

Supporting Information

High-efficiency removal of NO_x using a combined adsorption-discharge plasma catalytic process

Qinqin Yu[#], Hui Wang[#], Tong Liu, Liping Xiao, Xiaoyuan Jiang, Xiaoming Zheng^{*}

*Key Lab of Applied Chemistry of Zhejiang Province, Department of Chemistry, Zhejiang
University (XiXi Campus), Hangzhou 310028, China.*

^{}Corresponding author. Tel.: +86 571 88273417; Fax: +86 571 88273283*

E-mail: xmzheng@zju.edu.cn (Xiaoming Zheng)

[#]The authors contributed equally to this work

Number of figures: 6

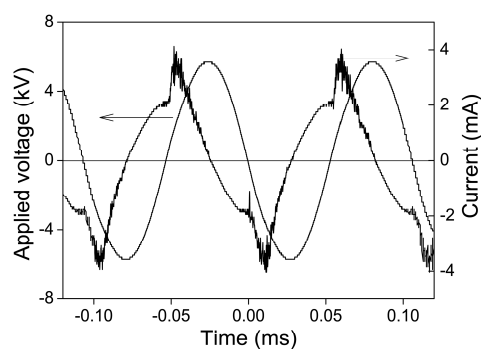


Figure S1. Waveforms of applied voltage and discharge current of the plasma.

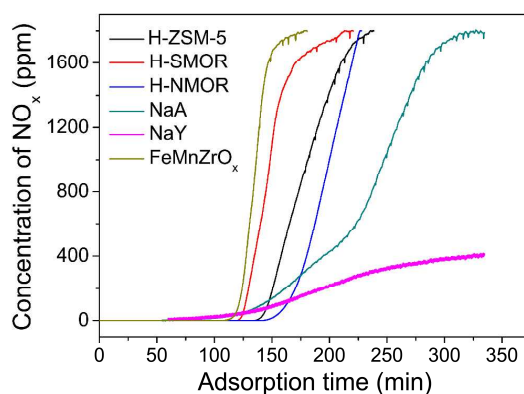


Figure S2. Breakthrough curves of NO_x on the catalysts.

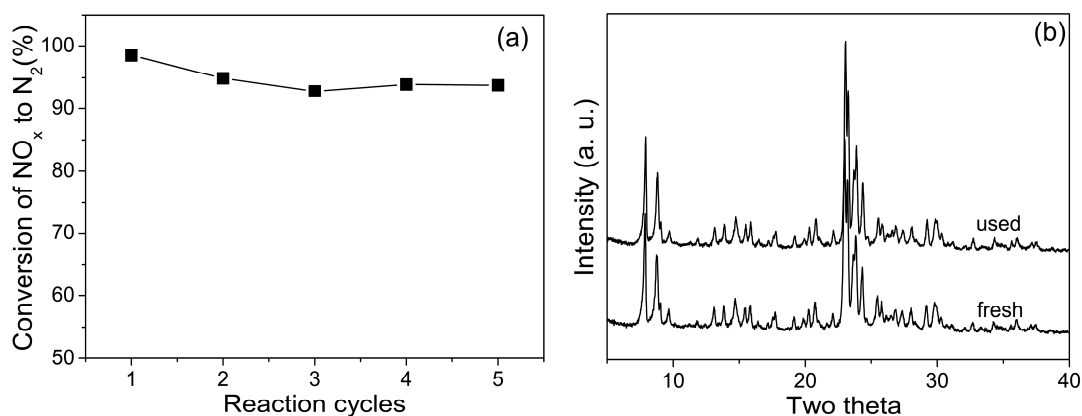


Figure S3. Cyclic performance of H-ZSM-5 in the adsorption-discharge process for NO_x removal (a) conversion of adsorbed NO_x (b) XRD patterns of H-ZSM-5 before and after 5-cycles' operation.

Conditions: in the adsorption stage, H-ZSM-5 adsorbed 0.645 mmol NO_x ; in the

discharge stage, 50 mL/min flow rate of Ar, discharge power is set at 3.6 W, total input energy of the plasma is fixed at 0.832 W·h.

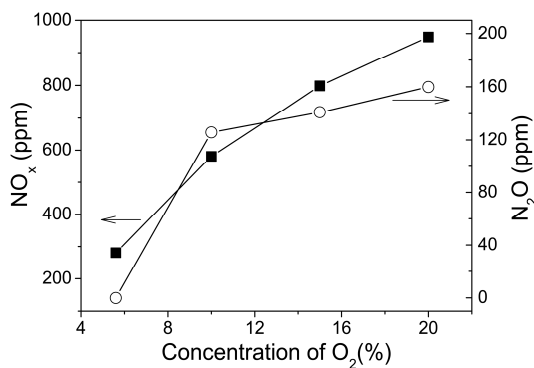


Figure S4. Reaction of O₂ with N₂ as a function of O₂ concentration in the DBD reactor.

Conditions: 50 mL/min of total flow rate, discharge power is set at 1.92 W.

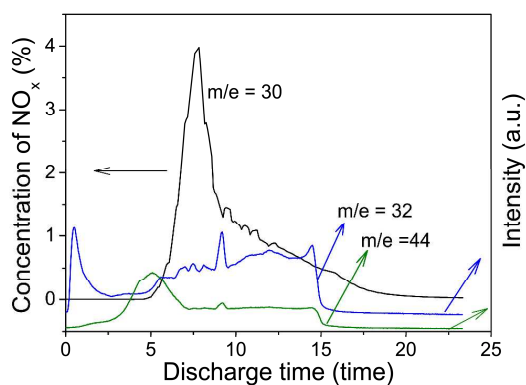


Figure S5. Effluents monitored by MS during decomposition of adsorbed NO_x on H-ZSM-5 in N₂ plasma at a discharge power of 3.6 W.

Notably, the MS signal $m/e = 30$ represents NO_x (NO + NO₂) in our MS

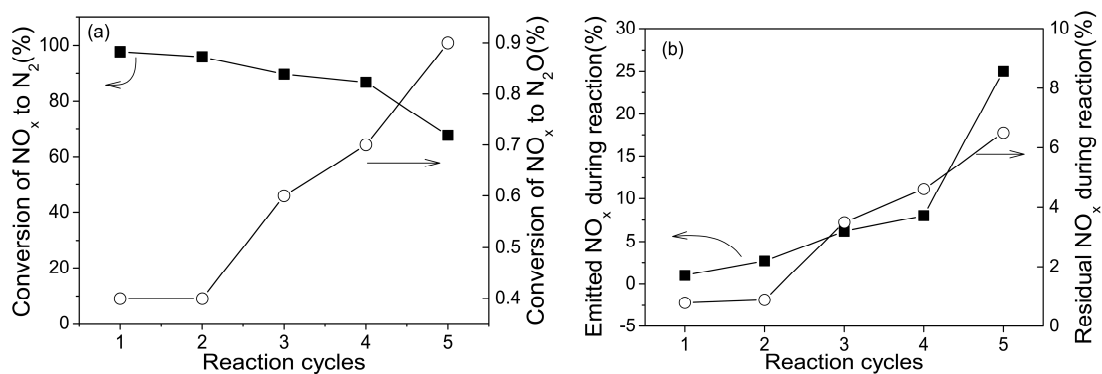


Figure S6. Cyclic operation of AC/H-ZSM-5 for the removal of NO_x (a) conversion of adsorbed NO_x (b) percentage of emitted NO_x during reaction and residual NO_x after reaction.

Conditions: in the adsorption stage, 0.645 mmol NO_x adsorbed; in the discharge stage, 50 mL/min flow rate of N₂, discharge power is 3.6 W, input energy of each cycle is 0.832W·h; between the intermission of two cycles, the AC/H-ZSM-5 should be treated in N₂ at 300 °C for 30 min, as without the procedure the conversion of NO_x is similar to that on bare H-ZSM-5. Notably, the percentage of NO_x remained on the catalyst after reaction is determined by the desorbed NO_x during the pretreatment process, as most adsorbed NO_x was desorbed in the temperature range (No other N-containing products were produced during the pretreatment process).