Oxidation of refractory sulfur-containing compounds with molecular oxygen catalysed by vanadoperiodate

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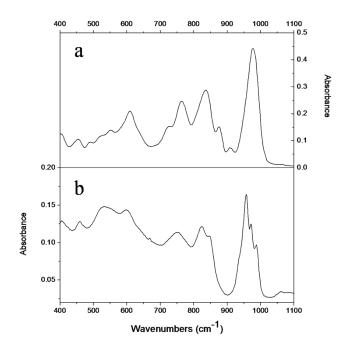
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 $\textbf{Fig. S1.} \ (a), \ FT\text{-IR spectrum of } (Q_8)_3HIV_9O_{28}; \ (b), \ FT\text{-IR spectrum of } (Q_8)_3H_3V_{10}O_{28}.$

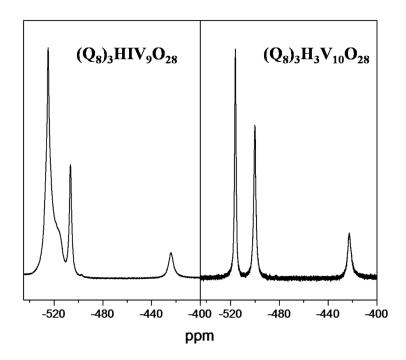


Fig. S2. 51 V NMR spectra of $(Q_8)_3$ HIV $_9$ O $_{28}$ and $(Q_8)_3$ H $_3$ V $_{10}$ O $_{28}$.

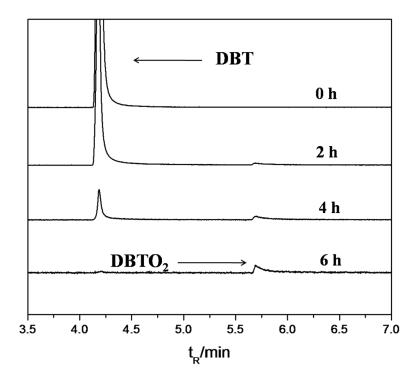


Fig. S3. Sulfur-specific GC-FPD chromatograms of the oxidation of DBT in decalin. Reaction conditions: $(Q_8)_3HIV_9O_{28}$ (40 mg), DBT(0.3 mmol) in 20 ml decalin, reaction temperature 90 °C, oxidant O_2 (1 atm).

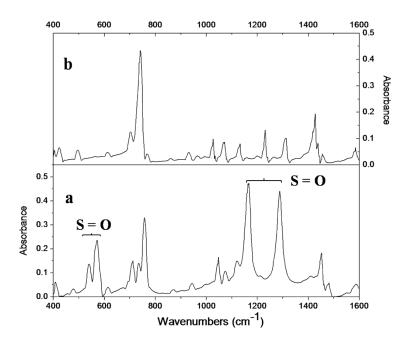


Fig. S4. FT-IR spectrum of DBT after reaction, wavenumbers : $400 \sim 1600$ cm⁻¹. (a), DBT after reaction; (b), DBT.

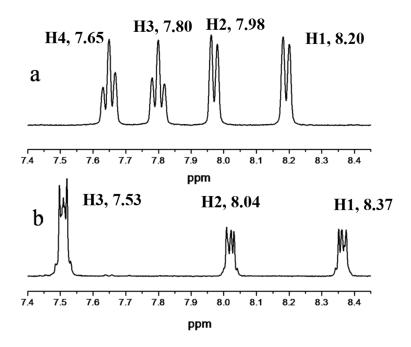


Fig. S5. (a), ¹H NMR spectrum of DBTO₂; (b), ¹H NMR spectrum of DBT.

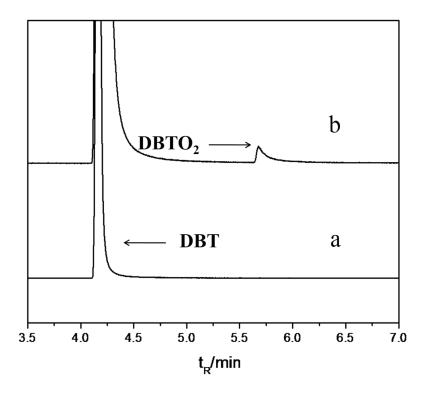


Fig. S6. Sulfur-specific GC-FPD chromatograms. (a) reaction product of experiment 1; (b) reaction product of experiment 2. Reaction conditions: (a) Reaction conditions: $(Q_8)_3$ HIV $_9$ O $_{28}$ (40 mg), DBT (0.3 mmol) in 20 ml decalin, reaction temperature 90 °C, Argon (1 atm); (b) Reaction conditions: $(Q_8)_3$ HIV $_9$ O $_{28}$ (40 mg), DBT (0.3 mmol) in 20 ml decalin, reaction temperature 90 °C, Argon (1atm), oxidant O $_2$ (1 atm).

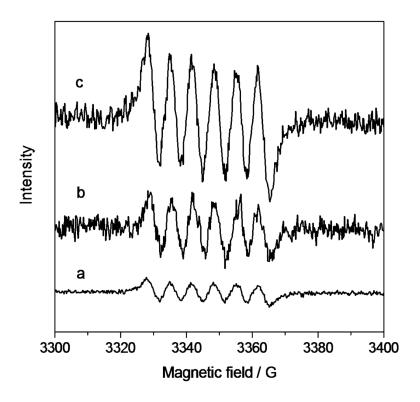


Fig. S7. In situ ESR spectra of DMPO- O_2 • adduct generated in the oxidative desulfurization process. (a) the sample tested with catalyst at 40 °C. (b) the sample tested with catalyst at 50 °C. (c) the sample tested with catalyst at 60 °C.

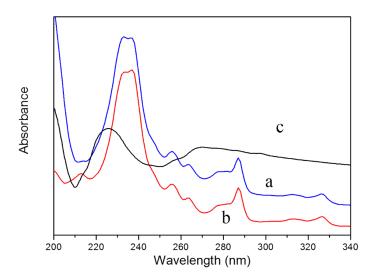


Fig. S8. UV-Vis spectra of DBT. All spectra were collected from 0.02 mmol L⁻¹ DBT and 30 mg L⁻¹ (Q_8)₃HIV₉O₂₈ solution. (a) DBT and (Q_8)₃HIV₉O₂₈ dissolved in decalin at 90 °C for 1 h under 1 atm Ar; (b) DBT dissolved in decalin at 90 °C for 1 h under 1 atm Ar; (c) (Q_8)₃HIV₉O₂₈ dissolved in decalin at 90 °C for 1 h under 1 atm Ar.