

## Supporting Information

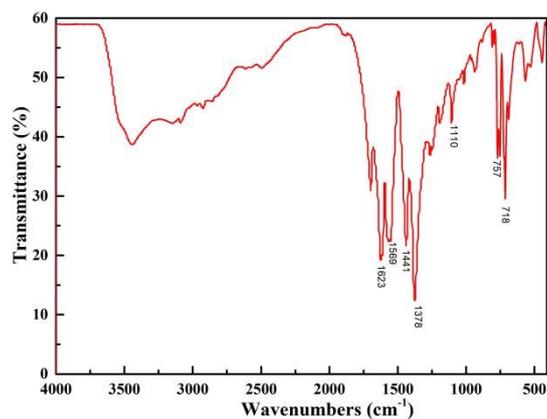
**Title:** Gas-Solid two-phase Flow (GSF) Mechanochemical Synthesis Dual-Metal-Organic Frameworks and Electrochemical Properties Research

**Author(s):** Jun Zhao, Bo Jin, Rufang Peng

**number of pages:** 5

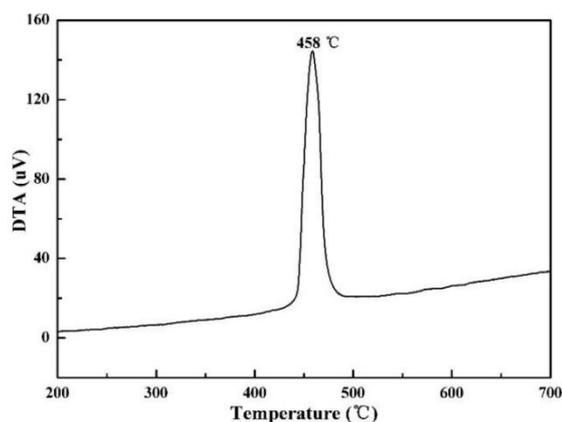
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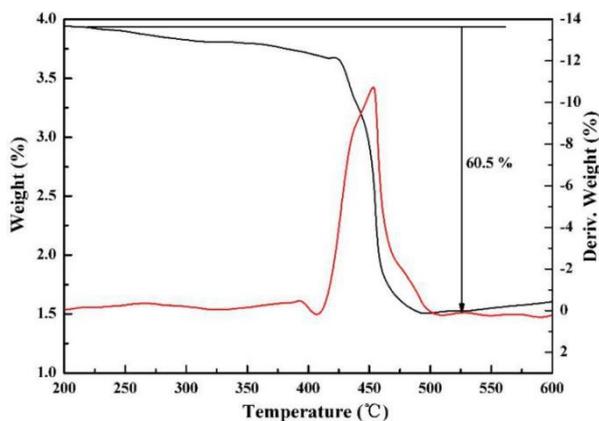


**Fig. S1** Fourier transform infrared spectroscopy (FT-IR) spectra of  $\text{CoMn}_2(\text{BTC})_2$

It can be seen from **Fig. S1** that the FT-IR spectra of  $\text{CoMn}_2(\text{BTC})_2$  synthesized by GSF was identical with the previously reported.<sup>1</sup> The asymmetric stretching vibrations and symmetric stretching vibrations of  $-\text{COO}-$  appeared in the regions  $1623\text{--}1569\text{ cm}^{-1}$  and  $1441\text{--}1378\text{ cm}^{-1}$ , respectively. The absorption at  $757\text{ cm}^{-1}$ ,  $1110\text{ cm}^{-1}$  were C-O-Mn and C-O-Co stretching vibration peaks, which suggested that the metal ions have been coordinated with the BTC successfully.

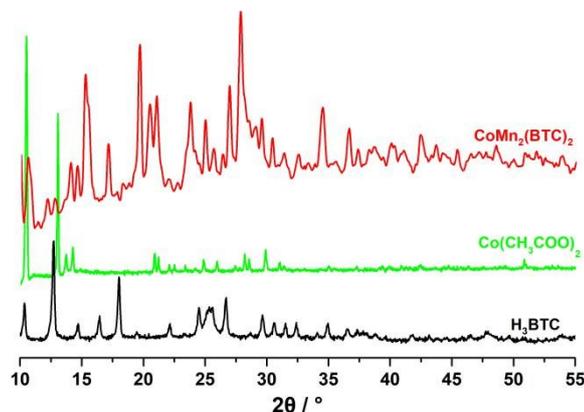


**Fig. S2** DTA of  $\text{CoMn}_2(\text{BTC})_2$



**Fig. S3** TG/DTG of  $\text{CoMn}_2(\text{BTC})_2$

In order to further verify the synthesized  $\text{CoMn}_2(\text{BTC})_2$ , DTA analysis was shown in **Fig. S2-S3**. According to the DTA, TG/DTG curves, the initial decomposition peak was about at  $458^\circ\text{C}$  and  $\text{CoMn}_2(\text{BTC})_2$  showed good thermal stability. The above information was completely consistent with the literature reports.<sup>1</sup>



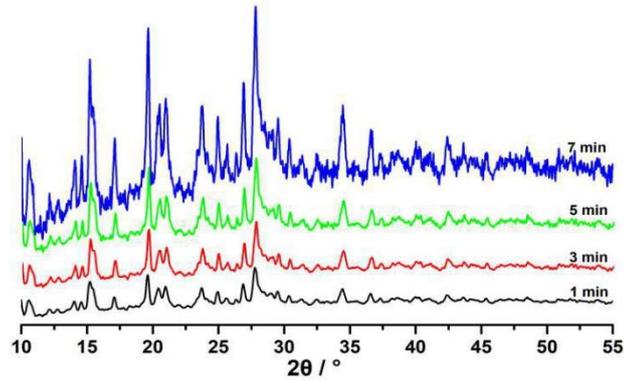
**Fig. S4** XRD comparison of  $\text{H}_3\text{BTC}$ ,  $\text{Co}(\text{CH}_3\text{COO})_2$ ,  $\text{CoMn}_2(\text{BTC})_2$

As shown in **Fig S4**, we also added the XRD comparison of  $\text{H}_3\text{BTC}$ ,  $\text{Co}(\text{CH}_3\text{COO})_2$ ,  $\text{CoMn}_2(\text{BTC})_2$ . It further explained that the raw materials ( $\text{H}_3\text{BTC}$ ,  $\text{Co}(\text{CH}_3\text{COO})_2$ ) reacted in the GSF and generated  $\text{CoMn}_2(\text{BTC})_2$ .



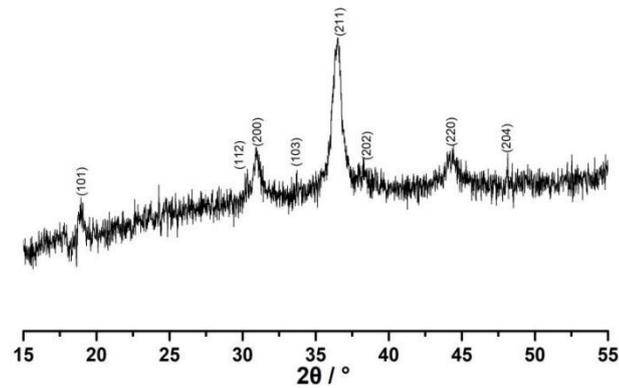
**Fig. S5** GSF reaction equipment with key parts highlighted.

The reaction process flow chart of GSF chemical synthesis was shown in **Fig S5**. Reactant particles were transferred into the impacting chamber through the feed port, firstly. Afterward, the particles were accelerated to high velocities (300 m/s) by compressed air (1.5 MPa).



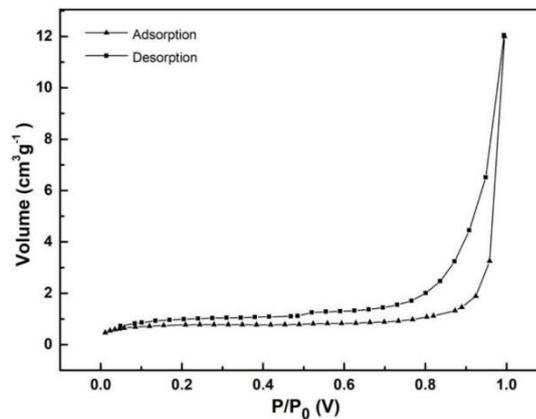
**Fig. S6** XRD of the product collected at different times

The XRD of  $\text{CoMn}_2(\text{BTC})_2$  at different times was shown in **Fig S6**. The characteristic diffraction peaks of the raw material completely disappeared after collision reaction 10 min. Meanwhile, new diffraction peaks assigned to  $\text{CoMn}_2(\text{BTC})_2$  appeared (**Fig S5**), which basically consistent with the characteristic diffraction peak reported in the related literature.<sup>1</sup>



**Fig. S7** XRD of the product collected after the TGA

As shown in **Fig S7**, The XRD results showed that the products after TGA still existed mainly in the form of  $\text{CoMn}_2\text{O}_4$ .



**Fig. S8**  $\text{N}_2$  adsorption/desorption isotherms of  $\text{CoMn}_2(\text{BTC})_2$

As shown in **Fig S8**, Nitrogen adsorption/desorption isotherms were conducted at 77 K to study the textural properties of the  $\text{CoMn}_2(\text{BTC})_2$  synthesized by GSF. The BET surfaces areas obtained from the nitrogen adsorption-desorption isotherms were  $2.45 \text{ m}^2/\text{g}$ .

## References

(S1) Hu, X. S.; Li, C.; Lou, X. B.; Yan, X. J.; Ning, Y. Q.; Chen, Q.; Hu, B. W. Controlled synthesis of  $\text{Co}_x\text{Mn}_{3-x}\text{O}_4$  nanoparticles with a tunable composition and size for high performance lithium-ion batteries. *RSC Advances*, **2016**, *6*(59), 54270-54276.