

Supporting Information for

Calcium-Doped Boron Nitride Aerogel Enables Infra-Red Stealth at High Temperature Up to 1300 °C

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Supplementary Figures

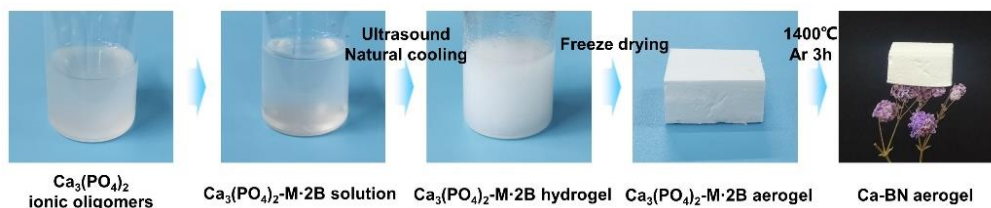


Fig. S1 Photographs of the synthesis of Ca-BN aerogel

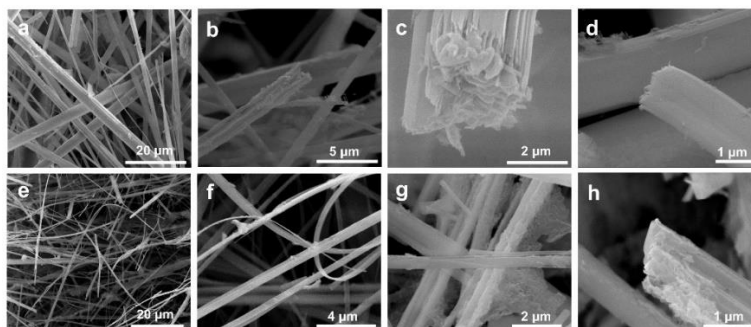


Fig. S2 SEM images of $\text{Ca}_3(\text{PO}_4)_2$ -M·2B aerogel

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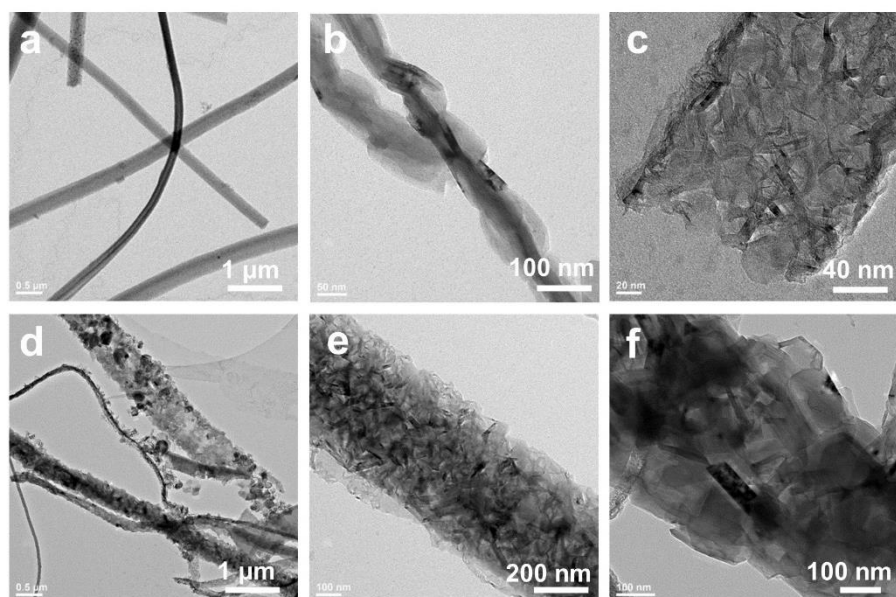


Fig. S3 TEM images of Ca-BN aerogel

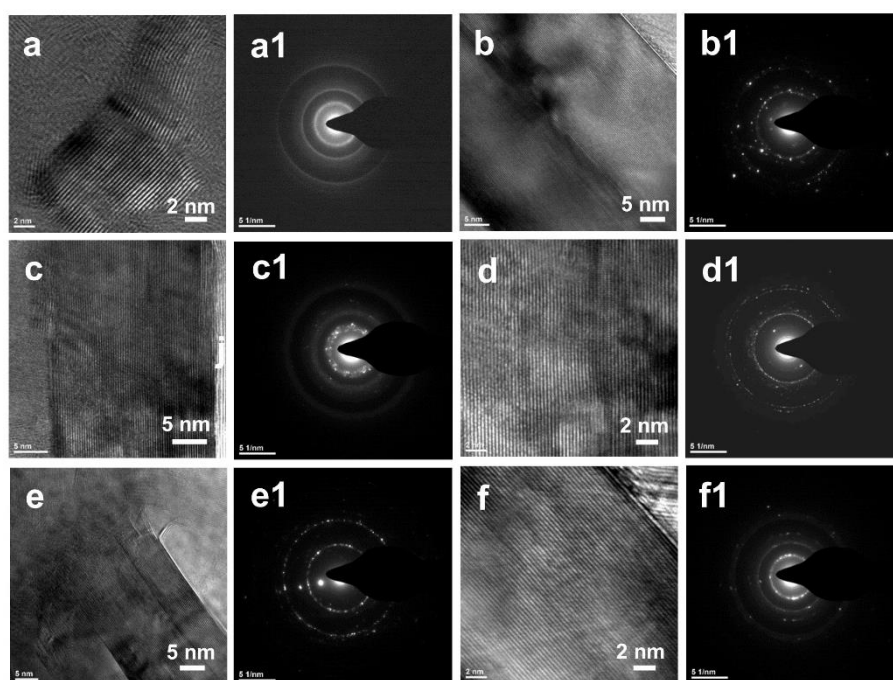


Fig. S4 HRTEM images and the corresponding selected area electron diffraction patterns of the Ca-BN aerogel

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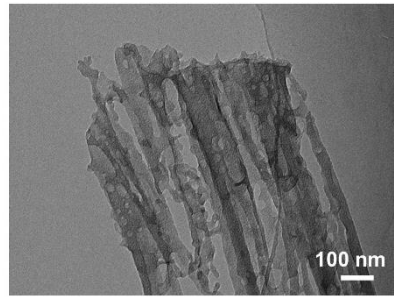


Fig. S5 TEM image of BN aerogel

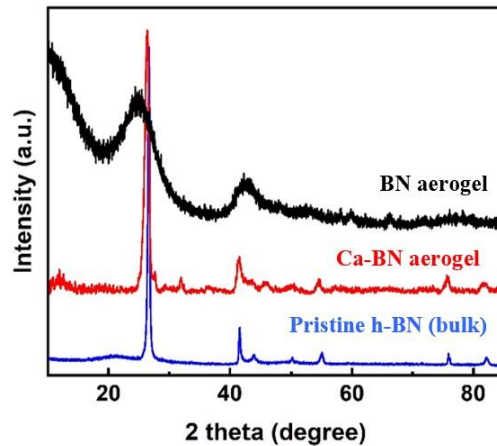


Fig. S6 XRD spectra of Ca-BN aerogel (red), bulk h-BN (blue) and BN aerogel (black). For Ca-BN aerogel, the (002) (peak at $2\theta=26.36^\circ$) interplanar spacing was about 0.338 nm, which is slightly larger than that of bulk h-BN (0.333 nm, $2\theta=26.74^\circ$), but is smaller than that of BN aerogel derived from M·2B (0.36 nm, $2\theta=24.64^\circ$)

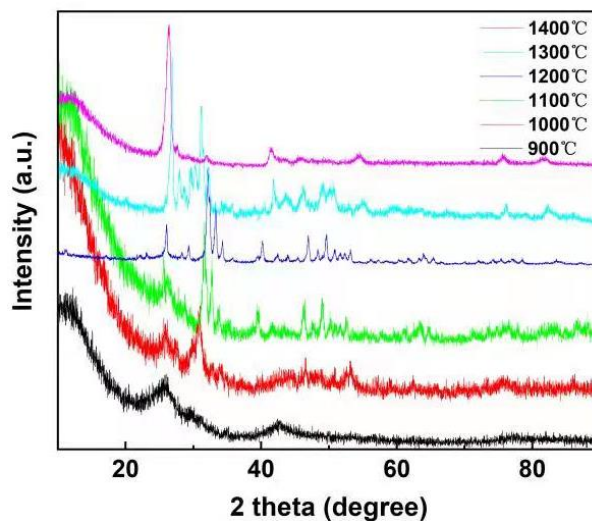


Fig. S7 XRD spectra of Ca-BN aerogels derived at different temperature. During the synthetic process, crystal structure of BN changes from amorphous to crystalline as temperatures increase from 900 °C

to 1400 °C. When synthetic temperature changes from 1000 °C to 1300 °C, other than diffraction peaks of BN, some diffraction peaks of $\text{Ca}_3(\text{PO}_4)_2$ still were observed in the XRD spectra. When synthetic temperature reaches to 1400 °C, nearly no obvious diffraction peaks of $\text{Ca}_3(\text{PO}_4)_2$ could be observed, mainly because $\text{Ca}_3(\text{PO}_4)_2$ occurred carbothermal reduction reaction.

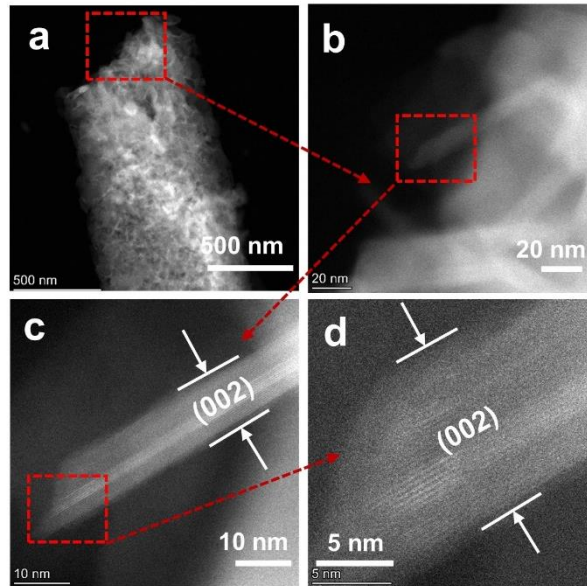


Fig. S8 STEM images of single ribbon of Ca-BN aerogel

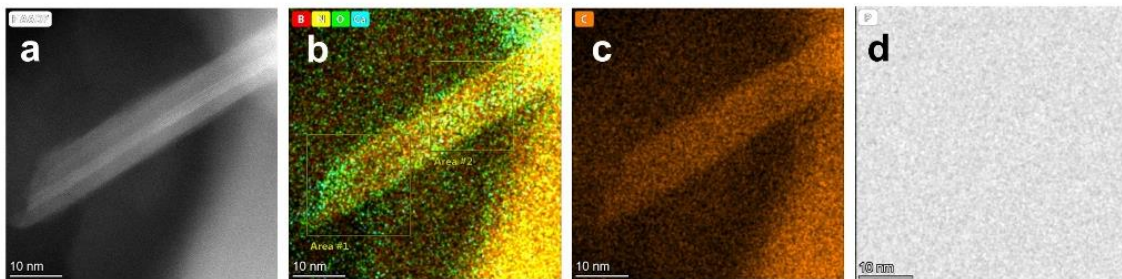


Fig. S9 (a) STEM image, (b) EDS overlaid mapping of selected elements (B, N, O, Ca), (c-d) EDS mapping of C, P of the resulting Ca-BN aerogel in the length of 10-20 nm

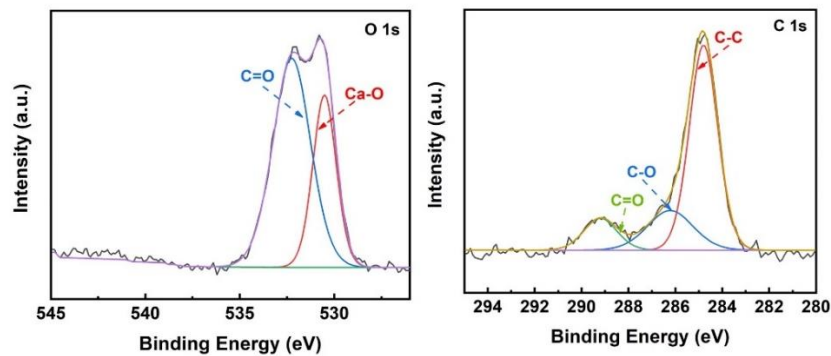


Fig. S10 High-resolution XPS spectra of O 1s and C 1s

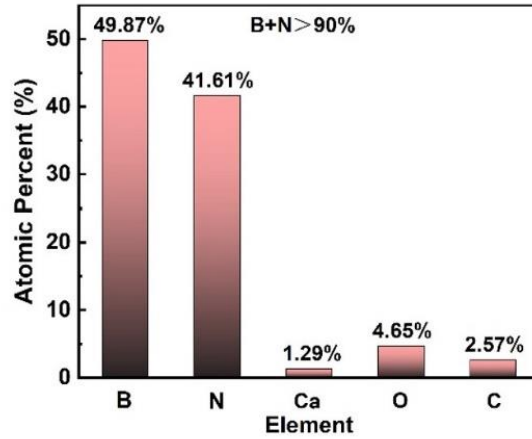


Fig. S11 Atomic ratio of B, N, Ca, O, C from XPS survey. Atomic ratio of B and N is close to 1:1, and total atomic ratio of B and N was about 92.4%, atom ratio of Ca was about 1.29%

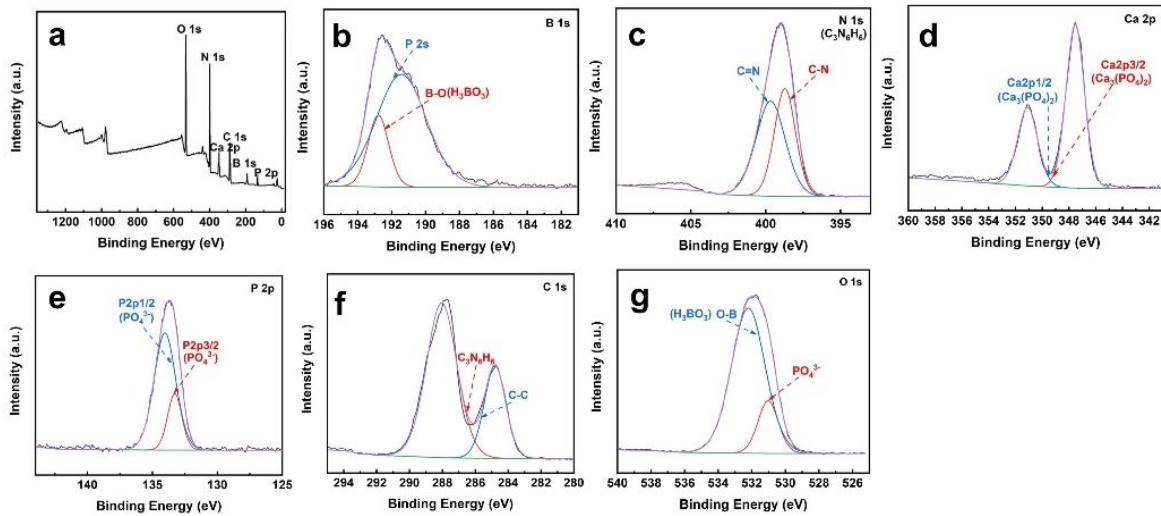


Fig. S12 (a) The XPS survey of $Ca_3(PO_4)_2$ -M·2B aerogel, (b-g) High-resolution XPS spectra of B1s, N1s, Ca 2p, P 2p, C1s, O1s

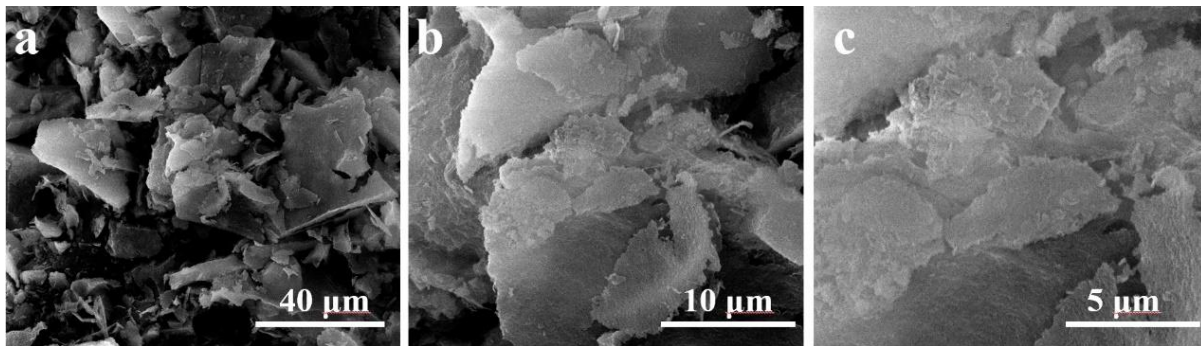


Fig. S13 SEM images of calcium phosphate solid fabricated from calcium phosphate ionic oligomer solution

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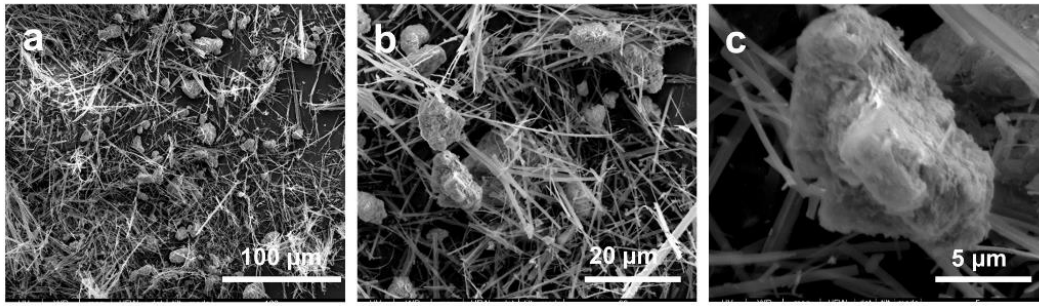


Fig. S14 SEM images of commercial $\text{Ca}_3(\text{PO}_4)_2$ powder-M·2B aerogel

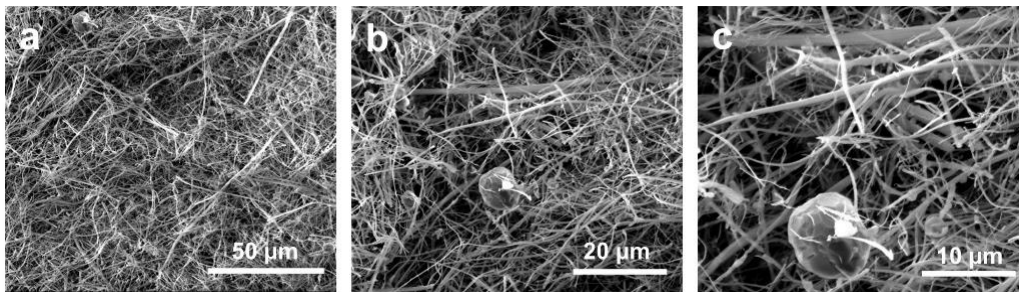


Fig. S15 SEM images of commercial $\text{Ca}_3(\text{PO}_4)_2$ powder directed Ca-BN aerogel

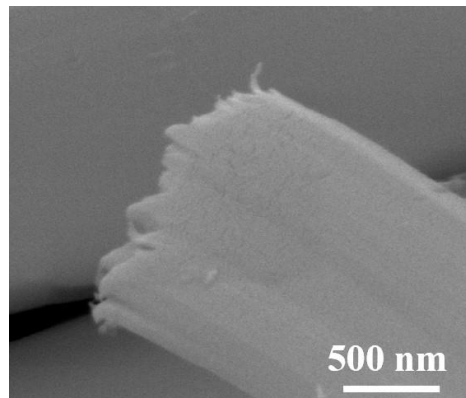


Fig. S16 SEM image of calcium phosphate doped M·2B ribbon

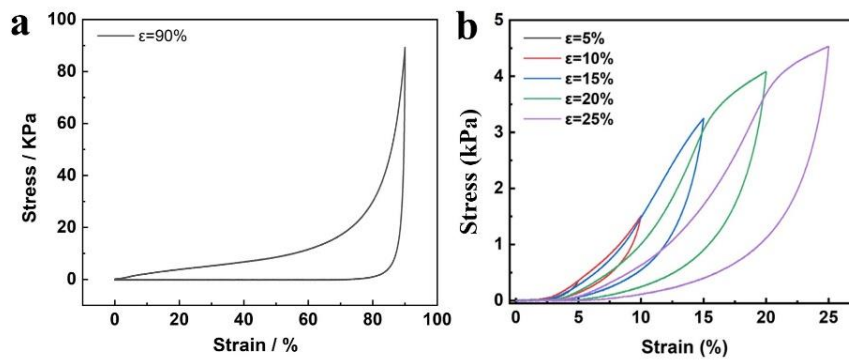


Fig. S17 Strain-stress curves of Ca-BN aerogel



Fig. S18 Photograph of infrared thermometer (ISR 6 Advanced, LumaSense IMPAC)

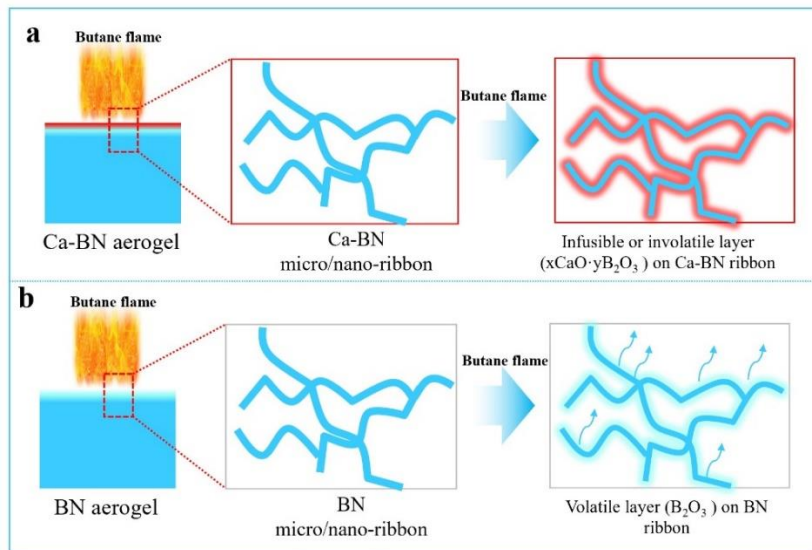


Fig. S19 Illustration of structural evolution of Ca-BN aerogel and BN aerogel when they heated by butane flame. For Ca-BN aerogel, Ca element could react to melted B_2O_3 and then form an involatile melted compound (e.g., $x\text{CaO}\cdot y\text{B}_2\text{O}_3$ system), the melted compound covered on the skeleton surface of the aerogel and prevented the further oxidization, pure BN aerogel was oxidized into B_2O_3 and then evaporated rapidly.

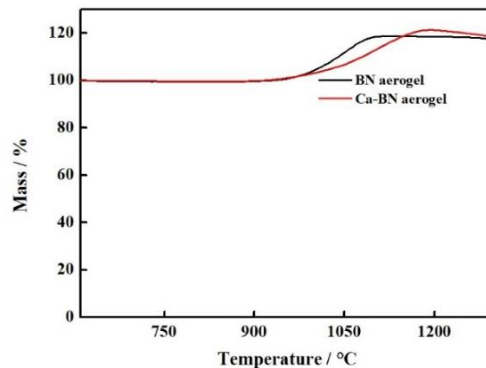


Fig. S20 TG curves of Ca-BN aerogel and BN aerogel in N_2/O_2 atmosphere. Compared to BN aerogel, Ca-BN aerogel exhibited a higher oxidizing temperature.

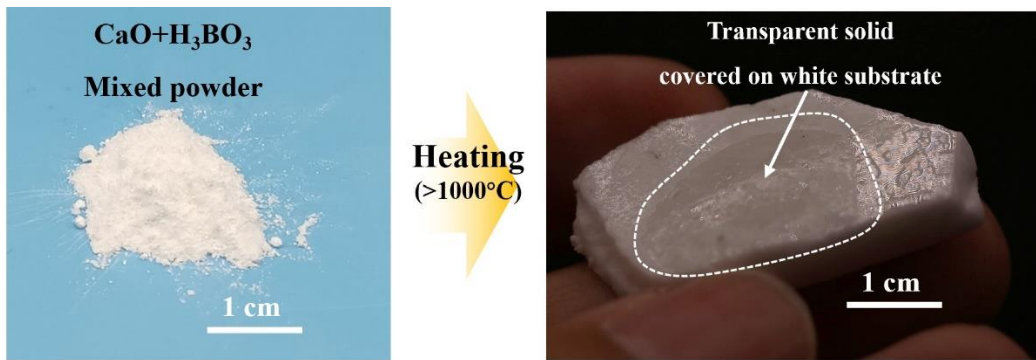


Fig. S21 Photograph of CaO and boric acid and their relevant product after high-temperature treatment

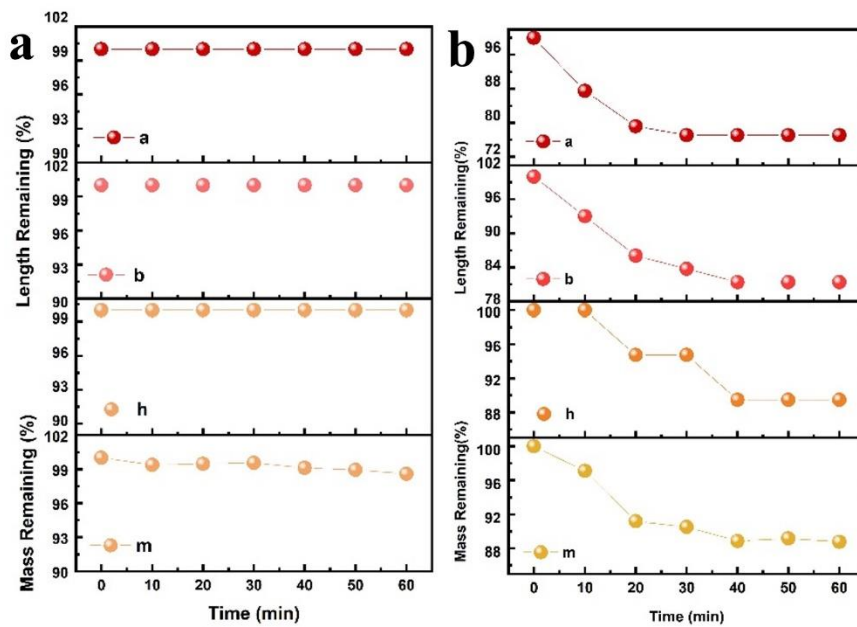


Fig. S22 Statistic diagram of the change rate of three side lengths and mass of Ca-BN (a) and BN aerogels (b)

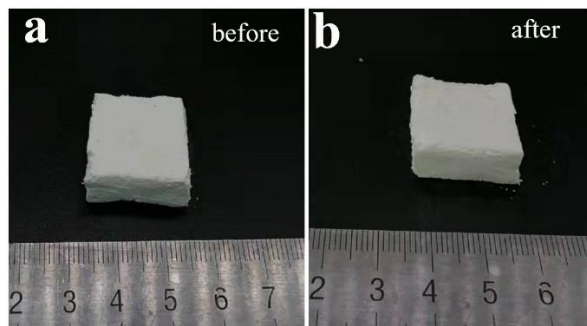


Fig. S23 Surface images of Ca-BN aerogels before (a) and after (b) being heated by butane flame

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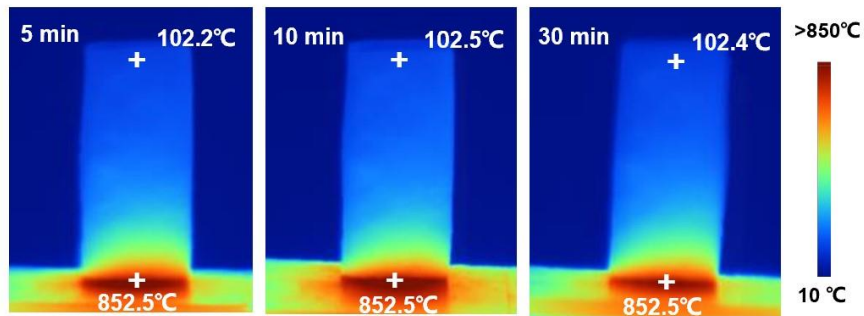


Fig. S24 Infrared images of Ca-BN aerogel on a hot Al_2O_3 plate for 30 min

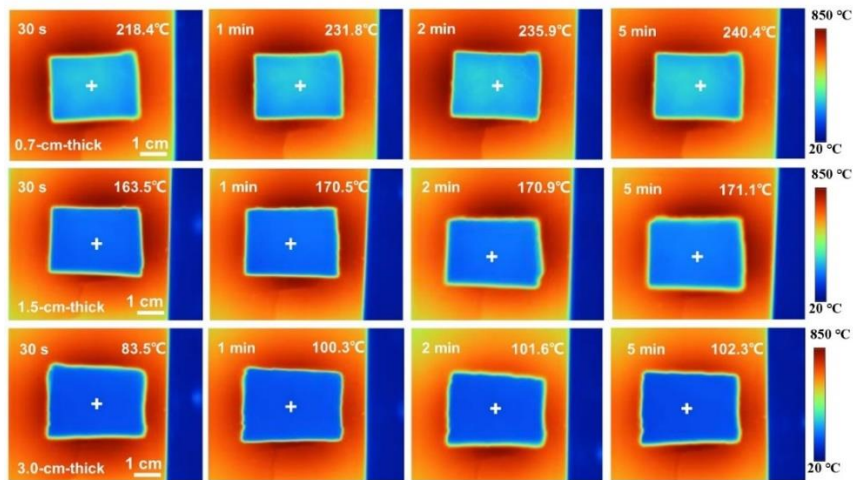


Fig. S25 Infrared images of hot target covered by Ca-BN aerogels with different thickness

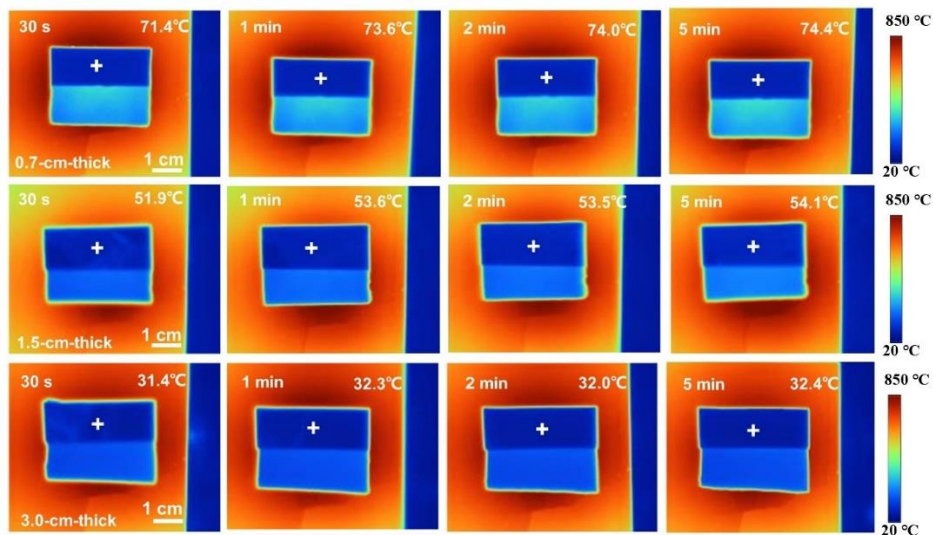


Fig. S26 Infrared images of hot target covered by Ca-BN aerogels (different thickness) together with Al foil

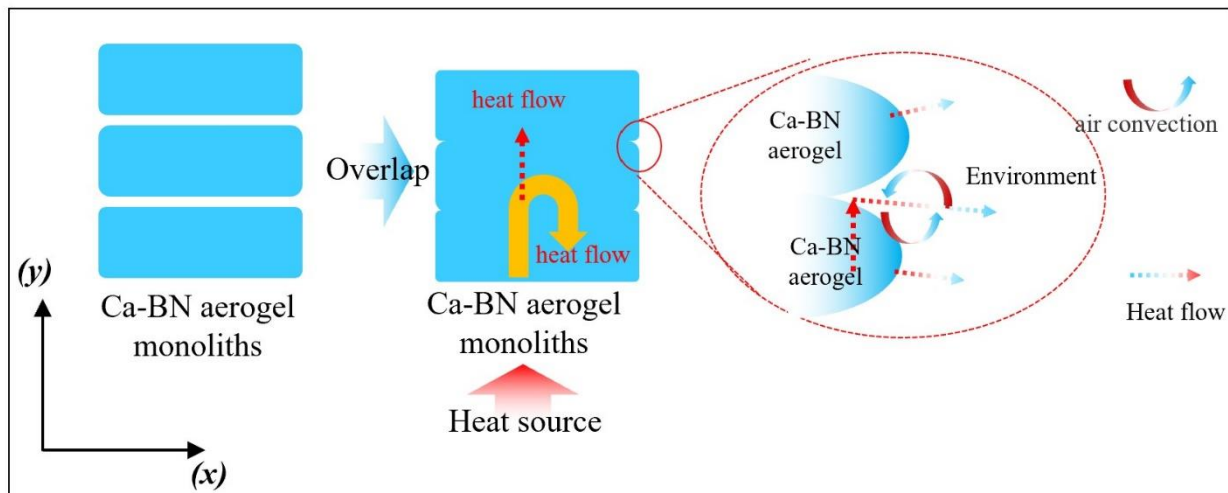


Fig. S27 Illustration of three layers of Ca-BN aerogels. Ca-BN aerogel monoliths were physically stacked together to restrain heat transport in vertical direction (oy), aerogel surfaces could touch with each other tightly excepted to their edge positions. At their edge positions, overlapping may bring small gaps because the surface of Ca-BN aerogels cannot be completely flat. Such gaps connected to surrounding environment, which could provide more locations for heat loss from Ca-BN aerogel to environment via air convection in horizontal direction (ox). In other words, heat transport along vertical direction (oy) was restrained.

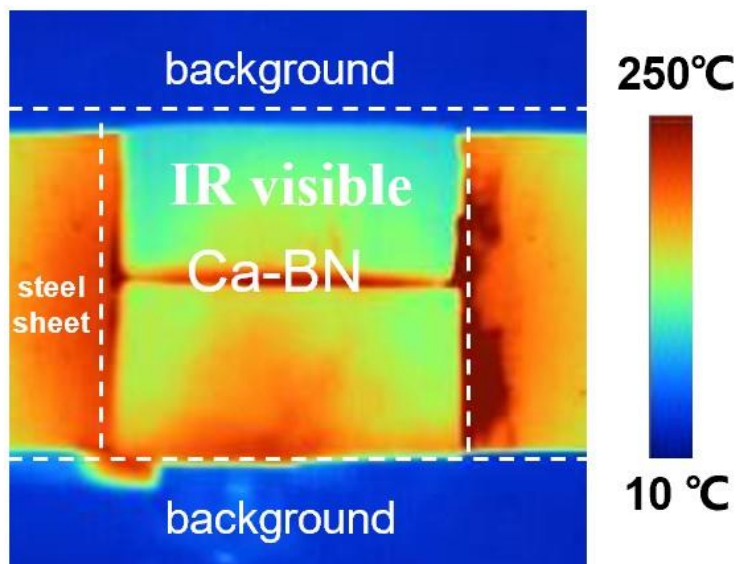


Fig. S28 Infrared image of hot target covered by Ca-BN aerogel