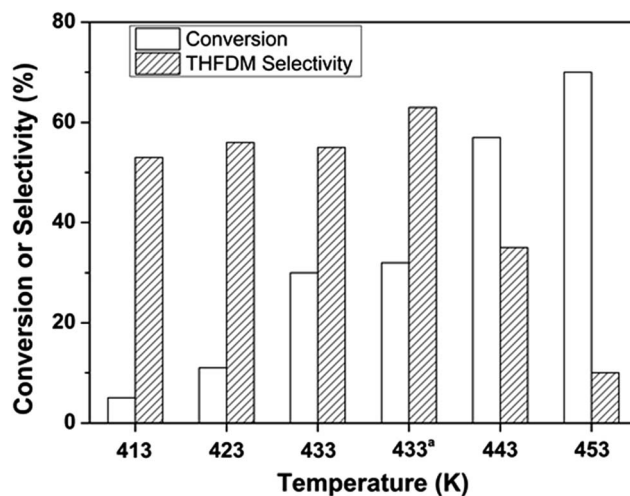


Fig. 4 Effect of temperature on the conversion of glucose to THFDM. ^aRepeated test for the temperature at 433 K. Reaction conditions: Nb_2O_5 -FP (40 mg), Ru/SiO₂-TM (30 mg), glucose aqueous solution (3 mL, glucose 1 mmol), cyclohexane (6 mL), H_2 (4 MPa), 433 K, 4 h.



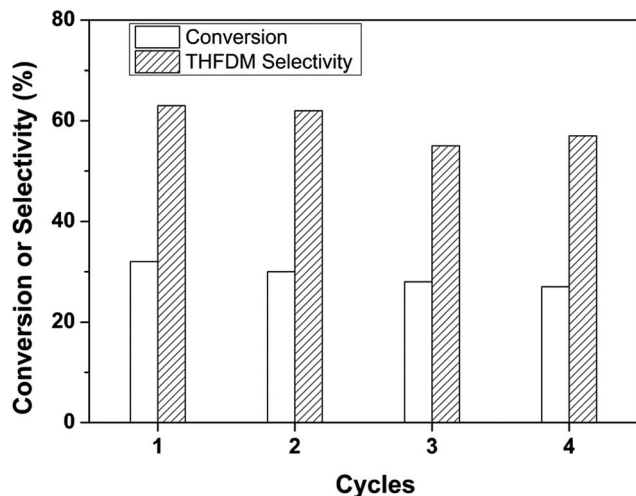


Fig. 5 Cycle experiments of the conversion of glucose to THFDM. Reaction conditions: Nb₂O₅-FP (40 mg), Ru/SiO₂-TM (30 mg), glucose aqueous solution (3 mL, glucose 1 mmol), cyclohexane (6 mL), H₂ (4 MPa), 433 K, 4 h.

We attempted to obtain THFDM directly from carbohydrates, containing glucose or fructose units, by the combination of Nb₂O₅-FP and Ru/SiO₂-TM. The fructose was almost completely converted under the reaction condition (Table 1, entry 2). However, the selectivity of THFDM was only 41% showed that the conditions adopted here were not appropriate for the conversion of fructose. As a result, the inulin, a polymer consisting of chain-terminating glucosyl moieties and a repetitive fructosyl moiety, gave 33% selectivity of THFDM although with 96% of conversion (Table 1, entry 3). The conversion of sucrose was higher than that of the other two kinds of disaccharides maltose and cellobiose (Table 1, entries 4–6). The reason was that the sucrose was a disaccharide comprised of glucose and fructose while the maltose and cellobiose were comprised of glucose only. The selectivity of THFDM for the three disaccharides was around 40%. The starch, the

Table 1 Conversion of carbohydrates to THFDM^a

Entry	Substrate	Conv. ^b (%)	Selectivity (%)			U. D. ^d
			THFDM	Fructose	Others ^c	
1	Glucose	32	63	9	15	13
2	Fructose	98	41	1	27	31
3	Inulin	96	33	4	31	32
4	Sucrose	61	42	4	25	29
5	Maltose	37	42	7	22	29
6	Cellobiose	33	32	8	19	41
7	Starch	26	34	11	19	36

^a Reaction conditions: Nb₂O₅-FP (40 mg), Ru/SiO₂-TM (30 mg), carbohydrate aqueous solution (3 mL, 1 mmol by monosaccharide), cyclohexane (6 mL), H₂ (4 MPa), 433 K, 4 h. ^b Conversion was calculated based on the amounts of monosaccharide. ^c Others referred to furfuryl alcohol, methyl furfuryl alcohol, γ -valerolactone, 1,2-pentanediol, 1,2-hexanediol, 1,2,5-hexanetriol, 1,2,6-hexanetriol. ^d Unidentified products.

polysaccharide of glucose, was also able to be converted to THFDM in one-step by this catalytic system. A conversion of 26% with 34% selectivity of THFDM was obtained under the same reaction conditions (Table 1, entry 7).

The physical properties of the Nb₂O₅-FP were characterized by BET and TEM. The results were shown in Fig. 6 and S3.† The BET surface of as-synthesized Nb₂O₅-FP was 62 m² g⁻¹. According to the IUPAC classifications, the sample had the characteristics of typical type IV physisorption isotherms, indicating the presence of mesoporous structure. There had a significantly increase in adsorbed volume at relatively high pressure which was the characteristic of the H3 type hysteresis loop indicating the existence of slit-shaped pores with a large pore size which was in accordance with the results of TEM. The distribution of pore size calculated by the non-local density functional theory (NLDFT) model showed there had very broad pore size distribution in the mesoporous range (2–50 nm) and macropore were also presented in the Nb₂O₅-FP.

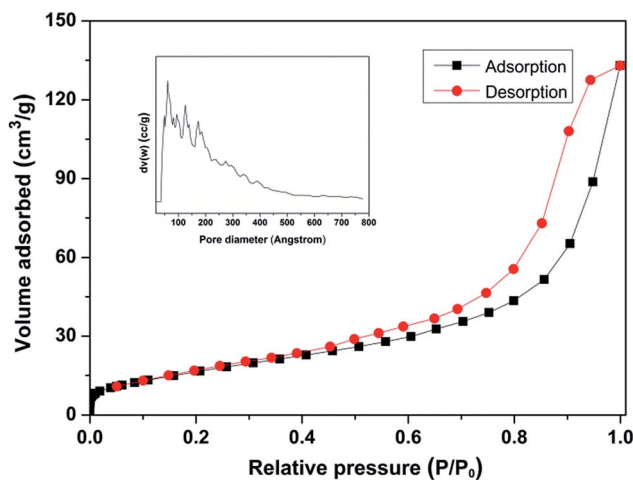


Fig. 6 N₂ adsorption/desorption isotherms of Nb₂O₅-FP.

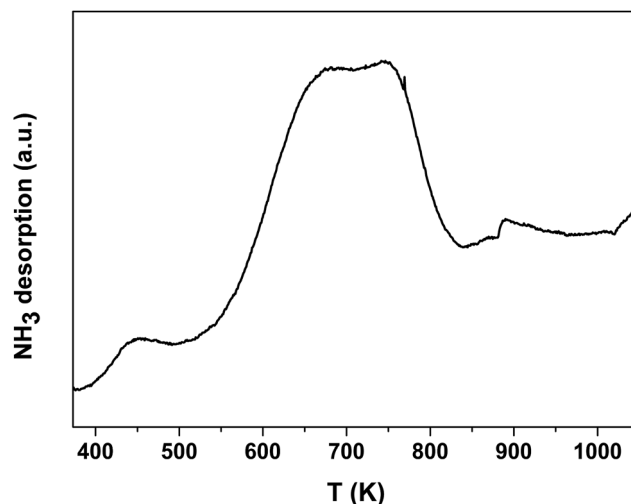


Fig. 7 NH₃-TPD profiles of Nb₂O₅-FP.



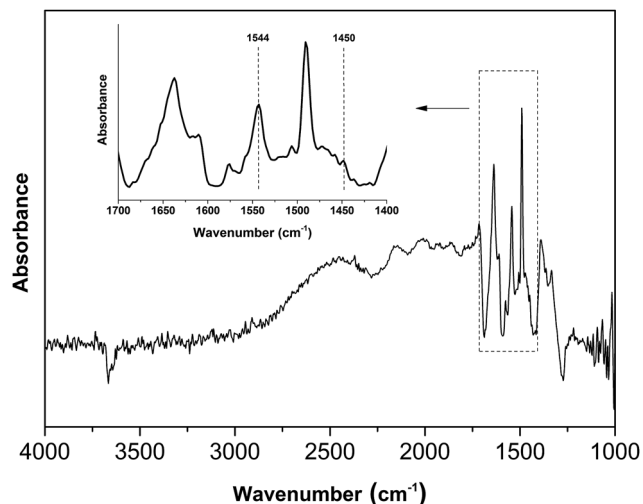


Fig. 8 FT-IR spectrum of Nb₂O₅-FP after pyridine adsorption and subsequent evacuation, the sample was treated in vacuum to remove adsorbed water.

The acid property of the Nb₂O₅-FP was characterized by temperature-programmed desorption of ammonia (NH₃-TPD) (Fig. 7) as well as Fourier transform infrared (FT-IR) spectroscopy of adsorbed pyridine (Fig. 8). There were two desorption peaks appeared in the NH₃-TPD profile. The weak desorption peak at low temperature (400–500 K) should be ascribed to the weakly held ammonia, probably the hydrogen-bonded ammonia according to the literature.⁶⁰ The strong broad desorption peak between 600 and 800 K corresponded to large amount of the medium and the strong acid sites in the sample. The FT-IR spectroscopy of adsorbed pyridine revealed the type of the acid sites.^{61,62} The medium peak at 1544 cm⁻¹ was attributed to pyridine protonated on Brønsted sites, and the weak peak at 1450 cm⁻¹ indicated the presence of Lewis acid sites. The peak intensity at 1544 cm⁻¹ and 1450 cm⁻¹ elucidated that the sample had a very high Brønsted/Lewis ratio. And the high Brønsted/Lewis ratio was favourable for high selectivity of HMF from glucose in the aqueous phase.⁶³

Conclusions

In conclusion, the direct conversion of a wide variety of carbohydrates to THFDM was achieved by the combination of Nb₂O₅-FP and Ru/SiO₂-TM in the water/cyclohexane biphasic system. The as-synthesized Nb₂O₅-FP, which was proved to have large amount of medium and strong acid sites with high Brønsted/Lewis ratio, was essential and effective for keeping high THFDM selectivity when glucose was used as substrate. This work provides a common strategy for the direct production of valuable chemicals from carbohydrates.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (21272109 and 21371089) and the

Intercollegiate Key Scientific Research Projects of Henan Province (16A150017). The authors express their gratitude for the help and guidance from Researcher Jie Xu at Dalian Institute of Chemical Physics, Chinese Academy of Sciences.

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