Exceptional Superionic Conductivity in Disordered Sodium Decahydro-closo-decaborate


Recently, complex hydride salts undergoing solid-state, entropy-driven, order–disorder transitions have been shown to exhibit impressive fast-ion conduction properties as a result of the appearance of vacancy-rich cation sublattices within networks of highly mobile, reorientationally disordered polyanions.[1] Initial interest has mainly focused on the light-metal Li and Na salts possessing tetrahydroborate (BH₄⁻) polyanions (see Figure 1), such as LiBH₄ and Na₂BH₄NH₂, as well as related derivative materials.[2–5] Very recently, it was discovered that Li and Na salts possessing the larger icosaedral dodecahydro-closo-dodecaborate (B₁₂H₁₀²⁻) anions (see Figure 1) also undergo order–disorder phase transitions,[6,7] with Na₂B₁₂H₁₂ exhibiting disorder-induced superionic conductivity (approaching 0.1 S cm⁻¹) above around 480 K.[8] This conductivity rivals that of traditional ceramic materials, Naβ'-alumina solid electrolyte (BASE), and Na superionic conductor (NASICON), as well as more recent Na₄PS₄-based glass ceramic electrolytes, all currently of considerable interest for use in Na-ion batteries.[9–11] The pronounced superionicity and relatively low conduction barrier for disordered Na₂B₁₂H₁₂ are probably due, in part, to the larger size (and roughly spherical shape) of the B₁₂H₁₀²⁻ anions compared with the substantially smaller BH₄⁻ anions (see Figure 1) present in other investigated fast-ion conductors.[12] These types of large polyanion compounds represent a potentially fertile area for discovering new materials with superionic conductivities, but with lower transition temperatures. Here we report our finding that sodium decahydro-closo-decaborate (Na₂B₁₀H₁₀), a related sodium salt containing large, ellipsoidal-shaped, B₁₀H₁₀²⁻ anions (see Figure 1), forms a disordered, face-centered-cubic (fcc) phase above ca. 360 K, possessing a vacancy-rich Na⁺ cation sublattice. This cation sublattice is highly mobile within the spacious corridors formed by the large B₁₀H₁₀²⁻ anions and exhibits remarkable superionic conductivity (e.g., σ ≈ 0.01 S cm⁻¹ at 383 K) to substantially lower temperatures than for Na₂B₁₂H₁₂. This conductivity is more than an order of magnitude higher than that of all other solid-state Na-based complex-hydrde materials investigated to date in this temperature region.[9] As such, this discovery represents a major advancement in the field of solid-state Na⁺ fast-ion conduction at technologically relevant device temperatures.

Figure 2 shows differential scanning calorimetry (DSC) results for Na₂B₁₀H₁₀ after several heating/cooling cycles. There is a clear reversible transition, first observed by Bonnet et al.,[12] with minor hysteresis. After a number of cycles to 500 K, the hysteresis decreases and the onset temperature stabilizes near 360 K upon both heating and cooling. A slow attenuation of the DSC features begins to occur upon repeated cycling to temperatures near 600 K.

Figure 3 shows the neutron powder diffraction (NPD) result for a partially deuterated Na₃⁴₁⁴¹B₁₀H₁₀ sample at 410 K above the phase transition. Table S1 in the Supporting Information lists the corresponding structural parameters. The Rietveld-refined model confirms transformation from the known, low-T, ordered monochinic structure[13] (not shown) to a high-T disordered structure with Na⁺ cations partially occupying a variety of interstitial sites within an fcc lattice of orientationally disordered anions. Neutron-scattering Fourier difference maps initially suggested that the broad distribution of cation positions and the inclusion of the three most intense positions were ultimately necessary to attain a good model fit to the data. Furthermore, the refinements suggested extensive anion orientational disorder, which could be represented in various ways via multiple B and H positions. The model reflects the simplest
representation of six approximately superimposed anion orientations, each equally probable. In particular, the six possible anion orientations are aligned in pairs with their long axes oriented along any one of the three orthogonal crystallographic axes. The members of each pair are azimuthally offset from each other by 45° about their long axes. Each B and H position in the structure is 1/3 occupied, being shared by two of the six possible orientations.

Anion dynamical behavior was probed by neutron elastic-scattering fixed-window scans (FWSs) of Na$_2$B$_{10}$H$_{10}$. The results in Figure 4 suggest that a dramatic change in B$_{10}$H$_{10}^{2-}$ anion reorientational mobility occurs upon phase transformation. In particular, the high elastic neutron counts in the low-T ordered phase suggest anion reorientational jump frequencies less than 10$^{8}$ s$^{-1}$, whereas the roughly 80% lower counts in the high-T disordered phase suggest an orders-of-magnitude enhancement in jump frequencies to greater than 10$^{10}$ s$^{-1}$. This is reminiscent of the FWS behavior observed for Na$_2$B$_{12}$H$_{12}$.[7,15] Indeed, the inset in Figure 4 showing a quasielastic neutron scattering (QENS) spectrum for the disordered phase at 375 K indicates a quasielastic component with a Lorentzian linewidth of about 41(1) µeV FWHM, which reflects a jump correlation frequency on the order of 3 × 10$^{10}$ s$^{-1}$. The ratio of elastic and total scattering intensities (which is also consistent with the ratio of FWS neutron counts in the disordered and ordered phases from Figure 4) suggests that, besides the B$_{10}$H$_{10}^{2-}$ reorientational jumps around the long axis, two-fold anion flips leading to exchanges of apical H atom positions are also occurring.

We also probed the Na$^+$ dynamical behavior in Na$_2$B$_{10}$H$_{10}$ by $^{23}$Na NMR measurements. Figure 5 shows the $^{23}$Na spin-lattice relaxation rate $R_1$ at the resonance frequency of $\omega/2\pi = 23$ MHz as a function of $T^{-1}$. The general features of the behavior of $R_1$ for Na$_2$B$_{10}$H$_{10}$ resemble those for Na$_2$B$_{12}$H$_{12}$ at the phase transition.[16] Here, $R_1$ exhibits a jump accompanied by a change in sign of its temperature dependence. Such behavior indicates that the transition from the ordered to

---

**Figure 1.** Geometries and approximate relative sizes of BH$_4^-$, B$_{10}$H$_{10}^{2-}$, and B$_{12}$H$_{12}^{2-}$ anions, each shown from top and side views. The boron and hydrogen atoms are denoted by the green and pink spheres, respectively.

**Figure 2.** Characteristic DSC measurements (2 K min$^{-1}$) for Na$_2$B$_{10}$H$_{10}$ after several heating and cooling cycles up to 410 K.

**Figure 3.** Experimental (circles), fitted (line), and difference (lower line) NPD profiles for Na$_2$B$_{10}(H_{0.73}D_{0.27})_{10}$ at 410K ($\lambda = 2.077$ Å; Fm-3m; $a = 9.8426(8)$ Å; $R_{wp} = 0.0123$; $R_p = 0.0104$; $\chi^2 = 0.817$). The vertical bars indicate the calculated positions of the Bragg peaks. Inset: Disordered structure. H atoms are omitted for clarity. B atoms are denoted by the green spheres. Anions are centered at 4a (0 0 0). Different Na$^+$ cation positions are denoted by the red [tetrahedral (t) sites at 8c (1/4 1/4 1/4)], yellow [octahedral (o) sites at 4b (1/2 0 0)], and blue [intermediate (i) sites at 24d (1/4 1/4 0) between two t sites] spheres. The sphere sizes are proportional to the partial occupancies of 0.58(1), 0.28(3), and 0.09(1) for the t, o, and i sites, respectively.

**Figure 4.** Neutron Counts (10$^7$) as a function of $2\theta$ for B$_{10}$H$_{10}^{2-}$ anions. The inset shows a quasielastic neutron scattering (QENS) spectrum for the disordered phase at 375 K.
the disordered phase is accompanied by an abrupt increase in the Na⁺ jump rate $\tau_+^{-1}$. The expected $R_1(T)$ maximum is ‘folded’; i.e., because of the abrupt increase in $\tau_+^{-1}$ at the phase transition, there is a jump directly from the low-$T$ slope to the high-$T$ slope of the $R_1(T)$ peak. On the low-$T$ slope, $R_1$ is proportional to $\tau_+^{-1}$; on the high-$T$ slope, $R_1$ is proportional to $\tau_+$. From the two slopes, we obtain activation energies for Na⁺ jumps in the ordered and disordered phases of 750(20) meV and 190(10) meV, respectively. Although we cannot reliably determine the absolute values of $\tau_+^{-1}$ due to the ‘folded’ nature of the $R_1(T)$ peak, the data do allow us to conclude that the Na⁺ jump rate exceeds $\omega \approx 1.5 \times 10^4$ s⁻¹ just above the phase transition. Moreover, the very small $^{23}$Na NMR linewidth (0.2 kHz full-width half-maximum (FWHM)) observed in the disordered phase confirms that the Na⁺ cations are undergoing long-range diffusion.

To characterize the Na$_2$B$_{10}$H$_{10}$ conduction behavior, we carried out AC impedance measurements between 295 K and 423 K using a pressed disk of polycrystalline Na$_2$B$_{10}$H$_{10}$ powder with either gold or (similarly performing) molybdenum foil contacts. Cross-sectional SEM images of the pelletized sample in either gold or (similarly performing) molybdenum foil contacts. Figure S1 of the Supporting Information confirms that intimate contacts among the particles were achieved. Na$_2$B$_{10}$H$_{10}$ can be easily pelletized without further sintering, as has been reported for LiBH$_4$ and other complex hydrides. The results are shown in Figure 6 for the gold contacts. The inset shows typical complex impedance plots at various temperatures. They consist of an arc in the high-frequency region and a spike in the low-frequency region due to contributions from the bulk/ grain boundaries and the electrode, respectively. The results suggest that Na$_2$B$_{10}$H$_{10}$, similar to Na$_2$B$_{12}$H$_{12}$, behaves like a typical ionic conductor. The temperature dependence of the conductivity indicates dramatic superionic conductivity above the hysteretic order–disorder transition near 373 K, rising two orders of magnitude higher than that in the low-$T$ phase. This behavior is superior to that of other investigated complex hydride materials. Indeed, the conductivity exhibits a value of about 0.01 S cm⁻¹ at 383 K, which is about 25× greater than that of NaBH$_4$ (4 × 10⁻⁴ S cm⁻¹). A conductivity above 0.1 S cm⁻¹ at 500 K is suggested by extrapolation of the lower-$T$ data. The activation energy for conduction is evaluated to be 0.47 eV, which is higher than that reported for Na$_2$B$_{12}$H$_{12}$ (0.21 eV) but lower than those reported for NaBH$_4$ (0.61 eV) and the high-$T$ phase of LiBH$_4$ (0.53 eV). Nevertheless, any rationalization of the differences based solely on anion size is complicated by the substantial differences in structure and conduction pathways.

It should be noted that the NMR-derived activation energies discussed above reflect the average microscopic barriers for all cation diffusional jumps between neighboring sites within the Na$_2$B$_{10}$H$_{10}$ lattice, some of which may have little effect on the macroscopic conductivity barrier. In contrast, the latter barrier is more reflective of an overall rate-limiting step, such as a particular type of cation jump within the material required to
maintain conduction pathways or cation transport, e.g., across grain boundaries (although such grain-boundary bottlenecks are believed to be small in the present system).

The superionic conductivity of disordered Na$_2$B$_{10}$H$_{10}$ is consistent with the relatively small activation energy for Na$^+$ diffusion within the liquid-like cation sublattice. Again, similar to disordered Na$_2$B$_{12}$H$_{12}$, the overly large size and spheroidal shape of the polyanions result in less restrictive interstitial pathways and, hence, reduced Na$^+$ diffusional bottlenecks between the various cation sites within the close-packed anion sublattice.

As for the other disordered complex hydrides, the reorientationally mobile anions associated with superionic Na$_2$B$_{10}$H$_{10}$ may also lower the cation diffusional barrier by providing a dynamically cooperative environment for cation jumps within the voids of the anion sublattice. Indeed, at least an order-of-magnitude-higher anion reorientational jump rate compared with the Na$^+$ diffusion jump rate provides a dynamic environment where the anions can behave as ‘lubricants’ for cation diffusive motions.

A comparison of the relative sizes of the B$_{10}$H$_{10}^2$ and B$_{12}$H$_{12}^2$ anions in Figure 1 indicates a similar maximum dimension for each anion. In fact, the lattice constants for the disordered fcc Na$_2$B$_{10}$H$_{10}$ and body-centered cubic (bcc) Na$_2$B$_{12}$H$_{12}$ structures[7] indicate that both disordered anions possess similar spherical packing radii of $\approx$3.5 Å. This makes the small 190 meV activation energy for Na$^+$ diffusion in Na$_2$B$_{12}$H$_{12}$ particularly noteworthy, since it is less than half that of Na$_2$B$_{12}$H$_{12}$ (410 meV).[16] Such a difference may be the result of the different natures of the diffusion saddle points inherent within the fcc and bcc structures, but may also signal a local geometric advantage that the less-spherical B$_{10}$H$_{10}^2$ anions have over their more spherical relatives. In particular, within a cubic structure, one might expect each of the more-ellipsoidal B$_{10}$H$_{10}^2$ anions to take up slightly less space in directions perpendicular to their long axes than the more spherical B$_{12}$H$_{12}^2$ anions. On a local level, this would allow more free space between anions for cation diffusion. The QENS results are consistent with a locally ellipsoidal anion, by suggesting that each anion retains a particular orientation of its long axis over at least a nanosecond timescale.

Much needs to be done to provide a better understanding of the superionic properties of this new class of conducting materials. For example, since Na$_2$B$_{10}$H$_{10}$ possesses a much lower order–disorder phase-transition temperature than Na$_2$B$_{12}$H$_{12}$, one might think that the lighter-metal analogue, Li$_2$B$_{10}$H$_{10}$, would also possess a lower order–disorder phase-transition temperature than Li$_2$B$_{12}$H$_{12}$,[6,7] thus enhancing the stability of the disordered, fast-ion-conducting structure. Our Li$_2$B$_{10}$H$_{10}$ DSC measurements suggest the contrary. Rather, Li$_2$B$_{10}$H$_{10}$ appears to possess a slightly higher transition temperature than Li$_2$B$_{12}$H$_{12}$ does, leading to an unstable disordered structure, which makes it unsuitable as a solid-state, Li$^+$-conducting electrolyte. However, the addition of other anions or cations to Li$_2$B$_{10}$H$_{10}$ and to Na$_2$B$_{10}$H$_{10}$ may lead to hybrid materials displaying even lower transition temperatures than seen here for pure Na$_2$B$_{10}$H$_{10}$. We are currently pursuing such potentially favorable modifications.

For all these disordered materials, a more thorough understanding of the relationship of structural disorder and anion reorientational mobility to cation diffusion and conductivity will benefit from future first-principles molecular-dynamics calculations,[1,17,18] which may in turn lead to a more rational pathway to develop improved modified materials.

In conclusion, the discovery of very high superionic conductivity in Na$_2$B$_{10}$H$_{10}$ that persists to temperatures as low as 360 K is a marked improvement over Na$_2$B$_{12}$H$_{12}$ and other complex hydrides. Although hygrosopic, Na$_2$B$_{10}$H$_{10}$ remains air-stable at room temperature with no noticeable degradation in its diffraction pattern up to at least 500 K. DSC indicates that the compound decomposes/polymerizes with some mass loss of presumably H$_2$ at around 850 K. Preliminary cyclic voltammetry measurements indicate that ordered Na$_2$B$_{10}$H$_{10}$ is electrochemically stable up to at least 4 V at 353 K and disordered Na$_2$B$_{10}$H$_{10}$ up to at least 5 V at 393 K (see Figure S2 in the Supporting Information). Its favorable properties and high conductivity warrant a serious investigation of Na$_2$B$_{10}$H$_{10}$’s applicability to next-generation solid-state Na-ion battery technologies. Based on these results, successful future searches for related materials with even better cation conductivity properties may be enhanced by the inclusion of similar- or even larger-sized polyanions compared with B$_{10}$H$_{10}^2$.

**Experimental Section**

**Synthesis:** 11Boron-enriched Na$_2$B$_{10}$H$_{10}$ (and partially deuterated Na$_2$B$_{10}$D$_{10}$) was synthesized as follows: the triethylammonium salt [(Et$_3$NH)$_2$B$_{10}$H$_{10}$] was synthesized via reaction of Na$_2$B$_{10}$H$_{10}$ and triethylamine in para-xylene at reflux. The crude product was recrystallized from water/EtOH and dried in vacuum (10 mTorr) at room temperature for 16 h. The (Et$_3$NH)$_2$B$_{10}$H$_{10}$ was then converted into the corresponding acid (H$_2$O)$_2$[(Et$_3$NH)$_2$B$_{10}$H$_{10}$] by ion exchange using an Amberlite resin in H+-form. Aqueous Na$_2$B$_{10}$H$_{10}$ was prepared by neutralization of (H$_2$O)$_2$[(Et$_3$NH)$_2$B$_{10}$H$_{10}$] with 0.1 M NaOH until a pH value of 7 was reached. The solvent was removed on a rotary evaporator at 323 K. Unlabeled Na$_2$B$_{10}$H$_{10}$ was synthesized using a similar approach. The resulting hydrated materials were dried under vacuum at 433 K for 16 h. For the partially deuterated sample, a single exchange treatment was performed by dissolution and stirring for 3 h of 1 g of Na$_2$B$_{10}$D$_{10}$ in 20 mL of D$_2$O slightly acidified by adding 50 µL of a saturated solution of deuterochloric (DCI) acid in D$_2$O. The resulting dried sample had a D/H ratio of only 27:73 as determined from refinement of the 20 K NPD pattern, yet led to some reduction of the incoherent neutron scattering background from the lighter H isotope.

**Measurement Details:** DSC measurements were made with a Netzsch (STA 449 F1 Jupiter) TGA-DSC under He flow with Al sample pans. The neutron-scattering measurements were performed at the National Institute of Standards and Technology Center for Neutron Research. NPD patterns were measured on the BT-1 High-Resolution Powder Diffractometer using the Ge(311) monochromator at a neutron wavelength of 2.077 Å. Horizontal divergences of 60°, 20°, and 7° of arc were used for the in-pile, monochromatic-beam, and diffracted-beam collimators, respectively. The sample was contained in a 6 mm-diameter V can inside a He closed-cycle refrigerator. FWSs were measured on the High-Flux Backscattering Spectrometer using 6.27 Å wavelength neutrons, with a resolution of 0.8 µeV FWHM. QENS spectra were collected at 270 K (resolution measurement) and 375 K on the Disk Chopper Spectrometer using 4.08 Å wavelength neutrons with a resolution of 79 µeV FWHM. 23Na NMR measurements were performed on the pulse spectrometer described earlier[9] at the frequency $\nu/2\pi = 23$ MHz. The nuclear spin–lattice relaxation rates were measured...
using the saturation–recovery method. NMR spectra were recorded by Fourier transforming the solid-echo signals. Ionic conductivities were determined in heating and cooling runs repeatedly in the temperature range between 303 K and 423 K by the AC complex-impedance method using an NF FRA5097 frequency response analyzer over a frequency range of 10 Hz to 10 MHz. All the measurements were performed under Ar. The powder sample was pressed into a pellet of 8 mm in diameter and 2 mm in thickness without sintering. The pellet density was about 1.17 g cm\(^{-3}\), which is more than 95% of the density calculated from the lattice parameters. Au or Mo foils were used as electrodes and were mechanically fixed on both faces of the pellet. The resistances of the sample were obtained by least-square fittings of a single arc in the high-frequency range using equivalent circuits of a parallel combination of a resistance and a capacitance. At high temperature, since only a spike was observed using equivalent circuits of a parallel combination of a resistance and a capacitance. At high temperature, since only a spike was observed with the observed scatter in the data, if not explicitly designated by the reference electrodes of Na or Na–In at 353 K and 393 K, respectively.

VersaSTAT4) with a Mo disk as the working electrode and counter/reference electrodes of a Mo disk as the working electrode and counter/reference electrodes of Na or Na–In at 353 K and 393 K, respectively.

Finally, for all the figures, standard uncertainties are commensurate with the observed scatter in the data, if not explicitly designated by the vertical error bars.

Supporting Information
Supporting Information is available from the Wiley Online Library or from the author.

Acknowledgements
This work was performed, in part, in collaboration between members of IEA HIA Task 32-Hydrogen-based Energy Storage. The authors gratefully acknowledge support from DOE EERE through Grant Nos. DE-EE0002978 and DE-AC04–94AL85000; the Russian Foundation for Basic Research under Grant No. 12–03–00078; the U.S. Civilian Research & Development Foundation (CRDF Global) under Award No. RUP1–7076-EK-12; the National Science Foundation (NSF) under Cooperative Agreement No. OISE-9531011; the Integrated Materials Research Center for the Low-Carbon Society (LC-IMR), Tohoku University; the Advanced Low Carbon Technology Research and Development Program (ALCA) from the Japan Science and Technology Agency (JST); and JSPS KAKENHI under Grant Nos. 25220911 and 26820311. This work utilized facilities supported in part by the NSF under Agreement No. DMR-0944772. The authors also thank Dr. Nina Verdal for assistance with the QENS measurements.

Received: July 14, 2014
Revised: August 30, 2014
Published online: October 13, 2014


The mention of all commercial suppliers is for clarity and does not imply recommendation or endorsement by NIST.

---

(References and Supporting Information are not included in this text)
Supporting Information

for Adv. Mater., DOI: 10.1002/adma.201403157

Exceptional Superionic Conductivity in Disordered Sodium Decahydro-closo-decaborate

Supporting Information

Exceptional Superionic Conductivity in Disordered Sodium Decahydro-closo-decaborate


NPD Results: The NPD patterns were analyzed by Rietveld refinement[1] using the GSAS package[2] assuming neutron scattering amplitudes of (3.63, 6.65, 6.67, and -3.74) fm for Na, $^{11}$B, D, and H, respectively. A D:H ratio of 27:73 was determined from the low-temperature monoclinic structure[3] model fit to the data at 20 K, assuming a random isotopic distribution of D and H atoms. Wavelength errors were not included in the standard deviations of the unit cells; i.e., the precisions reported in this paper for the structural parameters reflect the quality of the data and the corresponding refinement model, assuming a fixed neutron wavelength.

Table S1. Structure parameters of the high-T disordered phase (Fm-3m) of Na$_2$$^{11}$B$_{10}$(H$_{0.73}$D$_{0.27}$)$_{10}$ at 410 K (a=9.8426(8) Å).

<table>
<thead>
<tr>
<th>Atom</th>
<th>Site</th>
<th>x</th>
<th>y</th>
<th>z</th>
<th>$U_{iso}$ (Å$^2$)</th>
<th>Occupancy</th>
</tr>
</thead>
<tbody>
<tr>
<td>B1</td>
<td>24e</td>
<td>0.186</td>
<td>0</td>
<td>0</td>
<td>0.10</td>
<td>0.333</td>
</tr>
<tr>
<td>B2</td>
<td>96k</td>
<td>0.063</td>
<td>0.063</td>
<td>0.128</td>
<td>0.12</td>
<td>0.333</td>
</tr>
<tr>
<td>D1</td>
<td>24e</td>
<td>0.306</td>
<td>0</td>
<td>0</td>
<td>0.4</td>
<td>0.333</td>
</tr>
<tr>
<td>D2</td>
<td>96k</td>
<td>0.110</td>
<td>0.110</td>
<td>0.226</td>
<td>0.4</td>
<td>0.333</td>
</tr>
<tr>
<td>Na1</td>
<td>8c</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td>0.15</td>
<td>0.58(1)</td>
</tr>
<tr>
<td>Na2</td>
<td>4b</td>
<td>0.5</td>
<td>0</td>
<td>0</td>
<td>0.4</td>
<td>0.28(3)</td>
</tr>
<tr>
<td>Na3</td>
<td>24d</td>
<td>0.25</td>
<td>0.25</td>
<td>0</td>
<td>0.3</td>
<td>0.09(1)</td>
</tr>
</tbody>
</table>

The center of the B$_{10}$D$_{10}^2$ anion was fixed at the $4a$ site (0 0 0). B-D bond lengths were fixed at 1.173 Å. B-B bond lengths were fixed at those present in the low-T ordered Na$_2$B$_{10}$H$_{10}$ phase. $R_w=0.0123$, $R_p=0.0104$, and $\chi^2=0.817$. A neutron scattering amplitude of 0.73(-3.74) + 0.27(6.67) = -0.93 fm was used for the D1 and D2 atoms. Na positions were determined from a Fourier difference map, and Na site occupancies were refined using reasonable fixed $U_{iso}$ values. It should be noted that the thermal factors and occupancies for Na were correlated. Changes in thermal factors affected the occupancies, but the variations in occupancies were not significant. The magnitudes of the Na1 (t), Na2 (o), and Na3 (i) site occupancies followed the same trend, i.e., occ(t) > occ(o) > occ(i), and the total number of Na atoms was basically around 8 per unit cell. This was also an indication of the correct structure model.
Figure S1. Cross-sectional SEM images of the pelletized Na$_2$B$_{10}$H$_{10}$ sample. As can be seen, intimate contacts among particles are achieved.

Figure S2. Cyclic voltammogram of disordered Na$_2$B$_{10}$H$_{10}$ at 393 K using Mo disk as the working electrode and a Na-In alloy as the counter/reference electrode at a scan rate of 5 mV s$^{-1}$.