### Electronic Supplementary Information

# Oxidative conversion of lignin and lignin model compounds catalyzed by CeO<sub>2</sub>-supported Pd nanoparticles

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### 1. XPS spectra for Pd species loaded on several metal oxides



Fig. S1. XPS spectra of Pd 3d for several metal oxide-supported Pd catalysts.

#### 2. H<sub>2</sub> chemisorption results for supported Pd catalysts

Catalyst	Dispersion <sup>a</sup> (%)	Mean diameter of Pd nanoparticles <sup>b</sup> (nm)
Pd/SiO <sub>2</sub>	52	2.1
$Pd/Al_2O_3$	46	2.4
Pd/CeO <sub>2</sub>	52	2.1
Pd/MgO	27	4.2

**Table S1** The dispersion and the estimated mean particle diameters for Pd loaded on several metal oxides

<sup>a</sup>Obtained from  $H_2$  chemisorption measurements.

<sup>b</sup>Estimated by using the following equation: Pd diameter =1.12/Pd dispersion (nm).<sup>1</sup>

### 3. HPLC spectra of soluble products from the conversion of Organosolv lignin over the Pd/CeO<sub>2</sub> catalysts



**Fig. S2** HPLC spectra of soluble products from the conversion of Organosolv lignin over the Pd/CeO<sub>2</sub> catalyst. Reaction conditions: Organosolv lignin, 0.10 g; catalyst, 0.20 g; MeOH, 50 mL; O<sub>2</sub>, 0.1 MPa; 458 K; 24 h.

# 4. Effect of $O_2$ pressure on catalytic performances of Pd/CeO<sub>2</sub> for the conversion of 2-phenoxy-1-phenylethanol

**Table S2** Catalytic conversion of 2-phenoxy-1-phenylethanol under different O<sub>2</sub> pressures<sup>a</sup>

2-phenoxyl	$\frac{1}{5} = \frac{1}{5} = \frac{1}$	Catalyst methanol O <sub>2</sub>	PPone OCI MB	+ (		Phol	
Entry	O <sub>2</sub>	Conv.		Y	′ield (%)		
	pressure (MPa)	(%)		ОН	O OCH3	° ↓	HO
1 <sup><i>b</i></sup>	0	6.7	0	0	0	3.4	4.7
2 <sup>c</sup>	0.02	10	0	0	3.1	5.7	4.8
3	0.1	64	12	0.03	14	38	48

<sup>a</sup> Reaction conditions: PP-ol, 0.125 g (0.6 mmol); catalyst, 0.2 g; MeOH, 50 mL; 458 K; 24 h.  ${}^{b}N_{2}$ , 0.1 MPa.  ${}^{c}N_{2}$  was added to keep the total pressure at 0.1 MPa.

### 5. Catalytic performances of Pd/CeO<sub>2</sub> for the conversion of 2-phenoxy-1-phenylethanone under different atmospheres

	$\beta 0 4 3 2 5 6 1$	Catalyst methanol $N_2$ or $O_2$	benzoic ar BA	OH + cid aceto	O HO + phenone pl	henol Phol
2-phenoxyl-1-phenylethanone						
	PP-one					
				netnyi benzoate	МВ	
Entry	Atmosphere	Conv.		Yield	d (%)	
		(%)	ОН	OCH3	o C	HO
1	$N_2$	27	0	0	20	18
2	O <sub>2</sub>	53	0.1	17	28	42

Table S3 Catalytic conversion of 2-phenoxy-1-phenylethanone under N<sub>2</sub> and O<sub>2</sub><sup>a</sup>

<sup>a</sup>Reaction conditions: PP-ol, 0.125 g (0.6 mmol); catalyst, 0.2 g; MeOH, 50 mL; N<sub>2</sub> or O<sub>2</sub>, 0.1 MPa; 458 K; 2 h.

#### 6. Conversion of methanol over Pd/CeO<sub>2</sub> in the absence of O<sub>2</sub>

**Table S4** The gaseous component in the conversion of methanol over the Pd/CeO<sub>2</sub> catalyst under  $N_2^{a}$ 

Gaseous component <sup>b</sup> (mol%)					
N <sub>2</sub>	H <sub>2</sub>	CO	CO <sub>2</sub>		
93.9	4.0	2.0	0.10		

<sup>a</sup> Reaction conditions: MeOH, 50 mL; Pd/CeO<sub>2</sub>, 0.1 g; N<sub>2</sub>, 0.1 MPa; 458 K; 2 h.

 ${}^{b}$ H<sub>2</sub> was analyzed by an Agilent Micro 3000-GC equipped with a Molecular Sieve 5A column and a high-sensitivity thermal conductivity detector. CO and CO<sub>2</sub> were separated by a carbon molecular sieve (TDX-01), and were further converted to CH<sub>4</sub> by a methanation reactor, and were then analyzed by a flame ionization detector (FID).

# 7. CeO $_2$ -supported Pd, Au and Pt catalysts for the conversion of 2-phenoxy-1-phenylethanol

**Table S5** Catalytic performances of CeO<sub>2</sub>-supported Pd, Au and Pt catalysts for the conversion of 2-phenoxy-1-phenylethanol<sup>a</sup>

OH 2-phenoxyl-1 PF	- - - phenyletha <b>2</b> - 1 - phenyletha	Catalyst methanol O <sub>2</sub>	→ Products			
Catalyst	Conv.		Yield (%)			
	(%)				HO	
CeO <sub>2</sub>	9.1	0	3.2	0	0	
Pd/CeO <sub>2</sub>	64	12	14	38	48	
Au/CeO <sub>2</sub>	58	11	24	24	46	
Pt/CeO <sub>2</sub>	42	7.0	19	11	28	
<sup>a</sup> Reaction con MPa; 458 K; 24	ditions: PF h.	P-ol, 0.125 g	(0.6 mmol); catalyst,	0.2 g;	MeOH, 50 mL; (	D <sub>2</sub> , 0.1

Reference

1. N. Mahata and V. Vishwanathan, J. Catal., 2000, 196, 262.