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TOPIC(s) : Homogenous, heterogenous and biocatalysis

## Tailored Hydrogen Generation from Borohydride in Ionic Liquids

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## PURPOSE OF THE ABSTRACT

The continuous depletion of fossil fuels promotes the development of sustainable energy systems, such as H<sub>2</sub>, an environmentally benign energy carrier [1]. H<sub>2</sub> can be obtained via water electrolysis, using renewable energy sources. After temporary storage of H<sub>2</sub>, it is fed to a fuel cell, recovering the retained energy, e.g. to charge a phone [2]. The H<sub>2</sub> lifecycle is schematically displayed in Figure 1. Especially for portable fuel cell applications, the H<sub>2</sub> storage is a key issue [3]. Current H<sub>2</sub> storage technologies include pressure tanks, cryo storage or chemically bound H<sub>2</sub>, such as LiBH<sub>4</sub> or NaBH<sub>4</sub> with a H<sub>2</sub> capacity of 18.4 wt% and 10.6 wt% H, respectively [4, 5]. BH<sub>4</sub><sup>-</sup> hydrolyzes when in contact with H<sub>2</sub>O, forming H<sub>2</sub> and the corresponding borate (Eq. (1)) [6, 7].

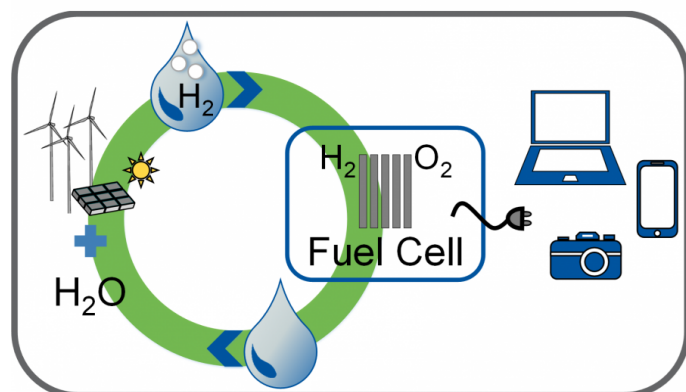


For the implementation of BH<sub>4</sub><sup>-</sup> in portable electronic applications, a hydrogen storage medium with a melting point below room temperature (RT) and the tailored catalytic H<sub>2</sub> release is mandatory. In the present study, we produce H<sub>2</sub> from liquid storage materials based on BH<sub>4</sub><sup>-</sup> in ionic liquids (ILs) via hydrolysis under ambient conditions, using various catalysts as releasing agents. For this purpose, various BH<sub>4</sub><sup>-</sup> ionic compounds were synthesized (Figure 2). Our findings indicate that EMIM BH<sub>4</sub><sup>-</sup> 2 and PMIM BH<sub>4</sub><sup>-</sup> 3 with a hydrogen storage capacity of max. 3 wt% H are promising storage media [1].

Subsequently, the BH<sub>4</sub><sup>-</sup> ILs were hydrolyzed with H<sub>2</sub>O using supported metal as well as acidic catalysts. All hydrolyses of EMIM BH<sub>4</sub><sup>-</sup> and PMIM BH<sub>4</sub><sup>-</sup> with metal/acid catalysts revealed saturation curves, whereas highest H<sub>2</sub> yields were obtained in a semi-batch process with continuous acid addition. Additionally, mass transport limitations as well as activation energies of the PMIM BH<sub>4</sub><sup>-</sup> hydrolysis with supported Co catalyst were investigated. A mechanism for the hydrolysis of PMIM BH<sub>4</sub><sup>-</sup> is proposed based on the <sup>11</sup>B NMR analysis of the H<sup>+</sup> catalyzed hydrolysis of BH<sub>4</sub><sup>-</sup> at various reaction times [1].

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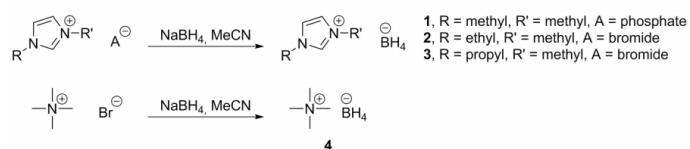
## FIGURES



**FIGURE 1**

### H<sub>2</sub> Lifecycle

Schematic lifecycle of H<sub>2</sub> as energy carrier, including the formation of H<sub>2</sub> via water electrolysis with renewable energy and subsequent application of hydrogen as a feedstock for fuel cells.



**FIGURE 2**

### Borohydride Ionic Liquid Synthesis Synthesis of BH<sub>4</sub><sup>-</sup> ILs.

## KEYWORDS

Borohydride | Ionic liquids | Hydrogen Storage | Hydrolysis

## BIBLIOGRAPHY

- [1] E. Klindtworth, I. Delidovich, R. Palkovits, *Int. J. Hydrogen Energy* 2018, 43(45), 20772-20782. [2] R. O'Hayre, S. W. Cha, F. B. Prinz, W. Colella, Wiley 2016. [3] L. Schlappbach, A. Züttel, *Nature* 2001, 414(6861), 353-358. [4] A. F. Dalebrook, W. Gan, M. Grasmann, S. Moret, G. Laurency, *Chem. Commun.* 2013, 46(78), 8735-8751. [5] M. Paskevicius, L. H. Jepsen, P. Schouwink, R. Cerny, D. B. Ravensbaek, Y. Filinchuk, M. Dornheim, F. Besenbecher, T. R. Jensen, *Chem. Soc. Rev.* 2017, 46(5), 1565-1634. [6] S. Eugenio, U. B. Demirci, T. M. Silva, M. J. Carmezin, M. F. Montemor, *Int. J. Hydrogen Energy* 2016, 41(20), 8438-8448. [7] H. I. Schlesinger, H. C. Brown, A. E. Finholt, J. R. Gilbreath, H. R. Hoekstra, E. K. Hyde, *J. Am. Chem. Soc.* 1953, 75(1), 215-219.