

Nitride materials as possible electrodes for NH₃ reversible fuel cells

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A principal aim of the EU 2030 climate & energy framework is the reduction of CO₂ emissions by the development of new technologies for industries and processes[1]. In this respect, ammonia (NH₃) is a key chemical that is produced in vast quantities worldwide as a precursor in a range of products that are vital to society, such as fertilizers and medicines. Moreover, due to its high energy density and the formation of only H₂O and N₂ upon its combustion, it also has been proposed as a potential carbon-free fuel [2].

The current industrial route for ammonia production is the Haber- Bosch process, which employs Fe-based catalysts to synthesize ammonia from nitrogen and hydrogen at high pressures (100-300bar) in the temperature range 300-550°C [3]. Despite this prevalence, the process has a very high energy consumption, with low equilibrium conversion (10-15%) and produces 300 million metric tons of CO₂ per year due to its continued reliance on hydrogen produced from natural gas [3].

New greener concepts for NH₃ production and utilisation are, therefore, urgently needed.

The current work aims to offer potential materials for a highly attractive alternative to mitigate this problem, based on the electrochemical synthesis of NH₃ directly from H₂O and N₂, with the input of renewable electricity, using a Proton Ceramic Electrolyser Cell (PCEC), Fig.1. Such a process can form NH₃ without CO₂ generation. Moreover, it can also provide a method for the chemical storage of renewable electricity, generating a route to balance renewable energy supply and demand, by the formation of a transportable chemical product of high energy density.

At this juncture, it is also important to highlight that the reverse operation of the electrochemical cell of Fig.1, would correspond to a fuel cell, where NH₃ would be combusted to form the products of H₂O and N₂, while also directly producing electricity and heat.

Despite this attraction, and the apparent similarity of these two devices, previous work on these concepts are scarce. For this reason, the current presentation aims to emphasise the challenges of these processes and the critical functionalities that are required in each case with respect the NH₃ electrode.

Discussion will then progress to describe some recent work of our group on the development of a range of nitride materials as possible NH₃-electrodes for these devices.

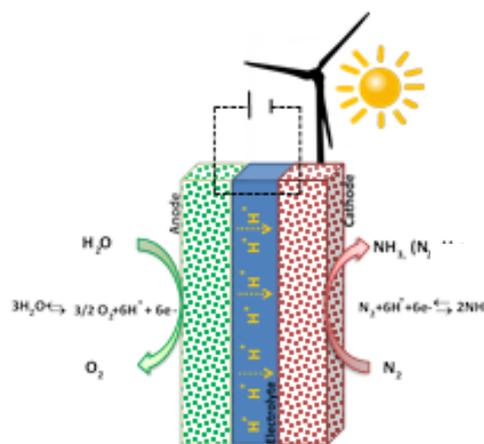


Figure 1. Electrochemical NH₃ production using a protonic membrane.

References

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