A hydrogen generator coupled to a hydrogen heater for small scale portable applications

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Abstract. This study aims to build and test a small scale portable device able to couple a hydrogen generation system (based on a NaBH₄ solution as liquid H₂ carrier) to a hydrogen heater (based on the exothermic catalytic combustion of the released H₂). The hydrogen generating system is based on the hydrolysis of stabilized solutions of NaBH₄ (fuel solutions) which are pumped into the hydrolysis reactor. The generated H₂ feeds the catalytic combustor. Two catalysts have been developed for the H₂ generation and the combustion reactions able to operate at room temperature without need of additional energy supply. For the NaBH₄ hydrolysis a Co-B catalyst was supported on a perforated and surface treated stainless steel (SS316) home-made monolith. For the flameless H₂ catalytic combustion a Pt catalyst was prepared on a commercial SiC foam. The device was automatized and tested for the on-demand production of heat at temperatures up to 100°C. In steady state conditions the NaBH₄ solution flow is controlling the H₂ flux and therefore the heater temperature. Once the steady-state is reached the system responds in a few minutes to up and down temperature demands from 80 to 100 ºC. The catalysts have shown no deactivation during the tests carried out in several days.

1 Introduction

For the implementation of the “hydrogen economy”, challenges related to its sustainable and low-cost production, transportation and storage are under continuous investigation [1–5]. The use of NaBH₄ (SBH) solutions as liquid hydrogen carrier constitutes an attractive strategy for hydrogen storage [6-8]. Solid SBH is stable in dry air, can be stabilized in basic solutions and produces hydrogen through its hydrolysis reaction (Eq.1). The borate by-product is non-toxic, and must be extracted from the spent fuel for further regeneration [9,10].

\[
NaBH₄ + 2H₂O \rightarrow 4H₂ + 2NaBO₂ \tag{1}
\]

The potential of SBH solutions for H₂ storage and release, coupled to PEM fuel cells for mobile or portable applications have been previously considered [11-14]. In this work we aim to couple a H₂ generation system to a reactor for the direct production of heat (e.g. cookers and heaters) based on the catalytic hydrogen combustion (CHC, Eq.2).

\[
H₂ + \frac{1}{2}O₂ \rightarrow H₂O \tag{2}
\]

The CHC is a key reaction in the “hydrogen economy” because it is safe, controllable and highly exothermic (286 kJ/mol) [15]. This reaction can be employed for heat production as well as for safety purposes for the elimination of undesired hydrogen [15-17]. The catalysts in this work have been selected to operate at room temperature for both H₂ generation and combustion reactions and have been deposited on selected structured supports according to the reactors’ requirements [18]. The design of the H₂ generation and combustion reactors and the final controlled heater device are described herein, together with the operation tests.

2 Catalysts and catalytic reactors

2.1. The H₂ generation by NaBH₄ hydrolysis

Co-B materials are the most investigated cobalt based catalysts for the SBH hydrolysis reaction and have been prepared on a wide range of conditions in powder as well as in supported form [12-14,19-21]. As catalyst’s structured support, a homemade cylindrical monolith was fabricated from commercially available perforated stainless steel (SS316) as shown in Fig.1a [13-14]. The SS support was calcined at 900°C to produce a well adhered oxide layer according to a previous work [13]. The supported catalyst was calcined at 900°C to produce a well adhered oxide layer according to a previous work [13]. Fig.1c shows SEM (scanning electron microscopy) images of the monolith oxidized surface at two magnifications. The Co-B catalyst was deposited by successive cycles of alternated immersions of the support on 30% CoCl₂·6H₂O and stabilized 19% SBH aqueous solutions [13]. Fig.1b shows the monoliths after deposition of the Co-B catalyst. A cylindrical reactor
with a total volume of 11 mL (hydrolysis reactor in Fig.3) can be made of PMMA or stainless steel. Catalytic monoliths are placed inside the reactor where the hydrolysis reaction will start spontaneously upon contact of the SBH stabilized solution with the catalysts.

Fig.1. Stainless steel monoliths before (a) and after (b) deposition of the Co-B catalysts. (c) SEM images of monolith surface modified after thermal treatment.

2.2 The catalytic H₂ combustion

For the flameless H₂ catalytic combustion Pt and Pd are reported as the most active catalysts which permit to initiate the reaction without ignition even at room temperature [15, 22-23]. In this work a Pt deposited catalyst was prepared on commercial SiC foam by impregnation with an ethanolic solution of H₂PtCl₆ for a final Pt loading of 1.0 wt%. The catalyst was dried in an oven and further calcined in static air at 400 °C during 4 h. Fig.2 shows the commercial disc of alumina bonded SiC foam (from Lanik s.r.o., 80 pores per inch) before (a) and after (b) deposition of the Pt catalyst. Fig.2c shows SEM (scanning electron microscopy) surface images of the monolith SiC structured supports at two magnifications.

Fig.2. SiC monoliths before (a) and after (b) deposition of the Pt catalysts. (c) SEM images of the bare SiC foam surface.

The reactor (combustor in Fig.3) was made of stainless steel for 30 mm diameter and 5 mm thickness size of catalytic monoliths. H₂ generated in the hydrolysis reactor will directly feed the combustor where the CHC reaction will start spontaneously in the presence of oxygen and the Pt catalyst.

3 Controlled heater device for portable applications

3.1. The experimental design

According to scheme in Fig.3 a prototype has been designed and constructed for a portable heater powered by H₂. Stainless steel was used to fabricate the two reactors according to machining availability in the laboratory. The hydrolysis reactor and the residues tank could however be constructed with polymeric materials. The tank for the SBH fuel solution was made of PP polymer.

Fig.3. Scheme of the complete experimental design for a portable H₂ heater: (1) Tank for the SBH fuel solution. (2) Micropump. (3) Hydrolysis reactor with catalytic monolith. (4) Tank for the borate residues solution. (5) CHC reactor with catalytic monolith. (6) Controller

The hydrolysis reactor consists of a tube of 105 mm long and 15 mm internal diameter. The reactor has three connections, the fuel inlet on the bottom, the waste outlet on the side and the gas outlet on the top. The waste outlet is connected to the waste tank (V ca. 300 mL), which is also connected to the gas outlet line (pressure balance in the system). Inside the reactor is the home-developed stainless steel monolith with the CoB catalyst of 60 mm length and 12 mm diameter. The fuel enters the reactor from down, and in contact with the catalyst begins to evolve H₂. The gas formed exits the reactor free from residues through the head of the reactor. At a height of 75 mm there is a connection (min. 1/8”) for the elimination
and separation of the reacted fuel to the waste tank. With this configuration the catalyst is wet during operation and the synergism between the high temperatures achieved and the continuous elimination of the spent fuel, prevent borates from crystallization on the surface of the CoB, which could easily cause deactivation [12,13]. Fig.4 shows the experimental realization of the portable H2 generator which also includes a thermocouple for reaction temperature measurements. The selected micro-pump was from HNP Microsysteme with reference mzr-2921.

![Fig. 4. Experimental realization of the portable H2 generator.](image1)

The catalytic combustion reactor is connected to the hydrolysis reactor (H2) and to the air supply from a compressor (for a final device a miniature air pump may be used). The reactor consists of a stainless steel body with an internal diameter of 30 mm and a height of 50 mm. The conical bottom with the gas inlet is 20 mm high and is filled with quartz wool for free volume reduction. On top it is a disc (5 mm thick) of the bare porous SiC foam as a diffuser. Above it is a free space where the air inlet is placed and is mixed with the H2. In the upper part the porous SiC plate with the deposited catalyst is located. The heat resulting from catalytic combustion is measured with a K-type thermocouple, which is in contact with the center of the catalytic plate. The reactor can be used in the open mode (for the final portable device) or with a cap including a gas outlet for gas chromatographic (GC) analysis. Fig.5 shows the experimental realization of the portable H2 combustor.

### 3.2. The operation tests

With the objective of controlling the heater temperature by the fuel addition rate, tests were performed with the following conditions:

(i) SBH hydrolysis reaction: Fuel (NaBH4) concentration 1-2 wt.%. Fuel flow between 0.5-2 mL/min. In these conditions the maximum H2 production is 30 mL/min. Maximum usage time: determined by the capacity of the waste tank (ca. 3h in this prototype).

(ii) Catalytic hydrogen combustion: Air flow 800 mL/min, fixed to ensure a concentration of H2 in air below 4 vol.%. H2 flow up to 30 mL/min. H2 conversion (determined by GC) > 99%

![Fig. 5. Experimental realization of the H2 combustor.](image2)

![Fig. 6. Relationship (as a function of experiment time) between the combustor temperature and the SBH fuel solution flow which is feeding the H2 generator. (a) 1wt.% NaBH4, (b) 2wt.% NaBH4.](image3)
First tests were devoted to determine the relationship between the fuel flow feeding the hydrolysis reactor and the temperature resulting in the combustor. Two SBH concentrations (1 and 2 wt.%) were compared. Results are shown in Fig.6a and 6b respectively. In both cases it is observed an increase in temperature, due to the hydrogen combustion, with an increased fuel flow. The reason is that there is a correlation between the fuel addition and the hydrogen generation rates [12,13]. With the 1 wt.% fuel a maximum temperature of 128 °C can be reached with 2.5 mL/min flow and 100 °C is reached with 1.5 mL/min fuel. Comparatively lower flows are needed and higher temperatures are reached for the 2 wt.% fuel as expected [12-13]. With a 1.2 mL/min flow, 142 °C can be reached while to achieve 100°C a flow of ca. 0.7 mL/min of fuel was needed. For the range of around 100 °C the 1 wt% SBH fuel was selected to test the application of an electronic control at these moderate temperatures.

Second tests were therefore done connecting the micro-pump to the electronic controller as shown in Fig. 3. The objective is to control the flow of the fuel in relation to the desired temperature of the combustor. Two set-point temperatures have been chosen, 100°C and 80°C. To avoid a long initial induction period, observed in preliminary experiment, a quantity of 2 mL of 19wt% SBH fuel has been initially injected, resulting in the production of ca. 600 mL of H₂. This will purge the complete system from air and will establish a base H₂ pressurization in the system. Once the combustor is at room temperature the experiments start. The set points temperatures given to the controller, along the experiment time, are indicated in red in Fig.7.

![Fig. 7. Evolution of the H₂ combustor temperature upon time as a response to the set-point temperatures given to the controller in an automatic mode operation](image)

Despite the previous injection of the concentrated fuel to purge and pressurize the system, the combustor starts to heat up slowly what results in a high flow demand of fuel. Due to this, an intense temperature increase can be observed at the beginning of the experiment until the system finally stabilizes at the predetermined temperature of 100 °C. This peak also results in high fuel consumption. Once in the range of this temperature, the system reaches the selected temperatures in a short time (2-3 min) (80°C-100°C-80°C-100°C-80°C), demonstrating the regulation effectiveness of the controller system. No deactivation of the catalysts has been observed, neither in this automated experiment, nor in the previous manual experiment.

4 Conclusions

In summary, in this work a small portable heater which operates at room temperature was designed and built. The system is based on coupling a hydrogen generator with a hydrogen combustor. The controlled addition of SBH in the form of stabilized solution (fuel solution) permits to generate hydrogen in a controlled rate to feed the combustor. In steady state conditions, the temperature of the combustor was tuned with the fuel addition rate, which was controlled with the micro-pump. The system operating in an automate mode, was able to respond fastly to the desired temperature from 80°C to 100°C and vice-versa. When the system is demanded to increase the temperature, the heating is slow, resulting in the increase of a fuel addition rate which produces a peak in the temperature of the combustor. On the contrary, the system responds more efficiently to a demand for decreasing temperature. The catalysts chosen for the hydrogen generator and the combustor demonstrated to be adequate to initiate the reactions with no extra energy supply and stable during each experiment and upon cycles. For a final mobile device the system requires the integration of a small battery (or fuel cell also fed by the H₂ generator) to power the small controller and both fuel and air micro-pumps. The designed (SS monolith) and selected (SiC disk) structured supports met the geometric requirements of each reactor and the necessary surface roughness for high catalyst anchorage.

In sum, the results presented herein constitute a proof of concept which can be adapted and scaled to different temperature demand. The operation time is limited by both the fuel addition and residues tanks which could be eventually replaced during operation.

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