

Supporting Information

Solvent Dielectricity-Modulated Helical Assembly and Morphologic Transformation of Achiral Surfactant-Inorganic Cluster Ionic Complexes

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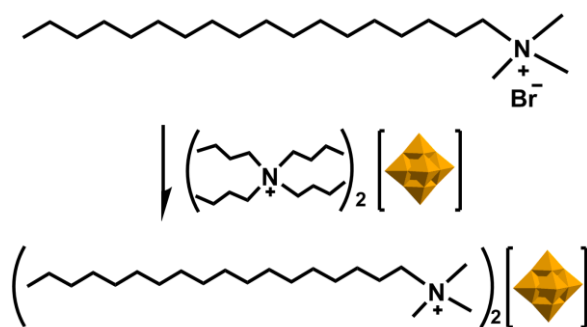


Figure S1. Synthesis route of $(\text{ODTA})_2[\text{Mo}_6\text{O}_{19}]$.

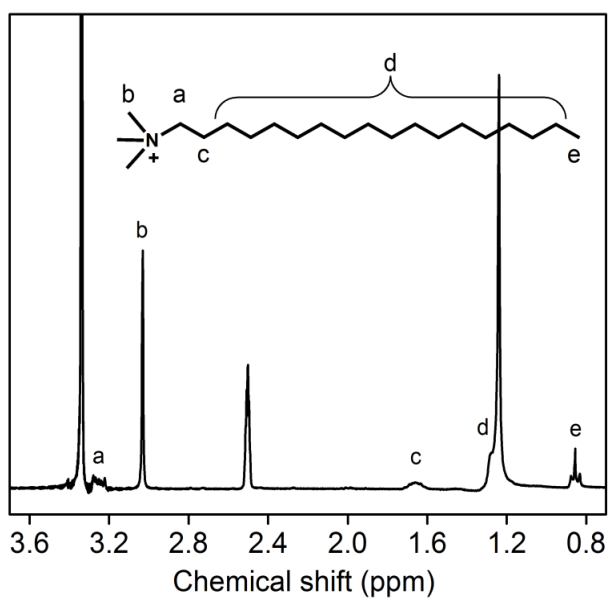


Figure S2. ^1H NMR spectrum of $(\text{ODTA})_2[\text{Mo}_6\text{O}_{19}]$ in $\text{DMSO}-d_6$.

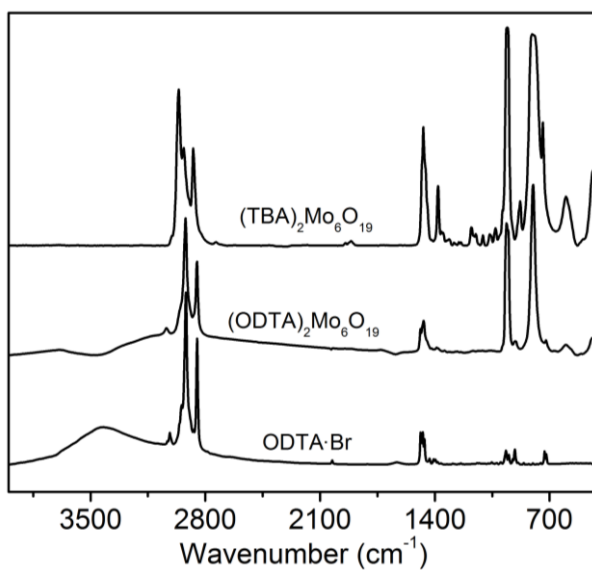


Figure S3. FT-IR spectra of $(\text{TBA})_2[\text{Mo}_6\text{O}_{19}]$, $(\text{ODTA})_2[\text{Mo}_6\text{O}_{19}]$, and $\text{ODTA}\cdot\text{Br}$.

Table S1. Assignments of characteristic infrared spectra of (TBA)₂[Mo₆O₁₉], (ODTA)₂[Mo₆O₁₉], and ODTA·Br.

(TBA) ₂ [Mo ₆ O ₁₉] (cm ⁻¹)	(ODTA) ₂ [Mo ₆ O ₁₉] (cm ⁻¹)	ODTA·Br (cm ⁻¹)	Assignments ^a
	3030	3433	O–H asymmetrical stretching
2962	2954	2956	CH ₃ asymmetrical stretching
2931	2920	2918	CH ₂ asymmetrical stretching
2873	2850	2850	CH ₂ symmetrical stretching
	1485		CH ₂ –N ⁺ scissoring
1468	1467	1467	CH ₂ scissoring
1378	1388		CH ₃ scissoring
956	962		$\nu_{\text{as}}(\text{Mo–O}_t)$
798	802		$\nu_{\text{as}}(\text{Mo–O}_b)$
597	605		$\delta(\text{O}_b\text{–Mo–O}_t)$
438	439		$\delta(\text{O}_b\text{–Mo–O}_t)$

Table S2. Elemental analysis for (ODTA)₂[Mo₆O₁₉].

Sample		C/%	H/%	N/%
(ODTA) ₂ [Mo ₆ O ₁₉]	Found	33.47	6.09	1.82
	Calcd.	33.52	6.16	1.86

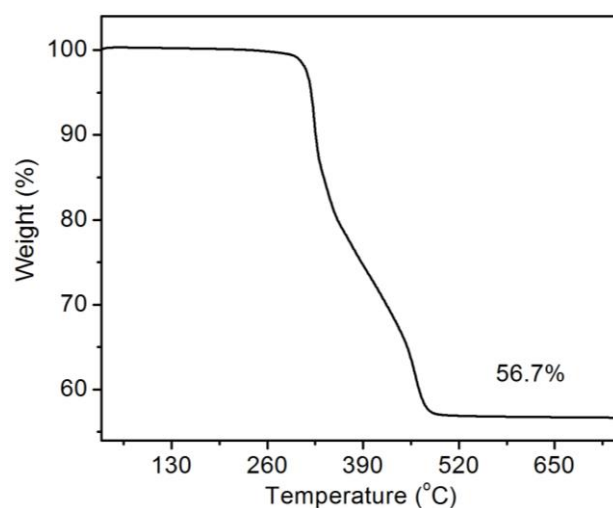


Figure S4. TGA curve of (ODTA)₂[Mo₆O₁₉] carried out in air, the measured residue of 56.7% at 700 °C is in agreement with the calculated value of 57.4% from the given formula C₄₂H₉₂N₂Mo₆O₁₉.

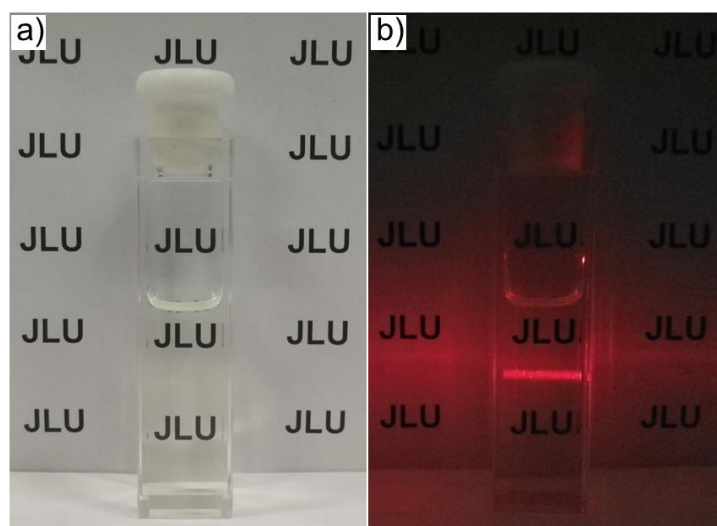


Figure S5. (a) Digital photo and (b) corresponding Tyndall scattering observation of (DODA)₂[Mo₆O₁₉] in the solution of dichloromethane/methanol (30:1 in v/v) at concentration of 1.0 mg mL⁻¹.

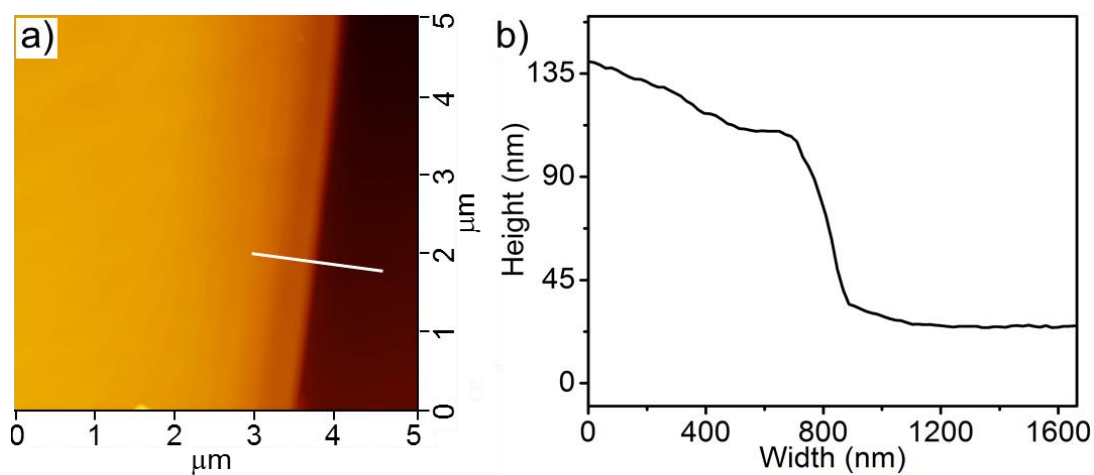


Figure S6. (a) AFM image and (b) the corresponding height profile of $(\text{DODA})_2[\text{Mo}_6\text{O}_{19}]$ self-assemblies in dichloromethane.

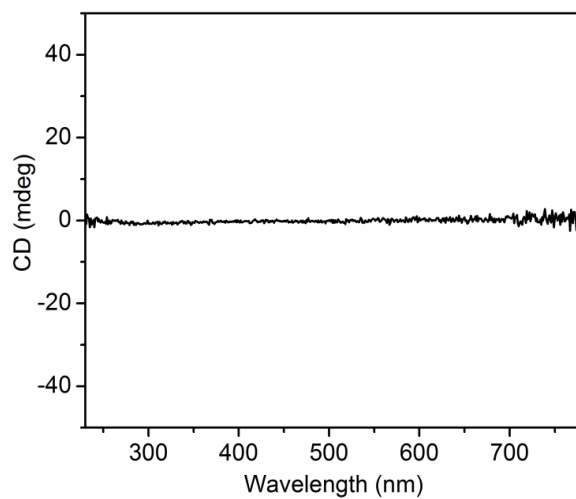


Figure S7. CD spectrum of $(\text{DODA})_2[\text{Mo}_6\text{O}_{19}]$ casting film on quartz substrate obtained from dichloromethane/methanol (30:1 v/v).

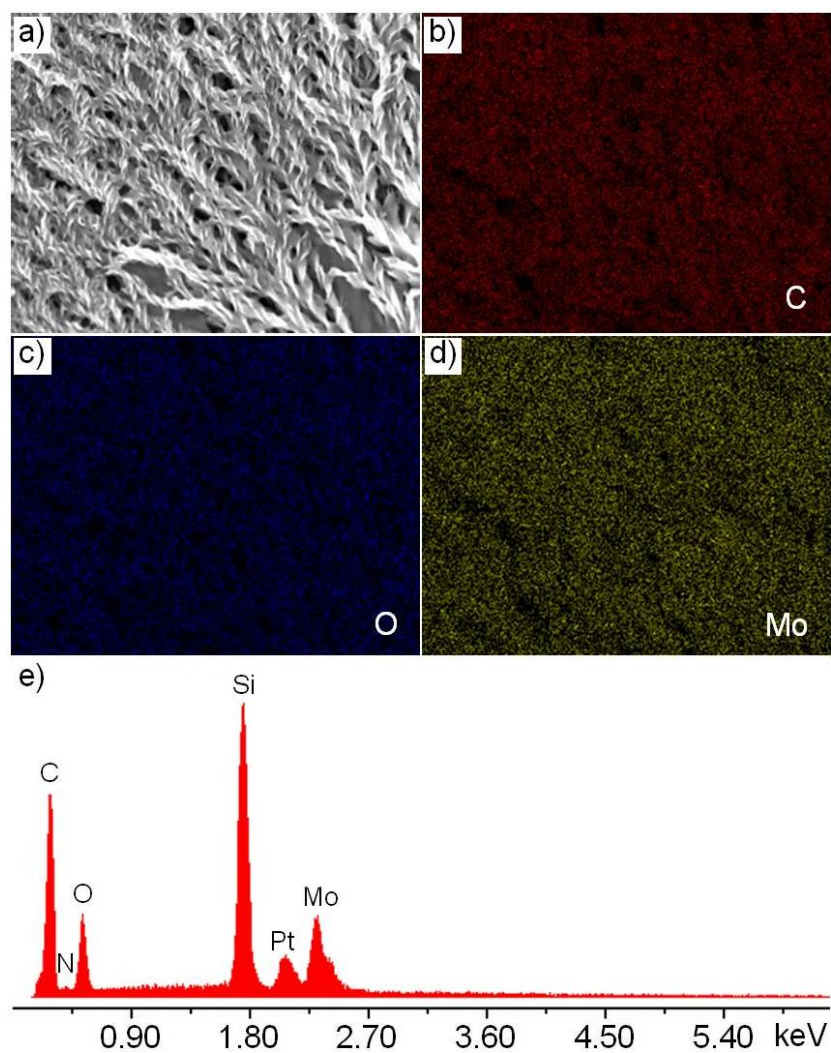


Figure S8. (a) HAADF-STEM image of the helical assemblies of $(\text{DODA})_2[\text{Mo}_6\text{O}_{19}]$ on silica substrate; (b), (c) and (d) the EDX-mapping image of C, O and Mo element distribution marked as red, blue and yellow dots, respectively; (e) the binding energy spectrum of the region of image (a). Conductive coating: platinum ultrathin coating of electrically-conducting material, deposited by low vacuum sputter coating of the sample.

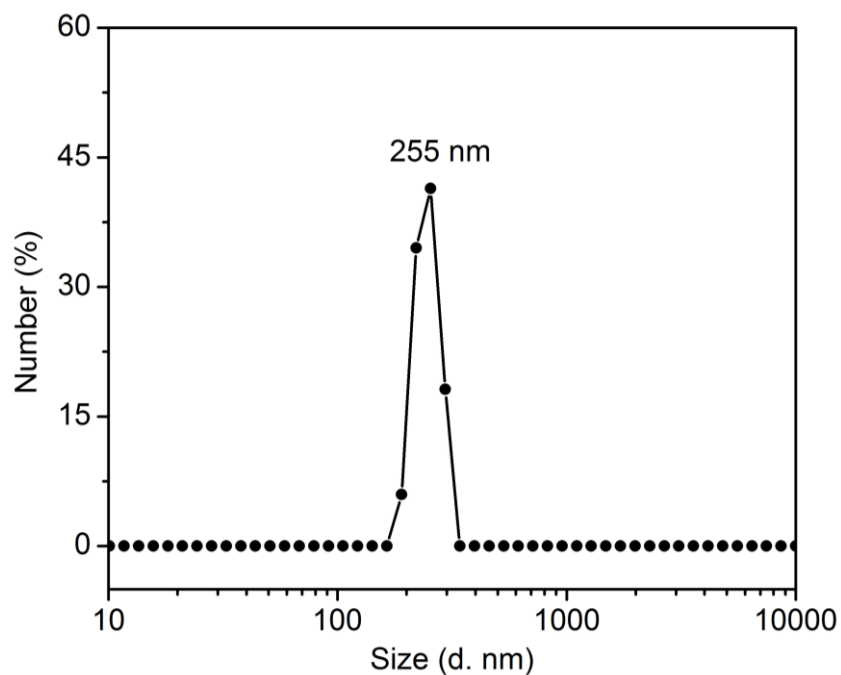


Figure S9. DLS of $(\text{DODA})_2[\text{Mo}_6\text{O}_{19}]$ in the solution of dichloromethane/methanol (30:1 in v/v) with concentration 1.0 mg mL^{-1} .

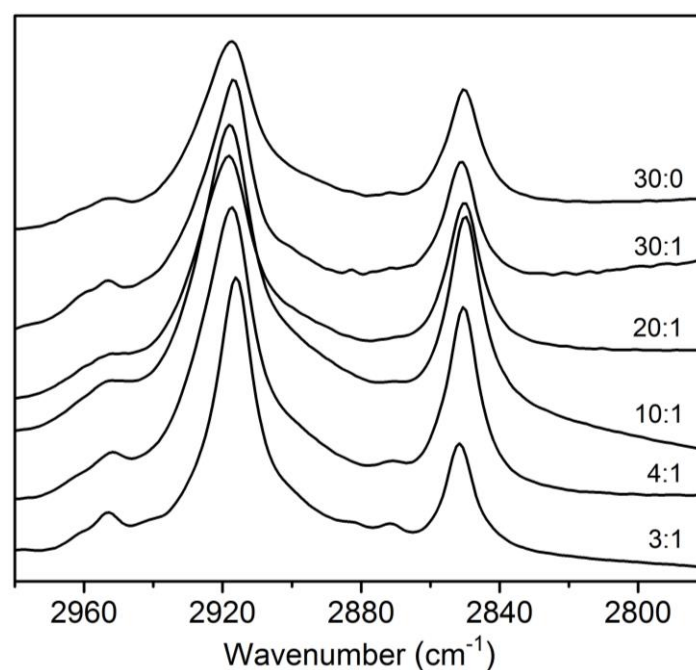


Figure S10. FT-IR spectra of $(\text{DODA})_2[\text{Mo}_6\text{O}_{19}]$ in the mixed solvent CH_2Cl_2 and CH_3OH with different volume ratios.

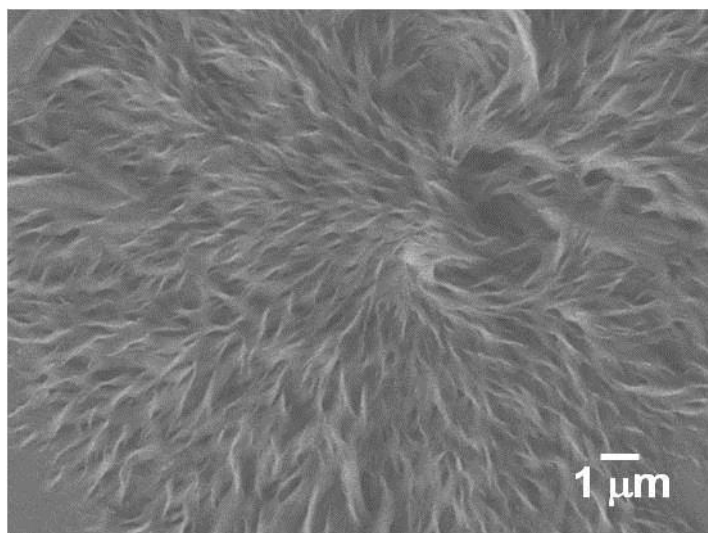


Figure S11. SEM image of the helical assembly of $(\text{DODA})_2[\text{Mo}_6\text{O}_{19}]$ in dichloromethane/methanol (20:1) solution at 30 °C with the concentration of 1 mg mL^{-1} .

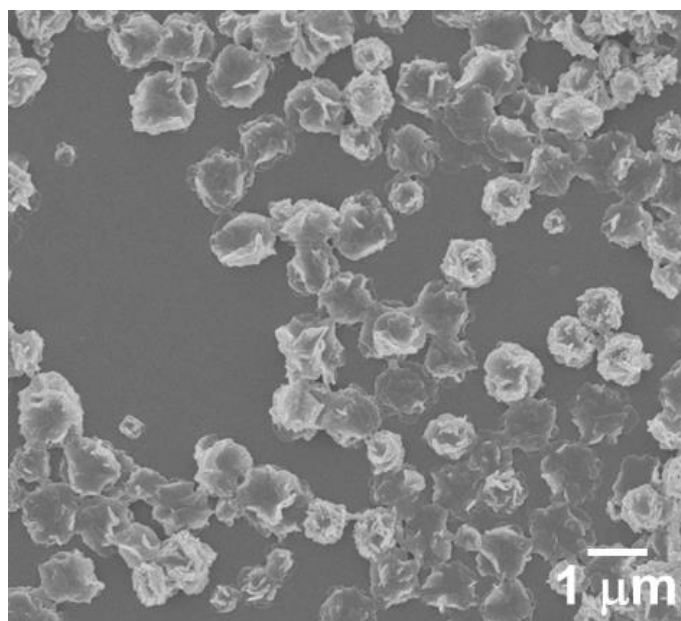


Figure S12. SEM image of $(\text{DODA})_2[\text{Mo}_6\text{O}_{19}]$ for the immediate state after sonication for 5 min, aging for 5 min, in dichloromethane/methanol (4:1 v/v) solution at 30 °C with the concentration of 1 mg mL^{-1} .

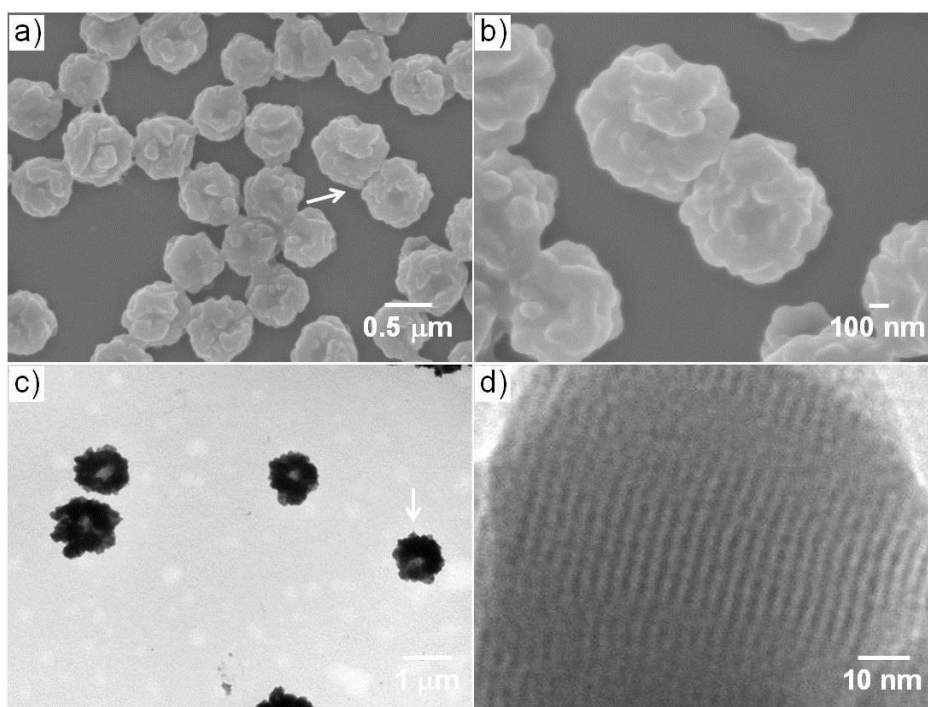


Figure S13. (a) SEM and (c) TEM images of $(\text{DODA})_2[\text{Mo}_6\text{O}_{19}]$ in dichloromethane/methanol (3:1 v/v) solution at 30 °C with the concentration of 1 mg mL⁻¹. Magnified (b) SEM and (d) TEM images of the area marked in (a) and (c).

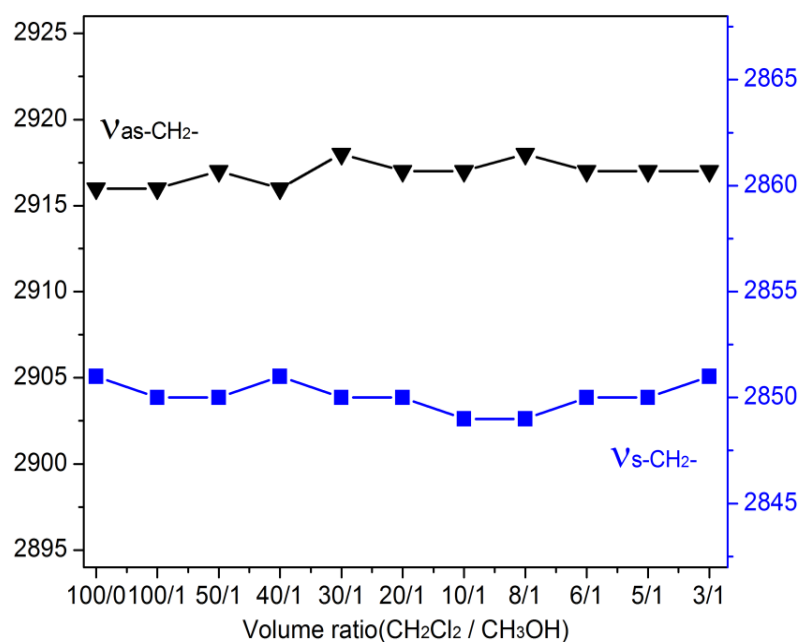


Figure S14. Solvent dependence of the positions of antisymmetric and symmetric CH₂ stretching mode of $(\text{DODA})_2[\text{Mo}_6\text{O}_{19}]$.