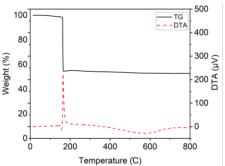
## Temperature-Dependent Carbonization of Pyridine-Stabilized Alkynylborane for Energy Storage Applications

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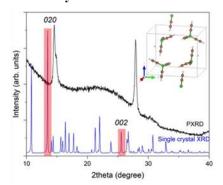
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Boron-doped carbon (BDC) has attracted attention as an alkali-ion storage anode material.<sup>1</sup> Previous studies have reported the synthesis of BDC materials using boric acid or boron oxide as starting materials.<sup>2</sup> However, these molecules require significant energy to cleave the B-O bonds and form the B-C bonds, making it challenging to control the bonding states of boron. Ethynyl groups are known as functional groups that promote carbonization through pyrolysis. Additionally, three-dimensional carbons are synthesized from ethynyl-containing organic compounds.<sup>3</sup> In this study, we designed ethynyl-containing boron molecules to serve as building blocks for BDC.

We focused on triethynylborane (TEB) as an ethynyl-containing boron molecule. Three-coordinated boron compounds like **TEB** are sensitive to oxygen because of their high Lewis-acidity. In this study, synthesized TEB-pyridine complex (TEB-Py), which is stabilized by coordinating Lewis base to the boron center. We investigated the thermal behavior of TEB-Py by TG-DTA (Figure 1). As a result, we confirmed endothermic and exothermic peaks at 157 and 160 °C, respectively. The first peak (157 °C) evidences the desorption of pyridine, and the second one (160 °C) indicates the polymerization of ethynyl groups. Based on the TGA study, we synthesized BDC at three different temperatures (200, 600, and 900 °C) and named them TEB- $C_X$ where x denotes temperature. We compared the experimental XRD pattern of TEB-C<sub>200</sub> with the simulated XRD pattern of TEB (Figure 2), which evidenced that carbonization proceeded by desorption of pyridine. Obtained BDCs were evaluated for their potential applications in lithium and sodium ion storage.



(Figure 1) TG-DTA analysis of **TEB-Py** 



(Figure 2) Simulated XRD pattern of **TEB** and measured XRD pattern of **TEB-C**<sub>200</sub>.

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