MODIFICATION OF LITHIUM BOROHYDRIDE AND MAGNESIUM HYDRIDE MIXTURE FOR HYDROGEN STORAGE

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ภัทรพร ศรีเคชประสาท : การปรับปรุงสารผสมระหว่างลิเทียมบอรอไฮไครค์และ แมกนีเซียมไฮไครค์ สำหรับการกักเก็บก๊าซไฮโครเจน (Modification of Lithium Borohydride and Magnesium Hydride for Hydrogen Storage) อ. ที่ปรึกษา : รศ. คร. ปราโมช รังสรรค์วิจิตร คร. สันติ กุลประทีปัญญา และ ผศ. คร. บุนยรัชต์ กิติยานันท์ 130 หน้า

งานวิจัยนี้ศึกษาคุณสมบัติการปลดปล่อย/ดูคซับก๊าซไฮโครเจนของสารผสมระหว่างลิ เชียมบอรอไฮไครค์และแมกนีเซียมไฮไครค์ ในอัตราส่วน 2:1 โคยโมล (สารผสม) คุณสมบัติที่ ตรวจสอบครอบคลุม อุณหภูมิของการปลดปล่อย กลไกการเกิดปฏิกิริยา และการผันกลับได้ ใน การศึกษาส่วนแรก เป็นการศึกษากลไกของการปลดปล่อย/ดูดซับก๊าซไฮโดรเจนของสารผสมที่ ผ่านการบดเป็นเวลา 1 ชั่วโมง ผลการทดลองแสดงให้เห็นว่า ที่อุณหภูมิต่ำกว่า 400 องศาเซลเซียส เกิดการปลดปล่อยก๊าซไฮโดรเจนบางส่วนจากลิเทียมบอรอไฮไดรด์และแมกนีเซียมไฮไดรด์ ในขณะที่อุณหภูมิ 450 องศาเซลเซียส เกิดการก่อตัวของแมกนีเซียมไดบอไรย์และลิเทียมไฮไดรด์ จากนั้นได้ศึกษาผลของการเติมไททาเนียมใตรคลอไรด์ ฮาฟเนียมเตตระคลอไรด์ และ เซอร์โคเนียมเตตระคลอไรด์ ในปริมาณร้อยละ 3 โคยโมล ลงไปในสารผสม ในการศึกษาส่วนที่ สอง ได้ศึกษาผลกระทบของเวลาในการบดต่อพฤติกรรมการปลดปล่อย/ดูดซับก๊าซไฮโดรเจนของ สารผสม ซึ่งพบว่าเมื่อเพิ่มเวลาของการบด อุณหภูมิและปริมาณของก๊าซไฮโครเจนที่ปลดปล่อย ออกมาลดลง สำหรับการศึกษาในส่วนที่สาม นั้นได้เติมตัวเร่งปฏิกิริยาไททาเนียม ไททาเนียมได ออกไซค์ และไททาเนียมไตรคลอไรค์ ลงไปในสารผสมและบคเป็นเวลา 5 ชั่วโมง ผลการศึกษา แสดงให้เห็นว่า การเติมไททาเนียม ให้ปริมาณก๊าซไฮโดรเจนที่ปลดปล่อยออกมาจากสารผสม สงสดที่ร้อยละ 5.0 โดยน้ำหนัก ในการศึกษาส่วนที่สี่ เป็นการศึกษาผลกระทบของความคันเริ่มต้น ต่อการกักเก็บก๊าซไฮโครเจนของสารผสมที่ผ่านการบดเป็นเวลา 5 ชั่วโมง จากผลการทคลอง พบว่า สารผสมที่สลายตัวภายใต้ความคันก๊าซไฮโครเจนเริ่มต้น 0.15 เมกะพาสคาล ปลดปล่อย ก๊าซไฮโครเจนออกมาสูงสุดที่ร้อยละ 5.0 โคยน้ำหนัก

ABSTRACT

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In this work, LiBH₄ was mixed with magnesium hydride (MgH₂) in a 2:1 molar ratio (mixture) in order to investigate the hydrogen desorption/absorption properties. In the first part, the mixture milled for 1 h was observed for its hydrogen desorption/absorption mechanism. The results showed that, when the temperature was lower than 400°C, the released hydrogen was from the partial decomposition of LiBH₄ and MgH₂. At 450°C, there was the formation of MgB₂ and LiH. 3 mol% of TiCl₃, HfCl₄, and ZrCl₄ was also added into the mixture. It was found that all the catalyzed mixtures had lower desorption temperatures. The effects of ball-milling time on the hydrogen desorption/absorption behaviors of the mixture were studied in the second part. It was found that the increase in the ball-milling time resulted in the decrease in the desorbed temperature as well as the hydrogen capacity. In the third part, Ti, TiO2, and TiCl3 were added into the mixture and milled for 5 h. The results showed that doping with Ti resulted in the highest amount of hydrogen, 5.0 wt%, released from the mixture. In the fourth part, the effects of initial pressure on the hydrogen storage properties of the mixture milled for 5 h were investigated. The results exhibited that the mixture decomposing under 0.15 MPa hydrogen had the highest reversible hydrogen capacity of 5.0 wt%.

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