

Shaping $\text{Mg}(\text{OH})_2$ crystals through hydrothermal treatment

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Magnesium hydroxide (MDH) is a compound widely employed in several industrial sectors, with increasing interest as halogen-free flame-retardant agent. Battaglia et al. [1] demonstrated a practical approach to recover MDH from waste brines via reactive crystallization, achieving high conversions and purities (about 100%). On the other hand, nanometric strongly aggregated crystals have been mainly obtained. These crystals do not comply with market requirements for flame-retardant applications: (i) a specific surface area (SSA) below $10 \text{ m}^2/\text{g}$, (ii) hexagonal plate-like morphology and (iii) a narrow particle size distribution without aggregation [2]. To enter the flame-retardant market, as part of the European project *LIFE23-ENV-IT-MareMag LIFE*, this work focuses on the hydrothermal modification of magnesium hydroxide powders obtained from waste bittern solutions. The hydrothermal treatment relies on a dissolution–recrystallization mechanism, promoting the rearrangement of the solid phase to minimize its surface energy and increase crystal size.

For the first time, a detailed comparative study was conducted exploring $\text{Mg}(\text{OH})_2$ suspensions and particles (after filtration and drying) with controlled particle sizes and morphologies. The influence of key operating parameters, namely temperature, reaction time, feed suspension concentration, and the presence of a mineralizing agent such as sodium hydroxide, was also analyzed.

A specific surface area of $8 \text{ m}^2/\text{g}$ and a hexagonal morphology were achieved at 200°C for reaction times exceeding 30 hours (using dried solids). Notably, similar performances were obtained under milder operating conditions (180°C and 5h) using suspensions with controlled initial nanoparticles distributions, leading to routes for industrial applications. Results will be adopted for scale-up and energy optimization studies.

Keywords: *Hydrothermal recrystallization, Magnesium hydroxide, Crystal morphology.*

References

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