Reversible proton-switchable fluorescence controlled by conjugation effect in an organically-functionalized polyoxometalate

Chunlin Lv1,*, Kun Chen1,*, Junjie Hu2, Jin Zhang1, Rao Naumaan Nasim Khan1 & Yongge Wei1,3

A novel monosubstituted organoimido hexamolybdate containing 6-nitroquinoline moiety has been successfully synthesized. This organically-functionalized polyoxometalate exhibits proton-induced switchable fluorescence property in aqueous acetonitrile solution at room temperature. Experimental and theoretical investigations of this reversible "on" and "off" switching mechanism have been carried out, and it is found that the protonation and deprotonation at the heterocyclic nitrogen atom within quinoline fragment leads to the breaking and reformation of the conjugation through strong d-π interaction between the hexamolybdate anionic cluster and the quinoline moiety, resulting in "on" and "off" luminescence signal.

Molecular switches constitute a variety of reporter molecules able to show response to external stimuli by the reversible molecular structural change such as cis-trans isomerization, ring-closing/ring-opening, molecular proton-transfer processes, behavior of assembly and so on1–3. As an efficient and highly sensitive method, fluorescent molecular switches have attracted substantial research interest in the past decades4–9. In particular, proton sensors which can indicate the presence of protons are of great use in the field of detectors response to environmental changes, memories or logic gates in nanotechnology, and especially life sciences10,11. Up to date, though numerous fluorescent systems exhibiting reversible response to protons have been reported, the new challenge of analysis constantly needs to develop more proton-responsive fluorescent sensors. Therefore, the development of molecular switches having fluorescent response to external stimuli represents a continuous research topic.

In this sense, polyoxometalates (POMs) which are a rich class of metal oxide anionic clusters with diversity of structure and unique photonic, electronic, and magnetic properties12–16 are good candidates for fabricating multifunctional fluorescent materials. Previously, one well understood synthetic method was to incorporate lanthanide ions, mainly Eu3+ and Tb3+, into POMs17–22. Alternatively, the introduction of organic fluorophore may be an efficient synthetic strategy and have been attracted much attention in the past decade23–32. Especially, the organic functionalization of POMs via covalent linkage to form organic-inorganic hybrid fluorescent materials is a very attractive and challenging research area. Much pioneer work has been done by Neumann and Peng27–30, however it is found that POM clusters exhibit efficient fluorescence quenching when the inorganic clusters directly linked to the organic conjugated polymers or moieties through Mo-N triple bond28–30. Recently, Liu and Wei and co-workers reported that surfactant functionalized hexavanadates with proton or sodium ions as the counter-ions have fluorescent property31. Meanwhile, Liu and Hill and co-workers also found this photoluminescence in inorganic-organic hybrid hexavanadate clusters containing pyrene moiety32. However, to the best of our knowledge, POMs-based single-molecular switches displaying reversible "on" and "off" photoluminescence response to protons have not been reported yet.

In this paper, we report the synthesis of a novel organoimido hexamolybdate containing 6-nitroquinoline moiety, which exhibits reversible stimuli-response to protons. Furthermore, the mechanism of this reversible fluorescent behavior based on electron delocalization breaking and reforming is also investigated in detail.

1Department of Chemistry, Tsinghua University, Beijing 100084, China. 2Solid Waste and Chemicals Management Center, Ministry of Environmental Protection, Beijing 100029, China. 3State Key Laboratory of Natural and Biomimetic Drugs, Peking University, Beijing 100191, China. *These authors contributed equally to this work. Correspondence and requests for materials should be addressed to Y.W. (email: yonggewei@mail.tsinghua.edu.cn)
Results

Synthesis and characterization. A reaction of the hexamolybdate salt (\((\text{Bu}_4\text{N})_2[\text{Mo}_6\text{O}_{19}]\)) with 5-amino-6-nitro quinoline in the ratio of 1:1 using the well-established DCC (N,N′-dicyclohexylcarbodiimide) protocol can conveniently give rise to the corresponding monosubstituted organoimido derivative, (\((\text{Bu}_4\text{N})_2[\text{Mo}_6\text{O}_{18}(\text{NC}_9\text{H}_5\text{N}-\text{NO}_2)]\) (Mo\(_6\)-Q-NO\(_2\)), which is crystallized as orange lamellar-like crystals when diethyl ether diffuses into their solution of acetonitrile. The IR spectrum of compound Mo\(_6\)-Q-NO\(_2\) exhibits similar characteristic features to those of the reported monosubstituted organoimido hexamolybdates (Supplementary Fig. S1). In addition, the ESI-MS of this compound shows clearly resolved peaks at 525.6 and 1293.9 which correspond to \([\text{Mo}_6\text{O}_{18}(\text{NC}_9\text{H}_5\text{N}-\text{NO}_2)]^{2-}\) (calcd. 525.4) and \((\text{Bu}_4\text{N})[\text{Mo}_6\text{O}_{18}(\text{NC}_9\text{H}_5\text{N}-\text{NO}_2)]^{-}\) (calcd. 1293.3) (Supplementary Fig. S2), respectively.

Furthermore, single-crystal X-ray diffraction investigation has also been performed. It is found that an asymmetric unit contains one independent cluster anion and two counter tetrabutylammonium cations as well as a half of acetonitrile molecule and one water molecule. The ORTEP drawings of the anionic cluster within compound Mo\(_6\)-Q-NO\(_2\) is presented in Fig. 1. As shown in Fig. 1, compound Mo\(_6\)-Q-NO\(_2\) displays the typical structural features as those of monofunctionalized hexamolybdates: one terminal oxo atom (Ot) of the hexamolybdate cluster has been replaced by an organoimido ligand. In addition, the formed Mo–N bond is found to be 1.730 Å, and the nearly linear Mo–N–C angle is refined as 167.1° (Supplementary Tables S1 and S2), exhibiting the typical triple bond features.

Photo-physical property. The UV-Vis spectrum of compound Mo\(_6\)-Q-NO\(_2\) is shown in Fig. 2. The lowest-energy electronic transition (LET) absorption band in 50% aqueous acetonitrile solution locates at 423 nm, which exhibits obviously bathochromic shift compared to the parent hexamolybdate due to the quinoline aromatic ring conjugated with the hexamolybdate framework through the strong d–π interaction. To assess the effect of acid and base, absorption changes of Mo\(_6\)-Q-NO\(_2\) in the presence of 10 equivalents of \(\text{H}^+\) and \(\text{OH}^-\) in 50% aqueous acetonitrile solution have been determined respectively. It is found that the LET absorption band
has no obviously change in the presence of 10 equivalents of OH\textsuperscript{−}, while the LET absorption band has significant hypsochromic shift from 423 nm to 383 nm in the presence of 10 equivalents of H\textsuperscript{+}. Interestingly, upon further addition of 0.5 mL dilute NaOH solution (10 equivalents) into the acidic solution to neutralize the HCl, the LET absorption band migrates back to 423 nm. Note that the decreased intensity of absorption peaks attributes to the dilution effect of the addition of 0.5 mL dilute NaOH solution. This reversible change of LET absorption band not only indicates that Mo\textsubscript{6}-Q-NO\textsubscript{2} is stable after these treatments but also implies that a corresponding reversible change of conjugate delocalization system within compound Mo\textsubscript{6}-Q-NO\textsubscript{2} takes place.

Theoretical investigation of the protonated structure. To explore the cause of reversible change of LET absorption band, we firstly need to determine the protonation site. Previous studies have pointed out that bridge oxygen (O\textsubscript{b}) is a preferable binding site to proton as compared to terminal oxygen\textsuperscript{36–39}. Therefore, within Mo\textsubscript{6}-Q-NO\textsubscript{2} there are four possible protonation sites which are bridge oxygen atoms bound to the molybdate atom at the trans site of the imido molybdate atom (p-O\textsubscript{b}), bridge oxygen atoms only bound to the molybdate atom at the cis site of the imido molybdate atom (m-O\textsubscript{b}), and bridge oxygen atoms bound to imido molybdate atom (o-O\textsubscript{b}) as well as the heterocyclic nitrogen atom. The corresponding structures are defined as p-OH, m-OH, o-OH and NH, respectively. By the optimization of their structures using DFT-B3LYP method\textsuperscript{40,41}, shown in Fig. 3, it was found that the structure corresponding to the protonation at the heterocyclic nitrogen atom is the most stable isomer. Based on this result, we can explain the reversible change of absorption band from the corresponding change of structure. The protonation at the heterocyclic nitrogen atom leads to the Mulliken charge of the hydrogen bound to the 4 site carbon in quinoline moiety increased from 0.362 to 0.555, resulting in much stronger hydrogen bonding between 4-H and the bridge atom O15. Combination of the effect of hindrance of 6-nitro, in addition, the angle of Mo-N-C decreased to 150.5°. Simultaneously, the bond length of Mo-N increased to 1.851 Å while that of N-C decreased to 1.315 Å (Supplementary Fig. S3). All these changes of structure indicate that the Mo-N bond exhibits much more like single bond, largely decreasing the d-\pi interaction between inorganic hexamolybdate cluster and organic quinoline aromatic ring. Therefore, the absorption band in UV-Vis spectrum shifts to 383 nm after protonation.

Reversible proton-switchable fluorescence. In previous investigations, it was found that the Mo-N triple bond linking POM clusters with conjugated aromatic moiety displayed significant fluorescence quenching because of the photo-induced electron transfer from the fluorophore to the clusters\textsuperscript{27–30}. Likewise, compound Mo\textsubscript{6}-Q-NO\textsubscript{2} also exhibits no fluorescence. However, compound Mo\textsubscript{6}-Q-NO\textsubscript{2} clearly displays proton responsive “on” fluorescence at 560 nm with excitation wavelength of 377 nm. As shown in Fig. 4a, upon gradual addition of
dilute HCl (0~20 equimolar) to the 50% aqueous acetonitrile solution of Mo6-Q-NO2, the intensity of the emission at 560 nm increased gradually. While upon further gradual addition of dilute NaOH (0~16 equimolar) to the acidic solution, the emission intensity exhibited gradual decreasing until the fluorescence almost “off”, (presented in Fig. 4b). To verify this reversible “on” and “off” fluorescence originated from the presence and absence of H+, further gradual addition of 8 and 16 equimolar dilute HCl have been performed, and the recovered fluorescence is observed. A control experiment confirmed that this proton-switchable fluorescence was due to Mo6-Q-NO2, as the organic ligand 5-amino-6-nitroquinoline showed almost no fluorescence with excitation wavelength of 377 nm (Supplementary Fig. S4). It is noteworthy that this is a first example of organoimido derivative of POMs with significant reversible proton-switchable fluorescent property.

Discussion

In light of the structural analysis as above, fluorescence enhancement upon gradual addition of dilute HCl into the 50% aqueous acetonitrile solution of Mo6-Q-NO2 is due to the protonation at the heterocyclic nitrogen atom leading to the Mo-N changing from triple bond to single bond, resulting in the effective channel of photo-induced electron transfer from the organic 6-nitroquinoline moiety to the hexamolybdate cluster that is blocked. Contrarily, the quenching of fluorescence upon addition of NaOH into this system attributes to subsequent deprotonation of the heterocyclic nitrogen atom, regenerating the effective photo-induced electron transfer channel, illustrated in Fig. 5.

In summary, we have successfully synthesized a novel mono-substituted organoimido hexamolybdate containing isoquinoline moiety which, as a first example of organoimido-POMs-based fluorescent single-molecule switch, exhibits proton-induced switchable fluorescence property in aqueous acetonitrile solution at room temperature. Moreover, experimental and theoretical investigations of reversible “on” and “off” switching mechanism have been carried out, and it is found that the protonation and deprotonation at the heterocyclic nitrogen atom within quinoline fragment leads to the breaking and reformation of the conjugation through strong d-π interaction between the hexamolybdate anionic cluster and the quinoline moiety, resulting in “on” and “off” luminescence signal. This proton-responsive switchable fluorescent activity provides a potential application in sensors using organic functionalized polyoxometalates. More importantly, this work paves a new way to design and synthesize POM-based organic-inorganic hybrid fluorescent single-molecule switches.
Methods

Materials. \((\text{Bu}_4\text{N})_2[\text{Mo}_6\text{O}_{19}]\) was synthesized according to the improved literature method\(^{42}\) and dried before use. 5-amino-6-nitroquinoline was purchased from Alfa Aesar. Acetonitrile was dried by refluxing in the presence of CaH\(_2\) and was distilled prior to use. N,N\('\)′-dicyclohexylcarbodiimide (DCC) and ethyl ether were directly used without further purification.

Spectroscopic Characterization. Infrared spectra were measured using the method of KBr pellets and recorded on a Perkin-Elmer FTIR spectrometer. The electrospray mass spectra (ESI-MS) were measured on a Bruker Apex IV FTMS Plus ion-trap mass spectrometer, and all experiments were carried out in the negative-ion mode using CH\(_3\)CN as solvent. UV-Vis absorption spectra of Mo\(_6\)-Q-NO\(_2\) in a mixed solution of acetonitrile and water were recorded on a UN-2100s spectrometer. Fluorescence measurements were recorded in 3 mL quartz cuvettes at room temperature using a Hitachi F-7000 FL Spectrophotometer equipped with a xenon lamp excitation source. All fluorescence spectra were measured at an excitation wavelength of 377 nm.

Crystallographic structural determinations. A suitable crystal of the compound Mo\(_6\)-Q-NO\(_2\) was covered with petroleum jelly, mounted onto glass fibers, and transferred directly to a Rigaku RAXIS-SPIDER IP diffractometer. Data collection, data reduction, cell refinement, and experimental absorption correction were performed with the software package of Rigaku RAPID AUTO (Rigaku, 1998, Ver2.30). Structures were solved by direct methods and refined against F\(^2\) by full matrix least squares. All non-hydrogen atoms, except disordered atoms, were refined anisotropically. Hydrogen atoms were generated geometrically. All calculations were performed using the SHELXTL V. 5.10 program\(^{43}\).

Theoretical calculations. All the calculations presented in this work were carried out with the Gaussian-09 program package\(^{44}\). The structures of each stationary point were fully optimized using the B3LYP method, in combination with the LANL2DZ basis set for molybdate atoms and the 6-31G(d) basis set for rest atoms. Vibrational frequencies of each stationary point were calculated at the same level of theory to characterize the nature of the stationary points and give the thermal correction to Gibbs free energy. In addition, to consider the solvent effect, a single point energy calculation was performed by using the integral equation formulism within the polarized continuum model (IEF-PCM)\(^{45}\) with the Bondi atomic radii for the united atom topological model at the B3LYP level based on each optimized geometry, with the LANL2DZ basis set for molybdate atoms and the 6–31+G(d, p) basis set for rest atoms denoted as IEFPCM-B3LYP/BS1. The dielectric constant of acetonitrile is the default value of the G09 program.

Synthesis of \((\text{Bu}_4\text{N})_2[\text{Mo}_6\text{O}_{19}(\text{NC}_9\text{H}_5\text{N-NO}_2)]\) \((\text{Mo}_6\text{Q-NO}_2)\). A mixture of \((\text{Bu}_4\text{N})_2[\text{Mo}_6\text{O}_{19}]\) (1.36 g, 1.0 mmol), 5-amino-6-nitroquinoline (0.189 g, 1.0 mmol), and DCC (0.23 g, 1.1 mmol) dissolved in 15 mL anhydrous acetonitrile and then heated to reflux at 110 °C for about 20 h. After being cooled to room temperature, the reaction solution was filtrated to remove the white precipitates of 1,3-dicyclohexylurea (DCU) and then the filtrate poured into ether, resulting in precipitates to come out. After the solution changing to clarify, the supernatant liquid was poured off, and the resulting small crystals were compound Mo\(_6\)-Q-NO\(_2\) (0.476 g, yield: 31%). By diffusion of Et\(_2\)O into their solution in acetonitrile, orange lamellar-like crystals which are suitable for the X-ray diffraction were obtained for compound Mo\(_6\)-Q-NO\(_2\). IR (KBr): 797, 953, 977, 1152, 1321, 1386, 1420, 1481, 1556, 2873, 2933, 2961 cm\(^{-1}\). UV/Vis (MeCN): \(\lambda_{\text{max}} = 423\) nm. ESI-MS(m/z): 1293.9 ([Mo\(_6\)O\(_{18}\)(NC\(_9\)H\(_5\)N-NO\(_2\))\(^+\), calcd 1293.3), 525.6 ([Mo\(_6\)O\(_{18}\)(NC\(_9\)H\(_5\)N-NO\(_2\))]\(^2\), calcd 525.4). Crystal data for compound (Mo\(_6\)-Q-NO\(_2\))

Figure 5. Schematic illustration of switchable fluorescence response to proton. Proposed mechanism for Mo\(_6\)-Q-NO\(_2\) displaying single-molecule switchable fluorescence response to proton.
References

1. Natali, M. & Giordani, S. Molecular switches as photocontrollable "smart" receptors. Chem. Soc. Rev. 41, 4010–4029 (2012).
2. Ray, D., Foy, J. T., Hughes, R. P. & Aprahamian, I. A switching cascade of hydrazine-based rotary switches through coordination-coupled proton relays. Nat. Chem. 4, 757–762 (2012).
3. Lee, L. K., Ginsburg, M. A., Crovace, C., Donohoe, M. & Stock, D. Structure of the torque ring of the flagellar motor and the molecular basis for rotational switching. Nature 466, 996–1000 (2010).
4. Kundu, P. K., Olsen, G. L., Kiss, V. & Klajn, R. Nanoporous frameworks exhibiting multiple stimuli responsiveness. Nat. Commun. 5, 3588 (2014).
5. Cui, X., Zhao, J., Zhou, Y., Ma, J. & Zhao, Y. Reversible photoswitching of triplet-triplet annihilation upconversion using dithienylethenophenanthroline. J. Am. Chem. Soc. 136, 9256 (2014).
6. Fukumaita, T. et al. Molecular design strategy toward diarylethenes that photoswitch with visible light. J. Am. Chem. Soc. 136, 17143–17154 (2014).
7. Amendola, V., Bergamaschi, G., Boiocchi, M., Fabbrizzi, L. & Mosca, L. The interaction of fluoride with fluorogenic ureas: an ON1–OFF–ON2 response. J. Am. Chem. Soc. 135, 6634–6655 (2013).
8. Biedermann, F., Elmaleh, E., Ghosh, I., Nau, W. M. & Scherman, O. A. Strongly fluorescent, switchable perylene bis(dicarboximide) host-guest complexes with cucurbit[8]uril in water. Angew. Chem. Int. Ed. 51, 7739–7743 (2012).
9. Kim, Y. et al. High contrast reversible fluorescent photoswitching of dye-crosslinked dendritic nanoclusters in living vertebrates. Angew. Chem. Int. Ed. 51, 2878–2882 (2012).
10. Auwärter, W. et al. A surface-anchored molecular four-level conductance switch based on single proton transfer. Nat. Nanotech. 7, 41–46 (2012).
11. Barile, C. J. et al. Proton switch for modulating oxygen reduction by a copper electrocatalyst embedded in a hybrid bilayer membrane. Nat. Mater. 13, 619–623 (2014).
12. Special thematic issue on polyoxometalates. Hill, C. L. Ed. Inorganic-organic hybrid vesicles with counterion- and pH-controlled fluorescent properties. Angew. Chem. Int. Ed. 49, 7702–7705 (2010).
13. Chen, L. et al. A basket tetradeceanadate cluster with blue luminescence. J. Am. Chem. Soc. 127, 8588–8589 (2005).
14. Stillman, M. J. & Thompson, A. J. Emission spectra of some lanthanoid decatungstate and undecatungstosilicate ions. J. Chem. Soc., Dalton Trans. 1138–1144 (1976).
15. Yamase, T., Kobayashi, T., Sugeta, M. & Naruke, H. Europium (III) luminescence and intramolecular energy transfer studies of polyoxometalatehydroporphosphates. J. Phys. Chem. A, 101, 5046–5053 (1997).
16. Blasse, G. & Dirksen, G. J. The luminescence of some lanthanide decatungstates. Chem. Phys. Lett. 83, 449–451 (1981).
17. Granadeiro, C. M., Ferreira, R. A. S., Soares-Santos, P. C. R., Carlos, L. D. & Nogueira, H. I. S. Lantanohexamolybdometallates as building blocks for multilength photoluminescent organic-inorganic hybrid materials. Eur. J. Inorg. Chem. 5088–5095 (2009).
18. Ritchie, C., Moore, E. G., Spel URich, M., Kogerler, P. & Boskovic, C. Terbium polyoxometalate organic complexes: correlation of structure with luminescence properties. Angew. Chem. Int. Ed. 49, 7702–7705 (2010).
19. Zhao, W.-F. et al. Synthesis of diamondoid lanthanide-polyoxometalate solids as tunable photoluminescent materials. Dalton Trans. 41, 10991–10996 (2012).
20. Wang, B., Bi, L.-H. & Wu, L.-X. Electroswitchable fluorescent thin film controlled by polyoxometalate. J. Mater. Chem. 21, 69–71 (2011).
21. Qin, B. et al. Reversible photoswitchable fluorescence in thin films of inorganic nanoparticle and polyoxometalate assemblies. J. Am. Chem. Soc. 132, 2886–2888 (2010).
22. Jin, L. et al. Polyoxometalate-based inorganic-organic hybrid film structure with reversible electroswitchable fluorescence property. Chem. Commun. 48, 2101–2103 (2012).
23. Zhang, H. et al. Hybrid microtubes of polyoxometalate and fluorescence dye with tunable photoluminescence. Chem. Commun. 48, 4462–4464 (2012).
24. Bar-Nahum, I., Narasimhulu, K. V., Weiner, L. & Neumann, R. Phenanthroline–polyoxometalate hybrid compounds and the observation of intramolecular charge transfer. Inorg. Chem. 44, 4900–4902 (2005).
25. Lu, M. et al. Synthesis of main-chain polyoxometalate-containing hybrid polymers and their applications in photovoltaic cells. Chem. Mater. 17, 402–408 (2005).
26. Xu, B. et al. Synthesis and optical properties of conjugated polymers containing polyoxometalate clusters as side-chain pendant. Chem. Mater. 17, 2841–2851 (2005).
27. Chakraborty, S., Keightley, A., Dusevich, V., Wang, Y. & Peng, Z. Synthesis and optical properties of a rod-coil diblock copolymer with polyoxometalate clusters covalently attached to the coil block. Chem. Mater. 22, 3995–4006 (2010).
28. Yin, P. et al. A double-tailed fluorescent surfactant with a hexavanadate cluster as the head group. Angew. Chem. Int. Ed. 50, 2521–2525 (2011).
29. Li, D. et al. Inorganic-organic hybrid vesicles with counterion- and pH-controlled fluorescent properties. J. Am. Chem. Soc. 133, 14010–14016 (2011).
30. Wei, Y. G., Xu, B. B., Barnes, C. L. & Peng, Z. H. An efficient and convenient reaction protocol to organoimido derivatives of polyoxometalates. J. Am. Chem. Soc. 123, 4083–4084 (2001).
31. Strong, J. B. et al. A new class of functionalized polyoxometalates: synthetic, structural, spectroscopic, and electrochemical studies of organoimido derivatives of [Mo6O19]4−. J. Am. Chem. Soc. 122, 639–649 (2000).
32. Xiao, F. et al. Polyoxometalatocyclonatophanes: controlled assembly of polyoxometalate-based chiral metallamarcocycles from achiral building blocks. J. Am. Chem. Soc. 132, 5956–5957 (2010).
33. Yao, J. P. Optical and electrochemical properties of polyoxometalates. J. Am. Chem. Soc. 123, 137–163 (2002).
34. López, X., Bo, C. & Poblet, J. M. Electronic properties of polyoxometalates: electron and proton affinity of mixed-Addenda Keggin and Wells–Dawson anions. J. Am. Chem. Soc. 124, 12574–12582 (2002).
35. Janik, M. J., Davis, R. J. & Neurock, M. Anhydrous and water-assisted proton mobility in phosphotungstic acid. J. Am. Chem. Soc. 127, 5238–5245 (2005).
36. Yang, J. et al. Location, acid strength, and mobility of the acidic protons in Keggin 12-HPW12O40−: a combined solid-state NMR spectroscopy and DFT quantum chemical calculation study. J. Am. Chem. Soc. 127, 18274–18280 (2005).
37. Becke, A. D. Density-functional thermodynamics. III. The role of exact exchange. J. Chem. Phys. 98, 5648–5652 (1993).

(MeCN)6[H2O]·(H2O): C42H39O43N42O21, M = 1574.26, Monoclinic, space group P-1 (No. 1), a = 12.6462(8), b = 12.7095(8), c = 19.7792(2), β = 101.3717(9), γ = 102.6574(9), V = 2936.64(4) Å3, T = 103 K, Z = 2, Dc = 1.780 g cm−3, m = 1.316 mm−1, 30329 collected reflections, 15122 independent (Rint = 0.0446), Goof = 1.068, Final R indices (I > 2σ(I)) R1 = 0.0647, wR2 = 0.1260.
41. Lee, C. T., Yang, W. T. & Parr, R. G. Development of the Colle-Salvetti correlation-energy formula into a functional of the electron density. *Phys. Rev. B* **37**, 785–789 (1988).
42. Hur, N. H., Klemperer, W. G. & Wang, R.-C. Tetrabutylammonium hexamolybdate (VI). *Inorg. Synth.* **27**, 77–78 (1990).
43. Sheldrick, G. M. SHELXTL v. 5.10, Structure determination software suite. *Bruker AXS*, Madison, Wisconsin (USA) (1998).
44. Gaussian 09, Revision A.1, Frisch, M. J. *et al.* Gaussian, Inc., Wallingford CT (2009).
45. Cossi, M., Barone, V., Mennucci, B. & Tomasi, J. Ab initio study of ionic solutions by a polarizable continuum dielectric model. *Chem. Phys. Lett.* **286**, 253–260 (1998).

**Acknowledgements**
This work is supported by the National Natural Science Foundation of China (NSFC Nos 21225103 and 21221062), THSJZ, Tsinghua University Initiative Foundation Research Program No. 20131089204 and the State Key Laboratory of Natural and Biomimetic Drugs.

**Author Contributions**
C.L. and K.C. conceived the idea, designed and performed the experiment, analyzed the results and wrote the manuscript. J.H. conducted the DFT calculations. J.Z. and R.N.N.K. assisted in writing the manuscript. Y.W. supervised the project, helped design the experiments and calculations, analyzed the data and wrote the manuscript. All the authors discussed the results and commented on the manuscript.

**Additional Information**
Supplementary information accompanies this paper at http://www.nature.com/srep

Competing financial interests: The authors declare no competing financial interests.

**How to cite this article**: Lv, C. *et al.* Reversible proton-switchable fluorescence controlled by conjugation effect in an organically-functionalized polyoxometalate. *Sci. Rep.* **6**, 27861; doi: 10.1038/srep27861 (2016).

This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/