Electronic Supplementary Information

Quick and selective synthesis of Li₆[α -P₂W₁₈O₆₂]·28H₂O soluble in various organic solvents

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Figure S1. The ⁷Li-NMR spectrum of the compound measured with deoxidized 0.3 M lithium chloride methanol solution as reference. There is only singlet peak at 0.006 ppm which signifies presence of solvated lithium ion.



Figure S2. The ³¹P-NMR spectrum of the product material; $[\alpha-P_2W_{18}O_{62}]^{6-}$ (-13.1 ppm) and $[\beta-P_2W_{18}O_{62}]^{6-}$ (-11.6 and -12.3 ppm). The integration ratio for $[\alpha-P_2W_{18}O_{62}]^{6-}$ and $[\beta-P_2W_{18}O_{62}]^{6-}$ was approximately 30 : 1.



Figure S3. The IR spectrum (KBr) of $Li_6[\alpha - P_2W_{18}O_{62}] \cdot 28H_2O$.



Figure S4. TG curve of $Li_6[\alpha\mathchar`-P_2W_{18}O_{62}]\mathchar`-28H_2O.$

Table S1. The comparison of this work and previous total experimental steps, time, yield and purity for the synthesis of highly-pure α -Dawson type POM.

entry	date	author	counter	total experimental	total experimental	yield / %	α -isomer
			cation	steps	time / days		purity / %
1	1997	Droege/Randall/frinke et al. ^{S1}	K+	5	10	82	≥97
2	2004	Nadjo et al.	K*	4-5	8-12	85-93	97-99
3	2012	This report	Li+	3	1-2	87	≥97

Reference

S1. C. R. Graham, R. G. Finke, Inorg. Chem., 2008, 47, 3679.3679.