Investigation of Heat Extraction Methods from Cuprous Chloride for Improving the Efficiency of the Thermochemical Copper-Chlorine Cycle

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ABSTRACT

In order to improve the efficiency of the thermochemical copper-chlorine cycle (Cu-Cl) for hydrogen production, a quench cell configuration for the quenching of cuprous chloride (CuCl) is being investigated. Initial testing has been performed in water to determine whether the molten cuprous chloride can be cooled enough to be quenched in hydrochloric acid (HCl (aq)). Quenching in HCl (aq) would reduce the number of components in the overall cycle, simplifying it and potentially making it more cost effective. A thermal camera was used to experimentally obtain the heat loss as the CuCl falls through the heat exchanger and into the quench cell. In addition, a FlowSense 2M shadow imaging camera was used to observe the behavior of the CuCl droplets interacting with the quench solution. It was observed that the CuCl droplet disintegrating into dust like particles as it enters the water pool and vapour was generated hence the temperature at the surface of the droplet as it entered the liquid water exceeded 100 °C.

Keywords: hydrogen production, copper chlorine cycle, quench cell, heat exchanger

INTRODUCTION

As the demand for energy increases, so do the challenges of producing enough fuel to meet the demand. Currently fossil fuels are the main source of energy in the world. Fossil fuels however are finite in supply, and contribute 79% of the total US greenhouse gas emissions (Marques et al., 2018). In order to meet future energy demands, alternative energy sources and means of energy production need to be considered.

One alternative fuel is hydrogen. Although hydrogen is not an energy resource itself, it is an energy carrier that can facilitate the use of many sustainable energy resources (Alanne and Cao, 2017; McDowall, 2012). Hydrogen is the most abundant element, making up roughly 90% of all the atoms in the universe. On earth, however, hydrogen is present mainly in water (H2O), as opposed to hydrogen gas (H2). Currently, hydrogen is mainly produced from natural gas using steam-methane reforming (McDowall and Eames, 2006). Since natural gas is a finite resource and since its use leads to environmental and other problems, it is necessary to explore other methods. Another method currently used for hydrogen production is water electrolysis. A challenge with this method is its efficiency, defined as the ratio of the energy in the hydrogen produced to the electrical energy input. The efficiency of water electrolysis is approximately 24% (Lewis et al., 2009). An alternative process to produce H2 in an efficient and a clean manner that exhibits significant potential is thermochemical water decomposition (Rosen et al., 2012; Naterer et al., 2010; Naterer et al., 2011; Dincer, 2012).
One thermochemical water decomposition process that has received increasing attention in recent years is a four-step thermochemical Cu-Cl cycle (Naterer et al., 2009). The overall Cu-Cl cycle is complex. A simplified version of it is shown in Figure 1, where it is seen to involve four main chemical reactions. In Table 1, the four main steps of the four-step Cu-Cl cycle are listed. The overall nature of the process can be observed from the chemical reactions taking place. An important step in the four-step Cu-Cl cycle involves converting the liquid CuCl in step 4 (Table 1) into an aqueous state by dissolving it in HCl (aq). The liquid CuCl after step 4 is in a molten state, being exposed to temperatures higher than its melting point. In order to prep the molten CuCl for step 1, the molten CuCl is quenched in a quench cell and then dissolved in HCl (aq). Given that the temperature of the CuCl is high when exiting step 4, the use of a heat exchanger can aid in heat recovery while cooling the molten CuCl from roughly 500 °C to a room temperature of 25 °C. In order to determine the most applicable heat extraction method, current heat extraction methods were explored. Heat transfers from high to low temperature, therefore it would be essential to ensure that the ambient temperature that the experiments conducted are as low as possible. Numerous studies have been conducted in the past regarding heat recovery methods that can be used in the Cu-Cl thermochemical cycle. These methods include atomization and steam generation with a quench bath (Ghandehariun et al., 2011), as well as a double-pipe heat exchanger (Ghandehariun et al., 2012).

An experiment is carried out to investigate this heat transfer. To observe the heat transfer during the experiment, a thermal camera is used. If a temperature of roughly 25 °C can be obtained with the heat exchanger, the CuCl can be quenched directly in HCl (aq), which implies the possible removal of the dissolution cell and ultimately reducing the cost of the overall setup. However, if a temperature of 25 °C is unattainable, the heat exchanger will be designed to achieve temperature of 100 °C and then quenched in water. In addition, how the molten CuCl behaves when it interacts with the surface of the quench liquid was examined. For example, does the molten CuCl form one solid mass, or break into many very small particles that easily dissolve in the quench solution. In order to observe this, a shadow imaging and fluid particle analysis software is used. The objective of this research paper is to improve understanding of waste heat extraction methods for the Cu-Cl thermochemical cycle and the behavior of molten CuCl in quench solution. Such an understanding is important for decision making on which measures to use to increase the efficiency of the cycle. Extracting the waste heat from the molten CuCl is beneficial to the overall system because the heat can be recycled into the cycle. This reduces the heat input required and thereby increases the efficiency of the cycle (Ghandehariun et al., 2016; Jaber et al., 2010).

Table 1. Main chemical reactions of the Cu-Cl cycle

| Step | Reaction | Temperature Range (°C) |
|------|----------|------------------------|
| 1    | \(2 \text{CuCl (aq)} + 2 \text{HCl (aq)} \rightarrow \text{H}_2\text{(g)} + 2 \text{CuCl}_2\text{ (aq)}\) | <100 (electrolysis) |
| 2    | \(\text{CuCl}_2\text{ (aq)} \rightarrow \text{CuCl}_2\text{ (s)}\) | <100 |
| 3    | \(2 \text{CuCl}_2\text{ (s)} + \text{H}_2\text{O (g)} \rightarrow \text{CuO}\ast\text{CuCl}_2\text{(s)} + \text{2HCl (g)}\) | 400 |
| 4    | \(\text{CuO}\ast\text{CuCl}_2\text{(s)} \rightarrow 2 \text{CuCl (l)} + \frac{1}{2}\text{O}_2\) | 500 |

Source (Naterer et al., 2010)
In order to conduct this experiment a quench cell assembly was proposed, designed and assembled. The experimental set up is illustrated in Figures 2 and 3. The quench cell assembly consists of a 25 L glass vessel, a heat exchanger, aluminum bars, and an insulated small glass tube wrapped in heating tape, as well as a thermal camera and a shadow imaging camera. The quench cell was equipped with valves at the water inlet, the nitrogen inlet, and the slurry outlet. The water inlet was connected to a water pump and a water reservoir and the nitrogen inlet was connected to a nitrogen tank. A pipe was added to connect the slurry outlet and a collection bucket. A pyrex Allihn condenser was installed between the molten salt reactor and the quench cell such that no air could enter the system to prevent oxidation of the CuCl. The condenser would ensure that the gasses rising during the quenching process are being condensed back into the cell so that the concentration is maintained constant and backflow into the decomposition reactor is prevented. The shadow imaging camera and the thermal camera were fixed onto the main frame by creating adjustable side supports from aluminum bars. After the installation of the cameras, they were configured in order to achieve accurate results. A small furnace was assembled to melt the CuCl and simulate the overflow from the decomposition reactor. The furnace consists of a glass tube that is open on one end to permit insertion of the CuCl powder, and an orifice on the other end for the molten CuCl to drip into the condenser. In order to obtain a temperature of 500 °C, a heat tape was wrapped around the tube and then insulated to reduce heat losses from the tube to the atmosphere.

The quench vessel was filled with water and the system was purged with nitrogen gas. The CuCl furnace was allowed to reach a steady state temperature of 500 °C. Once the temperature was achieved, 24.5 g of CuCl powder was poured into the open end of the mini CuCl furnace and then closed with insulation to prevent the melting CuCl from oxidizing in air (Jianu et al., 2018). Meanwhile the cameras were set to start recording as the molten CuCl drips out of the furnace. As each drop developed and detached, the thermal camera recorded the temperature change in the droplet whereas the shadow imaging camera recorded the behavior of the CuCl droplet as it collided with the surface of the water in the quench vessel. The droplet traveled a distance of 1.5 m from detachment to collision with water surface as shown in Figures 2 and 3.

To effectively determine the method that offers the most recovery of waste heat, multiple tests were conducted. In each said test, the rate of heat transfer from the molten CuCl and the interaction between the molten CuCl and the aqueous quench solution were observed. Overall, three variations for the quench cell assembly were or are to be examined. The first variation for the quench cell assembly consisted of dropping the CuCl into the heat exchanger and having the heat transfer occur with the ambient air and then quenching the CuCl in water at the
bottom. Using this variation, two runs of experiment were conducted and it was observed the CuCl cools to an average of 385 °C-410 °C. With heat transfer only to the air, it was observed that the only 30% of the waste heat was removed from the molten CuCl droplets. In order to increase the percentage of waste heat removed from the CuCl droplet and the ambient air, the distance between the tip of the furnace and the quench solution can be increased.

The shadow imaging camera recorded the interaction between the CuCl and the water. The obtained images were then analyzed using the shadow processing function in Dynamic Studio. It was observed that the CuCl dispersed into numerous small particles. An image of the interaction obtained from each run can be seen in Figure 4. The one on the left is run 1 and on the right is run 2. As seen in the images, the CuCl droplets behaved similarly both times when hitting the water. It was observed that the CuCl particles settled at the bottom of the water tank which is expected since it is not soluble in water (Jianu et al., 2013; Jianu et al., 2016).

The last variation for the proposed setup is to have a gas rise from the bottom of the heat exchanger to the top of the heat exchanger. This method can be beneficial in two ways. For example, depending on the choice of gas, its thermal conductivity can be higher than that of air and therefore can be coupled with a liquid flowing through the outer shell of the heat exchanger to further improve the heat extraction occurring. In addition, the air would be opposing the motion of the falling droplet, and increase the duration of the droplet in the air. Before any experimentation could occur with counter-flowing gas, it was important to identify the ideal gas for the scenario using a criteria matrix. Nitrogen was selected as the counter flow gas since it purges oxygen thus preventing the oxidation of CuCl to other undesirable complexes. An additional consideration is to observe if the counter-flow gas will affect the particles interaction with the quench solution. In order to achieve accurate results, multiple tests should be conducted. During experimentation a temperature drop from 450 °C to approx. 325 °C was observed at the surface of the droplet. The shadow imaging camera observed that the interaction of the CuCl droplet and the quench solution was unchanged.
The second variation uses the heat exchanger and involves running chilled water on the outside to extract heat from the droplet. This method however is inefficient at such a small scale. Since the thermal conductivity of the air is low and the size of the drop is small relative to the heat exchanger, the rate of heat transfer does not change significantly from variation 1. Therefore, in order to ensure that the obtained results are of use, changes to either the droplet size or diameter of the heat exchanger would be necessary.

CONCLUSIONS

A quench cell assembly that will improve the efficiency of the thermochemical Cu-Cl cycle was designed and assembled. In addition to improving efficiency, an investigation into methods of recovering the waste heat was conducted. If the heat can be extracted such that the CuCl cools to room temperature, then the CuCl can be quenched in HCl directly. The result can be removing the need of a dissolution cell in the cycle and reducing the cost of the overall cycle. Three variations were examined based on the same general design. The first variation involves natural convection of the heat to ambient air. The second variation involves the use of a chilled water blanket. This method however was unfavorable given the scale of the experiments, so the results obtained for this variation have little significance in determining the best heat extraction method. The last variation explored is floating nitrogen gas through the heat exchanger opposing the motion of the falling droplet. Based on results from the experimentation conducted, extracting the waste heat so the final temperature is about 25 °C is not possible with any heat exchanger; hence quenching in HCl would not be advisable as HCl vapour would be formed which would rise and potentially enter the decomposition reactor, causing a safety concern.

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