Sustainable production of benzene from lignin

Qinglei Meng, Jiang Yan, Ruizhi Wu, Huizhen Liu, Yang Sun, NingNing Wu, Junfeng Xiang, Lirong Zheng, Jing Zhang & Buxing Han

Benzene is a widely used commodity chemical, which is currently produced from fossil resources. Lignin, a waste from lignocellulosic biomass industry, is the most abundant renewable source of benzene ring in nature. Efficient production of benzene from lignin, which requires total transformation of C\(_{sp2}\)-C\(_{sp3}\)/C\(_{sp2}\)-O into C-H bonds without side hydrogenation, is of great importance, but has not been realized. Here, we report that high-silica HY zeolite supported RuW alloy catalyst enables in situ refining of lignin, exclusively to benzene via coupling Bronsted acid catalyzed transformation of the C\(_{sp2}\)-C\(_{sp3}\) bonds on the local structure of lignin molecule and RuW catalyzed hydrogenolysis of the C\(_{sp2}\)-O bonds using the locally abstracted hydrogen from lignin molecule, affording a benzene yield of 18.8% on lignin weight basis in water system. The reaction mechanism is elucidated in detail by combination of control experiments and density functional theory calculations. The high-performance protocol can be readily scaled up to produce 8.5 g of benzene product from 50.0 g lignin without any saturation byproducts. This work opens the way to produce benzene using lignin as the feedstock efficiently.
Nowadays, benzene is an indispensable commodity in chemical industry with world production of more than 61 million metric tons in 2019 (www.statista.com/statistics/1108114/global-benzene-capacity). The global benzene demand is anticipated to grow with a rate of 2.9% annually in the next decade (www.chemanalyst.com/industry-report/benzene-market-56). Especially in the manufacturing industry, benzene is widely used in many sectors, where it is combined and processed with other basic chemicals (e.g. ethylene and propylene) to produce valuable consumer goods, such as clothing, packaging, car parts, building materials, pharmaceuticals, cosmetics, flame retardant, compact discs, eyeglass lenses, carpet, medical implants, foam insulation, adhesives, footwear, contact lenses, dyes, agrochemicals (Supplementary Fig. 1) (www.statista.com/statistics/1108114/global-benzene-capacity), (www.chemanalyst.com/industry-report/benzene-market-56). Currently, benzene is dominantly produced from petroleum and coal via catalytic reforming, steam cracking and toluene disproportionation processes, as well as coal processing (Supplementary Note 1) (Fig. 1a)3–5. Besides, there is also ongoing research to convert methane to benzene (Fig. 1b)6–8. However, all of the above routes depend on fossil resource, and has several disadvantages, such as complicated and severe conditions, high energy consumption and serious environmental pollution9–11. Thus, more benign and sustainable strategy, such as utilization of renewable resources as raw materials to economically produce benzene, is highly desired, which can liberate us from the reliance on fossil resource and is of great industrial and social significance12,13. Such an attractive, cheap and non-edible material is lignocellulosic biomass, generated from forestry and agricultural activity worldwide14. As a main constituent of lignocellulose, lignin is the most abundant renewable source composed of aromatic building blocks in nature15, with an annual production of around 50 billion metric tons16. In terms of molecular structure, the aromatic character of lignin springs from the benzene ring structures, which renders itself a sustainable and desirable candidate feedstock for benzene production16–18.

In recent years, valorizing lignin into fossil-based chemicals has received extensive attention, and various valuable chemicals used for transport fuels and industrial manufactures have been
obtained employing lignin as the feedstock. For example, advances in the seminal work of Luterbacher and co-workers on aldehyde stabilization of lignin improved the lignin depolymerization efficiency and yielded the mixture of guaiacyl and syringyl monomers, such as alkyl-substituted guaiacol/syringol, hydroxyprefyl-substituted guaiacol/syringol and guaiacyl/syringyl propane dienes, via hydrogenolysis and oxidation approaches. In the pioneering work of Stahl and co-workers, lignin could be depolymerized into syringaldehyde and vanillin via successive oxidation-formylation-hydrolysis processes. Wang and co-workers conducted the direct hydrodeoxygenation of lignin to alkyl-substituted arenes and hydrocarbons. Sels and co-workers developed an integrated biofinery process that converted lignin into phenol with high yield. Wang and co-workers also obtained phenol product from lignin by a multistep oxidation and decarbonylation methodology. These findings place a premium on strategies that pursue benzene from lignin, however, in the currently reported methodologies, such as catalytic pyrolysis, hydrodeoxygenation and combined catalytic processing, benzene could only be detected in quite a low yield in the complex mixture containing the aforementioned phenolic hydroxyl, methoxyl and alkyl-substituted aromatic products. Although there is ongoing research on lignin valorization, efficient lignin-to-benzene route has not been reported so far. Obviously, the complex chemical bonding environment, especially the stable aromatic carbon–aliphatic carbon (Csp2–Csp3) and aromatic carbon–oxygen (Csp2–O) bonds functionalized on the benzene rings truly limited the abstraction of benzene from the lignin structure. In this context, further cleavage of the unbroken Csp2–Csp3/Csp2–O bonds in the above generated alkylbenzenes and alkylphens for desired benzene product can only be realized via dealkylation, dealkenylation and reductive cleavage processes under quite harsh conditions (for example, high temperature, etc.). Moreover, such required reaction conditions often lead to inevitable side reactions. For example, the concurrent hydrogenation of the benzene ring in the hydroprocessing stage of the lignin upgrading with exogenous hydrogen source always further reduces the benzene selectivity. Thus, a key and great challenge for efficient lignin-to-benzene route would be to design practical strategy for combining the complicated refinements of the Csp2–Csp3/Csp2–O bonds into Csp2–H bonds via friendly catalysis, without multistep technological procedure and extra hydrogenation of benzene ring. In our previous work, we reported the hydrogenolysis of the aromatic Csp2–O(CH3) bond without need of any exogenous hydrogen or other reductants, termed as self-supported hydrogenolysis (SSH, Supplementary Fig. 2), by which the Csp2–O bond can be transformed into Csp2–H bond using the in situ abstracted hydrogen from the methoxyl group, and the side hydrogenation of the benzene ring can be completely avoided. Besides, some catalysts, such as MoO3, have also been proven to be effective in hydrodeoxygenation process of the lignin model compounds or lignin oil for arene products with high selectivity, using exogenous hydrogen.

To develop desired lignin-to-benzene route, in this work, we propose an integrated catalytic strategy that in situ refines the Csp2–Csp3/Csp2–O bonds functionalized on the benzene ring structures into Csp2–H bonds over the RuW/HY30 catalyst without any saturation of the ring, thereby achieving a single production of benzene from lignin (Fig. 1c). The in situ refining mechanism of the Csp2–Csp3/Csp2–O bonds is disclosed, which effectively couples the direct deconstruction of the Csp2–Csp3 bonds in the phenylpropanol units on the local structure of the lignin molecule without any preceding reductive fractionation process, and the hydrogenolysis of the Csp2–O bonds using the local-access hydrogen source in lignin molecule. With water acting as the cheap and green reaction medium, no other reductant or reactant is needed. This efficient strategy affords mild-condition abstraction of benzene from lignin molecule structures with a benzene yield as high as 18.8% on lignin weight basis, and our high-performance protocol can be readily scaled up to produce 8.5 g of benzene product from 50.0 g lignin without any saturation byproducts.

Results
Optimization study. We focused our in situ refining strategy studies initially on the overall transformation of the Csp2–Csp3/Csp2–O bonds by using 1-(4-methoxyphenyl)-1-propanol (1a) as the model compound because of its lignin-mimetic phenylpropanol structure [(CH3O)Ph–Csp3(OH)–] with only one methoxy group [Csp2–O(CH3)] and one 1-hydroxypropyl group [Csp2–Csp3–O–]. In the lignin depolymerization study, the optimized catalytic system and reaction conditions shown in Table 1 and Supplementary Fig. 3, which employed cheap and commercially available hydrogen-type faujasite Y zeolite (HY30, Si/Al ratio = 30) with supported RuW alloy components using water as the reaction medium at 180 °C under nitrogen atmosphere (Table 1, entries 1–3, Supplementary Fig. 3a, b). As expected, the transformations of the Csp2–Csp3/Csp2–O bonds in 1a did not occur without the catalyst, despite the achievable dehydroxylation (1b) in the blank experiment (Table 1, entry 1). It is clear that HY30 zeolite efficiently catalyzed the deconstruction of the Csp2–Csp3 bond in 1a into Csp2–H bond, but unfortunately the Csp2–O bond in anisole (1d) was not reactive when only HY30 was used as the catalyst (Table 1, entry 2). With RuW alloy introduced into the catalytic system, these unreacted anisole molecules could be further converted into benzene (1b) via the SSH process of the Csp2–O bond (Table 1, entry 3). It can be found that the selectivity to benzene (1b) could be 97.3% with a remained anisole (1d) selectivity of 2.5% at 99.9% conversion of 1a over the RuW/HY30 catalyst under nitrogen atmosphere in water (Table 1, entry 3, Supplementary Fig. 3b, c), and according to the 1H nuclear magnetic resonance (NMR) analysis of the aqueous phase (Supplementary Fig. 3d), there was no water-soluble product. As the reaction medium in the catalytic system, water also served as an ideal booster for the SSH reaction of the Csp2–O bond by way of forming hydrogen bonds with the O atom in anisole, which was beneficial for the in situ refining strategy. A total conversion of 1a into benzene product could be achieved in 6 h without any saturated cyclohexane product (1c) (Table 1, entry 4). In contrast, a test with exogenous hydrogen source led to benzene and cyclohexane products with competitive selectivities of 43.3% and 16.9%, which was additionally coupled with n-propylbenzene (1e) and n-propylcyclohexane (1f) in respective selectivities of 35.8% and 3.9% (Table 1, entry 5), albeit in lowered selectivities of cyclohexane (1c), n-propylbenzene (1e) and n-propylcyclohexane (1f) under low pressure of exogenous H2 (Table 1, entry 6). This is rationalized by the inevitable competitions of the hydrogenation of the benzene rings and hydrogenolysis of the hydroxyl group substituted at the aliphatic α-C (Cα) position (namely the Csp3 carbon in the Csp2–Csp3 bond, Fig. 1c) in the traditional hydroprocessing processes using exogenous hydrogen as reductant, which have been successfully avoided via the HY30 zeolite and RuW NPs accomplished refining strategy for the in situ transformation of the Csp2–Csp3 and Csp2–O bonds. To confirm the key roles of RuW and HY30 in the catalysts, we prepared the reference Ru/HY30, W/HY30, RuW/Al2O3 catalysts, and studied their catalytic performances for the reaction of 1a (Table 1, entries 7–10). Under nitrogen atmosphere, the
Table 1 Screening of catalysts for in situ refining 1-(4-methoxyphenyl)-1-propanol (1a) over supported metal catalysts at different conditions.

| Entry | Catalytic systemb | t (h) | Conv.(%) | Select. (%) | Ye. of 1b (%) |
|-------|-------------------|-------|----------|-------------|---------------|
|       | Catalystb         |       |          | 1b  | 1c  | 1d  | 1e  | 1f  | 1g  | 1h  |       |
| 1c    | -                 | 5.5   | 99.9     | 0.0  | 0.0  | 0.0  | 0.0  | 0.0  | 0.0  | 99.9 | 0.0  |
| 2     | HY50              | 1.0   | 99.9     | 0.0  | 0.0  | 99.0 | 0.0  | 0.0  | 0.0  | 0.0  | 97.2 |
| 3     | RuW/HY50          | 5.5   | 99.9     | 97.3 | 0.0  | 2.5  | 0.0  | 0.0  | 0.0  | 0.0  | 99.5 |
| 4     | RuW/HY50          | 6.0   | 99.9     | 99.6 | 0.0  | 0.0  | 0.0  | 0.0  | 0.0  | 0.0  | 99.5 |
| 5d    | RuW/HY50          | 6.0   | 99.9     | 43.3 | 16.9 | 0.0  | 35.8 | 3.9  | 0.0  | 0.0  | 43.2 |
| 6e    | RuW/HY50          | 6.0   | 99.9     | 26.6 | 2.0  | 0.0  | 10.8 | 0.5  | 0.0  | 0.0  | 86.5 |
| 7     | Ru/HY30           | 5.5   | 99.9     | 0.0  | 0.0  | 98.8 | 0.0  | 0.0  | 0.0  | 0.0  | 0.0  |
| 8     | W/HY30            | 5.5   | 99.9     | 0.0  | 0.0  | 98.6 | 0.0  | 0.0  | 0.0  | 0.0  | 0.0  |
| 9f    | RuW/SiO2          | 5.5   | 99.9     | 0.0  | 0.0  | 0.0  | 94.5 | 0.0  | 0.0  | 0.0  | 49.0 |
| 10e   | RuW/Al2O3         | 5.5   | 99.9     | 0.0  | 0.0  | 0.0  | 72.2 | 0.0  | 0.0  | 0.0  | 9.5  |
| 11    | RuW/HY5            | 5.5 | 99.9 | 13.2 | 0.0 | 0.0 | 73.3 | 0.0 | 0.0 | 0.0 | 4.3 |
| 12    | RuW/HY6            | 5.5 | 99.9 | 31.9 | 0.0 | 0.0 | 61.8 | 0.0 | 0.0 | 0.0 | 21.0 |
| 13    | RuW/HY15           | 5.5 | 99.9 | 96.0 | 0.0 | 3.5 | 0.0 | 0.0 | 0.0 | 0.0 | 9.5 |
| 14h   | RuW/HZSM-518       | 5.5 | 99.9 | 10.0 | 0.0 | 0.0 | 75.0 | 0.0 | 0.0 | 0.0 | 3.5 |
| 15    | RuW/Beta           | 5.5 | 99.9 | 8.9  | 0.0 | 0.0 | 70.0 | 0.0 | 0.0 | 0.0 | 8.3 |
| 16    | RuW/HZSM-514       | 5.5 | 99.9 | 8.1  | 0.0 | 0.0 | 74.9 | 0.0 | 0.0 | 0.0 | 7.7 |
| 17    | RuW/HZSM-518       | 5.5 | 99.9 | 7.5  | 0.0 | 0.0 | 78.1 | 0.0 | 0.0 | 0.0 | 7.1 |
| 18    | RuW/HZSM-520       | 5.5 | 99.9 | 6.3  | 0.0 | 0.0 | 82.6 | 0.0 | 0.0 | 0.0 | 6.2 |
| 19    | RuW/HZSM-5235      | 5.5 | 99.9 | 5.2  | 0.0 | 0.0 | 87.6 | 0.0 | 0.0 | 0.0 | 4.9 |
| 20    | RuW/Mordenite      | 5.5 | 99.9 | 2.0  | 0.0 | 0.0 | 92.9 | 0.0 | 0.0 | 0.0 | 3.0 |
| 21    | RuW/SAPO           | 5.5 | 99.9 | 0.0  | 0.0 | 0.0 | 83.8 | 0.0 | 0.0 | 0.0 | 9.2 |
| 22d   | RuW/MCM-41         | 5.5 | 99.9 | 0.0  | 0.0 | 0.0 | 94.0 | 0.0 | 0.0 | 0.0 | 5.1 |

aReaction results are the averages of three experiments conducted in parallel. 1a (1.0 mmol), H2O (5.0 mL), 180 °C, 0.1 MPa N2, 800 rpm.
bRu/HY30 (0.18 g, 2.5 wt% Ru) and W/HY30 (0.20 g, 14 wt% W) catalysts (the content of metal is based on HY30 zeolite and determined by ICP); RuW/zeolite, RuW/SiO2 and RuW/Al2O3 catalysts (0.20 g, 2.5 wt% Ru, 14 wt% W, the content of metal is based on zeolite, SiO2 and Al2O3 materials, and determined by ICP).
cWithout catalyst.
d1.0 MPa H2.
e0.1 MPa H2.
fAmorphous SiO2.
gγ-Al2O3.
hThe catalyst dosage is 1.5 times that of the RuW/HY30 catalyst.
iHydrogen-type Beta, Si/Al ratio is 25.
jHydrogen-type Mordenite, Si/Al ratio is 10.
kHydrogen-type SAPO-34, Si/Al ratios is 0.25.
lAll-silicon MCM-41.
monometallic Ru/HY30 (Table 1, entry 7) and W/HY30 (Table 1, entry 8) catalysts performed the same function as HY30 zeolite, and just led to the anisole product. Likewise, the bimetallic MW/HY30 (M = Ni, Co, Fe, Mo, and Cu) catalysts (Supplementary Table 1, entries 1–5) also only catalyzed the Csp2–Csp3 bond deconstruction without reaction of Csp2–O bond, proving that the combination of Ru and W was necessary for the SSH reaction of the Csp2–O bond. Although the Csp2–O bond could be hydrogenolyzed in the presence of the exogenous hydrogen over the MW/HY30 catalysts (for example, NiW/HY30, CoW/HY30, and FeW/HY30), their selectivities of benzene were much lower than that over RuW/HY30 catalyst (Supplementary Table 1, entries 6–10). As the constituent of HY30 zeolite, silicon and aluminum oxides supported RuW catalysts (Table 1, entries 9 and 10) only promoted the conversion of 1a to n-propylbenzene (1e) without any benzene product, disclosing that the unique catalytic properties derived from the zeolitic aluminosilicate [–TO2–(T = Si, Al)] frameworks can offer a particular way to tailor the Csp2–Csp3 bond into Csp2–H bond under mild conditions. The above control experiments showed that the HY30 zeolite and RuW alloy are respectively essential to realizing the transformations of the Csp2–Csp3 and Csp2–O bonds into Csp2–H bonds for producing benzene product (1b).

To get insight into the structural origins of the observed excellent catalytic performance, we characterized the catalysts by several techniques. A typical x-ray diffraction (XRD) pattern of the as-synthesized RuW/HY30 catalyst (Fig. 2a) revealed well-defined diffraction peaks of the RuW alloy and HY30 zeolite structures. Based on the Rietveld refinement analysis of the XRD data (Fig. 2b), the random occupancies of Ru and W atoms are estimated to be 20.3 and 79.7% in the RuW nanoparticles (NPs) with a range of 1.0–4.0 nm (Supplementary Fig. 4a–c). The Ru K-edge extended x-ray absorption fine structure (EXAFS) fitting (Fig. 2c and Supplementary Table 2) exhibits a Ru-W first shell coordination number (CN) of 3.1, while the W-Ru CN was just 0.9 due to the high W/Ru molar ratio (Fig. 2d and Supplementary Table 2), directly proving the alloy properties between the homogeneously distributed Ru and W atoms in the HY30 supported RuW NPs (Supplementary Fig. 4d–i). Accordingly, X-ray photoelectron spectroscopy (XPS) analysis of the RuW/HY30 catalyst showed the opposite shifts for the Ru3p (towards lower energy, Fig. 2e) and W4f (towards higher energy, Fig. 2f) orbital peaks, as compared respectively to the Ru/HY30 and W/HY30 catalysts, which can be ascribed to that the higher electronegativity of Ru induced electron transfer from W to Ru in the RuW NPs. This interplay between Ru and W atoms could also be discovered from the slightly lower energy absorption threshold value in the Ru K-edge x-ray absorption near-edge structure (XANES) curve of the RuW/HY30 catalyst (Supplementary Fig. 5a), as compared with that of Ru foil, despite the shift tendency to higher threshold value was inconspicuous for W L2-edge (Supplementary Fig. 5b). The analyses of the acid and textural properties indicated that the RuW/HY30 catalyst was highly strong Bronsted acidic (Supplementary Table 3), and had substantial mesoporous structures which are favorable to the access of reactant molecules to the acid sites, and then the rapid deconstruction of the Csp2–Csp3 bonds (Supplementary Table 4, Supplementary Fig. 4b, c). Compared with RuW/HY30 catalyst, although the RuW/HY catalysts with lower zeolite framework Si/Al ratios (3, 5, and 15) possessed more acid sites, their strong Bronsted acid sites were in lower proportions (Supplementary Table 3) and actually distributed in the micropores with higher proportions (Supplementary Table 4), which heavily restricted the deconstruction efficiency of the Csp2–Csp3 bonds on the Bronsted acid sites due to the inferior diffusibility of the microporous structures, matching that only less than one-third of 1a were transformed into benzene, and the main product was n-propylbenzene (1e) (Table 1, entries 11–13). RuW/HY40 catalyst with a higher zeolite Si/Al ratio of 40 had the appropriate textural properties, which gave a high selectivity of benzene (Table 1, entry 14), but requiring 1.5 times catalyst dosage of RuW/HY30 catalyst, due to its lower quantity of Bronsted acid sites (Supplementary Table 3). The overall transformation of 1a could also be achieved over other types of zeolite supported RuW catalysts with inferior mesoporous structures (Table 1, entries 15–23), but similarly generate n-propylbenzene (1e) in quantity, albeit with limited selectivities of benzene over RuW/Beta (Table 1, entry 15) and RuW/HZSM-5 (Table 1, entries 16–21) catalysts. It is well known that pure silica zeolites do not contain any acid sites, thus RuW/MCM-41 catalyst was incapable of Csp2–Csp3 bond deconstruction, also with n-propylbenzene (1e) as the main product (Table 1, entry 24). As the comparison of the utility of the RuW/HY30 catalyst, it is the poor activities of the supports in the above catalysts (Table 1, entries 9–13 and 15–24) on the deconstruction of the Csp2–Csp3 bonds that offered RuW the opportunity to hydrogenolyze the hydroxyl group substituted at the aliphatic Csp3 position of the [Csp2–Csp3(OH)] motif using the active hydrogen derived from the formaldehyde molecules generated during the SSH reaction of the Csp2–O bonds (Supplementary Fig. 2), consequently leading to the n-propylbenzene byproduct (Supplementary Table 5). Similarly, the 1-methoxy-4-propylbenzene (1g) was also resulted from the above hydrogenolysis of the hydroxyl group in [Csp2–Csp3(OH)] motif of 1a, but without SSH reaction of the Csp2–O bond proceeding yet. Aside from the catalytic properties, RuW/HY30 catalyst also exhibited excellent stability in the reaction, which was confirmed by reusing the catalyst (Supplementary Fig. 6a) and characterizations of the catalyst before and after the reaction by XRD (Supplementary Fig. 6b) and TEM techniques (Supplementary Fig. 6c).

**Mechanistic study.** On the basis of the above comparison tests (Table 1), and given that traditional hydroprocessing with exogenous hydrogen source resulted not only in the unavoidable hydrogenation of the benzene ring to cyclohexane but also in the n-propylbenzene byproduct that could not be further converted into benzene under the mild condition (Table 1, entries 5 and 6), we proposed that the outstanding efficiency of the refining strategy for benzene should be attributed to the innovative matching of RuW NPs and HY30 zeolite in the catalyst, where RuW NPs catalyzed hydrogenolysis of the Csp2–O bonds with the in situ abstracted hydrogen from the reactant can not only completely avoid the saturation of the benzene ring, but more importantly, allow the Csp2–Csp3 bonds to deconstruct promptly over the HY30 zeolite without the competition from the hydrogenolysis of the hydroxyl group in [Csp2–Csp3(OH)] motif. The pathway for the SSH reaction of Csp2–O bond on the RuW centers has been established and illustrated in Supplementary Fig. 24. We here focus on the study of the unique catalysis of Csp2–Csp3 into Csp2–H bond by means of the HY30 zeolite. Based on our experimental characterizations, we constructed the Bronsted acid model to represent the Csp2–Csp3 Reaction center in the RuW/HY30 catalyst, and performed density functional theory (DFT) calculations to study the reaction path of the Csp2–Csp3 bond. As shown in Fig. 3a, initially, the hydroxyl group substituted at the aliphatic C position is easily protonated on the Bronsted acid site, leading to the oxonium ion (structure I) that then transforms into a carbonium ion (structure II) by eliminating a molecule of H2O. Thermodynamically, the γ-methyl on the side chain is then shifted to the Cα position under the catalysis of the Bronsted acid center, evolving to the chemically adsorbed
structure III with a migration barrier of 1.25 eV (TS-1), and subsequent proton abstraction by the Bronsted acid center delivers structure IV in a lower barrier of 0.89 eV (TS-2). Then, by a second protonation at the Cα position of structure V, the resulting carbonium ion V (structure V) would be formed by the trap of H₂O, which gives tertiary alcohol geometry (structure VI) and concomitantly regenerates the Bronsted acid center with the proton. When the benzene ring of the tertiary alcohol (structure VI) is turned endothermically toward the Bronsted acid center, an appropriate geometry (structure VII) is formed to enable the protonation of the Csp² position (structure VIII) with an effective barrier of 0.53 eV (TS-3). After that, the Csp²–Csp³ bond can be exothermically transformed into the desired Csp²–H bond via the ultimate β scission step³⁶.

Taken together, the mild-condition refining of Csp²–Csp³ bond into Csp²–H bond was proceeded via a stepwise protonated dehydroxylation, γ-methyl shift and Csp²–Csp³ β scission pathway. Although the dehydroxylation could also occur in the water-only system (Table 1, entry 1), 1-methoxy-4-(prop-1-en-1-yl) benzene (1h) was the sole product without other following steps, which provided compelling proof for the requisite catalysis worked in the desired refining of the Csp²–Csp³ bond. In terms of catalytic chemistry, highly active acid nature and chemical accessibility of the framework in the mesopore-rich HY₃₀ innately enabled the aliphatic γ-methyl C (Cγ) atom to be activated on the Bronsted acid center, which could be illustrated by the space-filling models (Fig. 3b), affording the migration of the γ-methyl⁴⁹,⁵⁰. Such a chemical bonding state should be

Fig. 2 Characterization of RuW/HY₃₀ catalyst. a XRD patterns of RuW/HY₃₀ (2.5 wt% Ru, 14 wt% W) catalyst (red), W/HY₃₀ (14 wt% W) catalyst (pink), Ru/HY₃₀ (2.5 wt% Ru) catalyst (blue) and HY₃₀ (black), standard patterns of RuW alloy and Y zeolite from ICDD PDF cards (65-6705 for RuW, 45-0112 for Y zeolite) are shown at the bottom. b Rietveld refinement of RuW/HY₃₀ catalyst (2.5 wt% Ru, 14 wt% W) using XRD pattern in a (inset: crystal structures of RuW alloy NPs and Y zeolite). Black circle marks (○) represent the observed intensities, and the red solid line is Rietveld-fit profile. The difference plot (blue) is shown at the bottom. The dark yellow (Y zeolite) and olive (RuW alloy) tick marks respectively indicate the positions of the Bragg reflections as obtained in the Rietveld refinement. The RuW alloy parameters are as follows: space group Im-3m, a = b = c = 3.1610 Å, Rp = 7.37% and Rw = 9.31%. EXAFS Fourier transformed (FT) k²-weighted χ(k) function spectra of the RuW/HY₃₀ catalyst, references, and corresponding EXAFS R-space fitting curves for Ru (c) and W (d) species respectively in the RuW/HY₃₀ catalyst. XPS spectra for Ru 3p (e) and W 4f regions (f).
accompanied by a change in the chemical environment of the Bronsted acid center and reactant, which were monitored by solid-state magic angle spinning nuclear magnetic resonance analyses of the quenched catalytic system, as well as the corresponding model compounds (Figs. 3c–e, Supplementary Figs. 7–10, Supplementary Tables 6 and 7). The $^{13}$C NMR showed a clear resonance signal at 55.1 ppm, characteristic of methyl C atom in the methoxy group, and the dehydroxylated $\text{C}_\alpha$ atom shift at 73.8 ppm instead of the initial $\text{C}_\alpha$ shift at 60.6 ppm (Fig. 3c, Supplementary Fig. 7 and Table 6), which are indicative of the catalysis-priority of the Bronsted acid center during the reaction of 1a, evidencing the trend of the product distribution on reaction time (Supplementary Fig. 3b). Although the $^{13}$C NMR signal for the $\text{C}_\gamma$ atom could still be detected, a new resonance signal at 30.5 ppm was also observed, together with the 25.3 and 21.8 ppm signals for the original $\beta_1$-C ($\text{C}_\beta$1, in Structure II) and $\beta_2$-C ($\text{C}_\beta$2, in Structure III) atoms, which could be ascribed to the $\beta_3$-C ($\text{C}_\beta$3, in Structure III) atom shifted from the $\text{C}_\gamma$ atom in reaction [41]. The above resolved signals were further supported by the solid-state two-dimensional (2D) $^{13}$C–$^1$H dipolar-mediated heteronuclear correlation (HETCOR) NMR analysis (Supplementary Fig. 8 and Supplementary Table 7). The original $\text{C}_\beta$1 and $\text{C}_\beta$2 atoms at 25.3 and 21.8 ppm are strongly correlated with the $^1$H signals at 1.4 and 1.8 ppm that should be associated with ($\text{C}_\beta$1H2–) and ($\text{C}_\beta$2H2) moieties in the intermediates II and III, respectively. The observed $^{13}$C and $^1$H signals at 13.5 and 1.0 ppm are assigned to the remained ($\text{C}_\beta$2H3) moiety in the intermediate II. In line with the $^{13}$C NMR analysis (Fig. 3c), 2D $^{13}$C–$^1$H HETCOR NMR also yields a well-resolved correlated signal at 31.0 ppm in the $^{13}$C dimension and at 3.0 ppm in the $^1$H dimension, which again reflects the transferred $\text{C}_\gamma$, namely, new $\text{C}_\beta$3 atom from the ($\text{C}_\beta$3H2–$\text{SiAlO}$) moiety in the intermediate III. It is the chemical bonding with the oxygen atom in the Bronsted acid [$\equiv\text{Al}–\text{O}–\text{Si}≡$] center that provided an inductive effect on the $\text{C}_\beta$3 atom, which shifted its resonance downfield from that of the original $\text{C}_\beta$1 and $\text{C}_\beta$2 atoms, precisely supporting the Bronsted acid-catalyzed mechanism of the y-methyl shift [52,53]. Echoing the chemical environment change of the reactant, such a chemical bonding in the [$\equiv\text{Al}–\text{O}(\text{C}_\beta$3)–$\text{Si}≡$] unit also fed back to the tetrahedrally coordinated Al [$\text{Al}(\text{VI})$] in the Bronsted acid center, which was further confirmed by $^{27}$Al multiple-quantum MAS (MQMAS) (Fig. 3d, e, Supplementary Fig. 9) and $^{27}$Al NMR (Supplementary Fig. 10) analyses. In the blank experiment (without reactant, Fig. 3d), the regular framework Al(VI) signal in the 2D plot of the boiled catalyst is clearly evident at 59.7 ppm (F2, Fig. 3d and Supplementary Fig. 9a) [54]. Besides the above regular Al(VI) signal, as expected, a resonance signal at 51.5 ppm (F2) was distinctly identified in the operating RuW/HY30 catalyst (Fig. 3e and Supplementary Fig. 9b), which had also been detected as a shoulder signal in the $^{27}$Al NMR results (Supplementary Fig. 10). This emerging Al environment could be assigned to the Al(VI) atom in the distorted [$\equiv\text{Al}–\text{O}(\text{C}_\beta$3)–$\text{Si}≡$] unit, where the charge compensation of the $\text{C}_\beta$3 atom weakened the inductive effect of the O atom on the neighboring Al(VI) atom, and then reduced the Al–O–Si bond angle and lengthened the Al–O and Si–O bonds (Fig. 3e and Supplementary Fig. 10b), shifting the resonance upfield from that of the framework Al(VI) atoms in the regular [$\equiv\text{Al}–\text{O}–\text{Si}≡$] environments [55]. The above identifications evidently described the key processes that occurred on the $\text{C}_\beta$2–$\text{C}_\beta$3 refining centers of the RuW/HY30 catalyst, highly agreeing with the DFT mechanism studies (Fig. 3a).
Scope of phenylpropanol derivatives. We proceeded to study the scope of the in situ refining strategy. As shown in Fig. 4, several structurally and substitutionally distinct Csp2–Csp3/Csp2–O bonds equipped phenylpropanol derivatives were refined effectively and singly into desired benzene product, including those G (guaiacyl)-monomeric lignin model compounds (Fig. 4a) bearing methoxyl (2a), hydroxyl (3a) and isopropoxyl (4a) groups, as well as the S (syringyl)-monomers (Fig. 4b, 5a, 6a and 7a). The in situ refining strategy is not limited to the monomers, a range of G and S monomer derived dimeric model compounds including those bearing the methoxyl groups (Fig. 4c, 8a and 9a) and the combinatorial groups of methoxyl and isopropoxyl (Fig. 4d, 10a–13a), were all viable in the reactions with benzene as the single product. The catalytic performance observed here represented an important proof of the multifunctional centers on the RuW/HY30 catalyst, where the interlaced Csp2–Csp3 and Csp2–O bonds in the complex compounds could be efficiently tolerated and transformed into the Csp2–H bonds, respectively, demonstrating the feasibility of the in situ refining strategy (Fig. 1c). As a further demonstration of the in situ refining protocol, we further performed the detailed analysis of the conversion of 1-(4-isopropoxy-3, 5-dimethoxymethyl)-2-(2-methoxynaphthalene)propane-1, 3-diol (11a) versus reaction time, which aimed to comprehensively understand the evolution of the Csp2–Csp3/Csp2–O bonds in the lignin-mimetic dimeric linkages (Supplementary Fig. 11). When 11a was subjected to our reaction conditions, the transformation proceeded readily to afford 2, 6-dimethoxyphenol and 2-methoxyphenol in sequentially and rapidly ascended yields, while the phenylpropanol analogues were not detected during the reaction, which evidenced that the deconstruction of the Csp2–Csp3 bond was occurred locally on the molecular structure of the reactant and accompanied by the HY30 catalyzed hydrolysis of the aliphatic carbon–oxygen (Csp3–O, Fig. 1c) bonds.56 At this stage, water was the only reactant besides 11a in the formation of the phenolic hydroxyl (–OH) groups, and no active hydrogen was needed. With the ongoing refining process, the methoxyl derived Csp2–O(CH3) bonds in the above monomeric intermediates were gradually transformed into the Csp2–H bonds. Meanwhile, the phenolic Csp2–O bonds with higher bond energy could also be cleaved along with the Csp2–O(CH3) bond, which should be attributed to the RuW catalyzed hydrogenolysis of the phenolic hydroxyl group using the active hydrogen derived from the formaldehyde molecules generated during the SSH reaction of the Csp2–O(CH3) bond (Supplementary Fig. 2). To explore the mechanism of dehydroxylation, we conducted the reaction of 4-(1-hydroxypropylyphenol over the RuW/HY30 catalyst in the formalddehyde solution (Supplementary Table 8), where thermostable paraformaldehyde was used as the source of formaldehyde to simulate the gradual supply of the formaldehyde generated from the SSH reaction. As expected, only the Csp2–Csp3 bond in 4-(1-hydroxypropylyphenol was deconstructed over the HY30 component with phenol as the sole product, and the cleavage of the phenolic Csp2–O bond did not occur without the source of formaldehyde (Supplementary Table 8, entry 1). With the introduction of formaldehyde, the benzene product was detected (Supplementary Table 8, entry 2), suggesting that the phenolic Csp2–O bond could be hydrogenolyzed over the RuW component using the active hydrogen derived from formaldehyde. Moreover, the yield of benzene was steadily increased with the increase of the source of formaldehyde (Supplementary Table 8, entries 2–8), which confirmed that the phenolic Csp2–O bonds can be efficiently hydrogenolyzed over the RuW component with the active hydrogen derived from the gradually increased formaldehyde molecules during the RuW catalyzed SSH reaction of the Csp2–O (CH3) bonds.
transformation from the interference of the reaction of the carbohydrate in wood powder. In the first step, lignin was extracted from wood powder by solid–liquid separation and solvent recuperation, during which the used wood powder was also recovered along with the organic solvent and then used in the continual extraction process. In the second step, the extracted lignin was fed to the catalytic reactor only with water, and exclusively converted into benzene product, where the catalyst could be recovered and reused. Meanwhile, as the only liquid product, benzene could be quite easily separated from the system without complex procedures. The sufficiently recyclable and highly selective features of the above processes are beneficial to producing benzene economically. From the perspective of atomic economy, the active hydrogen atoms in the lignin molecule could also be utilized successfully along with the abstraction of the benzene rings from lignin under the in situ refining strategy. Moreover, the lignin residue obtained in the lignin conversion process can be collected and further valorized into high value-added fuel products and chemicals. Given the above advantageous features, this lignin-to-benzene route has the potential of industrial application.

**Discussion**

In summary, we have developed an in situ refining strategy to the transformation of the $C_{sp2}$–$C_{sp3}$ and $C_{sp2}$–$O$ bonds for the sustainable production of benzene from lignin using RuW/HY$_{30}$ as the multifunctional catalyst and water as the reaction medium, which further draws attention to biomass valorization methods. It demonstrates that the HY$_{30}$ zeolite and RuW components in the

---

**Fig. 4 In situ refining strategy: portfolio scope of the $C_{sp2}$–$C_{sp3}$ and $C_{sp2}$–$O$ bonds.** The general reaction scheme is shown at the top. The chemical bonds colored with red is transformed, and the benzene ring colored with green is the desired product from the in situ refining strategy. Yields of benzene product provided are at full conversion of substrates, as averages of three experiments conducted in parallel. $^a$CH(CH$_3$)$_2$, isopropyl. $^b$C$_7$H$_7$O$_2$, 2-methoxyphenoxyl. $^c$C$_8$H$_9$O$_3$, 2, 6-dimethoxyphenoxyl. Reaction conditions: H$_2$O (5.0 mL), 0.1 MPa N$_2$, 800 rpm. **Conditions a:** Substrate (1.0 mmol), RuW/HY$_{30}$ catalyst (0.25 g, 3.0 wt% Ru, 17 wt% W), 190 °C, 7.0 h. **Conditions b:** Substrate (1.0 mmol), RuW/HY$_{30}$ catalyst (0.30 g, 3.0 wt% Ru, 17 wt% W), 200 °C, 7.5 h. **Conditions c:** Substrate (0.5 mmol), RuW/HY$_{30}$ catalyst (0.35 g, 3.0 wt% Ru, 17 wt% W), 210 °C, 8.0 h. **Conditions d:** Substrate (0.5 mmol), RuW/HY$_{30}$ catalyst (0.35 g, 3.5 wt% Ru, 20 wt% W), 210 °C, 8.0 h.
n = phenylpropanol structure. The formula is displayed in Supplementary information. Priorly deconstruct the Csp3–Csp2–Csp2 and Csp2–O bonds in lignin structure, by which benzene can be exclusively produced from lignin with a maximum benzene yield of 18.8% on the lignin weight basis, highlighting the significance of developing catalytic technologies for the aromatic building blocks in lignin. Innovatively, the RuW component cannot only catalyze the hydrogenolysis of the Csp2–O bond using the active hydrogen in situ abstracted from lignin molecule, but more importantly, allow Bronsted acid sites of HY30 zeolite to promptly deconstruct the Csp3–Csp2 bonds on the local structure of lignin molecule without any precedent reductive catalytic fractionation process and competition from the hydrogenolysis of the hydroxyl group in Csp2–Csp3(OH) motif. In the scale-up experiment, 8.5 g of benzene can be produced from 50.0 g of lignin without any saturation byproducts. This in situ lignin refining strategy liberates the trapped benzene rings from the molecular structure of lignin, and paves a new way for sustainable production of benzene using lignin as the feedstock, which has great potential of practical application.

Methods

Catalyst preparation. Supported Ru/W/Zeolite catalysts were prepared by commonly used wet impregnation method. Prior to the impregnation, the used zeolites were pretreated by calcination at 550 °C for 4 h. In a typical preparation, ammonium metatungstate (AMT) (1.25 g) and RuCl3·xH2O (0.40 g) were dissolved in 15 mL of deionized water, respectively. Then, the precursor solutions containing AMT and Ru3+ ion were successively added dropwise to 70 mL deionized water with 2.0 g of zeolite at room temperature. The obtained mixture was vigorously stirred for 48 h, evaporated, and dried at 473 K for 16 h in an oven. The as-prepared sample was reduced in a continuous 10% H2/Ar flow, from room temperature to 400 °C at 5 °C/min and then to 900 °C at 1 °C/min, and maintained at 900 °C for 1 h. After being cooled to room temperature under Ar atmosphere, the reduced catalyst was exposed to 1% O2/Ar atmosphere for 1 h to form a passivation layer to prohibit against bulk oxidation before exposure to air. The passivated catalyst was kept under an inert atmosphere before testing and characterization, and denoted as Ru/W/Zeolite (x, the framework Si/Al ratio of the zeolite). For comparison, supported Ru/Zelite, W/Zelite, and other catalysts were prepared by the same procedures for preparing the Ru/W/Zeolite catalyst.

Catalyst characterization. N2 adsorption-desorption of the samples were measured using a Micromeritics Tristar II 3020 at liquid nitrogen temperature. The specific surface areas were calculated by using the Brunauer–Emmett–Teller model. The pore size distribution of the sample was calculated using the Barrett–Joyner–Halenda pore size model. XRD measurements were conducted on an X-ray diffractometer (D/MAX-RC, Japan) operated at 40 kV and 200 mA with Cu Kα (λ = 0.154 nm) radiation. Rietveld refinements were performed applying the TOPAS program60 for the measured XRD profile of all the catalysts. The XPS measurements were carried out on an ESCALAB 220i-XL spectrometer at a pressure of ~3 × 10−9 mbar (1 mbar = 100 Pa) using Al Ka as the excitation source (hv = 1486.6 eV) and operated at 15 kV and 20 mA. The X-ray absorption fine structure (XAFS) spectra (Ru K-edge and W L3 edge) were collected at 1W1B station in Beijing Synchrotron Radiation Facility (BSRF). The storage ring of BSRF was operated at 2.5 GeV with a maximum current of 250 mA. Using Si (111) double-crystal monochromator, the data collection were carried out in transmission mode using ionization chamber. All spectra were collected in ambient conditions. The acquired EXAFS data were processed according to the standard procedures using the ATHENA method. The k2-weighted EXAFS software packages. The k2-weighted EXAFS spectra were obtained by subtracting the post-edge background from the overall absorption and then normalizing with respect to the edge jump step. Subsequently, k2-weighted k(ξ) spectra of Ru K-edge and W L3 edge were Fourier transformed to real (R) space using a hanning windows (dk = 1.0 Å−1) to separate the EXAFS contribution from different coordination shells. To obtain the quantitative structural parameters around central atoms, least-squares curve parameter fitting was performed using the ARTEMIS module of IFEFFIT software packages. TEM and HRTEM images were obtained on a JEOL-2011F electron microscope operating at 200 kV. The contents of supported metals on the catalysts were determined by ICP.

Lignin extraction and characterization. In a typical experiment, 70 g of wood powder, 490 mL of acetic acid and 210 mL of water were loaded into a 1-L autoclave (Weihai Xinyuan Chemical Machinery Co. Ltd.). The autoclave was sealed and purged with N2 to remove the air at room temperature and subsequently charged with 0.1 MPa of N2. Then, the autoclave was heated to 160 °C within 1 h. After that, the stirrer was started with a stirring speed of 600 rpm, and the reaction time was recorded. After 1 h, the autoclave was cooled down slowly, and the gas was released. The liquid was collected by a filtration process. The filter residue was washed three times with 200 mL of acetone/H2O (9:1), and the washing liquor was collected and combined with the filtrate liquid. After that, the liquid was concentrated under vacuum at 30 °C until the liquid became muddy and then dissolved again by 30 mL of acetone. The obtained lignin was dissolved in 550 mL of dimethyl sulfoxide (DMSO)—d6 and characterized by 1H/13C 2D HSQC NMR and quantitative 1H NMR analyses using Bruker Avance III 500WB and Bruker Avance 600, as described by other researchers.67

Catalytic performance. The reaction was carried out in a Teflon-lined stainless-steel reactor of 20 mL with a magnetic stirrer. In a typical experiment, a suitable amount of reactant, catalyst, and water were loaded into the reactor. The reactor was sealed and purged with N2 for three times to remove the air at room temperature and subsequently charged with desired gas. Then the reactor was placed in a furnace at desired reaction temperature. When the reactor reached the desired reaction temperature, the stirrer was started with a stirring speed of 800 rpm, and the reaction time was recorded. After the reaction, the reactor was cooled down in ice water, and the gas was released, passing through the ethyl acetate. The reaction mixture in the reactor was transferred into a centrifuge tube. Then the reactor was washed with the ethyl acetate used for the gas filtration, which was finally combined with the reaction mixture. After centrifugation, the catalyst was separated from the reaction mixture. The quantitative analysis of the liquid products in the organic phase was conducted using a GC (Agilent 6820) equipped with a flame ionization detector and HP-5MS/HP-INNOWAX capillary columns (0.25 mm in diameter, 30 m in length). Identification of the products and reactant was performed using a GC-MS (Agilent 5977A, HP-5MS capillary column [0.25 mm in diameter, 30 m in length]) and by comparing the retention time to respective standards in GC traces. Biphenyl was used as the internal standard to determine the conversions of substrates, selectivities and yields of the products. Identification of the products in the aqueous phase was conducted by 1H NMR analysis on a Bruker Avance III 400 HD with D2O as the solvent. The carbon balance for the reaction was determined by elemental analysis (Elementar vario EL III), which was given relative to the aromatic compounds.63 The Cnatable mass balances for the reaction of the model compounds were better than 99%. The recovered lignin residual solid article NATURE COMMUNICATIONS | https://doi.org/10.1038/s41467-021-24780-8 | www.nature.com/naturecommunications

NATURE COMMUNICATIONS | https://doi.org/10.1038/s41467-021-24780-8 | www.nature.com/naturecommunications
were analyzed by H/^{13}C 2D HSQC NMR and quantitative^{13}C NMR spectroscopy (Bruker Avance III 600 HD).

**Recycling of the catalyst.** The reusability of Ru/W/HY_{50} catalyst was tested using the reaction of the model compound 1-(4-methoxyphenyl)-1-propanol (1a). After the reaction, the reaction mixture in the reactor was transferred into a centrifuge tube. Then the reactor was washed with ethyl acetate, which was combined with the reaction mixture. Subsequently, the reaction mixture was centrifuged and the ethyl acetate layer was analyzed by GC. After that, the used Ru/W/HY_{50} catalyst was separated from the reaction mixture and successively washed with ethanol (5 × 10 mL) and water (5 × 10 mL). Then, the recovered catalyst was reused directly for the next run.

**Detection of intermediates.** The detection of intermediates in the dried sample was conducted by 1H and 13C NMR analyses (Bruker Avance III 400) and 2D MQMAS and 27Al NMR analyses on Bruker Avance III 400.

**Mass balance analysis of the lignin transformation.** After the reaction of lignin, the gas was released, passing through the ethyl acetate. Then, the reaction mixture in the reactor was transferred into a separatory funnel, and then the aqueous layer was removed. After the reaction of lignin, the gas was released, and the liquid layers were pooled and used for the GC analysis. The detection of the intermediates in the dried sample was conducted by 1H and 13C NMR analyses (Bruker Avance III 400) and 2D MQMAS and 27Al NMR analyses on Bruker Avance III 400.

**Scale-up transformation of the pine lignin.** The scale-up reaction was performed in a 1-L autoclave (Weihai Xinyuan Chemical Machinery Co. Ltd.). In the reaction, 50.0 g of lignin, 50.0 g of Ru/W/HY_{50} catalyst and 650 mL of H_{2}O were loaded into the autoclave. The autoclave was sealed and purged with N_{2} three times to remove the air at room temperature and subsequently charged with 0.1 MPa of N_{2}. Then, the stirrer was started with a stirring speed of 800 rpm, and the reaction time was recorded. After 10 min, the reactor was transferred to a bath of liquid nitrogen very quickly. When the reaction mixture was frozen, the gas was released immediately. The reaction mixture was freeze-dried under vacuum for 12 h to remove the water. The detection of the intermediates in the dried sample was conducted by 1H and 13C NMR analyses (Bruker Avance III 400).

**Data availability**

All data needed to evaluate the conclusions in this paper are present in the paper and/or the Supplementary Information. Additional data available from authors upon request.

Received: 24 March 2021; Accepted: 29 June 2021; Published online: 26 July 2021

---

**References**

1. Zhu, X. F. (Xin X.), Johnson, J. A., Ablin, D. W. & Ernst, G. A. Efficient petrochemical processes: technology, design and operation (Wiley, 2019).
2. Vaughan, B. A., Webster-Gardiner, M. S., Cundari, T. R. & Gunnne, T. B. A rhodium catalyst for single-step styrene production from benzene and ethylene. *Science* **348**, 421–424 (2015).
3. Davis, R. J. & Deroanne, E. G. A non-porous supported-platinum catalyst for aromatization of 2-naphthalene. *Nature* **349**, 313–315 (1991).
4. Squires, A. M. Chemicals from coal. *Science* **191**, 689–700 (1976).
5. Mochida, I., Okuma, O. & Yoon, S.-H. Chemicals from direct coal liquefaction. *Chem. Rev.* **114**, 1637–1672 (2014).
6. Guo, X. G. et al. Direct, nonoxidative conversion of methane to ethylene, aromatics, and hydrogen. *Science* **344**, 616–619 (2014).
7. Gao, J. et al. Identification of low-dimensional nanostructures on zeolites for natural gas conversion. *Science* **348**, 686–690 (2015).
8. Morejudo, S. H. et al. Direct conversion of methane to aromatics in a catalytic co-ionic membrane reactor. *Science* **353**, 563–566 (2016).
9. Kintisch, E. Making dirty coal plants cleaner. *Science* **317**, 184–186 (2007).
10. Chow, J., Kopp, R. J. & Portney, P. R. Energy resources and global development. *Science* **302**, 1528–1531 (2003).
11. Huber, G. W., Iborra, S. & Corma, A. Synthesis of transportation fuels from biomass: chemistry, catalysts, and engineering. *Chem. Rev.* **106**, 4044–4098 (2006).
12. Sutton, A. D. et al. The hydrodeoxygenation of bioderived furans into alkanes. *Nat. Chem.* **5**, 428–432 (2013).
13. Deng, W. P. et al. Catalytic amino acid production from biomass-derived intermediates. *PNAS* **115**, 5093–5098 (2018).
14. Sun, Z., Frdric, B., de Santi, A., Elangovan, S. & Barta, K. Bright side of lignin depolymerization: toward new platform chemicals. *Chem. Rev.* **118**, 614–678 (2018).
15. Sudarsanam, P., Peeters, E., Makshina, E. V., Parvulescu, V. I. & Sels, B. F. Advances in porous and nanoscale catalysts for viable biomass conversion. *Chem. Soc. Rev.* **48**, 2366–2421 (2019).
16. Li, C. Z., Zhao, X. C., Wang, A. Q., Huber, G. W. & Zhang, T. Catalytic transformation of lignin for the production of chemicals and fuels. *Chem. Rev.* **115**, 11559–11624 (2015).
17. Wu, X. J. et al. Solar energy-driven lignin-first approach to full utilization of ligninolulosic biomass under mild conditions. *Nat. Catal.* **1**, 772–780 (2018).
18. Zeng, H. Q., Cao, D. W., Qiu, Z. H. & Li, C. J. Palladium-catalyzed formal cross-coupling of diaryl ethers with amines: slicing the 4-O-5 linkage in lignin models. *Angew. Chem. Int. Ed.* **57**, 3752–3757 (2018).
19. Rakagaskas, A. J. et al. Lignin valorization: improving lignin processing in the bioeconomy. *Science* **344**, 709–719 (2014).
20. Shuai, L. et al. Formaldehyde stabilization facilitates lignin monomer production during biomass depolymerization. *Science* **354**, 329–333 (2016).
21. Lan, W., Amiriz, M. T., Hunston, C. M. & Luterbacher, J. S. Protection group effects during a γ, δ-diol lignin stabilization promote high-selectivity monomer production. *Angew. Chem. Int. Ed.* **57**, 1356–1360 (2018).
22. Lan, W., de Baeren, J. B. & Luterbacher, J. S. Highly selective oxidation and depolymerization of α, γ-diol-protected lignin. *Angew. Chem. Int. Ed.* **58**, 2649–2654 (2019).
23. Ragauskas, A. J. et al. Lignin valorization: improving lignin processing in the bioeconomy. *Science* **344**, 709–719 (2014).
24. Wang, M. et al. Dealkylation of lignin to phenol via oxidation-hydrogenation strategy. *ACS Catal.* **8**, 6837–6843 (2018).
25. Decus, P. J. et al. aromatic monomers by in situ conversion of reactive intermediates in the acid-catalyzed depolymerization of lignin. *J. Am. Chem. Soc.* **137**, 7456–7467 (2015).
26. Li, H. L. & Song, G. Y. Ru-catalyzed hydrogenolysis of lignin: base-dependent tunability of monomeric phenols and mechanistic study. *ACS Catal.* **9**, 4054–4064 (2019).
27. Zhou, G. F., Jensen, P. A., Le, D. M., Knudsen, N. O. & Jensen, A. D. Direct upgrading of fast pyrolysis lignin vapor over the HZSM-5 catalyst. *Green. Chem.* **18**, 1965–1975 (2016).
28. Zhou, S., Xue, X., Sharma, A. & Bai, X. L. Lignin valorization through thermochemical conversion: comparison of hardwood, softwood and herbaceous lignin. *ACS Sustain. Chem. Eng.* **4**, 6608–6617 (2016).
29. Mullen, C. A. & Boetang, A. A. Production of aromatic hydrocarbons via catalytic pyrolysis of biomass over Fe-modified HZSM-5 zeolites. *ACS Sustain. Chem. Eng.* **3**, 1623–1631 (2015).
30. Saraeian, A. et al. Evaluating lignin valorization via pyrolysis and vapor-phase hydrodeoxygenation for production of aromatics and alkenes. *Green. Chem.* **22**, 2513–2525 (2020).
31. Dong, L. et al. Breaking the limit of lignin monomer production via cleavage of interunit carbon-carbon linkages. *Chem. 5*, 1521–1536 (2019).
32. Huang, Y. B., Yan, L., Chen, M. Y., Guo, X. & Fu, Y. Selective hydrogenolysis of phenols and phenyl ethers to alkenes through direct C-O cleavage over ruthenium–tungsten bifunctional catalysts. *Green. Chem.* **17**, 3010–3017 (2015).
35. Vispute, T. P., Zhang, H. Y., Sanna, A., Xiao, R. & Huber, G. W. Renewable chemical commodity feedstocks from integrated catalytic processing of pyrolysis oils. Science 330, 1222–1227 (2010).
36. Toppi, S. et al. On the radical cracking of n-propylbenzene to ethylbenzene or toluene over SnAl$_2$O$_4$-Cl catalysts under reforming conditions. J. Catal. 230, 255–268 (2005).
37. Verboekend, D., Liao, Y., Schutyser, W. & Sels, B. F. Alkylphenols to phenol and aromatics by zeolite catalysed cracking of diesel raw and fossilized lignocellulose. Green. Chem. 18, 297–306 (2016).
38. Smaligo, A. J. et al. Hydrodealkenylative C(sp$^3$)-C(sp$^2$) bond fragmentation. Science 364, 681–685 (2019).
39. Murphy, S. K., Park, J. W., Cruz, F. A. & Dong, V. M. Rh-catalyzed C-C bond cleavage by transfer hydroformylation. Science 347, 56–60 (2015).
40. Chopp, K. et al. Reactive cleavage of the C$_2$=C$_2$-C(sp$^3$)-C(sp$^3$) bond of secondary benzyl alcohols: rhodium catalyst directed by N-containing groups. Angew. Chem. Int. Ed. 51, 9851–9855 (2012).
41. Ferrini, P. & Rinaldi, R. Catalytic biofining of plant biomass to non-pyrolytic lignin bio-oil and carbohydrates through hydrogen transfer reactions. Angew. Chem. Int. Ed. 53, 8634–8639 (2014).
42. McClelland, D. J. et al. Supercritical methanol depolymerization and hydrodeoxygenation of lignin and biomass over reduced copper porous metal oxides. Green. Chem. 21, 2988–3005 (2019).
43. Meng, Q. L. et al. Self-supported hydrolysis of aromatic ethers to arenes. Sci. Adv. 5, eaax8339 (2019).
44. Prasomsri, T., Shetty, M., Murugappan, K. & Roman-Leshkov, Y. Insights into the catalytic activity and surface functionality of MoO$_3$ during the hydrodeoxygenation of lignin-derived model compounds into aromatic hydrocarbons under low hydrogen pressures. Energy Environ. Sci. 7, 2660–2669 (2014).
45. Akiya, N. & Savage, P. E. Kinetics and mechanism of cyclohexanol dehydration in high-surface-area zeolites. Ind. Eng. Chem. Res. 40, 1832–1833 (2001).
46. Ribblett, J. W., Sinclair, W. E., Borst, D. R., Yi, J. T. & Pratt, D. W. High resolution electronic spectra of anisole and anisole-water in the gas phase: Hydrogen bond switching in the S$_1$ state. J. Phys. Chem. A 110, 1478–1483 (2006).
47. Sieve, B. et al. Cubic aluminum silicides RE$_2$Ru$_4$Al$_2$Si$_2$ (Al$_{12-x}$Si$_{x}$) (RE=$\text{Pr}$, Sm) from liquid aluminum. Empty (Si, Al)$_3$ cubocahedral clusters and assignment of the AES distribution with neutron diffraction. J. Am. Chem. Soc. 123, 7040–7047 (2001).
48. Bereciartua, P. J. et al. Control of zeolite framework flexibility and pore topology for separation of ethane and ethylene. Science 358, 1068–1071 (2017).
49. Blomma, E., Martens, J. A. & Jacobs, P. A. Reaction mechanisms of isomerization and cracking of heptane on Pd/H beta zeolite. J. Catal. 155, 141–147 (1995).
50. Zhou, X. et al. Observation of an oxonium ion intermediate in ethanol dehydration to ethene on zeolite. Nat. Commun. 10, 1961 (2019).
51. Gabrieenko, A. A., Arzumanov, S. S., Toktarev, A. V. & Stepanov, A. G. Solid-state NMR characterization of the structure of intermediates formed from olefins on metal oxides (Al$_2$O$_3$ and Ga$_2$O$_3$). J. Phys. Chem. C. 116, 21430–21438 (2012).
52. Zardkooohi, M., Haw, J. F. & Lunsford, J. H. Solid-state NMR evidence for the formation of carboxylates from propene in acidic zeolite-Y. J. Am. Chem. Soc. 109, 5278–5280 (1987).
53. Haoaas, M., Walspurer, S., Taulelle, F. & Sommer, J. The initial stages of solid acid-catalyzed reactions of adsorbed propane: A mechanistic study by in situ MAS NMR. J. Am. Chem. Soc. 126, 599–606 (2004).
54. Zhang, X. Y. et al. Synthesis of self-pillared zeolite nanosheets by repetitive branching. Science 336, 1684–1687 (2012).
55. van Bokhoven, J. A., Koningsberger, D. C., Kunkeler, P., van Bekkum, H. & Kentgens, A. P. M. Stepwise dealumination of zeolite beta at specific T-sites observed with $^{27}$Al MAS and $^{27}$Al MQ MAS NMR. J. Am. Chem. Soc. 122, 12842–12847 (2000).