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Long-term Assessment of Geochemical Reactivity of CO₂ Storage in Highly Saline Aquifers: Application to Ketzin, In Salah and Snøhvit storage sites

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Abstract

Saline aquifers are choice targets for geological storage of CO₂ because of their storage potential and because these formations are not suitable for other uses. Geochemical modeling is an interesting tool to assess the geochemical behavior of CO₂ in the saline aquifer, including its dissolution in the brine and its interactions with minerals. Two key parameters which determine the confidence one can have in the results of geochemical modeling are tested in this paper: i) the establishment of the conceptual model, including the selection of the primary and secondary minerals expected to react; and ii) the activity model and the associated thermodynamic databases to calculate the interaction energies within the saline solution. In this study, we performed an analysis of a large set of CO₂ storage natural analogues, which makes it possible to identify the minerals that are likely to precipitate and dissolve during CO₂ – brine – rock interactions. Interestingly, this analysis indicates a strong dependence of dawsonite precipitation on the initial sandstone mineralogy. Dawsonite can precipitate in lithic and feldspar rich sandstones but was not observed in quartz rich sandstones. These observations on mineral reactivity are used to establish reactivity conceptual models for three CO₂ storage case-studies in saline sandstone aquifers (Ketzin, In Salah and Snøhvit) and a methodology is proposed to evaluate the long-term geochemical reactivity of these saline aquifers as a result of CO₂ injection. Noticeable differences are obtained between the case-studies as a function of the initial mineralogy and chemical conditions in the sandstones, which highlight that CO₂ mineral trapping can take place in a given storage site but can be almost absent in other storage sites. Regarding the activity model and the database, the Pitzer interaction model is rarely used for simulating CO₂ geochemical behavior in saline aquifers despite the fact that more conventionally

used activity models are not valid for such salinities. A comparison between calculated mineral solubility evolution with salinity versus experimental data is performed here using both B-dot and Pitzer activity models as well as six different databases. This comparison exercise shows that chemical interactions within saline solutions can only be reproduced using the Pitzer model, even though Pitzer databases are still incomplete or are not coherent for a wide range of chemical species and temperatures. The geochemical simulations of CO₂ injection in Ketzin, In Salah and Snøhvit saline aquifers give divergent results using different activity models and databases. A high uncertainty on the simulation results is then linked to the database choice and this study clearly stresses the need for a Pitzer database that can be confidently used in all physical/chemical conditions found in deep sedimentary aquifers.

1 Introduction

Carbon capture and storage (CCS) is considered as a promising option to limit the effects of anthropogenic greenhouse gas emissions on climate change (IPCC, 2007). Deep saline aquifers are interesting targets for geological storage of carbon dioxide (CO₂) because of their storage potential and because these formations are not suitable for other uses. Indeed, waters from deep saline aquifers in sedimentary basins have salinities ranging from 5 to >350 g L⁻¹ and contain dissolved toxic and radioactive metals which limit their use as a resource, in particular for potable use (Kharaka and Hanor, 2003). Before implementing CCS technology on a large scale, the viability of this technology must be evaluated and uncertainties must be removed regarding injectivity, containment and long-term safety. Within this purpose of assessing the viability of CO₂ storage, it is essential to predict the geochemical behavior of the CO₂ in the reservoir and in the underground media as well as the chemical and mineralogical changes linked to CO₂ injection. Assessing the geochemical evolution is mainly required for predicting near-well interactions which can alter injectivity, CO₂ trapping in the reservoir, interactions with the caprock and leakage pathways and chemical impacts in overlying aquifers (Gaus, 2010).

Different complementary methods can be undertaken to assess the behavior of the CO₂ stored in saline sandstone aquifers and its interactions with water and the minerals of the host formation CO₂-fluid-rock in a storage site (Czernichowski-Lauriol et al., 2006; Gaus, 2010). The first method is the characterization of natural CO₂ storage sites, which makes it possible to consider geological timescales at the reservoir and basin scales. The second method consists in laboratory experiments, which provide direct observations of CO₂-fluid-rock interactions, but at the experiment duration and spatial scales. Field and demonstration tests are also a valuable way to obtain information on the interactions occurring during CO₂ injection. However, only changes in gas and water compositions

are generally monitored during these tests and not changes on the rock. The last method is numerical geochemical and reactive transport modeling, which can be used to interpret and simulate natural analogues, laboratory experiments and field tests or to predict the behavior of a CO₂ geological storage, including over large time and spatial scales. In order to make accurate simulations and predictions, modeling requires a good description of the processes taking place and a precise characterization of the parameters involved in the calculations.

In geochemical modeling, the quality of the calculations is determined by the constraint of some key parameters and processes: the thermo-physical properties of CO₂, the interactions within the brine, the selection of the reacting mineral phases and their solubility properties and the kinetic laws and parameters (Marini, 2006; Gaus et al., 2008). Constraining these parameters is a challenging but essential task to understand and accurately simulate the geochemical evolution of a CO₂ storage. This work is focused on the selection of the minerals to consider in a model and on the influence of the different options offered to a modeler for describing the interaction processes within the solution.

Many modeling studies on the geochemical evolution of CO₂ storages including mineral changes have been carried out over recent years (Xu et al., 2003, 2010; Knauss et al., 2005; Lagneau et al., 2005; Zerai et al., 2006; Audigane et al., 2007, 2009). However, the selection of the minerals involved in the interaction processes does not always rest on a solid conceptual model specific to the studied storage site. Indeed, it is essential to first interpret in terms of reactivity the mineral and pore solution initial states and, then, based on natural system observations and on laboratory interaction experiments, to establish what reactions are expected when CO₂ is introduced into this particular water – rock system. All primary minerals are sometimes considered in the simulation (Xu et al., 2003; Zhang et al., 2009), without any consideration of the minerals really expected to react, and the behavior of the system is controlled by the thermodynamics and the kinetics of the reaction, in spite of their uncertainties. This kind of simulation leads to a complicated and certainly erroneous behavior where some primary minerals replace in several years or hundreds of years other primary minerals because they are thermodynamically more stable, although these minerals can coexist in natural systems over millions of years. Furthermore, the selection of the secondary minerals resulting from CO₂ injection into sandstone is still under debate, in particular for dawsonite. Under storage conditions, this Na-Al carbonate is thermodynamically favored to precipitate but is not often observed in CO₂ natural storage analogues and never in laboratory experiments at low pH and temperatures relevant for CO₂ storage. The purpose is then to be able to establish a solid conceptual model clearly pointing out the minerals likely to react in a given sandstone, in coherence with the overall observations from natural analogues and experiments, as a function of the mineralogical and

chemical properties of the studied formation (Gaus, 2010; Dethlefsen et al., 2011; Hellevang et al., 2011). Such a good constraint on the conceptual model appears all the more important in that it has been demonstrated that mineral trapping is highly influenced by the selection of the initial and secondary minerals (Kihm et al., 2012).

When performing geochemical simulations in deep saline aquifers, it should be recalled that the high pressure, temperature and salinity characterizing these aquifers confer to the system a different chemical behavior than in surface conditions. Conventional activity models of Debye-Hückel type are frequently used in simulations of CO₂ storage in deep saline aquifers, even though these models are only valid for dilute solutions. Likewise, thermodynamic databases are also used out of their temperature validity range or for chemical species without interaction coefficients in Pitzer databases. On this topic, the purpose of this paper is to evaluate the uncertainty of the activity model and the thermodynamic database used on the simulation results. This paper investigates the conditions to assess the geochemical behavior of CO₂ storage in deep saline aquifers aiming at improving the establishment of the conceptual model and at evaluating the effect of the activity model and thermodynamic database selection on the simulation results. These subjects are illustrated and discussed on the basis of three case studies: Ketzin, In Salah and Snøhvit CO₂ storage sites.

2 Geochemical modeling approach for CO₂ storage in saline aquifers

2.1 Conceptual model of long-term CO₂ evolution in saline sandstone aquifers: insights provided by natural analogues

In a CO₂ geological storage, CO₂ is injected into the host formation in a supercritical state and will distribute according to four trapping mechanisms, namely structural, residual, solubility and mineral trapping, respectively to the increase of storage safety (Benson and Cole, 2008). The distribution of CO₂ in these four traps will evolve over time, distance from the injection point and physical conditions (P, T, salinity) in the host formation. From a geochemical perspective and over long time spans, perturbations linked to CO₂ will mainly concern the dissolved CO₂ and the interaction with the rock.

The dissolution of CO₂ in brines leads to an acidification and an increase in bicarbonate content, which have important consequences regarding the evolution of the geochemical system in the saline

sandstone aquifer. Information on the resulting interactions is given by laboratory experiments and by the study of natural CO₂ geological disposal analogues. During laboratory interaction experiments between CO₂ saturated brines and sandstones or rock-constituting minerals (e.g., Kaszuba et al., 2003, 2005; Hangx and Spiers, 2009; Ketzer et al., 2009; Luquot et al., 2012), only fast reactions can be observed but system conditions are well controlled, allowing a precise characterization of the observed interaction processes. Natural analogues make it possible to observe the results of CO₂ – water – rock interactions over geological periods, mainly regarding the dissolution and precipitation of minerals obeying slow reaction kinetics. However, in natural analogues uncertainties remain regarding the fluid and gas compositions during the interaction with CO₂, the initial mineralogy, the temperature during interaction and the interaction duration. A compilation of natural analogues of CO₂ storage in saline aquifers is presented in Table 1. It emphasizes the changes in mineralogy that occurred during CO₂ intrusion in deep saline sandstone aquifers. It is necessary to consider both fast and slow CO₂ – water – rock reactions given by laboratory experiments and natural analogues to establish an interaction model. These insights make it possible to select the minerals expected to react or to precipitate during the interaction with CO₂ saturated brines, calibrate the model and compare the simulation results with data.

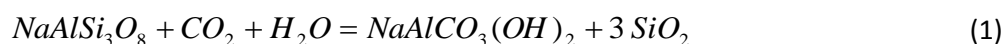
During the interaction of CO₂ saturated brines and sandstones calcite, dolomite and sulfate minerals dissolve rapidly and the solutions are close to equilibrium with these phases. Such fast reactions are observed during laboratory experiments and evidenced in natural analogues. Aluminosilicate primary sandstone minerals, such as feldspars, clay minerals, micas, chlorites or zeolites, also tend to dissolve because of the brine acidity. However these minerals have slow rates of dissolution and can persist over geological periods in thermodynamically unstable conditions. Natural analogues (Table 1) present dissolution footprints on primary aluminosilicate minerals but these minerals are still present even after several Ma of interaction. The dissolution of primary rock minerals induces a supply of cations in the brine, such as Ca, Mg, Na, Al, Fe and Si, which are available for precipitation of secondary minerals. Chalcedony or quartz are expected to precipitate relatively rapidly, since their precipitation is observed during CO₂ – sandstone interaction experiments accelerated using higher temperature and pressure than in the aquifers (Kaszuba et al., 2005) or even during experiments at *in situ* temperature and pressure (Luquot et al., 2012). With a few exceptions, quartz precipitation linked to CO₂ intrusion is observed in almost all natural analogues (Table 1). These exceptions are probably due to the difficulty of dating all quartz cementations in the rock in relation to their diagenesis event. Kaolinite is expected to precipitate as a result of primary aluminosilicate mineral

dissolution. Kaolinite precipitation is slow and is commonly observed in natural analogues but rarely during laboratory experiments (Ketzer et al., 2009; Luquot et al., 2012).

CO₂ mineral trapping occurs because of precipitation of carbonate minerals, in relation to the available dissolved cations in the brine. Commonly observed carbonate phases are calcite, dolomite, ankerite, siderite and dawsonite (Table 1). Precipitation of Ca, Mg and Fe carbonates are observed in all CO₂ storage analogues. Precipitation of dawsonite (NaAlCO₃(OH)₂) is not observed in all cases after interaction with CO₂ rich brines. The introduction of dawsonite into geochemical simulations of CO₂ storage in saline aquifers is controversial (Haszeldine et al., 2005; Hellevang et al., 2005, 2011; Kaszuba et al., 2011) since its precipitation is predicted by numerical simulations (Xu et al., 2003; Audigane et al., 2007) but dawsonite has rarely been observed in sedimentary basins (Smith and Milton, 1966; Baker et al., 1995). The thermodynamic properties of dawsonite were recently re-evaluated (Benezeth et al., 2007) and confirm that this mineral can be thermodynamically stable under CO₂ storage conditions (Cantucci et al., 2009). With regards to the thermodynamic database, it has also been pointed out that a consistent database for Al – phases and the Al system is required for a correct reproduction of dawsonite behavior (Kaszuba et al., 2011). Uncertainties regarding the kinetics of dawsonite precipitation and its nucleation have been suggested as a cause of the discrepancy between simulation predictions and field observations. The difference between rates measured during laboratory experiments and rates at the field scale is particularly uncertain to address (Zhu, 2005; Gannor et al., 2007). However, sensitivity analysis on the reaction rates and the nucleation rates (Hellevang et al., 2011) only indicate dawsonite precipitation is brought forward or postponed. After a long duration of interaction, dawsonite is predicted to precipitate. A dependence of the dawsonite precipitation rate on the aluminosilicate mineral dissolution rates has also been pointed out (Hellevang et al., 2010), because of a limited Al supply rate from aluminosilicates. It is difficult to estimate the time required for dawsonite to precipitate from the natural analogues. The only insight is given by the Bravo Dome analogue, where the absence of dawsonite can partly be explained by the too recent CO₂ intrusion in the arkosic sandstone, which occurred 8,000 to 100,000 years ago (Baines and Worden, 2004). However, the mineralogical assemblage of Bravo Dome analogue also suggests the sandstone underwent open-system alteration, leaching out the sources for dawsonite. Further natural CO₂ storage analogues have also been investigated in recent years and dawsonite precipitations were observed in various saline aquifers as a result of CO₂ rich brine – sandstone interaction (Moore et al., 2005; Worden, 2006; Gao et al., 2009; Wilkinson et al., 2009; Liu et al., 2011; Zalba et al., 2011). Dawsonite precipitation related to CO₂-rich fluid flooding has also been observed in carbonate veins in mudstone of the Cretaceous Izumi group (SW Japan),

which can be considered as a natural analogue of CO₂ storage caprock (Okuyama and Take, 2011). These studies on natural analogues indicate that dawsonite is more frequent than previously thought. It is also possible that dawsonite is missed in non-precise mineralogical and petrographical investigations since this mineral is characterized by a petrographical aspect similar to that of illites (Smith and Milton, 1966; Baines and Worden, 2004; Worden, 2006).

The influence of the initial composition of the sandstone can be considered to explain why dawsonite is sometimes observed in natural analogues and is sometimes absent. Dawsonite can precipitate from slightly acidic Na rich brines with high CO₂ contents and a source of Al (Worden, 2006; Benezeth et al., 2007; Hellevang et al., 2011). Only a relatively moderate CO₂ fugacity of some bars is required for the brine to be within the dawsonite stability zone (Worden, 2006). Studies on the reaction paths for dawsonite formation have shown that its precipitation follows the dissolution of an aluminosilicate mineral, such as a feldspar or a clay mineral (Johnson et al., 2001, 2004; Benezeth et al., 2007; Hellevang et al., 2011). Indeed, dawsonite is frequently observed to pseudomorphically replace feldspar crystals (Table 1). With regards to the temperature, dawsonite can form over a range from 25 to 120 °C (Table 1). An overall reaction is frequently proposed (Zerai et al., 2006; Audigane et al., 2007; Gaus, 2010) to represent the feldspar (NaAlSi₃O₈) evolution to dawsonite (NaAlCO₃(OH)₂) and chalcedony (SiO₂) in CO₂-rich brines. The reaction can be written as follows:



The compositions of the detrital phases of natural CO₂ storage analogue sandstones are shown in a Folk's diagram (Figure 1) with a distinction between dawsonite bearing sandstones and sandstones without dawsonite. The mineralogical composition of the total rock (i.e., not only including the detrital phases) is given in Table 1 for each analogue. Note that the Folk's diagram (Figure 1) shows some uncertainty regarding the composition of several analogues: i) the available mineralogical compositions of the Bravo Dome and Vert le Grand analogues are not detailed; ii) for the Miller Field analogue, relatively different mineralogical compositions are given by the different authors (Marchand et al., 2001; Baines and Worden, 2004); and iii) the Aldebaran sandstones of the Bowen, Gunnedah and Sydney basins present heterogeneities so that their compositions vary from litharenite to quartzarenite (Baker et al., 1995) but there are no details as to whether dawsonite is encountered or not in all facies.

From the Folk's diagram, which reports fourteen natural CO₂ geologic disposal analogues, a zonation of the sandstone compositions is highlighted regarding the occurrence or the absence of dawsonite. It appears that dawsonite bearing sandstones generally contain less quartz than sandstones without

dawsonite. Figure 1 also indicates that reactive minerals, such as clay minerals or volcanic or metamorphic derived phyllosilicates or tectosilicates (classified as lithics), may be necessary for dawsonite precipitation, in addition to feldspars. Indeed, these silicate rock fragments affect the silica activity and the pH, which are known to be crucial in dawsonite stability (Hellevang et al., 2011). On the other hand, it is observed that subarkose and arkose sandstones cannot lead to the precipitation of dawsonite consecutively to an interaction of the rock with CO₂-rich brines. In fact, mineralogical compositions of the whole rocks (Table 1) indicate that there are more feldspars in sandstones with dawsonite than in sandstones without dawsonite, in spite of being classified as subarkoses or arkoses. There are also more aluminosilicate minerals in dawsonite bearing sandstones. Immature sandstones (with a low SiO₂/Al₂O₃ ratio in aluminosilicates) seem consequently more adequate for dawsonite precipitation. Conversely, natural analogues suggest that no dawsonite can be expected in mature sandstones (with a high SiO₂/Al₂O₃ ratio).

Together with a diagram of the thermodynamic stability and the content of reactive minerals in the total rocks, this type of Folk's diagram compiling a set of natural CO₂ storage analogues seems interesting to evaluate the pertinence of introducing dawsonite to simulate the long-term interaction between a determined saline sandstone aquifer and CO₂. This diagram is further used for the selection of the minerals introduced in Ketzin, In-Salah and Snøhvit CO₂ storage site simulations.

2.2 Modeling aspects inherent to high salinity waters

2.2.1 Simulation code and activity models

In this study, the calculations were performed using the PHREEQC v2.18 code (Parkhurst and Appelo, 1999). PHREEQC is a computer program for simulating chemical reactions and transport processes in natural water systems. PHREEQC v2.18 allows high salinity solutions to be considered, such as sedimentary basin brines, using the Pitzer formalism as activity model. When concentrated solutions are considered, the solution ideality assumption is no longer valid and an activity coefficient is used to account for deviations from ideal behavior. Debye-Hückel (Debye and Hückel, 1923), extended Debye-Hückel, or Davies activity models are the most often used to calculate the activity coefficients. These models can provide an excellent description of the interactions occurring between solutes in solutions with an ionic strength lower than 0.5 to 0.7 M. In the case of higher concentrated solutions, the B-dot model (Helgeson et al., 1969), an extension of the Debye-Hückel model, can be used for NaCl dominant solutions. However, the more adequate formalism to consider the solution ideality deviation at high salinities is the Pitzer formalism (Pitzer et al., 1973; Pitzer, 1991). The Pitzer model considers both the electrostatic interactions between ions and the specific interactions between ion

pairs and triplets. These interactions amongst ions and solvent are introduced in Pitzer equations through linear combinations of parameters of a virial expansion of the excess Gibbs free energy. The determination of the Pitzer parameters is difficult, which limits the domain of application given the currently available thermodynamic databases.

2.2.2 Evaluation of thermodynamic databases

In the context of CO₂ geological storage in saline aquifers envisaged in this study, the range of ionic strengths is largely above the 1.0 M limit that “conventional” formalism can accept and the use of Pitzer equations is thus required to calculate the interaction energies within the solution. Considering Pitzer formalism is necessary to adequately predict the fluid rock interactions occurring during and after CO₂ injection. Very few applications of the use of Pitzer activity coefficient models for CO₂ geological storage can be found in the literature (Xu, 2008; Taberner et al., 2009). The purpose of the current study is to objectively evaluate the influence of databases and activity models on the simulation results based on the same initial data sets.

Before estimating this influence on three case-studies (Ketzin, Snøvit and In Salah), the efficiency of different thermodynamic databases to reproduce simple systems is evaluated. The solubility of simple minerals is calculated in saline solutions at different temperatures using different databases and compared with experimental data. Five thermodynamic databases are used for this comparison in simple geochemical systems. The purpose is to highlight the importance of considering Pitzer parameterization in highly saline solutions and to test the different databases including Pitzer coefficients. The six databases used in this study are: i) LLNL; ii) Thermoddem; iii) Pitzer.dat; iv) THEREDA; v) SandiaLab; and vi) SCALE2000 databases. The LLNL database corresponds to the “thermo.com.V8.R6.230” database (Johnson et al., 1992), produced by the Lawrence Livermore National Laboratory. Using the LLNL database, activity coefficients are calculated with the B-dot model. The Thermoddem database (Blanc et al., 2007, 2012), which is developed by BRGM, also uses the B-dot model to calculate the activity coefficients. The Pitzer.dat database is the default database provided with the PHREEQC code to calculate activity coefficients using Pitzer’s ion interaction theory. However, the use of the Pitzer.dat database is limited since few species are considered and the temperature is limited to 25 °C. The THEREDA database (Altmaier et al., 2011) includes Pitzer interaction coefficients but on few species. In particular, the THEREDA database cannot consider Al and Si species and C species for temperatures different to the standard temperature. The SandiaLab database corresponds to the “data0.ypf.R2” Pitzer database produced in the Yucca Mountain Project (USDOE, 2007) by the Sandia National Laboratory and reformatted to the PHREEQC format by Quintessa (Benbow et al., 2008). Finally, the SCALE2000 Pitzer database (Azaroual et al., 2004) has

been developed at BRGM for high saline solutions and brines (up to 300 g L⁻¹) at high pressure and temperature conditions. The last two Pitzer databases, i.e. the SandiaLab and SCALE2000 databases, contain most species of interest for performing calculations in deep saline sedimentary environments. However, the establishment of Pitzer databases is still on-going and there are no Pitzer interaction coefficients available for all species (e.g. for Al in the SCALE2000 database), or are valid only for a limited range of temperatures (e.g. many species have only interaction coefficients at 25 °C in the SandiaLab database), or incoherencies can be observed between the data. In particular, redox issues cannot be addressed using Pitzer databases since no interaction parameters describing the electron behavior in saline solutions are available at the present time. An attempt to deal with redox reactions in saline solutions using O₂ fugacity rather than electron activity has been undertaken by Marion et al. (2003). However, formalisms for redox reactions involving O₂ and related Pitzer interaction coefficients have not been introduced in the available databases. As a consequence, species which behavior is highly dependent to the redox, in particular Fe and S systems, cannot be considered in models when Pitzer formalism is used.

As a result, none of the currently available databases is totally appropriate for the purpose of the present study. Therefore, several databases are used to evaluate the uncertainties associated with the activity coefficient models. The comparison of the results calculated using these different databases with experimental data of simple mineral solubilities enables the accuracy of the different databases to be evaluated. Solubility measurements as a function of solution molarity were obtained from the literature for gibbsite in NaCl solutions from 30 to 70 °C (Palmer and Wesolowski, 1992), for silica in KCl, CaCl₂, MgSO₄ and MgCl₂ solutions at 25 °C (Marshall and Warakowski, 1980), for silica in NaCl solutions at 25 and 150 °C (Chen and Marshall, 1982), for calcite in NaCl solutions at 25 and 60 °C (Wolf et al., 1989) and for gypsum in NaCl solutions at 25 °C (Linke and Seidell, 1965). Solubilities were calculated for these minerals under the experimental conditions using the different databases previously presented. Calculations using the Pitzer.dat database were only performed for calcite and gypsum because this database does not contain Si and Al species.

As shown in Figure 2, calcite solubility as a function of solution salinity is well reproduced using the four Pitzer databases (only three databases at 60 °C). For the LLNL and Thermoddem databases, data reproduction is only satisfactory up to 0.5 and 1 mol NaCl L⁻¹, respectively. Similarly to calcite solubility, the gypsum solubility in saline NaCl solutions (Figure 3) can only be reproduced using the Pitzer databases. Reproduction is slightly improved using Pitzer.dat and THEREDA rather than the SCALE2000 and SandiaLab databases. In Figure 4, amorphous silica solubility is calculated and compared with experimental data in NaCl solutions at 25 and 150 °C, but also in KCl, CaCl₂, MgSO₄

and CaCl_2 saline solutions at 25 °C. Generally speaking, silica solubility experimental data are better reproduced using the SCALE2000 database. The range of dissolved silica and its trend with increasing salinity is reproduced satisfactorily with the SCALE2000 database in all solutions. Calculations using the SandiaLab database can provide satisfactory data reproduction, but strong discrepancies are also observed in some cases (in NaCl solutions at 150 °C and in KCl and CaCl_2 solutions). For salinities below 1 mol L^{-1} , experimental data are generally better reproduced using the Thermoddem database. At higher salinities, the Thermoddem database reproduces the trend of silica solubility with increasing salinity but significant discrepancies with the data are sometimes observed. Calculated silica solubility using the LLNL database remains constant with salinity. The reason of this constant silica solubility is due to the fact that the activity coefficient of uncharged species is calculated using the Setchenow equation and that the Setchenow parameter (b_i) has a fixed value in PHREEQC (Parkhurst and Appelo, 1999). However, the Setchenow parameter is dependent on the temperature and the salt in which the uncharged species dissolves (Marshall and Chen, 1982). When this salting-out effect is considered, using the b_i parameter determined for amorphous silica solubility in different salts and at different temperatures by Marshall and Chen (1982), the amorphous silica solubilities calculated using LLNL database as a function of the salinity are in much better agreement with the data (Figure 4). It illustrates the fact that the amorphous silica solubility evolution as a function of the salinity can be captured using a Debye-Hückel equation (Trudell and Jones, 1974). However, it requires an adjustment of the activity coefficient equation parameter for each condition of temperature and solution composition, which limits its application to natural systems. The SandiaLab database is the only database with Pitzer interaction coefficients for Al. However, as shown in Figure 5, experimental data reproduction is not necessarily better than with other databases. Nevertheless, the trend of gibbsite solubility with increasing salinity is reproduced only when using the SandiaLab database. The evolution of the Al^{3+} activity coefficient as a function of salinity shown in Figure 6 for the different databases in the same calculation conditions as in Figure 5-a indicates that Al behavior in solution depends on the presence or the absence of interaction parameters for Al. Indeed, in the Pitzer activity model formulation (Pitzer, 1991) part of the expression corresponds to an extended Debye-Hückel equation, including a dependence on the ionic strength, while the interaction parameters are involved in the other part of the expression. The differences between Al^{3+} activity coefficients calculated using the SandiaLab database, which includes Pitzer interaction parameters, and using the SCALE2000 database, without Pitzer interaction parameters, illustrates the influence of considering Al interaction parameters. Without interaction parameters, the calculated activity coefficient is similar to those obtained using the B-dot activity model (Figure 6).

The comparison of calculated and measured solubilities of simple minerals as a function of salinity highlights the importance of considering Pitzer interaction parameters in saline solutions. The SCALE2000 database provides the most coherent results including different temperatures and different ionic species, despite its lack of Pitzer interaction coefficients for Al species. The SandiaLab database contains a lot of elements with associated Pitzer interaction coefficients but results can be inconsistent, as observed for some ionic species and for temperatures different to 25 °C. Despite its aspect, the SandiaLab database cannot reliably be used for many problems, in particular those considering complex interactions and at temperatures different to 25 °C. The Pitzer.dat and THEREDA databases contain few species, which considerably limits their application range. In particular, the Pitzer.dat database is unable to consider geochemical systems encountered in siliciclastic sedimentary basins and the THEREDA database is also unable to consider C species as a function of temperature. These databases are not further used in this work for the case-studies of CO₂ storage in saline sandstone aquifers. Among the databases without Pitzer interaction parameters, the Thermoddem database provides the best results. The Thermoddem database is further considered in the case-studies, but not the LLNL database.

2.3 Proposed methodology

The following methodology is proposed to assess the fluid – rock interaction occurrences due to CO₂ injection in deep saline aquifers. A batch modeling approach is considered here, where no transport of elements is calculated and the geochemical system is assumed to be a “perfectly mixed reactor” in which every single chemical compound is homogeneously distributed. The modeling approach is based on 3 steps (Figure 7):

STEP 1: This first step consists in establishing the initial conditions in the aquifer by recalculating the brine composition in reservoir conditions. Once the charge balance of the sampled water has been controlled, the solution speciation is recalculated at formation temperature. In the case where an element of interest for the studied problem has not been analyzed, its content is adjusted to achieve equilibrium with a mineral phase showing evidence of equilibrium with the formation water (e.g. Al content adjusted to kaolinite equilibrium observed as the last cement formed in the rock). The saturation indices are checked to estimate what mineral phases are at equilibrium with the brine and whether these results are consistent with the mineralogical and petrological observations. The results obtained with the Thermoddem, SCALE2000 and SandiaLab databases are compared to evaluate the differences linked to databases. Note that the effect of pressure on the thermodynamic constants is not considered here to avoid introducing further uncertainties to the calculations, which can be difficult to constrain.

STEP 2: The second step consists in introducing the perturbation into the formation brine due to the CO₂ injection. The perturbation is introduced by solubilizing CO₂ in the brine at a given fugacity of CO₂. The fugacity of CO₂ is calculated using an equation of state (Duan and Sun, 2003) for 1.5 times the initial formation fluid pressure. Indeed, the CO₂ is planned to be injected at a pressure 1.5 times higher than the formation fluid pressure, at least in large reservoirs with high permeability. A mixing approach between a water acidified at the CO₂ injection fugacity end-member and a non-perturbed water end-member is then used to mimic the CO₂ impact at various distances from the injection well. Considering the different databases, the effect on pH is checked. The saturation indices are also computed to identify the mineral phases destabilized by the CO₂-induced acidification.

STEP 3: This last step consists in calculating the geochemical and mineralogical evolution of the system due to the CO₂ perturbation. For this purpose, a water-rock interaction model is proposed for each case-study. The goal is to quantify the mineral changes induced by the injection of CO₂ and, in particular, those contributing to CO₂ mineral trapping. Special attention is paid to the influence of the database on the calculation results. These interaction models are based on the characterization of the geochemical system for the different saline sandstone aquifers (step 2) and the identification of the minerals expected to dissolve or to precipitate, in agreement with the conceptual model previously presented (cf. section 2.1). The reactions introduced into the interaction model are summarized in Table 3. Dissolution of primary aluminosilicate minerals is kinetically controlled while thermodynamic equilibrium governs the precipitation of secondary phases. Kinetics are introduced for aluminosilicate dissolution because alteration of these mineral phases may last over very long time periods and equilibrium assumptions are not realistic. On the contrary, precipitation can be considered at equilibrium over long time periods for the sake of calculation simplicity (Helgeson, 1968). Precipitation is then controlled by the release of cations by dissolution of primary minerals (see section 2.1). Kinetics are introduced through a general transition state theory law using Arrhenius-type rate equations (Lasaga et al., 1994), which may be expressed as:

$$\frac{dm}{dt} = -A(1-\Omega)m \left(k_{acid}^{25} e^{-\frac{E_{acid}}{R} \left(\frac{1}{T} - \frac{1}{298.15} \right)} a_{H^+}^n + k_{neutral}^{25} e^{-\frac{E_{neutral}}{R} \left(\frac{1}{T} - \frac{1}{298.15} \right)} \right) \quad (2)$$

where, A is the reactive surface area ($m^2 mol^{-1}$), k^{25} are the reaction rate constants for acid and neutral mechanisms at 25 °C ($mol m^{-2} s^{-1}$), E are the activation energies ($J mol^{-1}$), R is the gas constant, T is the temperature (K), a_{H^+} is the proton activity, n is an empirical exponential term, m is the amount of considered mineral (mol) and Ω is the saturation state corresponding to the ratio between the ion activity product and the thermodynamic equilibrium constant. The reaction rates and

activation energies used in these calculations come from the compilation by Palandri and Kharaka (2004). Table 4 summarizes these parameters together with the reactive surface areas used for the different minerals. Important differences are usually observed between the dissolution rates measured in laboratory experiments, such as those reported by Palandri and Kharaka (2004), and long-term field dissolution rates (Steefel et al., 2005; Zhu, 2005; Gannor et al., 2007). In practice, this difference is often corrected by introducing a scaling factor of 0.001 on the reactive surface area measured on individual minerals at the laboratory scale (Gaus, 2005). Such a scaling factor is also used in the present calculations for the different case-studies.

Calculations were carried out over 1000 years considering the interaction between Ketzin, In Salah and Snøhvit sandstone and brines with acidified brine mixing ratios of 0, 5, 20, 50, 70 and 100 %. The storage aquifer temperature of 36 °C was considered and the SCALE2000, SandiaLab and Thermoddem databases were used. Calculations with a non-perturbed brine correspond to a baseline case without CO₂, making it possible to distinguish the brine-mineral interactions due to sandstone natural evolution from the interactions due to the CO₂ intrusion.

3 Case-studies for evaluating the long-term geochemical evolution of CO₂ in saline aquifers

3.1 Ketzin

3.1.1 Site presentation

Ketzin is an onshore site located in the Northeast German Basin for pilot injection of CO₂ into an 80 m thick saline sandstone aquifer (Förster et al., 2006). The Ketzin injection project was funded by the European Commission under the FP6 framework (CO2SINK Project). A modest amount of CO₂ was injected (60,000 tons) into an anticlinal structure of the Triassic Stuttgart Formation located at a depth of about 650 m above a salt pillow situated deeper (Figure 8). The Stuttgart Formation is of fluvial origin and exhibits a heterogeneous lithology showing alternations of mudstones, siltstones and fluvial sandstones (Förster et al., 2006, 2010). This formation has good reservoir properties, with an average porosity of 23 % and a permeability of 50 to 100 mD (Wiese et al., 2010). Reservoir temperatures of 33 and 36 °C were measured at depths of 600 and 700 m, respectively. Formation pressure at 700 m depth is determined to range between 70 and 75 bars. The brines sampled in the formation have a salinity of about 230 g L⁻¹ (Würdermann et al., 2010). The Stuttgart formation is overlaid by mudstones of the Weser and Arnstadt formations, which form a 210 m thick caprock section (Förster et al., 2006). At Ketzin experimental site, food grade CO₂ (purity 99.9 %) and CO₂ from an oxyfuel pilot plant (purity 99.7%) were injected (Martens et al., 2012).

3.1.2 STEP 1: Initial reservoir conditions at the Ketzin site

A detailed characterization of the Ketzin site has been carried out before injecting CO₂ (Förster et al., 2006). In particular, the evaluation of the water – rock interactions at the initial state in the Stuttgart Fm can be confidently addressed since several water sampling analyses (Würdermann et al., 2010) and a detailed mineralogical and petrological analysis (Förster et al., 2010) are available. Formation water was sampled (Würdermann et al., 2010) after pumping several times the borehole volume to obtain a water sample representative of the formation. The composition of water once the borehole volume was pumped 6.8 times (KTZI 202, No 26) is reported in Table 2 and used in this study. Waters sampled after 5 times or more the borehole volume was pumped have similar compositions. Fluorescein was used as a tracer to ensure the sampled waters were not perturbed by the drilling mud (Würdermann et al., 2010). Stuttgart Fm water has a total dissolved solids (TDS) content of about 235 g L⁻¹ and is mainly composed of Na and Cl, with noticeable amounts of Ca, SO₄ and Fe. Note that Al content was not determined. Charges are balanced within the solution, which has a 1.4 % error.

At Ketzin, the Stuttgart Fm presents channel and overbank facies, respectively corresponding to interbedded sandstones and thin mudstone and coal layers. Mineralogy composition is obtained from Förster et al. (2010). Quartz content ranges from 30 to 40 %. The sandstone has a high feldspar content (from 20 to 40 %), mainly plagioclase but also K-feldspar. Mg-, Fe-, K-sheet minerals, including chlorite, illite, mixed layered illite/smectite and micas are also present in the sandstone at relatively important amounts (between 10 and 15 %). These phyllosilicates can be detrital or authigenic minerals. Amongst the authigenic minerals, analcime (a Na-zeolite) and anhydrite are the most frequent and most abundant phases. Further authigenic minerals observed at low amounts, but not in all samples, are feldspar, quartz, ferric iron oxides, clay minerals and dolomite. Petrological observations, in particular with regards to the mineral crystallinity state or the presence of dissolution footprints and overgrowths, give reliable information on the equilibrium of the mineral phases respectively to the brine in the formation conditions. At Ketzin (Förster et al., 2010), feldspar grains are partially or totally altered and replaced by illite or by other authigenic cements. An albitization of K-feldspar and plagioclase also occurred as an early diagenetic process. Lithics are strongly altered, mainly the illite and chlorite-rich grains and the feldspar-rich rock fragments. The first minerals to precipitate during diagenesis were quartz, albite and iron oxides. These precipitations were followed by precipitations of clay minerals and analcime. Dolomite then precipitated, followed by anhydrite as the latest authigenic mineral.

The last aluminosilicate minerals to precipitate were illite and analcime and, even if these phases are not the last phases to precipitate during diagenesis, they do not show dissolution footprints. Illite can then be considered at equilibrium with the actual fluids circulating in the formation. Based on this hypothesis, Al content is adjusted to achieve solution equilibrium with illite (Mg-illite for the SCALE2000 and Thermoddem databases and Mg, K-illite for the SandiaLab database). When the solution speciation is recalculated at the formation temperature (36 °C), pH values of 6.40, 6.36 and 6.32 are obtained with the SCALE2000, SandiaLab and Thermoddem databases, respectively. These values are close to the 6.40 pH value measured during sampling (Table 2). Carbon dioxide fugacity log values of -1.86, -1.92 and -1.86 are calculated using the different databases. This narrow range of CO₂ fugacities can be compared with the CO₂ fugacity values measured in sedimentary basins and reported as a function of temperature by Coudrain-Ribstein et al. (1998). Calculated CO₂ fugacity values for the Ketzin injection site at a temperature of 36 °C are in the range of typical CO₂ fugacity in sedimentary formations for the corresponding temperature. This comparison suggests that the measured pH value is reliable and that the carbonate system is reliably constrained for this case-study. Indeed, the pH and carbonate system can be easily altered during sampling in a deep formation, mainly because of CO₂ outgassing.

Figure 9 shows the saturation indices of calcite, dolomite (medium-ordered dolomite), microcline, albite (low-ordered albite), anorthite, a Mg-chlorite (clinochlore), a Na-zeolite (analcime), quartz, anhydrite, kaolinite and dawsonite calculated using the SCALE2000, SandiaLab and Thermoddem databases. Stuttgart Fm porewater, with Al content adjusted to achieve illite equilibrium, is close to equilibrium with microcline, albite, analcime, quartz, anhydrite and dawsonite. A good agreement between databases is observed for these phases. Dolomite is slightly oversaturated when using the SCALE2000 and SandiaLab databases but undersaturated using the Thermoddem database. A similar difference in saturation state is obtained for kaolinite, which is oversaturated according to the SandiaLab and Thermoddem databases but is undersaturated using the SCALE2000 database. Anorthite and clinochlore are strongly undersaturated with the three databases. However, an important difference is observed between clinochlore saturation indices calculated with the SCALE2000 database and using the SandiaLab and Thermoddem databases. One can then expect different evolutions of the system after CO₂ perturbation, depending on the database.

3.1.3 STEP 2: Acidification of Ketzin brine by CO₂

In this second step, the effect of CO₂ on the formation brine is simulated, without considering interactions with the rock at this stage. Under the injection pressure (1.5 times the formation pressure of about 75 bars) and temperature (36 °C) conditions at the Ketzin injection site, CO₂

fugacity is 56.8 bars. The initial formation brine is equilibrated with such a CO₂ fugacity to calculate the brine composition and speciation with maximal CO₂ perturbation. Several mixings between CO₂-perturbed (100 %) and non-perturbed (0 %) end-member brines are then computed, leading to brines acidified at different levels.

Speciation, pH and saturation indices are calculated for the different resulting solutions using the SCALE2000, SandiaLab and Thermoddem databases (Figure 10). pH calculations show a decrease from about 6.4 for the non-perturbed brine to about 3.1 with the SCALE2000 and Thermoddem databases or 4.39 with the SandiaLab database for the acidified brine. pH strongly decreases even for solutions with low percent of acidified brines. Indeed, for a 5 % acidified brine ratio the pH is calculated at 4.21, 4.13 and 6.07 using the SCALE2000, Thermoddem and SandiaLab databases, respectively. This decrease is to be compared with the increase in CO₂ fugacity observed for mixed solutions. In Figure 10, calcite has been elected to show the carbonate mineral saturation index evolution and microcline for the aluminosilicate minerals. Saturation indices of these different minerals are calculated to strongly decrease when acidified brines are considered, because of the pH decrease. A decrease in saturation index is observed both for silicate and carbonate minerals, in spite of the increase in dissolved carbonate content. Differences in saturation index evolution are observed between the different databases. Higher saturation indices are calculated using the SandiaLab database for which higher pH are calculated. Differences in saturation index are also related to the considerations of ion interactions, since differences are observed between the SCALE2000 and Thermoddem databases, although pH are similar in both calculations.

3.1.4 STEP 3: Evaluation of CO₂ – water – rock interactions at Ketzin by batch modeling

This step aims at estimating the long-term geochemical and mineralogical interactions between the CO₂ perturbed brine and Ketzin injection site sandstone. An interaction model is proposed for the Ketzin site, based on the sandstone aquifer mineralogy and the observations derived from CO₂ storage natural analogues and laboratory experiments, presented in section 2.1. Ketzin sandstone has an important content in feldspars and aluminosilicate minerals such as chlorite, illite and zeolite (Förster et al., 2010). As indicated by the feldspar alteration observations, feldspars are unstable under present-day conditions in the formation. Feldspars are all the more unstable when the brine is acidified as a consequence of CO₂ introduction (Figure 10) and microcline saturation indices indicate that microcline is slightly more unstable than plagioclase. Chlorite (clinochlore) also shows alteration indications and is calculated to be strongly undersaturated in initial Ketzin brine (Figure 9) and in acidified brines (Figure 10). Illite and analcime precipitated during sandstone diagenesis and are

considered at equilibrium with the aquifer porewater (Figure 9 and section 3.1.2). These aluminosilicates were identified in section 2.1 as reactive phases likely to dissolve as a result of interaction with CO₂-rich water and saturation indices calculated for the CO₂ perturbed Ketzin brine indicate that these phases are unstable when a CO₂ perturbation is considered (Figure 10). Microcline has been selected as reactive feldspar because this phase is abundant in Ketzin sandstone and is already partially altered and, consequently, more sensitive to another perturbation. Microcline, clinocllore, analcime and Mg-illite are then introduced in the interaction model as phases that can dissolve (Table 3). With regards to the minerals that can precipitate as a result of high CO₂ content in water and the dissolution of primary minerals, kaolinite, quartz or chalcedony and carbonate minerals were identified in section 2.1 to precipitate in almost all natural analogues and during laboratory experiments. Kaolinite, chalcedony and dolomite are then introduced into the interaction model for Ketzin as phases allowed to precipitate. Chalcedony has been preferred over quartz because it can form more easily and is often observed as secondary silica phase. Dolomite has been selected because precipitations of Ca, Mg (and Fe) carbonates are generally observed in the natural analogues rather than pure Ca or Mg carbonates. Furthermore, Mg is expected to be released in solution if clinocllore and Mg-illite dissolve. In addition to these secondary phases, dawsonite is also introduced into the CO₂-water-rock model for Ketzin as a potential secondary phase. This choice has been made because Ketzin sandstone has large amounts of reactive aluminosilicate minerals and sandstones with similar composition that have been subject to CO₂ perturbation exhibit dawsonite precipitation. In particular, Ketzin sandstone has a composition similar to the sandstone of the Songliao Basin analogue where dawsonite has been observed (Liu et al., 2011). At Ketzin reservoir temperature (36 °C), dawsonite precipitation can perhaps be slow (Hellevang et al., 2011), but dawsonite precipitation has been observed in natural analogues for temperatures ranging for 25 to 120 °C and can then take place at Ketzin.

Calculation results for the Ketzin injection site are shown in Figure 11, where the cumulated mineral evolution over 1000 years is expressed as variations in mineral %_{wt} within the rock. Analcime is the main mineral to dissolve as a result of sandstone interaction with CO₂-rich brine. Up to 2 % of analcime is calculated to dissolve for the acidified brine. Analcime dissolution is less important when a mix of initial and acidified brines is considered and its dissolution increases regularly with the ratio of acidified brine. Other minerals allowed to dissolve evolve at low and non-representative amounts. A maximum of 0.016 % of illite is predicted to dissolve while less than 0.0001 % of clinocllore and 0.000008 % of microcline are calculated to dissolve (not shown in Figure 11). The main simulated precipitations are chalcedony and dawsonite (up to 1 % each for the acidified brine). Dissolved and precipitated amounts increase regularly with the ratio of acidified brine. Some differences in the

results are observed between the different databases. A little more analcime is calculated to dissolve using the SandiaLab database and a little more chalcedony also precipitates. Using the SandiaLab database, limited amounts of dolomite and kaolinite are also calculated to precipitate, while these minerals are not observed (or at very small amounts) with the two other databases. More noticeable differences between databases then occur for minerals precipitating or dissolving at very limited amounts. When the initial brine is considered, mineralogy does not evolve or does so to a very limited extent.

Regarding the evolution of the modeled system over time, in the present batch conditions without sustaining the CO₂ perturbation, aluminosilicate saturation indices become positive or null after relatively few dissolution steps. Dissolution then stops or becomes very limited after several tens of years. The time needed to reach system equilibrium tends to increase when more acidified brines are considered. Brine pH values increase during system evolution and these values are similar once system equilibrium is reached, whatever the acidified brine mixing ratio of the solution. Final pH values are 6.30, 7.46 and 6.66 using the SCALE2000, SandiaLab and Thermoddem databases, respectively. Solute component contents remain relatively stable over system evolution with a small increase in Al and Mg, but most components released by aluminosilicate dissolution are calculated to reprecipitate. Even the decrease in carbonate content due to dawsonite and dolomite precipitation tends to be limited.

3.2 In Salah

3.2.1 Site presentation

In Salah is located in the Sahara desert in Algeria and the methane production and CO₂ injection project is developed jointly by BP, Statoil and Sonatrach at the Carboniferous Krechba gas field. The produced gas contains up to 10 % of CO₂ while export sales require concentrations less than 0.3 %. To achieve this limit, 17 Mt of CO₂ have been separated and geologically stored over 20 years since 2004. At the In Salah site (Figure 8), a 20 m thick sandstone of Tournaisian age known as 'C10' sandstone forms the principal reservoir in an anticlinal structure for gas production but also for CO₂ injection. It is overlaid by over 950 m of Carboniferous mudstones forming an efficient seal. The producing wells are located in the hydrocarbon gas column on the crest of the structure and CO₂ is injected through 3 wells on the flanks of the reservoir, below the gas – water contact. The reservoir lies at depths of around 1800 m below the surface and comprises fine-grained quartzose sandstones. According to the site description given within the CO₂ReMoVe FP6 European project (Wildenborg et al., 2009; Vong et al., 2010), porosities range from 13 to 20 % and permeabilities are highly variable

as a function of the chlorite content, but average around 10 mD. Temperature and pressure in the reservoir are about 96 °C and 180 bars, respectively. Brine salinity is about 170 g L⁻¹, with Na, Ca and Cl as main components. The injected gas is composed of CO₂ at concentrations higher than 99 % and by trace amounts of methane, nitrogen and other components (including heavy alkanes, benzene and toluene).

3.2.2 STEP 1: Initial reservoir conditions at the In Salah site

The composition of a water sample taken in November 2004 at the bottomhole of one of the CO₂ injection wells (Kb502) has been provided through the CO2ReMoVe project as representative of the 'C10' Tournaisian sandstone porewater (Table 2). This water has a TDS of about 175 g L⁻¹ and is mainly composed of Na, Ca and Cl, with noticeable amounts of Mg. Note that the Al content is missing for this water. Solution electroneutrality is achieved, with an error of less than 0.1 %.

The Tournaisian sandstone at the In Salah injection site is mainly composed of quartz (between 69.9 and 88.5 %). Further detrital phases are mica and illite (about 5 %). The main authigenic minerals are Fe-rich chlorite (from 5 to 10 %) and siderite (from 1.2 to 11.7 %). Small amounts (less than 1 % each) of pyrite, dolomite and kaolinite also compose the In Salah reservoir. Diagenesis consisted first in cementation of chlorite, siderite, Fe-dolomite and pyrite. This first cementation was followed by chalcedony precipitation and, then, by a dissolution event which mainly affected the detrital clays, chlorite and pyrite. The last observed diagenetic event was a minor precipitation of kaolinite.

As the last authigenic mineral to precipitate, kaolinite can be assumed to be at equilibrium with present-day formation water. Therefore, the missing Al content is adjusted to achieve kaolinite equilibrium. Solution speciations recalculated at reservoir temperature (96 °C) with the SCALE2000, SandiaLab and Thermoddem databases give pH values of 5.55, 5.16 and 5.33, while the pH value measured after sampling is 5.20 (Table 2). Calculated carbon dioxide fugacities are similar, with log values of -0.57, -0.51 and -0.49 using the SCALE2000, SandiaLab and Thermoddem databases, respectively. These values are in the range of CO₂ fugacities measured in sedimentary basins for a temperature corresponding to the In Salah reservoir temperature (Coudrain-Ribstein et al., 1998), suggesting pH and carbonate system are reliable for the In Salah formation water.

Figure 12 shows the saturation indices calculated at 98 °C with Al content set at kaolinite equilibrium using the different databases for calcite, dolomite, illite, clinocllore, quartz, anhydrite and dawsonite. According to these calculations, calcite, dolomite, quartz and anhydrite are close to equilibrium in the brine. Illite, clinocllore and dawsonite equilibrium states are uncertain because of differences between databases. A good agreement between databases is obtained for quartz and

anhydrite saturation indices. For calcite and dolomite, slight differences are observed but minerals are oversaturated using the SCALE2000 database and undersaturated using the SandiaLab and Thermoddem databases. Strong differences in aluminosilicate saturation indices are calculated according to the database used. The same reaction constants for illite and clinocllore are used in the SCALE2000 and Thermoddem databases but not in the Sandialab database. These differences in saturation indices are then partly due to reaction constant differences but mostly to differences related to the ion interaction treatments, as illustrated by the differences in the Al^{3+} activity coefficient calculated in Figure 6 depending on whether Pitzer interaction parameters for Al are considered or not. A difference in saturation indices is also observed for dawsonite, because of the differences in Al behavior in solution when the different databases are used.

3.2.3 STEP 2: Acidification of In Salah brine by CO₂

Acidification of the In Salah injection site brine is made considering a CO₂ fugacity of 143.8 bars, corresponding to the fugacity calculated for 1.5 times the 180 bars formation pressure at a temperature of 96 °C. In Figure 13, pH is shown to decrease from 5.55 to 3.43 using the SCALE2000 database, from 5.16 to 2.80 using the SandiaLab database and from 5.33 to 3.06 using the Thermoddem database. pH decreases significantly even for low acidified brine ratios. Saturation indices are also calculated to decrease with respect to their initial values in the reference brine. Noticeable differences in saturation indices for acidified brines are obtained with the different databases. For 100 % acidified brine, calcite saturation index is -1.2 with the SCALE2000 database, -2.4 with the SandiaLab database and -2.2 with the Thermoddem database. For illite, this difference is even more pronounced with a saturation index of around -11 with the SCALE2000 and Thermoddem databases and a saturation index of -18.5 using the SandiaLab database. Differences in saturation index increase with the percent of acidified brine. Carbon dioxide fugacities of the mixed solutions are similar using the different databases and notably increase even for solutions with the lowest acidified brine mixing ratios.

3.2.4 STEP 3: Evaluation of CO₂ – water – rock interactions at In Salah by batch modeling

The interaction model for simulating interactions between CO₂-rich brine and the sandstone at the In Salah injection site is based on the rock mineralogy. In Salah sandstone is mainly composed of quartz, which is a phase with a relatively weak reactivity. The rock forming aluminosilicate minerals expected to dissolve as a consequence of interaction with CO₂-rich brine are illite, mica and chlorite. Note that there are no feldspars in this sandstone. One of the latest events during rock diagenesis was dissolution of clays and chlorite. Therefore, these phases are not at equilibrium with the brine prior

to CO₂ injection. Because of database differences, calculated saturation indices (Figure 12) cannot confirm these petrological observations. Furthermore, these phases are likely to dissolve in contact with acidified brines (Figure 13 and section 2.1). Mg-illite and clinochlore are then considered as the phases able to dissolve in the interaction model for In Salah (Table 3). Carbonate minerals (siderite and dolomite) precipitated during the rock evolution and can precipitate in CO₂-rich brine when cations are released by aluminosilicate dissolution. Dolomite is then considered in the model as a mineral allowed to precipitate. Chalcedony and kaolinite are also allowed to precipitate as a consequence of illite and clinochlore dissolution (section 2.1). Dawsonite precipitation seems unlikely in In Salah sandstone. Indeed, this sandstone has little amounts of reactive minerals and the study of CO₂ storage natural analogues showed that dawsonite is observed to precipitate in sandstone with elevated relative contents in aluminosilicates (section 2.1). In Figure 1, the In Salah composition is covered in the composition range of the Bowen-Gunnedah-Sydney Basin analogue. This analogue contains dawsonite but a heterogeneous composition is reported for it (Baker et al., 1995) without being precise whether dawsonite is observed for all sandstone types. Given the observations provided by other analogues, it is not very probable that dawsonite precipitates in a sandstone with In Salah sandstone composition and, therefore, it will not be considered in the interaction model.

Results of mineralogy changes calculated with the interaction model are shown in Figure 14. Important differences are observed depending on the database. Up to 3 % of illite is expected to dissolve using the SandiaLab database, while approximately half this amount dissolved using the Thermoddem database and less than 0.1 % of illite is found to dissolve using the SCALE2000 database. The second mineral allowed to dissolve, namely clinochlore, dissolves at limited amounts, 0.15 % at a maximum, for all the databases. Precipitations are influenced by the noticeable differences in dissolution between databases. Indeed, up to 2.5 % of kaolinite and up to 0.6 % of chalcedony precipitate using the SandiaLab database, while about half this amount precipitates using the Thermoddem database. Using the SCALE2000 database, no kaolinite is calculated to precipitate and less than 0.1 % of chalcedony precipitates. Precipitation of dolomite is also predicted, with less difference between databases. The evolution of dolomite precipitation with the increase in the acidified brine mixing ratio seems irregular, contrary to illite dissolution evolution and kaolinite and chalcedony precipitation evolutions, which are regular. No significant mineralogical modifications are simulated when the initial brine is considered in the interaction model.

In the case of In Salah CO₂ injection simulation, very different results are obtained as a function of the databases. While limited mineralogical modifications are expected using the SCALE2000 database, relatively important changes are observed using the SandiaLab database. However, the

most important differences do not concern dolomite precipitation, which contributes to CO₂ mineral trapping.

The water evolution shows a pH increase during the simulations (Figure 14) but, contrary to the Ketzin case, different final pH values are obtained depending on the acidified brine mixing ratio of the initial solution. The pH value further increases using the database predicting the most reactive system (i.e. the SandiaLab database), up to being buffered at non-perturbed brine pH values for the solutions with the lowest acidified brine mixing ratios.

3.3 Snøhvit

3.3.1 Site presentation

Snøhvit is an offshore CO₂ injection site located in the south western Barents Sea in Norway in the central part of the Hammerfest Basin. CO₂ is re-injected after its separation from methane produced at the Snøhvit, Albatross and Askeladd fields, which contain 5 to 8 % of CO₂. The Snøhvit production and injection site is operated by Statoil and owned by Statoil, Petoro, Total E&P, GDF Suez, Hess and RWE-Dea. Injection started in 2008 and about 23 Mt of CO₂ are planned to be injected over the 30 year lifespan of the project. The reservoir lithostratigraphic formations are late Triassic – Middle Jurassic formations called Fruholmen, Tubåen, Nordmela and Stø and are mainly composed of sandstones interbedded with thin shale layers (Linjordet and Grung Olsen, 1992). They are overlaid by Jurassic shale and thick Cretaceous shale caprocks (Figure 8). Natural gas is extracted from the Stø Fm reservoir and CO₂ is re-injected into the Tubåen saline aquifer and into the Stø Fm since 2011 after unexpected pressure increase in the Tubåen Fm (Hermanrud et al., 2013). The Tubåen aquifer has a thickness of about 45 to 75 m and is situated below the Stø Fm, at a depth of 2600 m. The Nordmela formation is sandwiched by the Tubåen and Stø formations and a shale-rich level of the Nordmela Fm acts as a caprock for the CO₂ storage. The Tubåen Fm is dominated by sandstones with subordinate shales and minor coals. The Tubåen aquifer porosity and permeability range from 10 to 15 % and 185 to 883 mD, respectively. Reservoir pressure and temperature are 285 bars and 98 °C, respectively. A high salinity of about 160 g L⁻¹ is measured in the Tubåen aquifer. The injected gas is mainly composed of CO₂ (99.8 %) with trace amounts of methane and heavy alkanes (Vong et al., 2010).

3.3.2 STEP 1: Initial reservoir conditions at the Snøhvit site

Relatively little information is available on the Tubåen Fm at Snøhvit, where CO₂ is injected. However, sufficient data for its geochemical characterization can be deduced from the overlying Stø Fm exploited gas reservoir, which is better characterized. Some information from the Stø Fm can be used

for the Tubåen Fm because no sedimentary hiatuses are observed between these two formations and because X-ray, neutron porosity and sonic logs (Linjordet and Grung Olsen, 1992) show similar trends for both formations.

The composition of a brine sampled by Statoil in the Tubåen Fm (well 7121/4-1) prior to CO₂ injection is reported in Table 2. Contents in minor components such as Ni, Fe and SiO₂ were deduced by Rochelle et al. (2007) from CO₂ – rock interaction experiments using Tubåen Fm sandstone. However Al content is missing in the water analysis. Tubåen Fm brine has a TDS of about 160 g L⁻¹ and is mainly composed of Na and Cl, with subordinate content in Ca. Brine is charge balanced, with a 0.2 % error.

No precise mineralogical and petrological descriptions of the Tubåen Fm sandstone are available. However the Stø and Tubåen formations have similar lithologies (Linjordet and Grung Olsen, 1992), with more argillaceous layers in the Tubåen Fm than in the Stø Fm. The mineralogy of the Stø Fm (Linjordet and Grung Olsen, 1992) is consequently considered here for the Tubåen Fm. Detrital minerals are mainly quartz with 80 to 90 % of monocrystalline quartz, polycrystalline quartz and chert. Further detrital phases are mica, K-feldspar, plagioclase, metamorphic and sedimentary rock fragments and heavy minerals. Quartz is the main cementing material. Cement is also composed of ankerite, dolomite, calcite, kaolinite, illite and pyrite.

In the absence of a precise diagenetic evolution description, illite is assumed to be at equilibrium with the actual formation water. Indeed, kaolinite and K-feldspar are expected to react together to form illite, which is observed as an authigenic mineral in the sandstone. Missing Al content is then adjusted to achieve solution equilibrium with Mg-illite. Solution speciation is recalculated at reservoir temperature (98 °C) using the SCALE2000, SandiaLab and Thermoddem databases. Calculated pH at reservoir temperature (6.19, 6.11 and 6.21, with the respective databases) is similar to the pH measured at surface conditions (6.20). Carbon dioxide fugacity log values of -0.42, -0.44 and -0.40 are calculated using the SCALE2000, SandiaLab and Thermoddem databases, respectively. These CO₂ fugacity values are in the range of measured and predicted CO₂ fugacity values in sedimentary basins at Tubåen Fm temperature (Coudrain-Ribstein et al., 1998). This observation suggests that outgassing and perturbation of the carbonate system equilibrium were limited during brine sampling. This brine can consequently be considered as representative of the Tubåen Fm porewater.

Saturation indices of calcite, dolomite, microcline, albite, kaolinite, Mg-chlorite, quartz, anhydrite and dawsonite calculated using the SCALE2000, SandiaLab and Thermoddem databases are shown in Figure 15. Saturation indices of aluminosilicate minerals (kaolinite and chlorite) and dawsonite calculated for Snøhvit injection reservoir water show noticeable differences, depending on the database used for the calculations. In particular, saturation indices calculated with the SCALE2000

database are different to saturation indices calculated using the SandiaLab and Thermoddem databases. Similar saturation indices are calculated with the three databases for further minerals shown in Figure 15 (calcite, dolomite, quartz and anhydrite). These differences concern only Al-bearing minerals, suggesting they are related to differences in Al behavior using the SCALE2000, SandiaLab and Thermoddem databases. Aluminosilicates are, depending on the database, slightly oversaturated to strongly oversaturated in the Tubåen Fm porewater. Calcite and dolomite are also oversaturated, while quartz is at equilibrium and anhydrite is slightly undersaturated.

3.3.3 STEP 2: Acidification of Snøhvit brine by CO₂

For the Snøhvit injection site case-study, a CO₂ fugacity of 207.9 bars is calculated considering 1.5 times the 300 bars formation pressure at a temperature of 98 °C. As a consequence of brine equilibrium with such a CO₂ fugacity, the pH value is calculated to decrease from about 6.3 to about 3.5 (Figure 16). pH values calculated with the three different databases show small differences. Small differences between databases are also noted for the calcite saturation index (Figure 16), which decreases to a minimal value of -1.6. Stronger differences are observed for aluminosilicate mineral saturation indices. For example, microcline saturation index for 100 % acidified brine is -7.2 and -9.7 using the SCALE2000 and SandiaLab databases, respectively, while it is only -3.9 using the Thermoddem database. Same CO₂ fugacities are calculated for a same mixing factor using the three databases.

3.3.4 STEP 3: Evaluation of CO₂ – water – rock interactions at Snøhvit by batch modeling

The same approach as in previous case-studies is undertaken to evaluate the long-term interactions between sandstone and CO₂-rich brine at the Snøhvit injection site. The Snøhvit target reservoir for CO₂ injection is mainly composed of quartz, both as detrital component and as cement. Rock-forming aluminosilicate minerals likely to react with CO₂-rich brine are feldspars, mica, chlorite (as metamorphic and sedimentary fragment) and illite (section 2.1). Furthermore, these minerals are undersaturated in acidified Snøhvit brine (Figure 16). In the interaction model for Snøhvit, microcline and Mg-chlorite are introduced as minerals allowed to dissolve (Table 3), following kinetic laws. Ca-, Mg- and Fe-carbonates, kaolinite and quartz are already present in the sandstone reservoir and these minerals can precipitate as a consequence of rock interaction with CO₂-rich brine (section 2.1). Dolomite, kaolinite and chalcedony are allowed to precipitate in the interaction model, at equilibrium. Dawsonite is not expected to form because the Snøhvit sandstone initial composition, with little amounts of reactive minerals, does not correspond to any sandstone composition in which dawsonite is observed, according to CO₂ storage natural analogues (section 2.1 and Figure 1).

Relatively limited amounts of minerals are calculated to dissolve and precipitate in the Snøhvit case (Figure 17). At most, for the acidified brine, 0.15 % of microcline and 0.07 % of clinocllore dissolve. Chalcedony, dolomite and kaolinite precipitate in a similar range of proportions and at limited amounts; i.e. less than 0.1 % at a maximum for each mineral. Note that no kaolinite precipitates using the SCALE2000 database, while the most important dolomite precipitations are calculated with this database. Largest microcline dissolutions are obtained using the SandiaLab database, as well as the most important chalcedony and kaolinite precipitations.

During system evolution, brine pH increases but its variation is lower than for the Ketzin and In Salah cases. This low pH buffering by the Snøhvit geochemical system is related to the lower reactivity of the system compared with the other case-studies. Using the different databases, pH values range in approximately 0.5 pH units.

4 Discussion

The characterization of the baseline geochemical systems of Ketzin, In Salah and Snøhvit CO₂ injection sites has highlighted three cases that are relatively different regarding their reactivity. Ketzin sandstone has high contents of reactive minerals while the two other sandstones are mainly composed of quartz. Amongst the two quartz-rich sandstones, the In Salah initial porewater is more acidic than the Snøhvit porewater, suggesting that dissolution of minerals in In Salah sandstone is likely to be more pronounced than in Snøhvit sandstone. Based on the mineralogical and petrological observations, a reactive model has been proposed for each case-study in the presence of CO₂. For dissolution, these models consider mineral phases present in the rock and expected to be altered by a CO₂-rich brine according to the general conceptual model based upon the laboratory experiments and natural analogue observations. Several aluminosilicates were selected for dissolution for each site but these minerals did not necessarily dissolve because the dissolution of an aluminosilicate can prevent another aluminosilicate from dissolving by switching its thermodynamic state to oversaturation. Dolomite, chalcedony and kaolinite were selected in all cases because these phases are commonly observed as secondary minerals in CO₂ storage natural analogues for all sandstone compositions. Dawsonite has been selected only for the Ketzin storage site, the reservoir of which contains large proportions of reactive minerals such as zeolite, feldspars and clay minerals. The conceptual model has highlighted that the possibility of dawsonite precipitation in a sandstone consequent to CO₂ seepage is strongly dependent on the sandstone composition (Figure 1). Indeed, dawsonite precipitation is rarely observed in natural analogues with high quartz contents and low clay contents but can precipitate in rocks with higher reactive mineral contents.

These differences in apparent reactivity between case-studies leading to the establishment of different models are logically observed in the simulation results. First of all, the Snøhvit case-study was identified as having potentially a low reactivity and presents very limited mineralogical changes after 1000 years of interaction for both acidified brine mixing ratios of the reacting solutions. pH buffering also seems limited. The most reactive case is effectively Ketzin, for which zeolite (analcime) is the most unstable primary aluminosilicate mineral under these conditions and up to 1 % of dawsonite is simulated to precipitate. Such a dawsonite precipitation after zeolite alteration is reported for the Cerro Barcino Formation CO₂ storage natural analogue, where up to 25 % of dawsonite is observed to precipitate (Zalba et al., 2011). A buffering of pH was also obtained for all degrees of CO₂ perturbation. For the In Salah site using the SandiaLab database, higher amounts of primary silicates are simulated to dissolve than for the Ketzin case, but this higher dissolution is not observed with the other databases. At In Salah, the precipitation of carbonates (dolomite) is also more restricted than at Ketzin.

Chlorite (clinochlore) is observed to be the main phase to dissolve at the In Salah and Snøhvit injection sites. Chlorite has already been shown in previous studies to preferentially dissolve in thermodynamic conditions corresponding to CO₂ disposal in sandstone layers (Khim et al., 2012). However, when present in the sandstone, zeolite is even more reactive than chlorite, as observed for the Ketzin site. Considering the most reactive case, only 1 % of carbonate minerals are calculated to precipitate. This suggests that mineralogical changes are limited in the presence of an acidic CO₂-rich brine. However, it is not possible to quantify CO₂ mineral trapping with such batch calculations considering only an input of CO₂ as dissolved CO₂ at the beginning of the calculations (Fig. 6). Indeed, here the perturbation in CO₂ is not maintained over time whereas, in a storage, the gas phase maintains high aqueous CO₂ contents when dissolved CO₂ is consumed by mineral reactions. In the present calculations, partial return to system equilibrium tends to consume a large proportion of the initially introduced CO₂, but the reactivity is kinetically controlled and the carbonation rate will not necessarily be higher if the CO₂ perturbation is maintained. Hydrodynamics also play a crucial role in CO₂ behavior in a deep aquifer, including on the system reactivity. In particular, chemical dynamics can establish within the reservoir with carbonate dissolution in some parts of the reservoir and precipitation in other parts (Audigane et al., 2007; Neufeld et al., 2010). Therefore, an extrapolation of the contribution of the different trapping mechanisms from these calculations appears hazardous.

In our calculations, mineral precipitation was assumed to occur at equilibrium, without kinetic constraints, for calculation time reasons and because of uncertainties in the kinetic parameters to be used for precipitation reactions. This assumption can be made since the objective of this work is to

consider long-term reactivity (1000 years or more). Moreover, the precipitations are limited by the dissolution of the primary minerals, including during the first calculation time-steps because secondary minerals are initially undersaturated in most acidified brines, as shown in Figure 10, Figure 13 and Figure 16.

Regarding the calculation method, it is also important to point out that the dissolution kinetics introduces a correction of 0.001 on the reactive surface area. This correction is in the range of corrections commonly applied in modeling studies using laboratory kinetic data at the natural system scale, where only part of the mineral surface is involved in the fluid-rock reaction (Gaus, 2005; Zerai et al., 2006; Xu et al., 2010). The reactive surface area always shows a strong uncertainty in a calculation and the selection of the correction factor can lead to noticeable overestimations or underestimations (White and Brantley, 2003; Ganor et al., 2007).

In order to evaluate the possible effect of the thermodynamic database on the simulation results, the calculations were performed for the three case-studies using the Thermoddem, SCALE2000 and SandiaLab databases. These three databases were selected from six databases after comparing calculated mineral solubility with experimental data. This preliminary comparison exercise highlighted the importance of using the Pitzer equations (Pitzer et al., 1973) as activity model when saline solutions are considered. The B-dot activity model, such as used with the Thermoddem database, is not sufficient to represent system behavior at high salinity (typically, beyond that of standard seawater). Between the three case-studies, using a given database does not tend to systematically favor or inhibit the system reactivity with the exception that mineral changes seem perhaps more pronounced when the SandiaLab database is used. However, fundamental differences in the geochemical system behavior are observed for the In Salah and Snøhvit case-studies as a function of the database used in the calculations. For the In Salah case, an important dissolution of illite and an important precipitation of kaolinite are calculated using the SandiaLab database while no changes in the amounts of these minerals are obtained using the SCALE2000 database and the half of illite dissolution and kaolinite precipitation is calculated using the Thermoddem database. The same discrepancy is obtained for the Snøhvit case regarding microcline dissolution and kaolinite precipitation, although dissolution and precipitation occur in more limited amounts. The evaluation of the different databases presented in section 2.2.2 suggested that the SCALE2000 database generally provides the most coherent results, including for the Si system at temperatures different to 25 °C. However, it should be recalled that parameters for Al species are not considered, impeding a correct representation of Al behavior in highly saline systems. The SandiaLab database contains Pitzer interaction parameters for Al allowing Al behavioral trends in solution to be reproduced, but

with a lack of accuracy that was evidenced in our tests. This lack of accuracy in the representation of the Si and Al chemical systems in saline solutions is most likely the cause of the differences between the results obtained using the different databases. It is difficult to conclude which reactivity case obtained with the SCALE2000 and SandiaLab databases is the most relevant since both database present limitations. However, it clearly illustrates the need to use a Pitzer database which includes all relevant chemical species over large ranges of temperature. Such a database is not available at the present time and requires further developments including the determination of many missing interaction parameters and the assessment of the global coherence of the database.

Another limitation related to the use of the Pitzer activity model concerns the redox. Redox reactions cannot be considered because the electron interaction has not been parameterized in saline solutions. As a consequence, an Fe or S system including Fe- or S-bearing minerals cannot be considered in these calculations because the behavior of these species is strongly dependent on redox.

Therefore, the calculations carried out in this study show uncertainties regarding some parameters and limitations in the representativeness of the results at the reservoir scale. However, these calculations make it possible to assess the possible reactivity of the storage system and evaluate what reactions are expected over long-term periods. These calculations consist in an essential first step before performing reactive-transport calculations considering both gas and aqueous phases. Pitzer formalism has been implemented in various reactive-transport codes such as TOUGHREACT (Zhang et al., 2006) or BIO-CORE^{2D} (Zhang et al., 2005). Using an adequate database, accurate simulations of geochemical evolution due to CO₂ injection can then be carried out, including multiphase-flow in the geological media geometry.

5 Conclusion

The aim of this work was to improve the constraint on two aspects determining the confidence in geochemical modeling of CO₂ storage in deep saline sandstone aquifers, namely, the way to calculate the interactions within the brine and the selection of the reactive mineral phases. Main results include:

- The synthesis of observations of mineralogical changes in sandstones applied to a CO₂ perturbation obtained from natural analogues and from laboratory experiments makes it possible to determine with good confidence the primary minerals expected to react and the minerals that are going to precipitate.

- The proposition of a way to identify whether dawsonite, a mineral which inclusion in models is under debate, can be formed as a secondary mineral during interaction with CO₂ from the initial mineralogy of the sandstone. Indeed, it has been shown from a review of observations in natural analogues that dawsonite is only observed in lithic and feldspar rich sandstones and is not observed in quartz rich sandstones. These observations can be related to the conditions of silica activity and pH which influence dawsonite stability.
- A method was established to assess the geochemical reactivity of the Ketzin, In Salah and Snøhvit case-studies and to evaluate the changes in mineralogy over a long-time period. Because of the mineralogical and chemical differences between these three CO₂ storage sites in saline sandstone aquifers, clear differences in geochemical behavior are predicted for these case-studies, with a higher mineral trapping expected for the Ketzin site than for In Salah and Snøhvit. This method constitutes an indispensable first step before considering reservoir geometry, mass transport and more realistic gas phase behavior.
- Two kinds of aqueous speciation models and several thermodynamic databases associated with these speciation models were tested. First in a comparison with experimental data, and then for the three CO₂ storage case-studies.
- Calculations using the B-dot model, an extension of the Debye-Hückel activity model, failed in reproducing simple mineral solubilities for salinities higher than 1 mol L⁻¹ even though the saline waters are of NaCl type. Two B-dot model databases were tested and better results were obtained using the Thermoddem database rather than the LLNL database.
- Calculations using the Pitzer interaction model clearly improved experimental data reproduction and allowed mineral solubility to be captured as a function of salinity in a relatively good manner. However, this study also points out that few Pitzer thermodynamic databases are currently available and the two databases containing interaction coefficients for a large number of ionic species, the SCALE2000 and SandiaLab databases, cannot confidently calculate the aqueous interaction energies and reproduce mineral solubility in all the physical/chemical conditions expected to be encountered in deep sedimentary aquifers.
- More experimental developments are clearly still needed to establish coherent databases for temperature ranges from 25 to 150 °C and to perform robust geochemical simulations involving saline solutions. It is all the more important to have reliable databases given that geochemical reactivity calculations for the Ketzin, In Salah and Snøhvit case-studies show strong differences as a function of the database used in the calculations.

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References

- Altmaier, M., Brendler, V., Bube, C., Marquardt, C., Moog, H. C., Richter, A., Scharge, T., Voigt, W., and Wilhelm, S., 2011. THEREDA. Thermodynamic reference database. Final Report, GRS-265, <https://www.thereda.de/>.
- Audigane, P., Gaus, I., Czernichowski-Lauriol, I., Pruess, K., and Xu, T., 2007. Two-dimensional reactive transport modeling of CO₂ injection in a saline aquifer at the Sleipner site, North Sea. *American Journal of Science* 307, 974-1008.
- Audigane, P., Lions, J., Gaus, I., Robelin, C., Durst, P., Van der Meer, B., Geel, K., Oldenburg, C. M., and Xu, T., 2009. Geochemical modeling of CO₂ injection into a methane gas reservoir at the K12-B Field, North Sea. In: Grobe, M., Pashin, J. C., and Dodge, R. L. Eds.), *Carbon dioxide sequestration in geological media - State of the science*.
- Azaroual, M., Kervévan, C., Durance, M.-V., and Brochot, S., 2004. SCALE2000 (v3.1), manuel utilisateur. BRGM.
- Baines, S. J. and Worden, R. H., 2004. The long-term fate of CO₂ in the subsurface: natural analogues for CO₂ storage. Geological Society, London, Special Publications 233, 59-85.
- Baker, J. C., Bai, G. P., Hamilton, P. J., Golding, S. D., and Keene, J. B., 1995. Continental-scale magmatic carbon dioxide seepage recorded by dawsonite in the Bowen-Gunnedah-Sydney Basin system, eastern Australia. *Journal of Sedimentary Research* 65, 522-530.

- Benbow, S. J., Metcalfe, R., and Wilson, J. C., 2008. Pitzer databases for use in thermodynamic modeling. *Quintessa*.
- Benson, S. M., and Cole, D. R., 2008. CO₂ sequestration in deep geological formations. *Elements* 4, 325-331.
- Bénézech, P., Palmer, D. A., Anovitz, L. M., and Horita, J., 2007. Dawsonite synthesis and reevaluation of its thermodynamic properties from solubility measurements: Implications for mineral trapping of CO₂. *Geochimica et Cosmochimica Acta* 71, 4438-4455.
- Blanc, P., Lassin, A., and Piantone, P., 2007. Thermoddem a database devoted to waste minerals. <http://thermoddem.brgm.fr>. BRGM, Orléans, France.
- Blanc, P., Lassin, A., Piantone, P., Azaroual, M., Jacquemet, N., Fabbri, A., and Gaucher, E. C., 2012. Thermoddem: A geochemical database focused on low temperature water/rock interactions and waste materials. *Applied Geochemistry* 27, 2107-2116.
- Cantucci, B., Montegrossi, G., Vaselli, O., Tassi, F., Quattrocchi, F., and Perkins, E. H., 2009. Geochemical modeling of CO₂ storage in deep reservoirs: The Weyburn Project (Canada) case study. *Chemical Geology* 265, 181-197.
- Chen, C.-T. A. and Marshall, W. L., 1982. Amorphous silica solubilities-IV. Behavior in pure water and aqueous sodium chloride, sodium sulfate, magnesium chloride, and magnesium sulfate solutions up to 350°C. *Geochimica et Cosmochimica Acta* 46, 279-288.
- Coudrain-Ribstein, A., Gouze, P., and de Marsily, G., 1998. Temperature-carbon dioxide partial pressure trends in confined aquifers. *Chemical Geology* 145, 73-89.
- Czernichowski-Lauriol, I., Rochelle, C., Gaus, I., Azaroual, M., Pearce, J., and Durst, P., 2006. Geochemical interactions between CO₂, pore-waters and reservoir rocks. *Advances in the Geological Storage of Carbon Dioxide*. In Lombardi, S., Altunina, L. K., and Beaubien, S. E. (Eds.), *NATO Science Series: IV: Earth and Environmental Sciences* 65, 157-174.
- Debye, P. and Hückel, E., 1923. The theory of electrolytes. I. Lowering of freezing point and related phenomena. *Physikalische Zeitschrift* 24, 185-206.
- Dethlefsen, F., Haase, C., Ebert, M., and Dahmke, A., 2011. Uncertainties of geochemical modeling during CO₂ sequestration applying batch equilibrium calculations. *Environmental Earth Sciences*, 1-13.
- Duan, Z. and Sun, R., 2003. An improved model calculating CO₂ solubility in pure water and aqueous NaCl solutions from 273 to 533 K and from 0 to 2000 bar. *Chemical Geology* 193, 257-271.

- Förster, A., Norden, B., Zinck-Jørgensen, K., Frykman, P., Kulenkampff, J., Spangenberg, E., Erzinger, J., Zimmer, M., Kopp, J., Borm, G., Juhlin, C., Cosma, C.-G., and Hurter, S., 2006. Baseline characterization of the CO₂SINK geological storage site at Ketzin, Germany. *Environmental Geosciences* 13, 145-161.
- Förster, A., Schöner, R., Förster, H. J., Norden, B., Blaschke, A. W., Luckert, J., Beutler, G., Gaupp, R., and Rhede, D., 2010. Reservoir characterization of a CO₂ storage aquifer: The Upper Triassic Stuttgart Formation in the Northeast German Basin. *Marine and Petroleum Geology* 27, 2156-2172.
- Ganor, J., Lu, P., Zheng, Z., and Zhu, C., 2007. Bridging the gap between laboratory measurements and field estimations of silicate weathering using simple calculations. *Environmental Geology* 53, 599-610.
- Gao, Y., Liu, L., and Hu, W., 2009. Petrology and isotopic geochemistry of dawsonite-bearing sandstones in Hailaer basin, northeastern China. *Applied Geochemistry* 24, 1724-1738.
- Gaus, I., 2010. Role and impact of CO₂-rock interactions during CO₂ storage in sedimentary rocks. *International Journal of Greenhouse Gas Control* 4, 73-89.
- Gaus, I., Azaroual, M., and Czernichowski-Lauriol, I., 2005. Reactive transport modelling of the impact of CO₂ injection on the clayey cap rock at Sleipner (North Sea). *Chemical Geology* 217, 319-337.
- Hangx, S. J. T. and Spiers, C. J., 2009. Reaction of plagioclase feldspars with CO₂ under hydrothermal conditions. *Chemical Geology* 265, 88-98.
- Haszeldine, R., S., Quinn, O., England, G., Wilkinson, M., Shipton, Z., K., Evans, J., P., Heath, J., Crossey, L., Ballentine, C., J., and Graham, C., M., 2005. Analogues géochimiques naturels pour le stockage du dioxyde de carbone en réservoir géologique poreux profond : perspective pour le Royaume-Uni. *Oil & Gas Science and Technology - Rev. IFP* 60, 33-49.
- Helgeson, H. C., 1968. Evaluation of irreversible reactions in geochemical processes involving minerals and aqueous solutions I. Thermodynamic relations. *Geochimica et Cosmochimica Acta* 32, 853-877.
- Helgeson, H. C., Brown, T. H., and Leeper, R. H., 1969. Handbook of theoretical activity diagrams depicting chemical equilibria in geologic systems involving an aqueous phase at one Atm and 0 to 300 °C. Freeman, Cooper & Company, 253 pp.
- Hellevang, H., Aagaard, P., Oelkers, E. H., and Kvanne, B., 2005. Can Dawsonite Permanently Trap CO₂? *Environmental Science & Technology* 39, 8281-8287.

- Hellevang, H., Declercq, J., Kvamme, B., and Aagaard, P., 2010. The dissolution rates of dawsonite at pH 0.9 to 5 and temperatures of 22, 60 and 77 °C. *Applied Geochemistry* 25, 1575-1586.
- Hellevang, H., Declercq, J., and Aagaard, P., 2011. Why is Dawsonite Absent in CO₂ Charged Reservoirs? *Oil Gas Sci. Technol. – Rev. IFP Energies nouvelles* 66, 119-135.
- Hermanrud, C., Eiken, O., Hansen, O.R., Nordgaard Bolaas, H., Simmenes, T., Teige, G., Hansen, H. and Johansen, S., 2013. Importance of pressure management in CO₂ storage. 2013 Offshore Technology Conference, May 06 - 09, 2013 2013, Houston, TX, USA
- IPCC, 2007. Special report on carbon dioxide capture and storage. Special report of the Intergovernmental Panel on Climate Change – Summary for Policymakers and technical summary. Intergovernmental Panel on Climate Change.
- Johnson, J. W., Nitao, J. J., and Knauss, K. G., 2004. Reactive transport modelling of CO₂ storage in saline aquifers to elucidate fundamental processes, trapping mechanisms and sequestration partitioning. *Geological Society Special Publications* 233, 107-128.
- Johnson, J. W., Nitao, J. J., Steefel, C., and Knauss, K. G., 2001. Reactive transport modelling of geological CO₂ sequestration in saline aquifers; the influence of intra-aquifer shales and the relative effectiveness of structural, solubility and mineral trapping during prograde and retrograde sequestration. First Annual Conference on Carbon Sequestration, Washington.
- Johnson, J. W., Oelkers, E. H., and Helgeson, H. C., 1992. SUPCRT92: A software package for calculating the standard molal thermodynamic properties of minerals, gases, aqueous species, and reactions from 1 to 5000 bar and 0 to 1000°C. *Computers & Geosciences* 18, 899-947.
- Kaszuba, J. P., Janecky, D. R., and Snow, M. G., 2003. Carbon dioxide reaction processes in a model brine aquifer at 200 °C and 200 bars: implications for geologic sequestration of carbon. *Applied Geochemistry* 18, 1065-1080.
- Kaszuba, J. P., Janecky, D. R., and Snow, M. G., 2005. Experimental evaluation of mixed fluid reactions between supercritical carbon dioxide and NaCl brine: Relevance to the integrity of a geologic carbon repository. *Chemical Geology* 217, 277-293.
- Kaszuba, J. P., Viswanathan, H. S., and Carey, J. W., 2011. Relative stability and significance of dawsonite and aluminum minerals in geologic carbon sequestration. *Geophysical Research Letters* 38, L08404.
- Ketzer, J. M., Iglesias, R., Einloft, S., Dullius, J., Ligabue, R., and de Lima, V., 2009. Water–rock–CO₂ interactions in saline aquifers aimed for carbon dioxide storage: Experimental and numerical

- modeling studies of the Rio Bonito Formation (Permian), southern Brazil. *Applied Geochemistry* 24, 760-767.
- Kharaka, Y. K. and Hanor, J. S., 2003. 5.16 - Deep Fluids in the Continents: I. Sedimentary Basins. In: Heinrich, D. H. and Karl, K. T. (Eds.), *Treatise on Geochemistry*. Pergamon, Oxford.
- Kihm, J.-H., Kim, J.-M., Wang, S., and Xu, T., 2012. Hydrogeochemical numerical simulation of impacts of mineralogical compositions and convective fluid flow on trapping mechanisms and efficiency of carbon dioxide injected into deep saline sandstone aquifers. *Journal of Geophysical Research* 117, B06204.
- Knauss, K. G., Johnson, J. W., and Steefel, C. I., 2005. Evaluation of the impact of CO₂, co-contaminant gas, aqueous fluid and reservoir rock interactions on the geologic sequestration of CO₂. *Chemical Geology* 217, 339-350.
- Lagneau, V., Pipart, A., and Catalette, H., 2005. Modélisation couplée chimie-transport du comportement à long terme de la séquestration géologique de CO₂ dans des aquifères salins profonds. *Oil & Gas Science and Technology - Rev. IFP* 60, 231-247.
- Lasaga, A. C., Soler, J. M., Ganor, J., Burch, T. E., and Nagy, K. L., 1994. Chemical weathering rate laws and global geochemical cycles. *Geochimica et Cosmochimica Acta* 58, 2361-2386.
- Linjordet, A. and Grung Olsen, R., 1992. The Jurassic Snøhvit gas fields, Hammerfest Basin, offshore Northern Norway. In: Halbouty, M. T. (Ed.), *Giant oil and gas fields of the decade 1978-1988, Proceedings of the Conference held in Stavanger, Norway, September 9-12, 1990*. American Association of Petroleum Geologists.
- Linke, W. F. and Seidell, A., 1965. *Solubilities of inorganic and metalorganic compounds*. American Chemical Society, Washington D.C.
- Liu, N., Liu, L., Qu, X., Yang, H., Wang, L., and Zhao, S., 2011. Genesis of authigenic carbonate minerals in the Upper Cretaceous reservoir, Honggang Anticline, Songliao Basin: A natural analog for mineral trapping of natural CO₂ storage. *Sedimentary Geology* 237, 166-178.
- Luquot, L., Andreani, M., Gouze, P., and Camps, P., 2012. CO₂ percolation experiment through chlorite/zeolite-rich sandstone (Pretty Hill Formation – Otway Basin–Australia). *Chemical Geology* 294–295, 75-88.
- Marchand, A. M. E., Haszeldine, R. S., Smalley, P. C., Macaulay, C. I., and Fallick, A. E., 2001. Evidence for reduced quartz-cementation rates in oil-filled sandstones. *Geology* 29, 915-918.

- Marini, L., 2006. Geological sequestration of carbon dioxide. Thermodynamics, kinetics and reaction path modeling. Elsevier.
- Marion, G. M., Catling, D. C. and Kargel, J. S., 2003. Modeling aqueous ferrous iron chemistry at low temperatures with application to Mars. *Geochimica et Cosmochimica Acta* 67, 4251-4266.
- Marshall, W. L. and Warakomski, J. M., 1980. Amorphous silica solubilities—II. Effect of aqueous salt solutions at 25°C. *Geochimica et Cosmochimica Acta* 44, 915-924.
- Marshall, W. L. and Chen, C-T. A., 1982. Amorphous silica solubilities—V. Predictions of solubility behavior in aqueous mixed electrolyte solutions to 300°C. *Geochimica et Cosmochimica Acta* 46, 289-291.
- Martens, S., Kempka, T., Liebscher, A., Lüth, S., Möller, F., Myrntinen, A., Norden, B., Schmidt-Hattenberger, C., Zimmer, M. and Kühn, M., 2012. Europe's longest-operating on-shore CO₂ storage site at Ketzin, Germany: a progress report after three years of injection. *Environmental Earth Sciences* 67, 323-334.
- Michael, K., Arnot, M., Cook, P., Ennis-King, J., Funnell, R., Kaldi, J., Kirste, D., and Paterson, L., 2009. CO₂ storage in saline aquifers I—Current state of scientific knowledge. *Energy Procedia* 1, 3197-3204.
- Moore, J., Adams, M., Allis, R., Lutz, S., and Rauzi, S., 2005. Mineralogical and geochemical consequences of the long-term presence of CO₂ in natural reservoirs: An example from the Springerville–St. Johns Field, Arizona, and New Mexico, U.S.A. *Chemical Geology* 217, 365-385.
- Neufeld, J. A., Hesse, M. A., Riaz, A., Hallworth, M. A., Tchelepi, H. A., and Huppert, H. E., 2010. Convective dissolution of carbon dioxide in saline aquifers. *Geophys. Res. Lett.* 37, L22404.
- Okuyama, Y. and Take, S., 2011. Dawsonite-aragonite association in the Cretaceous Izumi Group, SW Japan: Evidence of CO₂-rich fluid invasion in the area of classical study. *Journal of Mineralogical and Petrological Sciences* 106, 79-84.
- Palmer, D. A. and Wesolowski, D. J., 1992. Aluminum speciation and equilibria in aqueous solution: II. The solubility of gibbsite in acidic sodium chloride solutions from 30 to 70°C. *Geochimica et Cosmochimica Acta* 56, 1093-1111.
- Parkhurst, D. L. and Appelo, C. A. J., 1999. User's guide to PHREEQC (Version 2)--a computer program for speciation, batch-reaction, one-dimensional transport, and inverse geochemical calculations. U.S. Geological Survey.

- Pauwels, H., Gaus, I., le Nindre, Y. M., Pearce, J., and Czernichowski-Lauriol, I., 2007. Chemistry of fluids from a natural analogue for a geological CO₂ storage site (Montmiral, France): Lessons for CO₂-water-rock interaction assessment and monitoring. *Applied Geochemistry* 22, 2817-2833.
- Pearce, J. M., Holloway, S., Wacker, H., Nelis, M. K., Rochelle, C., and Bateman, K., 1996. Natural occurrences as analogues for the geological disposal of carbon dioxide. *Energy Conversion and Management* 37, 1123-1128.
- Pearce, J. M., Kirby, G. A., Lacinska, A., Bateson, L., Wagner, D., Rochelle, C. A., and Cassidy, M., 2011. Reservoir-scale CO₂ -fluid rock interactions: Preliminary results from field investigations in the Paradox Basin, Southeast Utah. *Energy Procedia* 4, 5058-5065.
- Pinti, D. L. and Marty, B., 1995. Noble gases in crude oils from the Paris Basin, France: Implications for the origin of fluids and constraints on oil-water-gas interactions. *Geochimica et Cosmochimica Acta* 59, 3389-3404.
- Pitzer, K. S., 1973. Thermodynamics of electrolytes, 1, Theoretical basis and general equations. *Journal of Physical Chemistry* 77, 268-277.
- Pitzer, K. S., 1991. Activity coefficients in electrolyte solutions. Boca Raton, CRC Press.
- Pruess, K. and García, J., 2002. Multiphase flow dynamics during CO₂ disposal into saline aquifers. *Environmental Geology* 42, 282-295.
- Rochelle, C., Pearce, J. M., Shaw, R., Taylor, H., Turner, G., and Williams, C., 2007. Geochemical interactions between CO₂ and host rocks at the Snøhvit field. Results of fluid-rock interaction experiments. British Geological Survey Commissioned Report CR/07/218.
- Smith, J. W. and Milton, C., 1966. Dawsonite in the Green River Formation of Colorado. *Economic Geology* 61, 1029-1042.
- Taberner, C., Zhang, G., and Catwright, L., 2009. Injection of Supercritical CO₂ into Deep Saline Carbonate Formations, Predictions from Geochemical Modeling. 2009 SPE EUROPEC/EAGE Annual Conference and Exhibition, Amsterdam, The Netherlands.
- Truesdell, A.H. and B. F. Jones, 1974. WATEQ a computer program for calculating chemical equilibria of natural waters. *Journal of Research of the U.S. Geological Survey* 2, 233-248
- USDOE, 2007. In-Drift Precipitates/Salts Model. United States Department of Energy (USDOE).

- Vong, C. Q., Jacquemet, N., Audigane, P. and Frykman, P., 2010. Compilation of geochemical data and injected gas compositions for modeling of fluid-rock reactions in the In Salah, Snøhvit, Ketzin and Sleipner CCS sites. CO2REMOVE, Deliverable D2.2.2C+D, restricted access.
- Watson, M. N., Zwingmann, N., and Lemon, N. M., 2004. The Ladbroke Grove–Katnook carbon dioxide natural laboratory: A recent CO₂ accumulation in a lithic sandstone reservoir. *Energy* 29, 1457-1466.
- White, A. F. and Brantley, S. L., 2003. The effect of time on the weathering of silicate minerals: why do weathering rates differ in the laboratory and field? *Chemical Geology* 202, 479-506.
- White, A. F. and Brantley, S. L., 2003. The effect of time on the weathering of silicate minerals: why do weathering rates differ in the laboratory and field? *Chemical Geology* 202, 479-506.
- Wiese, B., Böhner, J., Enachescu, C., Würdemann, H., and Zimmermann, G., 2010. Hydraulic characterisation of the Stuttgart formation at the pilot test site for CO₂ storage, Ketzin, Germany. *International Journal of Greenhouse Gas Control* 4, 960-971.
- Wildenborg, T., Bentham, M., Chadwick, A., David, P., Deflandre, J.-P., Dillen, M., Groenenberg, H., Kirk, K., Le Gallo, Y., 2009. Largescale CO₂ injection demos for the development of monitoring and verification technology and guidelines (CO2ReMoVe), GHGT 9, *Energy Procedia* 1, 2367-2374.
- Wilkinson, M., Haszeldine, R. S., Fallick, A. E., Odling, N., Stoker, S. J., and Gatliff, R. W., 2009. CO₂–Mineral Reaction in a Natural Analogue for CO₂ Storage—Implications for Modeling. *Journal of Sedimentary Research* 79, 486-494.
- Wolf, M., Breitkopf, O., and Puk, R., 1989. Solubility of calcite in different electrolytes at temperatures between 10° and 60°C and at CO₂ partial pressures of about 1 kPa. *Chemical Geology* 76, 291-301.
- Worden, R. H., 2006. Dawsonite cement in the Triassic Lam Formation, Shabwa Basin, Yemen: A natural analogue for a potential mineral product of subsurface CO₂ storage for greenhouse gas reduction. *Marine and Petroleum Geology* 23, 61-77.
- Worden, R. H., Coleman, M. L., and Matray, J. M., 1999. Basin scale evolution of formation waters: a diagenetic and formation water study of the Triassic Chaunoy Formation, Paris Basin. *Geochimica et Cosmochimica Acta* 63, 2513-2528.
- Würdemann, H., Möller, F., Kühn, M., Heidug, W., Christensen, N. P., Borm, G., and Schilling, F. R., 2010. CO2SINK—From site characterisation and risk assessment to monitoring and verification:

- One year of operational experience with the field laboratory for CO₂ storage at Ketzin, Germany. *International Journal of Greenhouse Gas Control* 4, 938-951.
- Xu, T., 2008. TOUGHREACT testing in high ionic strength brine sandstone systems. Lawrence Berkeley National Laboratory, Berkeley, California, USA.
- Xu, T., Apps, J. A., and Pruess, K., 2003. Reactive geochemical transport simulation to study mineral trapping for CO₂ disposal in deep arenaceous formations. *J. Geophys. Res.* 108, 2071.
- Xu, T., Kharaka, Y. K., Doughty, C., Freifeld, B. M., and Daley, T. M., 2010. Reactive transport modeling to study changes in water chemistry induced by CO₂ injection at the Frio-I Brine Pilot. *Chemical Geology* 271, 153-164.
- Zalba, P. E., Conconi, M. S., Morosi, M., Manassero, M., and Comerio, M., 2011. Dawsonite in tuffs and litharenites of the Cerro Castaño member, Cerro Barcino formation, Chubut group (Cenomanian), Los Altares, Patagonia, Argentina. *The Canadian Mineralogist* 49, 503-520.
- Zerai, B., Saylor, B. Z., and Matisoff, G., 2006. Computer simulation of CO₂ trapped through mineral precipitation in the Rose Run Sandstone, Ohio. *Applied Geochemistry* 21, 223-240.
- Zhang, G., Spycher, N., Xu, T., Sonnenthal, E., and Steefel, C., 2006. Reactive Geochemical Transport Modeling of Concentrated Aqueous Solutions: Supplement to TOUGHREACT User's Guide for the Pitzer Ion-Interaction Model.
- Zhang, G., Zheng, Z., and Wan, J., 2005. Modeling reactive geochemical transport of concentrated aqueous solutions. *Water Resour. Res.* 41, W02018.
- Zhang, W., Li, Y., Xu, T., Cheng, H., Zheng, Y., and Xiong, P., 2009. Long-term variations of CO₂ trapped in different mechanisms in deep saline formations: A case study of the Songliao Basin, China. *International Journal of Greenhouse Gas Control* 3, 161-180.
- Zhu, C., 2005. In situ feldspar dissolution rates in an aquifer. *Geochimica et Cosmochimica Acta* 69, 1435-1453.

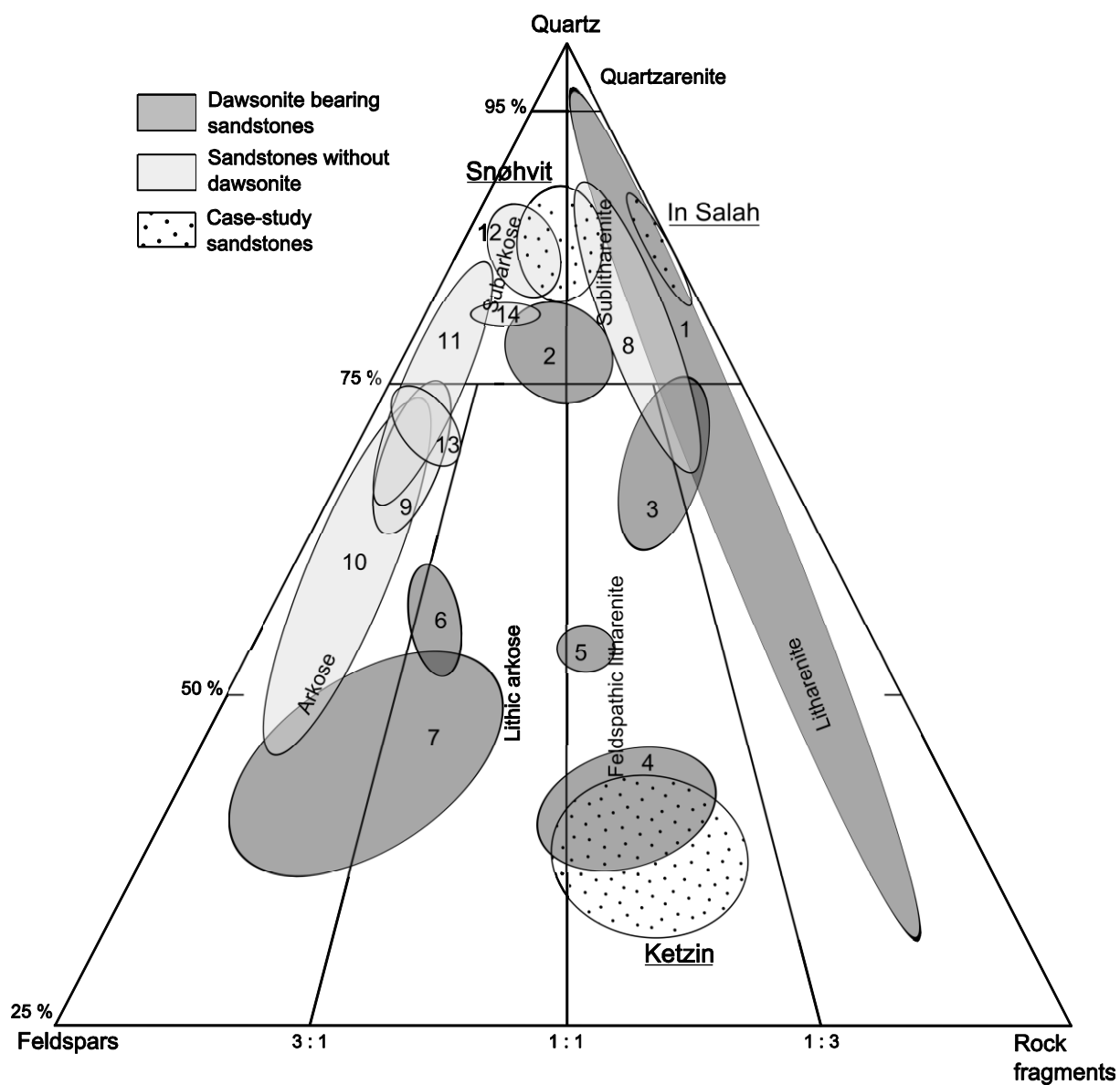


Figure 1 : Folk's diagram of CO₂ geologic disposal natural analogue compositions where the presence or absence of dawsonite is shown. Numbers correspond to natural analogues: 1) Bowen-Gunnedah-Sydney Basin; 2) Lam Formation; 3) Fizzy accumulation; 4) Songliao Basin; 5) Cerro Barcino Formation; 6) Springerville – St Johns Field; 7) Hailer Basin; 8) Miller Field; 9) Magnus Field; 10) Bravo Dome; 11) Vert le Grand; 12) Ladbroke Grove Field; 13) Montmiral reservoir; 14) Paradox Basin. Ketzin, In Salah and Snøhvit case-study sandstone compositions are also reported.

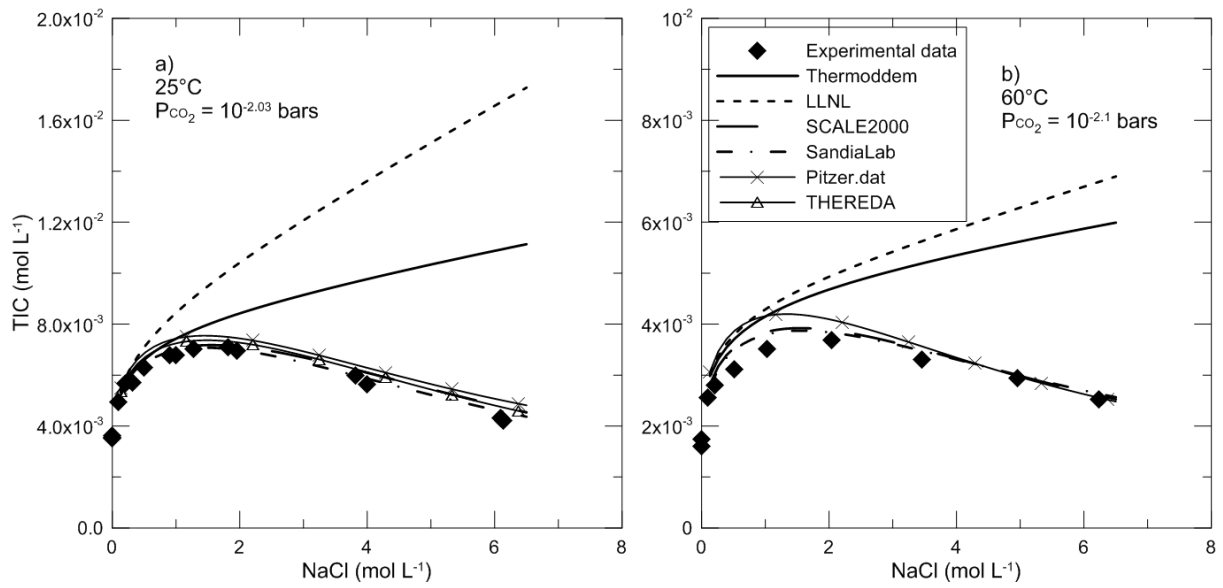


Figure 2: Measured and calculated calcite solubilities in NaCl solutions at a) 25 °C and b) 60 °C. Experimental data from Wolf et al. (1989).

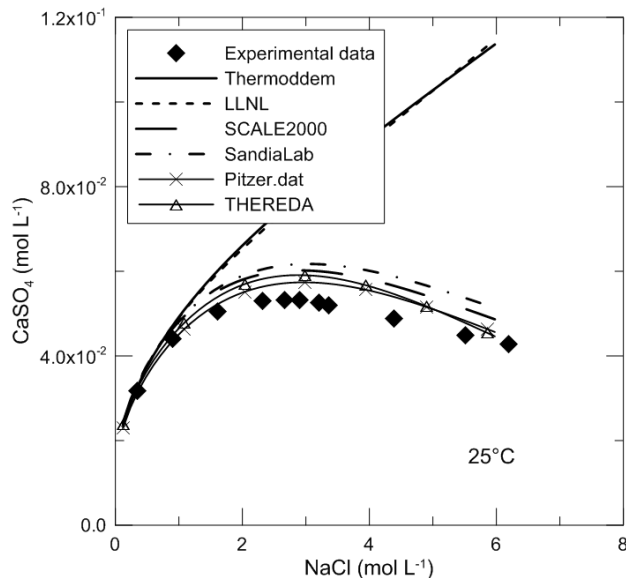


Figure 3: Measured and calculated gypsum solubilities in NaCl solutions at 25 °C. Experimental data from Linke and Seidell (1965).

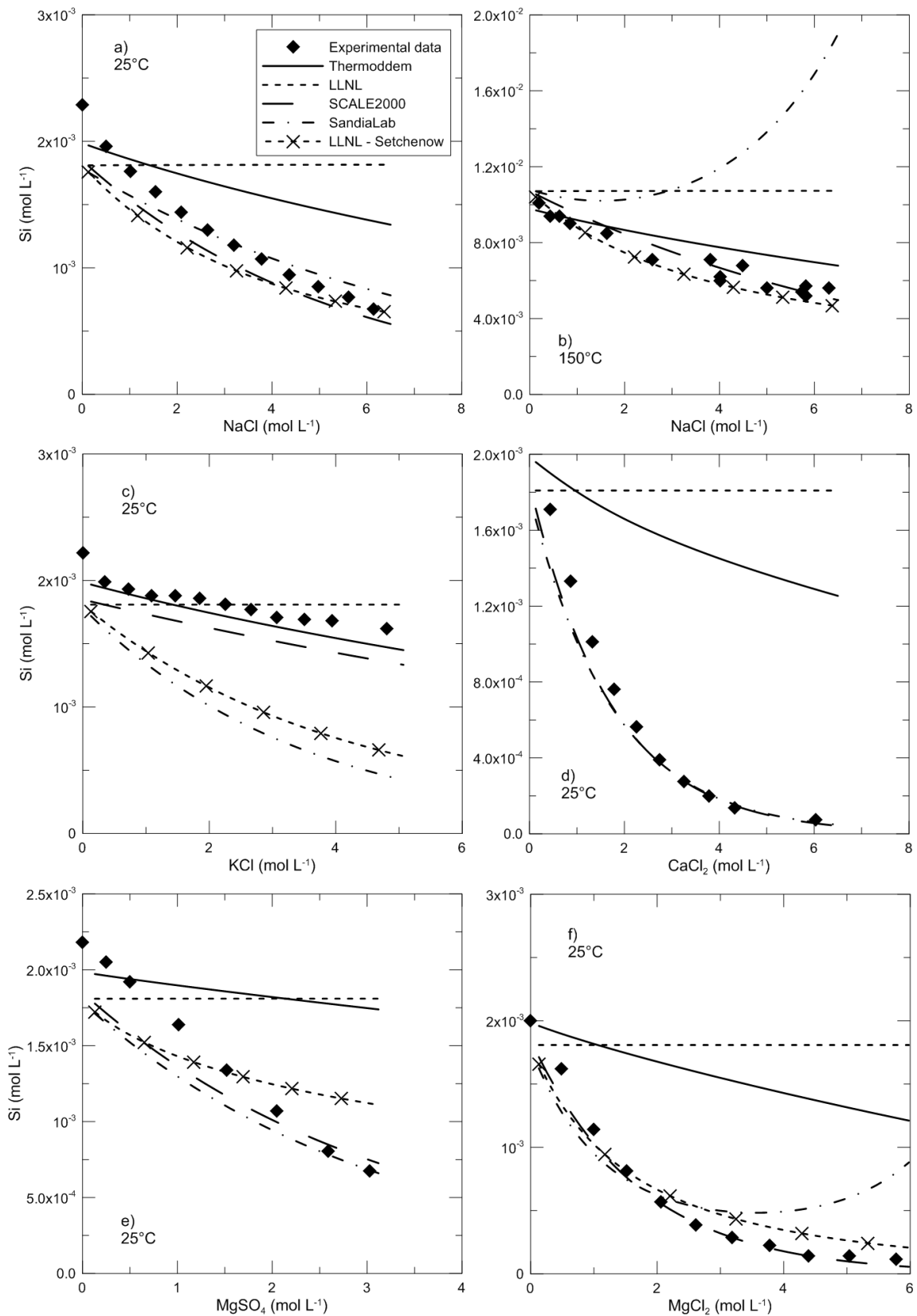


Figure 4: Measured and calculated amorphous silica solubilities in a) NaCl solutions at 25 °C, b) NaCl solutions at 150 °C, c) KCl solutions at 25 °C, d) CaCl_2 solutions at 25 °C, e) MgSO_4 solutions at 25 °C and f) CaCl_2 solutions and 25 °C. Experimental data in NaCl solutions from Chen and Marshall (1982) and further experimental data from Marshall and Warakomski (1980).

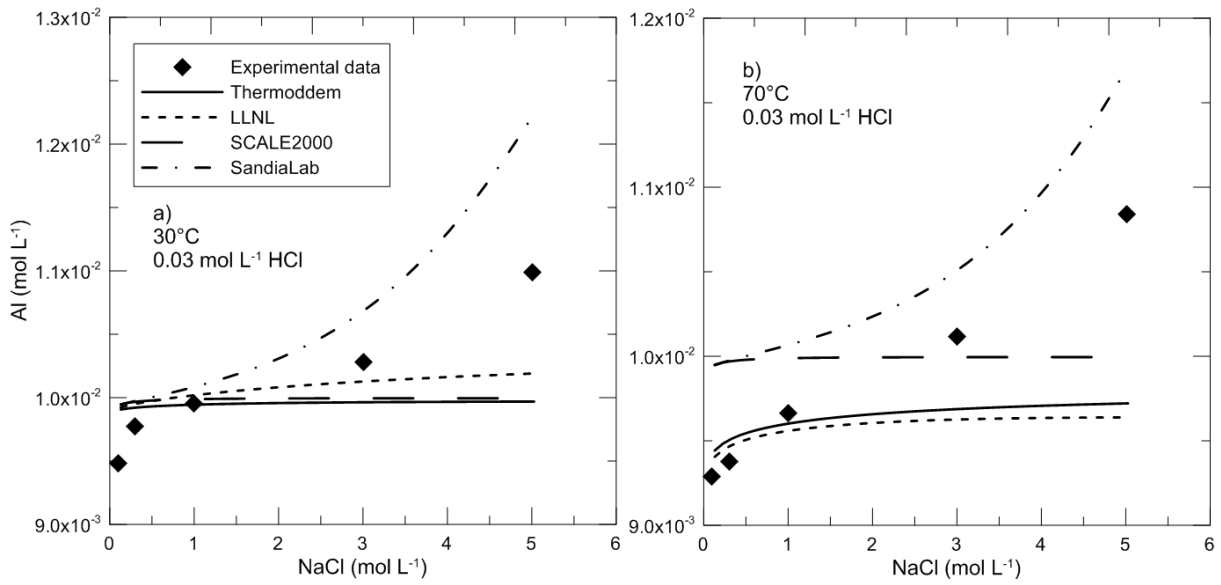


Figure 5: Measured and calculated gibbsite solubilities in NaCl solutions at a) 30 °C and b) 70 °C. Experimental data from Palmer and Wesolowski (1992).

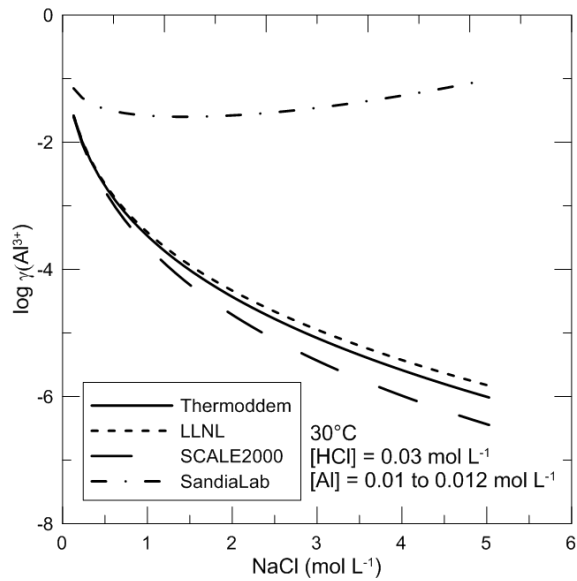


Figure 6: Al^{3+} activity coefficient evolution as a function of the ionic strength for different databases during gibbsite solubility calculations shown in Figure 5-a.

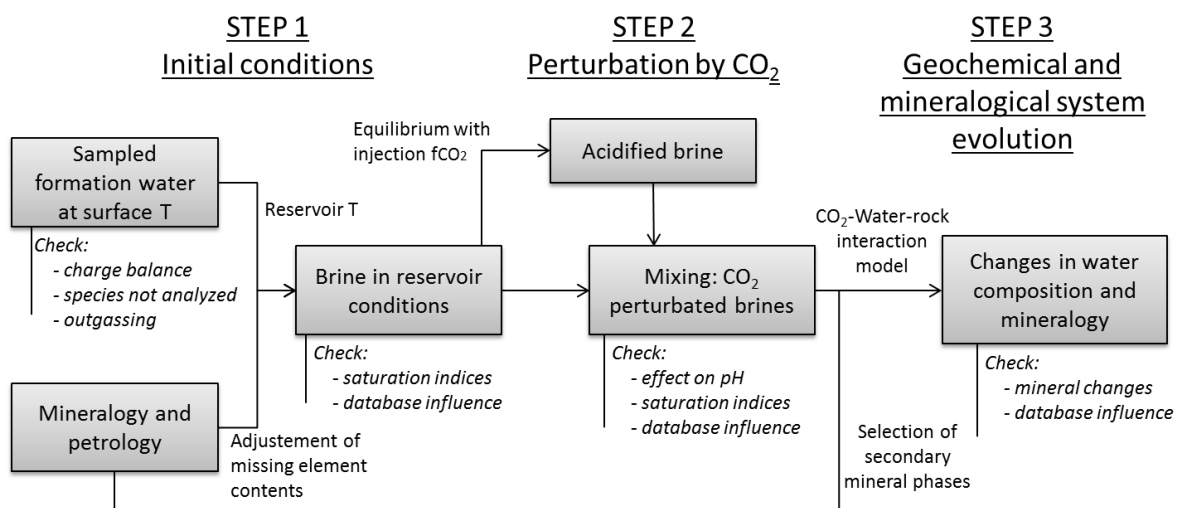


Figure 7: Workflow of the modeling approach proposed to study geochemical fluid – rock interactions induced by CO₂ injection in deep saline aquifers.

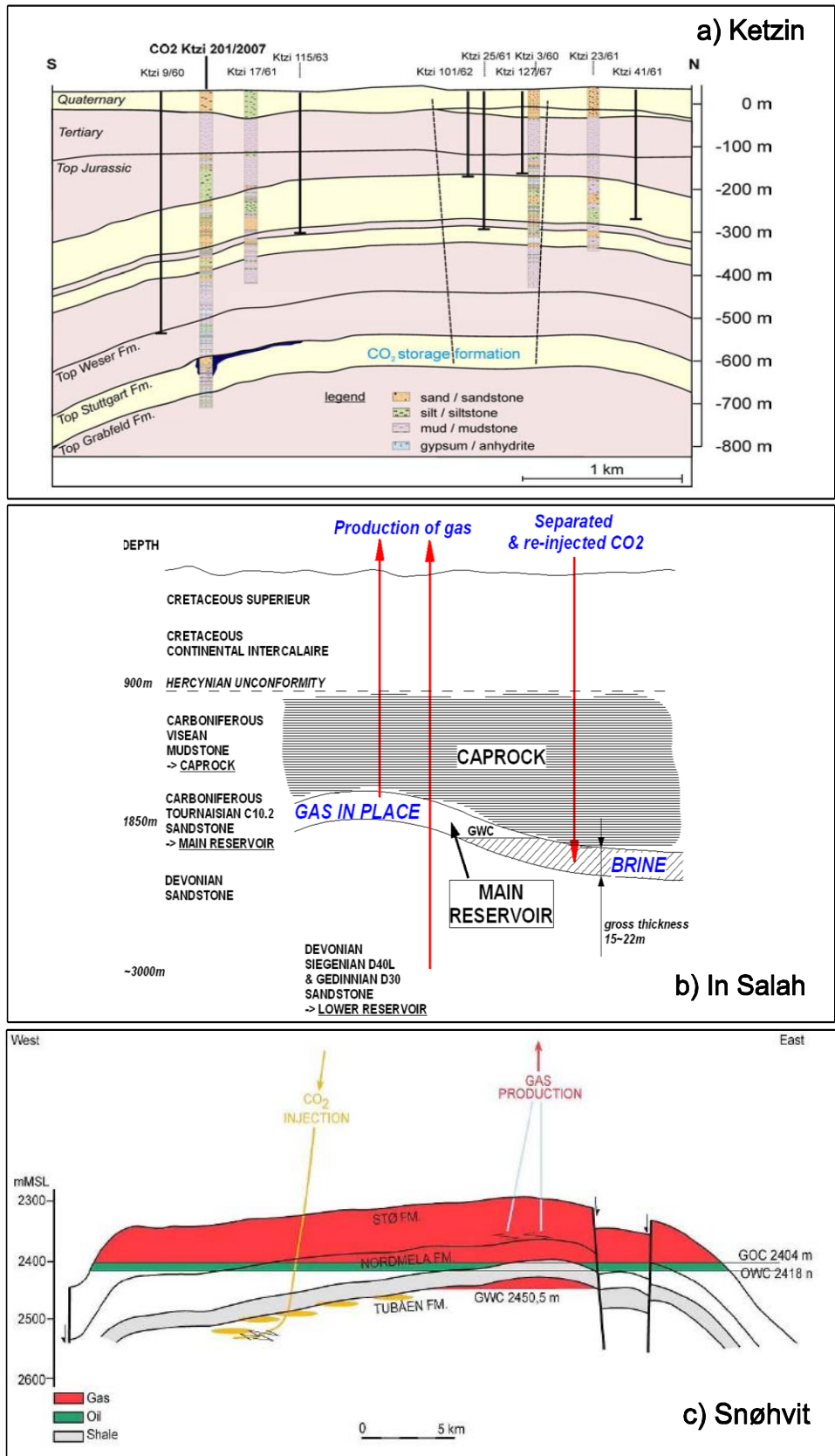


Figure 8: Schematic cross-section of a) Ketzin (Förster et al., 2008), b) In Salah and c) Snøhvit CO₂ injection site case-studies.

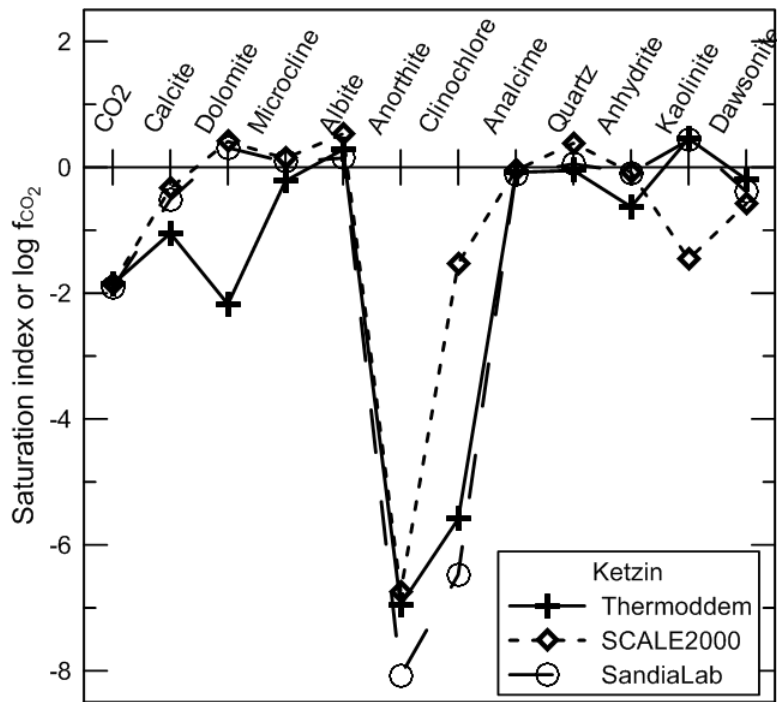


Figure 9: Mineral saturation indices and CO₂ fugacities of Ketzin site porewater calculated using SCALE2000, SandiaLab and Thermoddem databases.

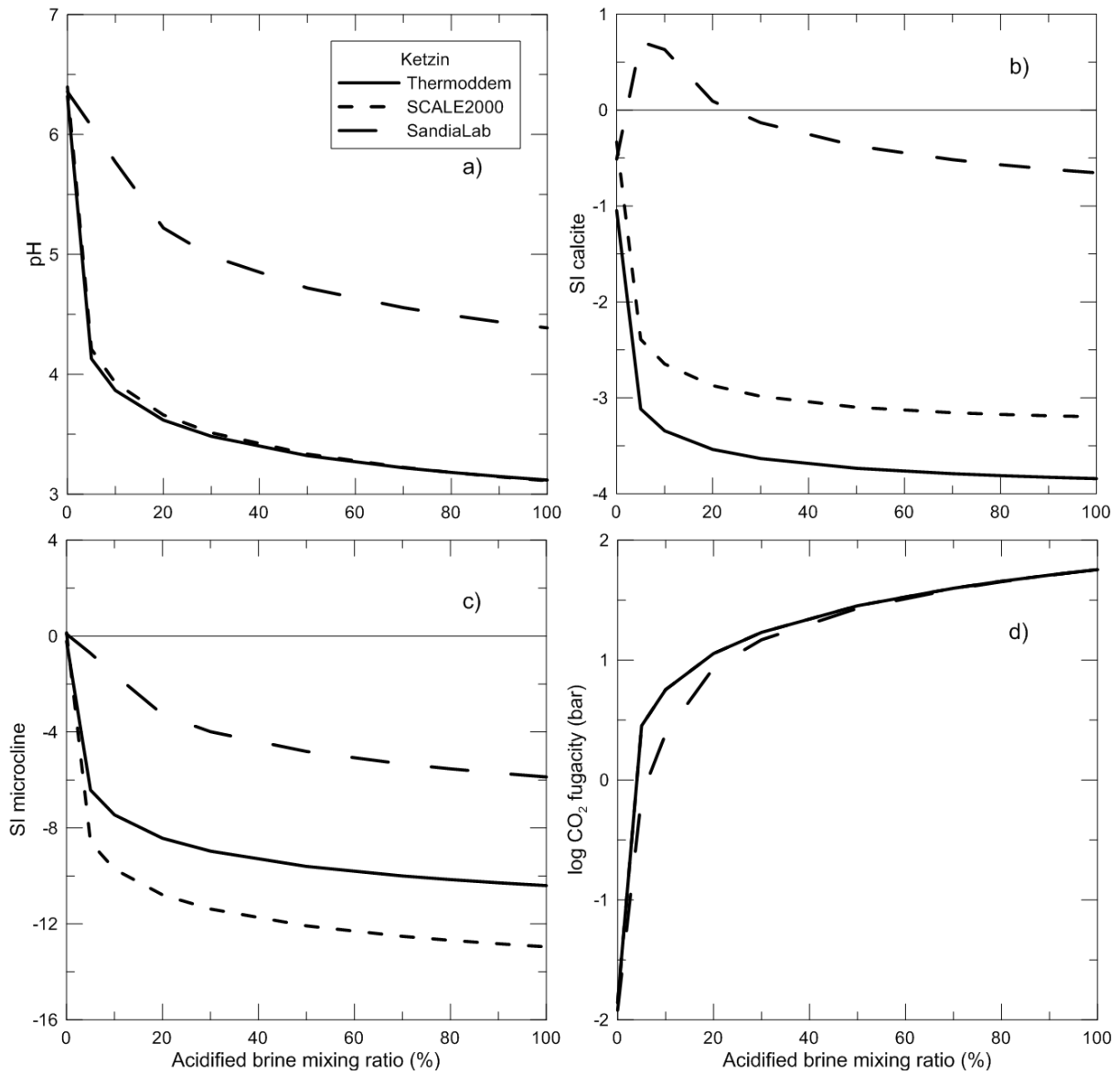


Figure 10: Evolution of a) pH, b) calcite saturation index, c) microcline saturation index and d) CO₂ fugacity of Ketzin water as a function of the acidified brine mixing ratio (0 % for Ketzin initial formation water to 100 % for Ketzin water acidified at injection CO₂ fugacity) calculated using the SCALE2000, SandiaLab and Thermoddem databases.

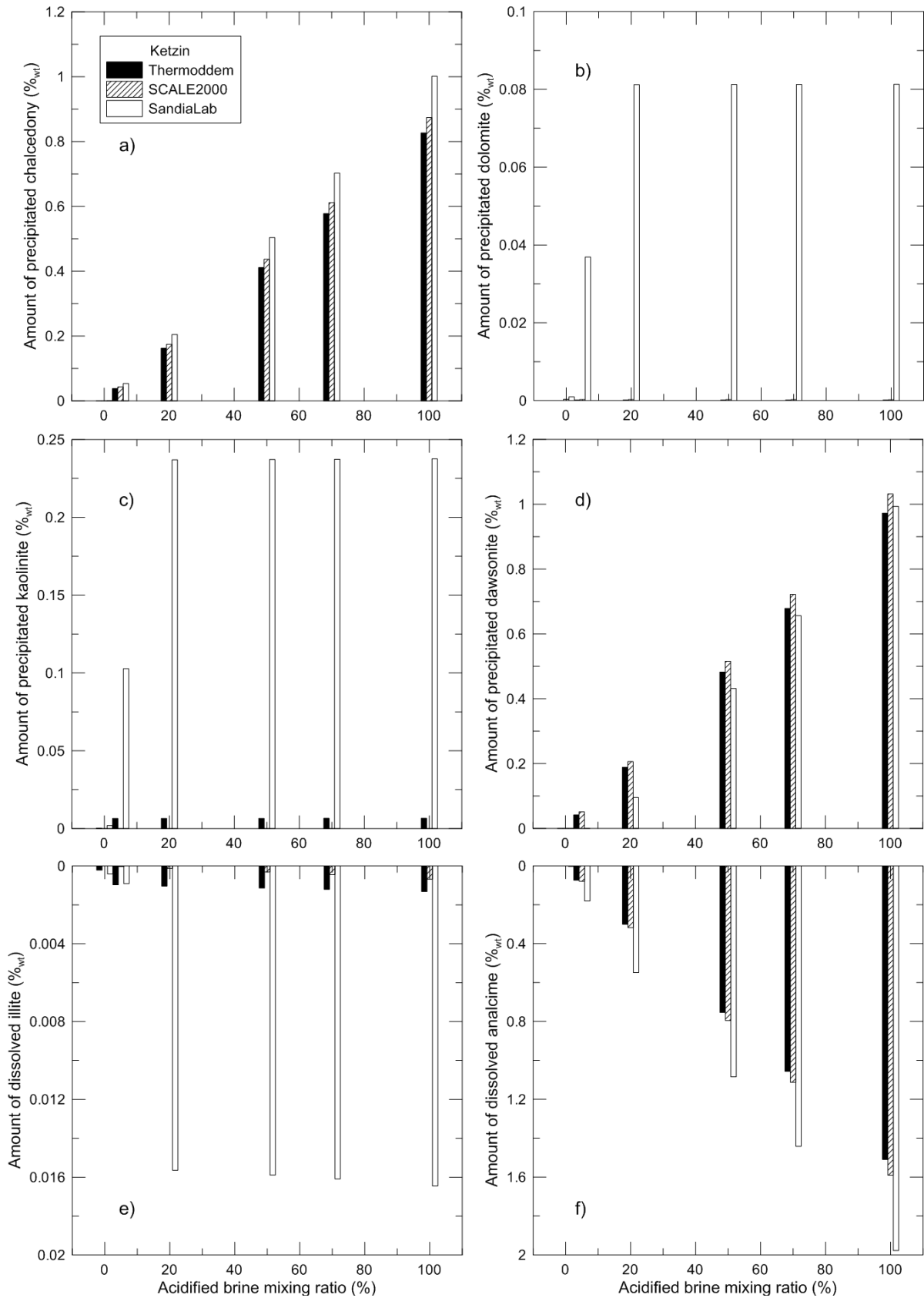


Figure 11: Amounts of a) chalcedony, b) dolomite, c) kaolinite, d) dawsonite, e) illite and f) analcime precipitating or dissolving during 1000 years for the Ketzin case-study for different acidified brine mixing ratios (0, 5, 20, 50, 70 and 100 %) calculated using the SCALE2000, SandiaLab and Thermoddem databases.

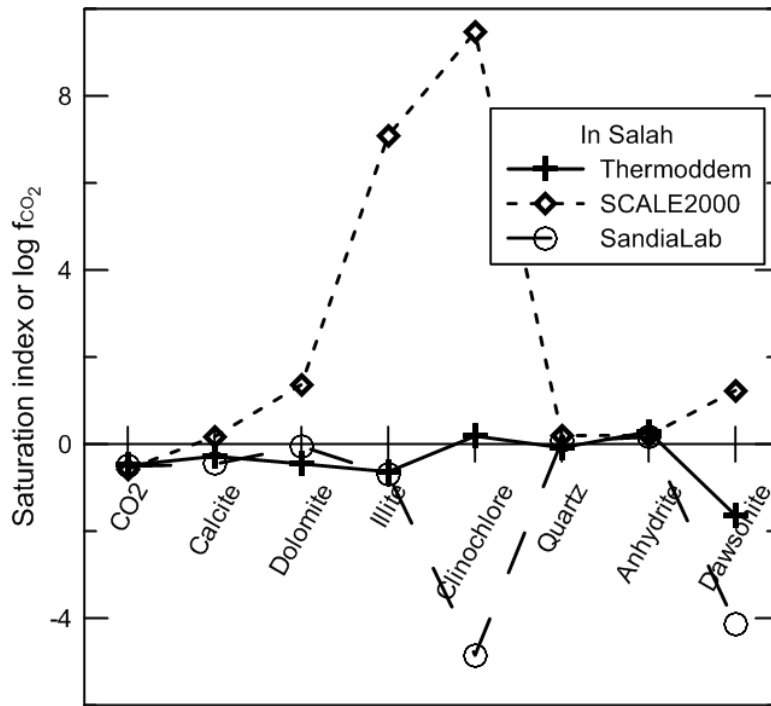


Figure 12: Mineral saturation indices and CO₂ fugacity of In Salah site porewater calculated using the SCALE2000, SandiaLab and Thermoddem databases.

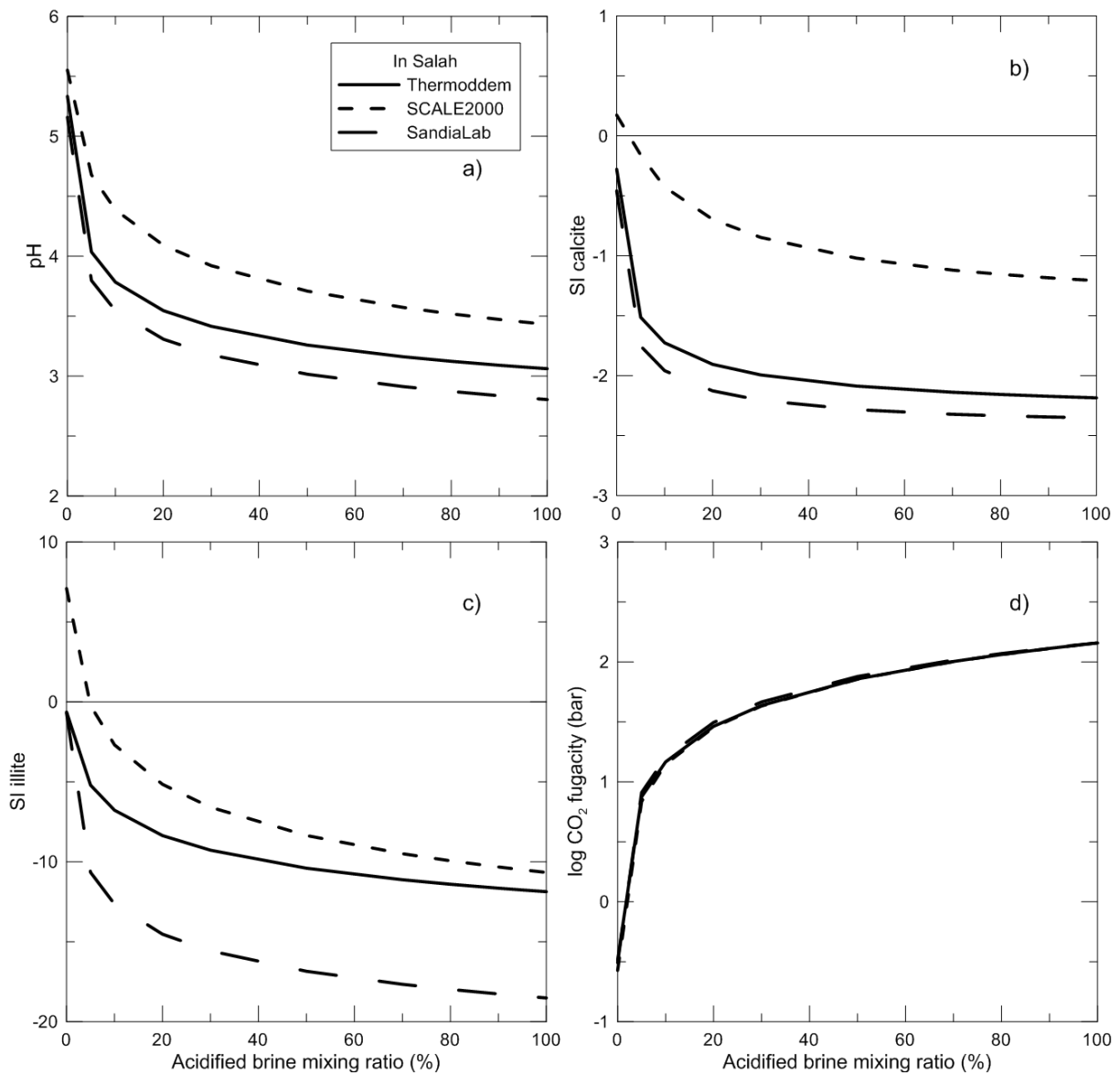


Figure 13: Evolution of a) pH, b) calcite saturation index, c) illite saturation index and d) CO₂ fugacity of In Salah water as a function of the acidified brine mixing ratio (0 % for In Salah initial formation water to 100 % for In Salah water acidified at injection CO₂ fugacity) calculated using the SCALE2000, SandiaLab and Thermoddem databases.

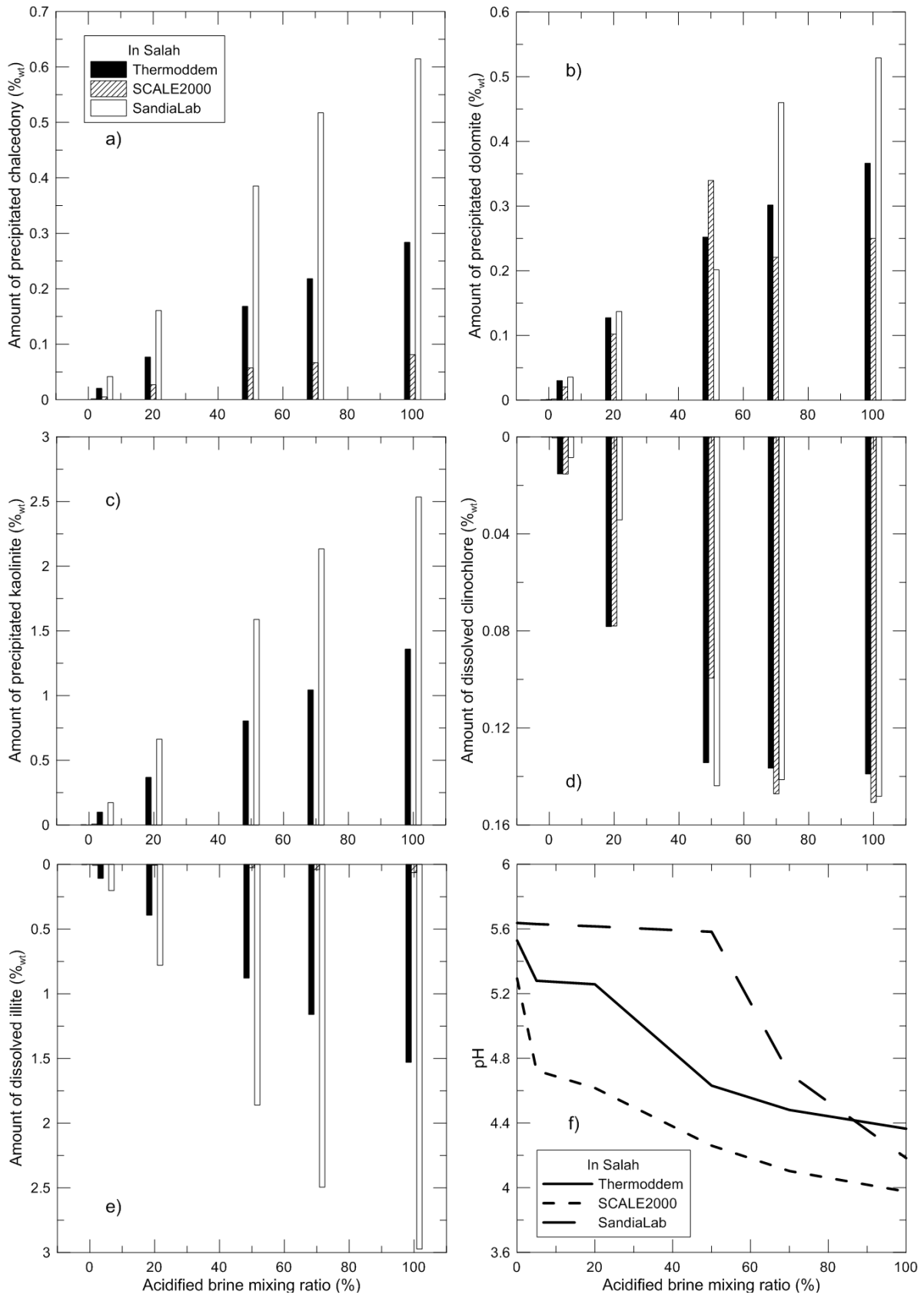


Figure 14: Amounts of a) chalcedony, b) dolomite, c) kaolinite, d) clinochlore and e) illite precipitating or dissolving during 1000 years and f) final pH value for the In Salah case-study for different acidified brine

mixing ratios (0, 5, 20, 50, 70 and 100 %) calculated using the SCALE2000, SandiaLab and Thermoddem databases.

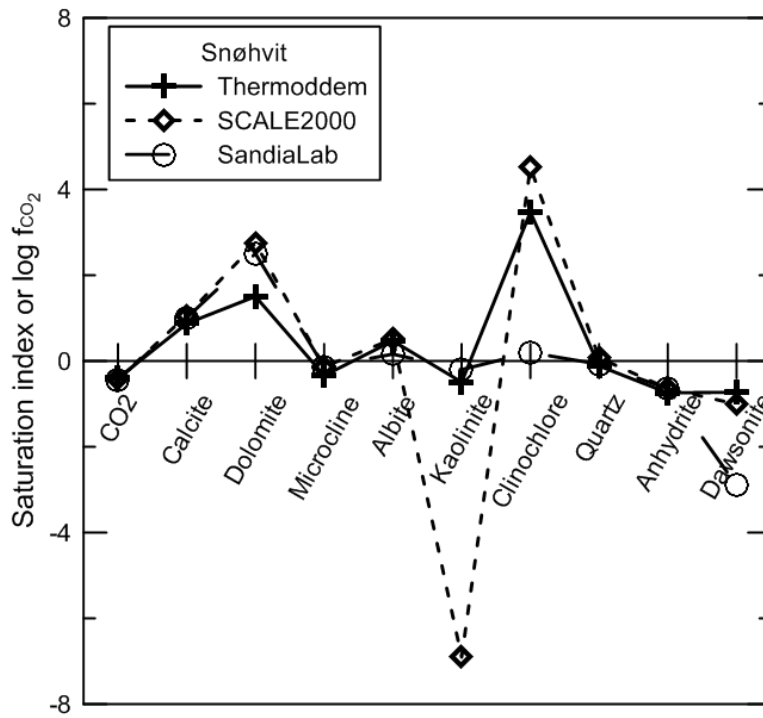


Figure 15: Mineral saturation indices and CO₂ fugacities of Snøhvit site porewater calculated using the SCALE2000, SandiaLab and Thermoddem databases.

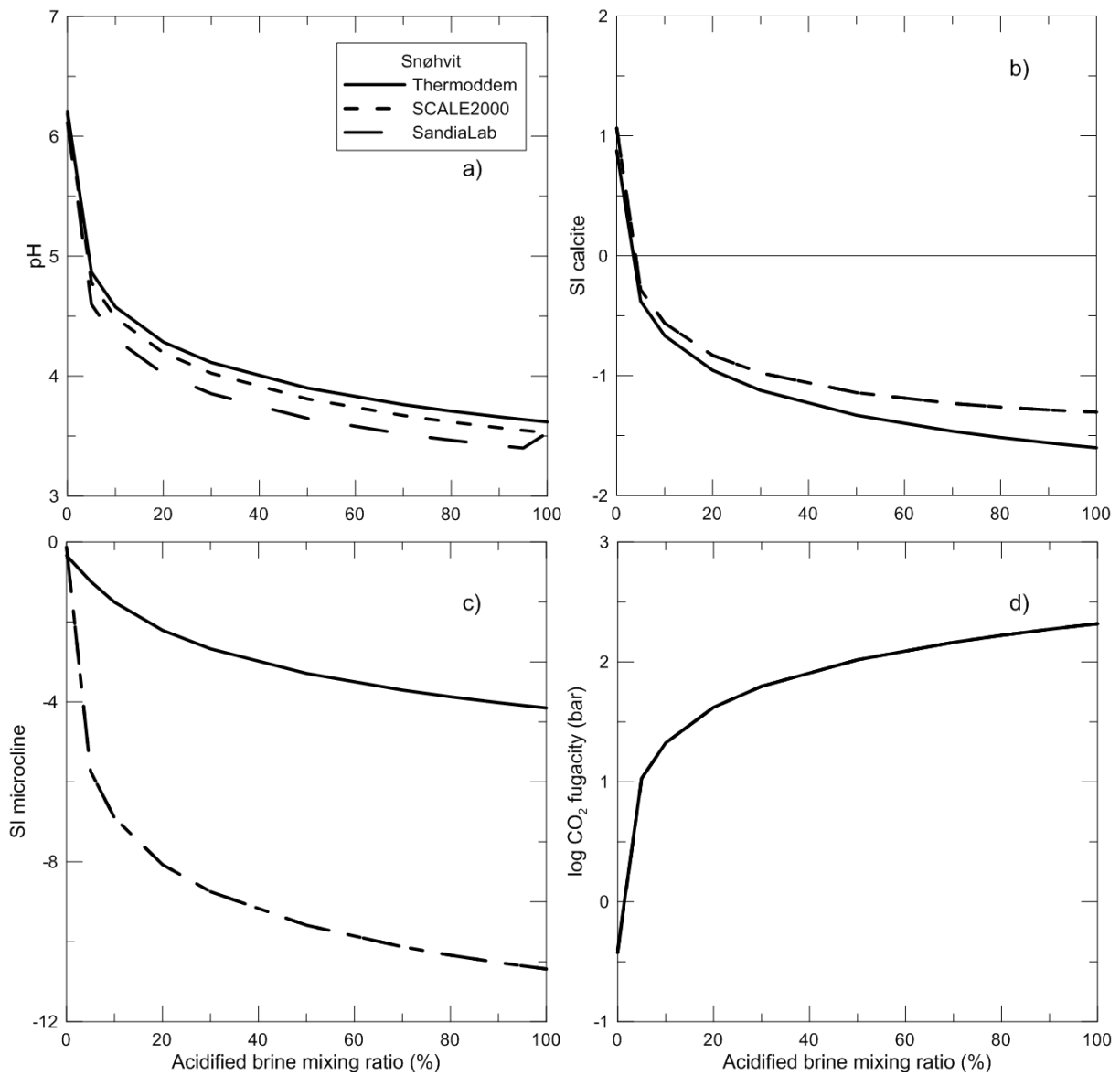


Figure 16: Evolution of a) pH, b) calcite saturation index, c) microcline saturation index and d) CO₂ fugacity of Snøhvit water as a function of the acidified brine mixing ratio (0 % for Snøhvit initial formation water to 100 % for Snøhvit water acidified at injection CO₂ fugacity) calculated using the SCALE2000, SandiaLab and Thermoddem databases.

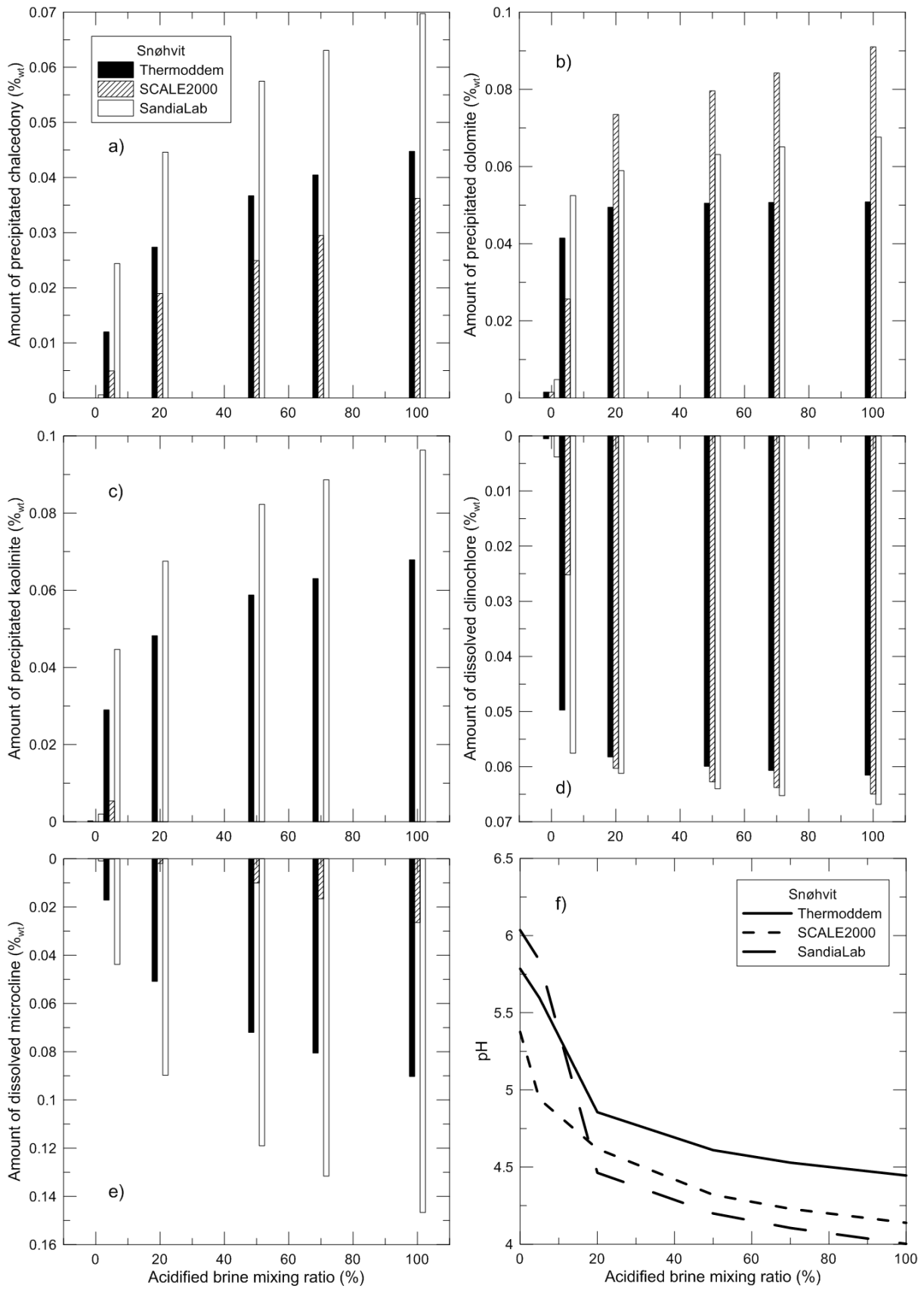


Figure 17: Amounts of a) chalcedony, b) dolomite, c) kaolinite, d) clinocllore and e) microcline precipitating or dissolving during 1000 years and f) final pH value for the Snøhvit case-study for different acidified brine

mixing ratios (0, 5, 20, 50, 70 and 100 %) calculated using the SCALE2000, SandiaLab and Thermoddem databases.

Table 1: Synthesis of natural analogues of CO₂ injection in deep sandstone aquifers

Name of CO₂ geological storage analogue	Bowen-Gunnedah-Sydney Basin	Cerro Barcino Formation	Miller Field
Geological formation and location	Permian Aldebaran Sandstone, Freitag Fm, Cattle Creek Fm, Wilton Fm and Black Jack Fm. Bowen, Gunnedah and Sydney Basins, eastern Australia	Cenomanian Cerro Castaño Member, Cerro Barcino Fm, Chubut Group. Los Altares, Patagonia, Argentina	Upper Jurassic Brae Fm at Miller oil and gas Field, South Viking Graben. North Sea, UK
References	Baker et al., 1995	Zalba et al., 2011	Baines and Worden, 2004; Marchand et al., 2001
CO₂ intrusion	Magmatic CO ₂ seepage at continental scale during Tertiary	CO ₂ seepage related to a regional intrusion of alkaline basic dykes during Eocene	CO ₂ accumulation mainly from inorganic crustal source. Accumulation persists at present-day with 15.95 to 64.3 % of CO ₂ in the gas phase
Temperature during rock-CO₂ interaction	Between 25 and 35 °C	Undetermined	About 120 °C, at present-day
Mineralogical composition	Quartz rich to rock-fragment rich sandstones: Quartz, feldspars (K-feldspars and feldspars from albite to anorthite series), rock fragments (volcanics, siliciclastics, metamorphics and granitics), dawsonite (up to 13.4 % with an average of 3.5 %), quartz, calcite, ankerite, siderite, kaolinite, illite, illite/smectite, alumohydroxides and alumohydrocalcite	Zeolitized and silicified tuffs and feldspathic litharenite sandstones: Quartz (40 to 80 %), analcime (2 to 52 %), mica (4 to 7%), sanidine (2 to 20 %), oligoclase (1 to 13 %), calcite (about 3 %), dolomite, dawsonite (up to 25 % and averages about 13 %), hematite and halite	Quartz arenite sandstones: Quartz (detrital and authigenic, 60 to 90 %), clays (about 2 %), muscovite (1 %), feldspars (2 to 6 %), calcite (2 to 11 %) and pyrite (about 2 %)
Brine composition	NaHCO ₃ -rich brines, with average concentrations of 6090 and 13360 mg L ⁻¹ , respectively	Na-rich solutions	Na, K, Cl and HCO ₃ -rich brines, with average contents of 27365, 1310, 43960 and 2495 mg L ⁻¹ , respectively
Diagenetic features linked to the CO₂ intrusion	Dissolution of feldspars and illitic clays. Precipitation of dawsonite, which fills pores and fractures and replaces plagioclases. Dawsonite postdates quartz,	Dissolution of oligoclase and analcime. Precipitation of dawsonite formed from analcime dissolution products and replacing pseudomorphically oligoclase	Extensive dissolution of feldspars (almost entirely dissolved), limited dissolution of calcite and creation of secondary porosity (6 to 15 %). Precipitation of minor

	calcite, ankerite, siderite and kaolinite precipitations, also probably related to CO ₂ intrusion	minerals. Precipitation of quartz, related to the transformation of oligoclase and analcime to dawsonite	volumes of ferroan dolomite and calcite
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Name of CO₂ geological storage analogue	Magnus Field	Bravo Dome	Vert le Grand
Geological formation and location	Upper Jurassic Magnus Sandstone member from Kimmeridge Clay Fm, North Viking Graben. North Sea, UK	Permian Wolfcampian-Leonardian Tubb Formation of the Bravo Dome reservoir. Northeastern New Mexico, USA	Upper Triassic Keuper Chaunoy Fm at Vert le Grand oil field, Paris basin. France
References	Baines and Worden, 2004	Pearce et al., 1996; Baines and Worden, 2004	Baines and Worden, 2004; Worden et al., 1999; Pinti and Marty, 1995
CO₂ intrusion	CO ₂ from thermal maturation of organic matter. 2 % of CO ₂ in the actual gas phase	CO ₂ accumulation linked to recent volcanic activity (100,000 to 8,000 years ago)	CO ₂ from organic matter maturation. 2 to 5 % of CO ₂ in the present-day gas phase. CO ₂ addition ceased 25 Ma ago
Temperature during rock-CO₂ interaction			Between 88 and 117 °C, at present-day
Mineralogical composition	Subarkosic sandstones: Quartz (52 to 62 %), feldspars (K-feldspars and plagioclases, 8 to 16 %) and lithic fragments (illite and glaucony grains, about 4 %), calcite (about 8 %) and pyrite	Arkosic sandstones: 25 to 65 % of K-feldspars and plagioclases, quartz, clays, zeolite, carbonates (dolomite, ferroan dolomite and siderite), hematite	Arkosic sandstones: several tens of percent of feldspars (K-feldspars, plagioclases, albite), quartz, carbonate cements (ferroan dolomite, dolomite and ferroan calcite)
Brine composition	Na, Cl and HCO ₃ -rich brines, with average concentrations of 11090, 17375 and 1040 mg L ⁻¹ , respectively		Na, Ca and Cl-rich brines, with average concentrations of 22000, 3600 and 40950 mg L ⁻¹ , respectively
Diagenetic features linked to the CO₂ intrusion	Partial dissolution of K-feldspars, associated with kaolinite and illite precipitation. Late-precipitation of ferroan dolomite and ankerite cements	Dissolution of dolomite, sulfate and aluminosilicates and precipitation of ferroan dolomite, siderite and small amounts of kaolinite possibly related to the CO ₂ intrusion. Large amounts of plagioclases, zeolite and clays remain in the sandstone	Dissolution of feldspars. Precipitation of quartz, ferroan dolomite, dolomite and calcite

Name of CO₂ geological storage analogue	Lam Formation	Songliao Basin	Hailer Basin
Geological formation and location	Upper Triassic Tithonian Lam Fm, Shabwa Basin. Yemen	Upper Cretaceous reservoir in Albian Qingshankou and Yaojia Fm, Honggang anticline, Songliao Basin. China	Lower Cretaceous CO ₂ gas reservoir in Namtum Fm, Hailer Basin. China
References	Worden, 2006	Liu et al., 2011	Gao et al., 2009
CO₂ intrusion	Elevated CO ₂ fugacity due to an influx of deep magmatic CO ₂	Accumulation of CO ₂ from mantle magmatic source	90 % of CO ₂ in present-day gas phase from magmatic source. Atmospheric CO ₂ also present during dawsonite precipitation
Temperature during rock-CO₂ interaction	85 to 105 °C	Between 100 and 120 °C	Between 85 and 105 °C
Mineralogical composition	Subarkosic to sublithic arenite sandstones. Detrital components: Quartz, feldspars (mainly K-feldspars, but also plagioclases and perthite), detrital calcite grains (trace to 10 %), micas and clays; cement: mainly carbonates (ferroan calcite, ferroan dolomite, dawsonite (up to 8 %) and siderite), quartz and traces of K-feldspars	Lithic arkose to feldspathic litharenite sandstones: Quartz (23 to 36 %), feldspars (14 to 25 %, mainly K-feldspars but also plagioclases and perthite), rock fragments (16 to 35 % including clasts from granite, micas, bioclasts and pyrite). Cement: dawsonite (2 to 27 %), calcite (up to 7 %), ankerite (3 %), illite, kaolinite, illite-smectite, quartz and feldspars (1 %)	Arkose to lithic arkose sandstones: Feldspars (14 to 48 %, mainly K-feldspars but also plagioclases and perthite), quartz (24 to 39 %), debris of granite, tuff and micas (2 to 26 %), dawsonite (2 to 22 % with an average of 10.8 %), siliceous cement (about 6 %), clay minerals (about 2.7 %) and ankerite (about 2.3 %)
Brine composition	Salinity of 180000 to 250000 mg NaCl L ⁻¹ with an average of 210000 mg NaCl L ⁻¹	Salinity ranges between 17000 and 144000 mg NaCl L ⁻¹ with an average of 56000 mg NaCl L ⁻¹	NaHCO ₃ -dominated brines
Diagenetic features linked to the CO₂ intrusion	Dissolution of K-feldspars and albite. Precipitation of dawsonite, which replaced K-feldspars and perthite grains and quartz, followed by precipitations of ferroan dolomite and siderite	Dissolution of feldspars and primary calcite. Precipitation of dawsonite, kaolinite, quartz, ankerite and late-generation calcite	Dissolution of feldspars and muscovite. Precipitation of quartz, kaolinite, dawsonite and ankerite as last cement

Name of CO₂ geological storage analogue	Ladbroke Grove Field	Montmiral reservoir	Fizzy accumulation
Geological formation and location	Early Cretaceous Pretty Hill Fm, Penola Trough at Ladbroke Grove, Otway Basin. Australia	Triassic sandstone, Sud-Est Basin. Montmiral, southeastern France	Early Permian Rotliegend Group, Fizzy accumulation. Southern North Sea, UK
References	Watson et al., 2004	Pauwels et al., 2007	Wilkinson et al., 2009
CO₂ intrusion	26 to 57 % of CO ₂ in the gas phase, related to recent volcanic activity (<1 Ma)	97.2 % of CO ₂ in the gas phase. CO ₂ accumulated since 35 Ma	About 50 % of CO ₂ in the gas phase, for at least 50 Ma
Temperature during rock-CO₂ interaction		103 °C at present-day	80 to 85 °C
Mineralogical composition	Lithic-rich sandstones: Quartz (45 to 70 %), feldspars (5 to 9 %), volcanic rock fragments (2 to 7 %), kaolinite (6 to 16 %), Mg/Fe-carbonates (2 to 8 %), siderite (1 to 8 %), calcite (<3 %) and chlorite (<2 %)	Feldspathic arenite sandstones: Quartz (62 to 79 %), K-feldspars (20 to 24 %), mica (1 to 12 %), ankerite (1 to 2 %) and traces of sphalerite and rutile	Sublitharenite sandstones: Quartz (49 to 62 %), fragments of quartz-rich sediments and muscovite (12 to 27 %), K-feldspars (2 to 10 %), dolomite (1 to 14 %), kaolinite (1 to 5 %), clay minerals (about 0.5 %), gypsum and anhydrite (about 1 %), dawsonite (0.1 to 0.8 %)
Brine composition	Na, Ca, Cl and HCO ₃ -rich brines with a salinity of 15000 mg NaCl L ⁻¹	Na, Cl and SO ₄ -rich brines with a salinity of 85000 mg NaCl L ⁻¹	
Diagenetic features linked to the CO₂ intrusion	Massive dissolution of feldspars, chlorite and calcite. Precipitation of quartz, kaolinite (replacing chlorite), ankerite and Fe/Mg-carbonates (replacing plagioclases). Precipitation of carbonates mainly in the gas phase zone, where the HCO ₃ content is 10 times higher than in the water zone of the reservoir	Dissolution of K-feldspars, ferroan dolomite and halite. Precipitation of kaolinite	Dissolution of K-feldspars. Precipitation of dolomite, little amounts of dawsonite and probably of quartz

Name of CO₂ geological storage analogue	Springerville – St Johns Field	Paradox Basin
Geological formation and location	Permian Supai Fm, Springerville – St Johns gas Field. Colorado Plateau, Arizona, USA	Jurassic Entrada sandstones, Paradox Basin. Colorado Plateau, Utah, USA
References	Moore et al., 2005	Pearce et al., 2011
CO₂ intrusion	About 90 % of CO ₂ (g). CO ₂ accumulation related to Pio-Pliocene volcanic activity	Circulation of CO ₂ -rich fluids related to Pio-Pliocene volcanic activity
Temperature during rock-CO₂ interaction		
Mineralogical composition	Siltstones: Quartz (26 to 46 %), plagioclases and K-feldspars (6 to 13 %, each), 5 to 18 % of illite, muscovite and biotite, kaolinite (2 to 6 %), dolomite (10 to 15 %), calcite (1 to 2 %), dawsonite (5 to 17 %), gypsum (1 to 9 %), anhydrite (1 to 2 %) and hematite (4 to 8 %)	Quartz (79 %), K-feldspars, albite, mica, pyrite (0.5 %) and hematite (0.5 %)
Brine composition	Na, Mg, SO ₄ and HCO ₃ -rich brines with a salinity of 4210 mg NaCl L ⁻¹	
Diagenetic features linked to the CO₂ intrusion	Dissolution of K-feldspars, plagioclases and calcite. Precipitation of dawsonite and kaolinite	Dissolution of lithic clasts, K-feldspars and albite. Precipitation quartz, carbonates and kaolinite

Table 2: Aquifer brine compositions prior to CO₂ injection at Ketzin (Sample KTZI 202, No 26 from Würdermann et al., 2010), In Salah and Snøhvit.

Components	Ketzin mg L⁻¹	In Salah mg L⁻¹	Snøhvit mg L⁻¹
Na	90400	35500	56418
Ca	2090	22400	4628
Mg	842	5276	477
Sr	48.9	506.0	207
Fe	5.56	262	14
K	282	225	496
Ba	0.082	10.0	4
Cl	139000	110250	96418
SO ₄	3744	656	210
HCO ₃	58.7	178	482
I	---	6.6	---
Br	44.9	---	---
Si	8.8	38	36
NH ₄	18.9	---	---
Mn	1.4	34	---
Li	1.8	13	---
B	36.2	8	---
Zn	---	1.5	---
Ni	---	---	1
Al	---	<3	---
P	---	<3	---
TDS (g L ⁻¹)	236.5	175.4	159.4
pH	6.4 (at 27°C)	5.2 (at 20°C)	6.2 (at 20°C)
Density (g cm ⁻³)	1.151 (at 27°C)	1.1279 (at 20°C)	1.104 (at 20°C)

Table 3: Mineral phases allowed to dissolve and precipitate in CO₂-brine-sandstone interaction models for Ketzin, In Salah and Snøhvit case-studies.

Case study	Minerals considered for kinetics dissolution	Minerals considered for equilibrium precipitation
Ketzin	Analcime Mg-Illite Microcline Clinochlore	Chalcedony Dolomite Kaolinite Dawsonite
In Salah	Mg-Illite Clinochlore	Chalcedony Dolomite Kaolinite
Snøhvit	Microcline Clinochlore	Chalcedony Dolomite Kaolinite

Table 4: Kinetic parameters used in the calculations: reaction rates at 25 °C (k_{25}), activation energies (E_a) exponential terms on H^+ activity (n) and reactive surface areas (A). Analcime kinetic parameters (k_{25} , E_a and n) taken as leucite parameters.

Mineral	A ($m^2 g^{-1}$)	Neutral mechanism		Acid mechanism		
		log k^{25} ($mol m^{-2} s^{-1}$)	E_a (KJ mol^{-1})	log k^{25} ($mol m^{-2} s^{-1}$)	E_a (KJ mol^{-1})	n
Analcime	35.5	-9.2	75.5	-6	132.2	0.7
Mg-Illite	30	-12.78	35.0	-10.98	23.6	0.34
Microcline	0.12	-12.41	38.0	-10.06	51.7	0.5
Clinochlore	0.0027	-12.52	88.0	-11.11	88.0	0.5