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# Pyrohydrolysis of CaCl<sub>2</sub> waste for the recovery of HCl acid upon the synergistic effects from MgCl<sub>2</sub> and silica

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A n efficient HCl acid recovery method from the mixture of alkaline earth metal chlorides waste was demonstrated via co-pyrohydrolysis in a lab-scale horizontal furnace at a temperature range of 700-1000°C and fixed additions of SiO<sub>2</sub> and steam. The synergistic effect of MgCl<sub>2</sub> on the HCl recovery from CaCl<sub>2</sub> was explored intensively. A double-sided effect was revealed. For the reaction temperatures below 1000°C, the MgCl<sub>2</sub> addition delayed the HCl release through competing with CaCl<sub>2</sub> for the inclusion into silica matrix. In contrast, once the chloride mixtures were subjected to 1000°C with a noticeable residence time (e.g. 2 hours) and at a minimum molar ratio of 0.5 of MgCl<sub>2</sub> to CaCl<sub>2</sub>, the MgCl<sub>2</sub> addition promoted the HCl release remarkably, via promoting the conversion of Ca<sub>3</sub>(SiO<sub>4</sub>)Cl<sub>2</sub> into Ca<sub>8</sub>Mg(SiO<sub>4</sub>)<sub>4</sub>Cl<sub>2</sub>. A portion of Mg<sup>2+</sup> derived from the early decomposition of MgCl<sub>2</sub> substituted the Ca(I) site in Ca<sub>3</sub>(SiO<sub>4</sub>)Cl<sub>2</sub>, thereby resulting in the formation of weak Mg-Cl bond that is in favor of the HCl release. Additionally, the remaining Mg<sup>2+</sup> consumed the excessive SiO<sub>2</sub> so as to cause the skeleton of [SiO<sub>4</sub>]<sup>4</sup> to be fully affiliated and balanced by cations to form Ca<sub>8</sub>Mg(SiO<sub>4</sub>)<sub>4</sub>Cl<sub>2</sub>, in which the weaker ionic polarization between Ca<sup>2+</sup> and adjacent anions further enhanced the breakage of the Ca-Cl bonds.

#### Biography

Song Zhou is currently a PhD student at Monash University. He is a Member of Clean Solid Fuel Laboratory (CSFL). His present research interest is mainly on chloride waste treatment and atomic structure analysis based on X-ray Adsorption Fine Spectroscopy (XAFS) analysis.

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