

Supporting Information

General Synthesis of Rare-Earth Orthochromites with Quasi-Hollow Nanostructures and Their Magnetic Properties

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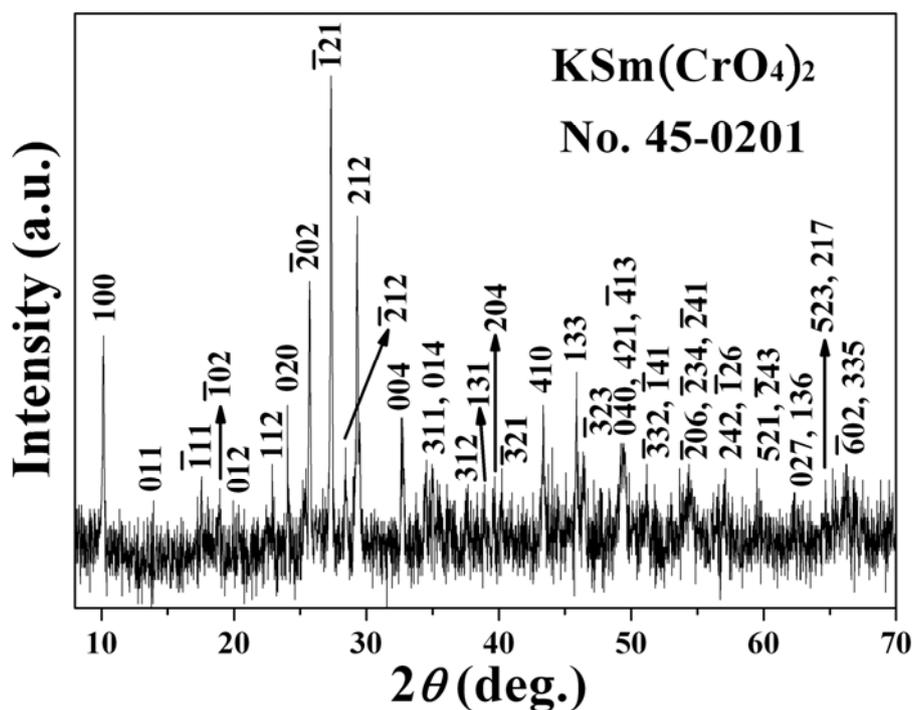


Figure S1. XRD patterns of the sample prepared by the direct reaction of K₂CrO₄ and Sm(NO₃)₃ aqueous solution at room temperature followed with annealing at 400 °C in nitrogen ambient.

All the diffraction peaks can be indexed to the monoclinic structure of KSm(CrO₄)₂ with lattice constants of $a = 8.682 \text{ \AA}$, $b = 7.351 \text{ \AA}$, $c = 10.873 \text{ \AA}$ and $\beta = 91.83^\circ$, which are in good agreement with the literature values (JCPDS Card File No. 45-0201, $a = 8.673 \text{ \AA}$, $b = 7.346 \text{ \AA}$, $c = 10.903 \text{ \AA}$ and $\beta = 91.89^\circ$). No characteristic diffraction peaks due to other impurities can be observed.

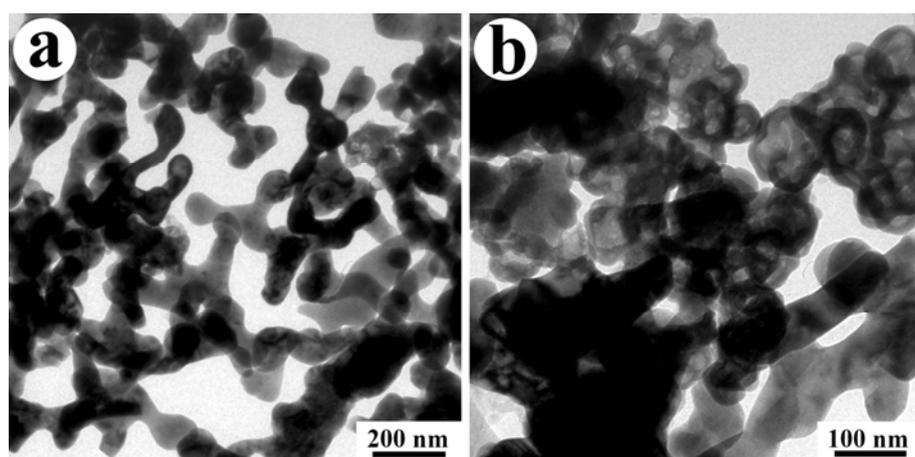


Figure S2. TEM images of the EuCrO_3 sample after annealing at 1200 °C in N_2 atmosphere.

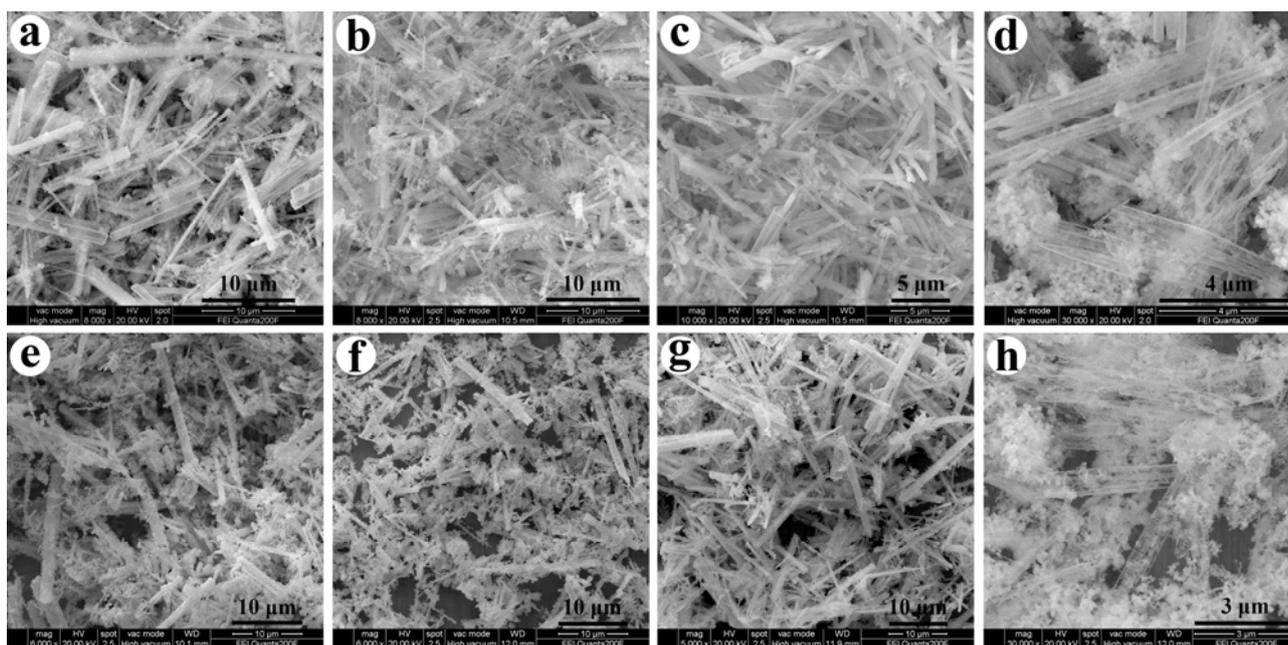


Figure S3. SEM images of the as-synthesized samples for (a) LaCrO₃, (b) PrCrO₃, (c) NdCrO₃ and (d) SmCrO₃ before annealing, and the corresponding 800 °C annealing samples for (e) LaCrO₃, (f) PrCrO₃, (g) NdCrO₃ and (h) SmCrO₃, respectively. Inset in S3e gives a close-up TEM image at high magnification of the annealing LaCrO₃ sample.

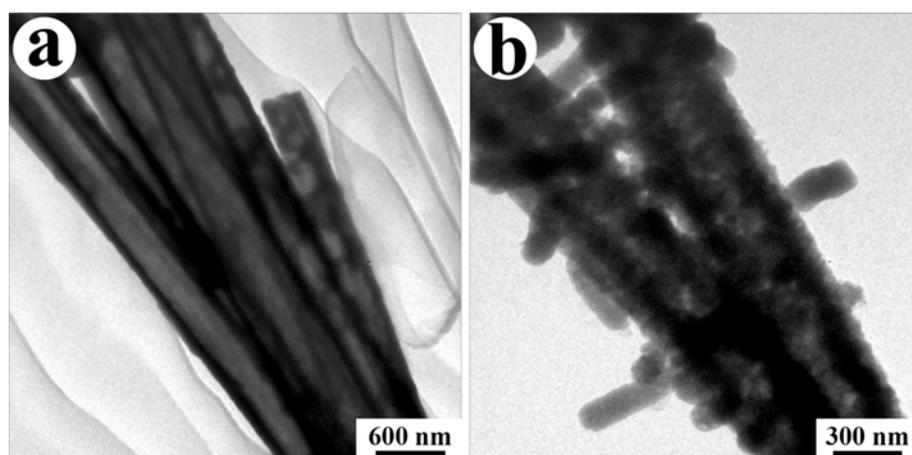


Figure S4. TEM images of the as-prepared NdCrO₃ samples (a) before and (b) after annealing.

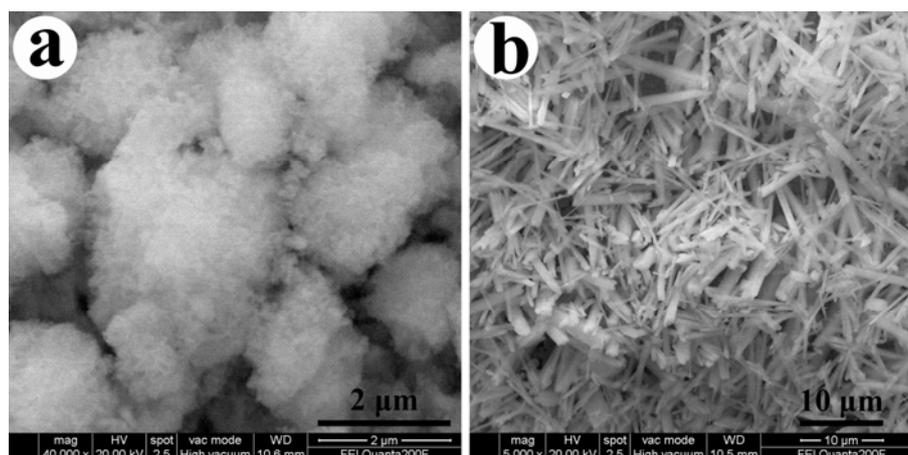


Figure S5. SEM images of the representative (a) $\text{KEu}(\text{CrO}_4)_2$ and (b) $\text{KNd}(\text{CrO}_4)_2$ precursors synthesized at room temperature by the direct precipitation between $\text{Eu}(\text{NO}_3)_3/\text{Nd}(\text{NO}_3)_3$ and K_2CrO_4 aqueous solution.

Table S1. Summary of the magnetic data based on the temperature dependence of the magnetization for all the as-prepared rare-earth orthochromites samples via a general solvothermal *in-situ* reduction route

RCrO ₃	T_{N1} (K)	$ \Theta $ (K)	C	Experimental $\mu_{\text{eff}} (\mu_B)^a$	Theoretical $\mu_{\text{eff}} (\mu_B)^b$	$\mu_{R^{3+}} (\mu_B)^c$	$\mu_{Cr^{3+}} (\mu_B)^d$
LaCrO ₃	288	947	3.11	5.00	3.87	0	3.87
PrCrO ₃	239	316	5.99	6.95	5.27	3.58	3.87
NdCrO ₃	224	324	4.68	6.14	5.30	3.62	3.87
SmCrO ₃	192	502	3.89	5.60	3.96	0.84	3.87
EuCrO ₃	181	1139	22.46	13.46	3.87	0	3.87
GdCrO ₃	167	37	8.16	8.11	8.83	7.94	3.87
DyCrO ₃	145	30	14.19	10.70	11.31	10.63	3.87
HoCrO ₃	140	25	14.10	10.67	11.28	10.60	3.87
YCrO ₃	140	327	2.35	4.35	3.87	0	3.87
ErCrO ₃	134	106	20.83	12.96	10.34	9.59	3.87
TmCrO ₃	124	45	8.06	8.06	8.50	7.57	3.87
YbCrO ₃	117	145	4.67	6.14	5.97	4.54	3.87
LuCrO ₃	110	369	3.00	4.92	3.87	0	3.87

^aThe effective magnetic moment (μ_{eff}) in units of the Bohr magneton (μ_B) is evaluated using the formula: $\mu_{\text{eff}} = (3k_B C/N_A)^{1/2} = 2.84 \cdot C^{1/2}$, where k_B is the Boltzmann constant, C is the Curie constant and N_A is Avogadro's number. ^bThe theoretical value of μ_{eff} is obtained by $[(\mu_{R^{3+}})^2 + (\mu_{Cr^{3+}})^2]^{1/2}$. ^c $\mu_{R^{3+}}$ can be calculated by $g[J(J+1)]^{1/2}$. ^d $\mu_{Cr^{3+}}$ can be calculated by $2[S(S+1)]^{1/2}$, where $S = 3/2$.