Supporting Information

Cascade and One Pot Dehydrative Amination of Glycerol to Oxazoline

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Scheme S1. Reaction mechanism for the glycerol amination.



Table S1. Effect of catalyst on acetol conversion to oxazoline.								
Sr. No.	Con.	Selectivity (%)						
	(%)	Oxazoline	5-methyl	1,2-dimethyl	Dialkyl	Other		
			imidazole	imidazole	Pipirazine			
1 ^a	99	95	3	2	00	00		
2 ^b	88	92	2	1	3	2		

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Reaction conditions: a With catalyst – Distillate (5 g), 30% aq. NH₃ = 15 mL, 50 °C, catalyst (0.01 g), 2 h. b Without catalyst - Distillate (5 g), 30% aq. NH_3 = 15 mL,, 50 °C, 2 h.

Figure S1. SEM images of a) Ru/AC, b) Ru/SiO₂ and c) Ru/Al₂O₃ catalyst.







Figure S3. ¹³C-NMR and ¹H- NMR of Oxazoline.



¹³C NMR (200MHz, CHLOROFORM-d) δ 14.89, 21.40, 66.6, 76.26, 110.40, 169.14 ¹HMR (200MHz, CHOLOROFORM-d) δ 1.39(CH₃,s,3H), 2.08 (CH₃,s,3H), 3.02(OH,s,1H), 3.64(CH₂,d,2H), 4.58(CH₂,s,2H).

Figure S4. IR spectra of oxazoline.









Figure S6. NH₃–TPD and Py-IR profiles of different activated copper catalysts.





Reaction conditions: 0.8 g Catalyst , 220 °C, 15 mL 30% aq. $NH_3,\,5$ h.

Figure S8. Effect of catalyst loading on dehydrative cyclised amination reaction.



Reaction conditions: 20 wt% glycerol aqueous solution, 220 °C, 15 mL 30% aq. NH₃, 5 h.

Figure S9. Reactive distillation set up.

