



Study of dolomite dissolution at various temperatures – Evidence for the formation of nanocrystalline secondary phases at dolomite surface and influence on dolomite interactions with other minerals

Mathieu Debure, Pascal Andreazza, Sylvain Grangeon, Catherine Lerouge,
German Montes-Hernandez, Benoît Madé, Christophe Tournassat

► To cite this version:

Mathieu Debure, Pascal Andreazza, Sylvain Grangeon, Catherine Lerouge, German Montes-Hernandez, et al.. Study of dolomite dissolution at various temperatures – Evidence for the formation of nanocrystalline secondary phases at dolomite surface and influence on dolomite interactions with other minerals. 2015 AGU Fall meeting, American Geophysical Union, Dec 2015, San Francisco, United States. hal-01182905

HAL Id: hal-01182905

<https://brgm.hal.science/hal-01182905>

Submitted on 5 Aug 2015

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Study of dolomite dissolution at various temperatures –Evidence for the formation of nanocrystalline secondary phases at dolomite surface and influence on dolomite interactions with other minerals

M. Debure^{1,*}, P. Andreazza², S. Grangeon¹, C. Lerouge¹, G. Montes-Hernandez³, B. Madé⁴, C. Tournassat¹

1. BRGM, French Geological Survey, D3E/SVP, 3, avenue Claude Guillemin - BP 36009, 45060 Orléans Cedex 2, France

(m.debure@brgm.fr, s.grangeon@brgm.fr, c.lerouge@brgm.fr, c.tournassat@brgm.fr)

2. ICMN - UMR 7374 CNRS - Université d'Orléans 1b rue de la Férollerie CS 40059, 45071 Orléans cedex 2, France (Pascal.ANDREAZZA@univ-orleans.fr)

3. ISTerre, Maison des Géosciences, Université de Grenoble I, CNRS, Grenoble, France (german.montes-hernandez@ujf-grenoble.fr)

4. Andra, 1 – 7 rue Jean Monnet, 92298 Châtenay-Malabry, France (benoit.made@andra.fr)

*Corresponding author

In most clay-rock geological formation studied for the storage of nuclear waste, pore water compositions are expected to be at equilibrium with carbonate minerals, which are always included in predictive models for pore water composition calculations [1]. Among the carbonates known to be present, dolomite may be problematic in the pore water composition calculation because its solubility spans a large range of values as a function of its crystallinity in thermodynamic databases. In addition, the composition of dolomite minerals observed in clay-rock formations such as Callovian-Oxfordian or Opalinus clay formation differs from this of a pure dolomite: the Ca/Mg stoichiometry is not ideal, and the minerals contain minor amounts of Fe and traces of many other elements [2].

To understand the influence of secondary phases precipitation during dolomite dissolution on pore water chemistry, the dissolution of monocrystals of dolomite were investigated at 25 °C and at 80 °C in a pH range 3 to 8 for various time periods (30 minutes to 21 days) in sealed PTFE reactors. Solution analyses evidenced a stoichiometric release of Ca and Mg in solution during dolomite dissolution. Scanning Electron Microscopy (SEM), Raman and X-Ray Diffraction (XRD) analyses did not evidence secondary Mg-bearing minerals precipitation, but revealed the formation of Fe-bearing particles on the dolomite surface. Morphological characterizations performed with Small-angle X-ray scattering (SAXS) evidenced that the precipitation occurs along a specific crystallographic plane of the dolomite monocrystal. Thus, the precipitated nanoparticles clustered on specific surface sites, and are made of Fe-rich phases poorly crystallized (carbonates, oxides and hydroxides).

[1] Tournassat et al. 2015. Ch. 3: Chemical Conditions in Clay-Rocks. *Natural and Engineered Clay Barriers*, Elsevier.

[2] Lerouge et al. 2011. Geochim. et Cosmoch. Acta, 2011, 75, 2633-2663.