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# Temperature dependence of hydrogen diffusion in reservoir rocks: implications for hydrogen geologic storage†

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Hydrogen (H<sub>2</sub>) has recently gained momentum as a promising clean energy alternative to fossil fuels. The intermittent nature of renewable energy, as the source of green H<sub>2</sub>, necessitates temporary H<sub>2</sub> storage in subsurface geologic formations. To quantify storage potential and leakage risk, it is crucial to fully characterize subsurface H<sub>2</sub> transport behavior. This work aims to measure the diffusion of H<sub>2</sub> through relevant reservoir rocks, including two sandstones (Amherst Grey and Birmingham) and a limestone (Indiana). Breakthrough as a function of temperature is measured and used to calculate the effective diffusion coefficients and activation energy for diffusion at three different temperatures between 20 and 75 °C. Calculated diffusion coefficients are then used to estimate the subsurface plume size during storage in sandstone and limestone reservoirs. We observe that diffusive flow slightly expands plume size by up to 7%, and this effect is most pronounced in formations with low water saturation. While the use of cushion gas can maintain reservoir pressure and enhance injection efficiency, it can also enlarge H<sub>2</sub> plume and hinder the recovery process due to molecular diffusion if the cushion gas differs from H<sub>2</sub>.

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## 1. Introduction

Underground hydrogen storage (UHS) provides an affordable and viable option for large-scale energy storage.<sup>1–3</sup> In light of the rapid depletion of fossil fuels and progressively worsening environmental concerns caused by the extensive use of traditional energy, there is an imperative need to develop alternative clean energy sources for sustainable economic and societal development.<sup>4</sup> Clean energy sources generally include nuclear power and renewable energy sources. In either case, the strongly fluctuating nature of energy production has been noted as the primary barrier to widespread deployment.<sup>5–8</sup> Specifically, due to the surplus in energy demand, nuclear stations have evolved from being base load plants to providing power during short periods of peak demand.<sup>9</sup> Considering the high capital cost of nuclear power plants, it is not optimal to leave some of them idle during off-peak demand periods and have them only operate during peak demand periods. If energy is stored *via*

economic means at times of excess energy supply, then the required number of nuclear stations can be reduced. Similarly, renewable sources such as wind and solar energy also exhibit variability in energy supply as they are heavily dependent upon weather conditions.<sup>10,11</sup> To overcome the variability of both energy source and market demand, hydrogen (H<sub>2</sub>), generated by electrolysis (power-to-gas), is projected to be the most favorable energy carrier by 2030 and associatively, large-scale economical and reliable hydrogen storage is at the heart of the clean hydrogen economy.<sup>2,12–14</sup>

Geologic storage of H<sub>2</sub> in subsurface porous formation, due to its immense capacity and relatively low cost (compared with surface storage), has gained momentum in academia and the energy industry to facilitate the fossil fuel to renewable energy revolution.<sup>15,16</sup> Based on decades of cumulative research efforts, we have gained extensive knowledge of underground natural gas storage (UGS) in naturally occurring formations (hydrocarbon reservoirs, aquifers) and artificial cavities (salt caverns).<sup>17,18</sup> UGS generally comprises permeable rock layers which serve as reservoirs, overlain by strata with ultra-low permeability, serving as cap rocks for trapping reservoir fluid.<sup>14,19</sup> In theory, if H<sub>2</sub> molecules behave the same as natural gas under geologic conditions, UGS should satisfy the requirements for storing H<sub>2</sub> (*e.g.*, engineering, operation, safety and capacity). However, there are limited field demonstrations of UHS due to the high cost to

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produce H<sub>2</sub> (at least \$900 per ton vs. average oil cost of \$385 per ton) and uncertainty surrounding the injection and production of H<sub>2</sub> from the subsurface formations.<sup>2,3</sup> Two projects associated with UHS in subsurface porous media include the German Hydrogen to Store (H2STORE) project<sup>20</sup> and the Austrian Sun Storage project (Underground Sun Storage Project final report, 2017).<sup>21</sup> The H2STORE project<sup>20</sup> explored the feasibility of utilizing UHS in siliciclastic depleted gas reservoirs. It specifically focused on the potential impacts of hydrogen on the petrophysical, geochemical, and mineralogical properties of siliciclastic reservoir rocks and caprocks within large, depleted gas storage sites. The results showed that, despite interactions between hydrogen and reservoir, these interactions do not significantly degrade the purity of the recovered hydrogen,<sup>22,23</sup> thereby demonstrating the feasibility of utilizing UHS in depleted gas reservoirs. However, this project is restricted to synthetic reservoir simulation, and more extensive field studies are needed for long-term monitoring. On the other hand, the Underground Sun Storage Project<sup>21</sup> injected and recovered a mixture of hydrogen (~10 vol%) and natural gas from a litharenitic sandstone reservoir and revealed a recovery factor of 82% for H<sub>2</sub> injected. However, only a small volume of H<sub>2</sub> was involved in this field trial, significantly reducing the scale of H<sub>2</sub> storage capacity and efficiency, and the feasibility of large-scale UHS remains unclear. To upscale UHS applications, safe and economic injection and production of H<sub>2</sub> needs to be guaranteed, which necessitates a solid understanding of the hydro-, bio-, fluid-dynamics of UHS. Subsurface transport of H<sub>2</sub> is controlled by the hydrodynamic behavior of H<sub>2</sub>, which includes rock properties, fluid properties (density, viscosity, molecular diffusion coefficient, fluid–fluid interfacial tension, solubility), and rock–fluid interactions (wettability, solid–fluid interfacial tension, relative permeability). This behavior can be mathematically represented using a compositional two-phase flow model that includes water and gas as phases, with all pertinent chemical species as components.<sup>22</sup> Advective flow is dominant during active injection and withdrawal periods, affecting the displacement of formation fluids and hydrogen.<sup>24,25</sup> Feldmann *et al.*, 2016<sup>22</sup> conducted a numerical simulation research to investigate subsurface H<sub>2</sub> transport and storage in a depleted gas reservoir for five years of seasonal cyclic operation, focusing on hydrodynamic and gas mixing processes. They found that mechanical dispersion, due to the movement of fluids in the porous medium, combined with molecular diffusion, can amplify the gas mixing process, which is crucial for understanding H<sub>2</sub> behavior in the subsurface. Lysy *et al.*, 2021<sup>26</sup> developed a site-specific numerical simulation study to investigate the feasibility of UHS in depleted gas reservoir of Norne field,

offshore Norway. After several injection and withdrawal cycles, the final recovery factor obtained from a prolonged withdrawal period was 87%. During idle periods with no flow, molecular diffusion played a significant role in mixing and spreading of H<sub>2</sub> in the reservoir, leading to a 30% H<sub>2</sub> gas mixture. Similar conclusions were drawn by Hagemann *et al.* 2016,<sup>27</sup> who found molecular diffusion of H<sub>2</sub> strongly controls gas mixing process. A thorough understanding of H<sub>2</sub> molecular diffusion is essential for accurately estimating plume migration dynamics and optimizing the overall storage and recovery process.

In the present work, H<sub>2</sub> permeation (diffusion) behaviors in various reservoir rocks (sandstone and limestone) were measured using a modified two-bulb diffusion cell. Based on the observed diffusion rate, we calculate H<sub>2</sub> diffusion coefficients at various temperatures. By examining the temperature dependence of H<sub>2</sub> diffusion behavior, we investigated the diffusion mechanism of H<sub>2</sub> in reservoir rocks from the pore scale, which is rarely discussed in the literature. Additionally, the impact of diffusion on H<sub>2</sub> plume size during H<sub>2</sub> injection is analyzed. This work provides fundamental data for H<sub>2</sub> permeation in subsurface reservoir rocks at geologic conditions.

## 2. Background

To date, the Sun project<sup>21</sup> is the only completed field trial that has resulted in a full field dataset to demonstrate UHS feasibility in depleted hydrocarbon reservoirs. One primary objective of this project was to investigate fundamental mechanisms controlling H<sub>2</sub> loss in subsurface reservoirs subjected to repeated injections and extraction cycles. Because of the concentration contrast between the injection stream and reservoir fluid and high H<sub>2</sub> diffusion coefficients, the field trial<sup>21</sup> shows that about 4 to 5% of H<sub>2</sub> was diffused and dissipated into the reservoir and could not be recovered. It should be noted that the injected fluid only contained 10 vol% of H<sub>2</sub> and underwent a relatively short storage period (~4 months). H<sub>2</sub> diffusive loss is expected to be considerably larger for longer-duration, pure H<sub>2</sub> storage. There have been limited experimental studies<sup>28–32</sup> to measure H<sub>2</sub> diffusion in minerals and some reservoir rocks, as compiled in Table 1. Except for diffusion measurements on coal, other existing works<sup>28–30,33–35</sup> were conducted at low-temperature and low-pressure conditions. Even for coal, the diffusion coefficients were quantified through gas adsorption kinetics that inherently assumed a strong adsorbent-adsorbate system.<sup>31,32</sup> This may be

**Table 1** Literature survey of hydrogen diffusion in reservoir rocks

| Rock type                                      | $D$ (m <sup>2</sup> s <sup>-1</sup> )                | $T$ (°C)  | $P$ (MPa) | Sample type | Porosity     | Source                                       |
|------------------------------------------------|------------------------------------------------------|-----------|-----------|-------------|--------------|----------------------------------------------|
| Sandstone                                      | $6.5 \times 10^{-6}$ – $1.9 \times 10^{-5}$          | 25        | 0.1       | Core        | 0.2–0.4      | Currie 1960 <sup>34</sup>                    |
| Sandstone (Berea, Bandera, <i>etc.</i> )       | $4 \times 10^{-6}$ – $1 \times 10^{-5}$              | 25        | 0.1       | Core        | 0.18–0.19    | Donaldson <i>et al.</i> , 1976 <sup>36</sup> |
| Berea sandstone                                | $2.1 \times 10^{-7}$ – $1.5 \times 10^{-6}$          | 28        | Up to 4   | Core        | 0.19–0.25    | Arekhov <i>et al.</i> , 2023 <sup>33</sup>   |
| Sandstone (Bentheimer, Chattian, <i>etc.</i> ) | $5 \times 10^{-9}$ – $2 \times 10^{-7}$              | Up to 100 | Up to 30  | Core        | 0.17–0.32    | Michelson <i>et al.</i> , 2023 <sup>37</sup> |
| Gypsum                                         | $1.5 \times 10^{-5}$                                 | 22        | 0.1       | Sheet       | Not measured | Yang <i>et al.</i> , 2013 <sup>35</sup>      |
| Opalinus claystone                             | $(1.2\text{--}1.8) \times 10^{-9}$ (water saturated) | 20        | 0.1       | Core        | 0.237        | Strauch <i>et al.</i> , 2023 <sup>30</sup>   |
| Anthracite coal                                | $(1\text{--}6.8) \times 10^{-9}$                     | Up to 60  | 1.3       | Powder      | Not measured | Keshavarz <i>et al.</i> , 2022 <sup>31</sup> |
| Coal in different ranks                        | $(0.5\text{--}1.8) \times 10^{-10}$                  | 30        | Up to 10  | Powder      | Not measured | Liu and Liu 2023 <sup>32</sup>               |



true for the  $H_2$  – coal system (as coal is a microporous media) but is unlikely to be valid for other non-sorptive reservoir rock types. For this reason, the present work aims to provide a benchmark dataset on  $H_2$  diffusion in natural rock samples at elevated temperatures using an improved diffusion cell design. Additionally, by systematically exploring the temperature effects on diffusion, we can determine activation energies that can be used to extrapolate diffusion behavior for a wide range of reservoir temperature conditions.

### 3. Methods

#### 3.1 Experimental diffusion apparatus and procedure

This work adopts the two-bulb method to measure the self-diffusion coefficient of  $H_2$  through the reservoir rocks at dry conditions (samples are filled with air). Fig. 1 shows the experimental apparatus that includes two bulb volumes and a connecting bridge of 316 stainless steel for holding the rock sample. The cell was constructed using ConFlat® parts with

copper gaskets. The rock cores used for experiments were 1" long by 1" in diameter. The cores were sealed on the outside using a UV-curing resin (UV15X-6NONMED-2, Master Bond, Inc., Hackensack, NJ, USA). The cores were then placed in the center of the bridging pieces, which were 1.5" in diameter. The axial space around the core was filled with Room Temperature Vulcanization (RTV) silicone (SILPAK, Inc. Pomona, CA, USA) to prevent diffusion around the outside of the rock core. This silicone was previously shown to be impermeable to gases.<sup>38</sup> The void volume of the system's components (including connecting pipe, bulb, bridge, ball valve) is measured through water displacement. The system dimensions are provided in Table 2.

To begin a diffusion experiment, the bottom bulb as depicted in Fig. 1 was spiked with a 5%  $H_2$  mixture to reach an injected  $H_2$  volume of 2 mL (approximately 15 800 ppmv). Each bulb was connected to a mass spectrometer (Pfeiffer Vacuum OmniStar GSD 320 mass spectrometer, Pfeiffer Vacuum, Aßlar, Germany) through one outlet port, which was connected to a dead-ended multiposition selector valve (VICI Valco Instruments, Houston, TX). The second port was connected to a length of capillary tubing open

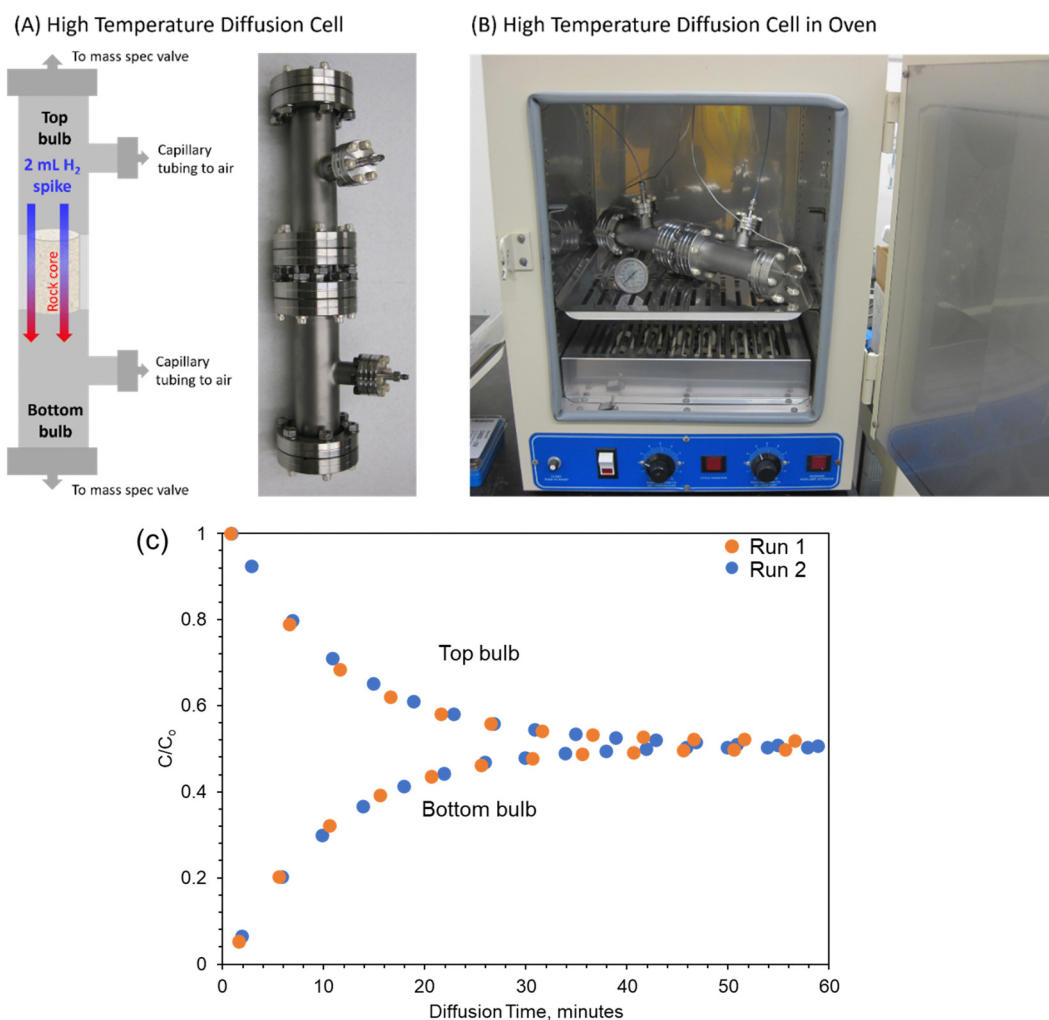


Fig. 1 (A) The two-bulb apparatus used in this study to measure  $H_2$  diffusion through reservoir rocks; (B) the laboratory oven for high-temperature experiments; (C) replicate experimental runs for Birmingham sandstone at 45 °C.



Table 2 System dimension

|                                                          |                         |
|----------------------------------------------------------|-------------------------|
| Length of bridge                                         | 2.54 cm                 |
| Cross-sectional area of bridge                           | 5.067 cm <sup>2</sup>   |
| Volume of top bulb (including connecting pipe, valve)    | 126.305 cm <sup>3</sup> |
| Volume of bottom bulb (including connecting pipe, valve) | 126.305 cm <sup>3</sup> |

to the atmosphere. This tubing prevents the formation of a vacuum during gas sampling, which would affect diffusion rates, while also preventing extensive back diffusion of air. This port allows for the replacement of sampled gas with air, which will result in some dilution of the spiked and diffusion chambers with air. The multi-position valve was used to sample from both bulbs for one minute every four minutes, and measured air the rest of the time to prevent oversampling. The sampling rate of gas from the mass spec is 0.2 mL minute<sup>-1</sup>. Therefore, the total gas sampled from the system over the course of the diffusion experiment is approximately 4 to 5 mL for an 80-to-100-minute experiment. The volume of the spiked and diffusion chambers is 126.3 cm<sup>3</sup>, so this volume is equivalent to 3.2–4.0% of the chamber volume being removed and replaced by air. Diffusion experiments were run until the two bulb concentrations equilibrated. Experiments were conducted at ambient temperatures up to 60–75 °C. Between diffusion runs, samples were placed in a vacuum oven to remove residual hydrogen gas. The impact of dilution due to sampling can be found in Fig. S1 in the ESI† Dilution did have a slight impact on the  $c/c_0$  values. However, this impact was very small (0.2–3.7% standard deviation in  $c/c_0$  values) over the considered diffusion timescale. Additionally, because sampling rates are equivalent between spiked/breakthrough chambers, rock types, and temperatures, we do not anticipate a significant change in the concentration gradient, which is the driving force for diffusion. Since the diffusion is normalized to  $c_\infty$  (see Section 3.3), changes in the equilibrium concentration are accounted for in diffusion coefficient calculation.

### 3.2 X-ray diffraction (XRD) analysis

For mineral analysis, quantitative X-ray diffraction (QXRD) analysis was carried out to identify the mineralogy of the studied rock samples.<sup>39</sup> Powdered samples were used, and the generated diffraction patterns were utilized to visualize diffraction measurements and identify the proportion of different minerals contained in each rock sample.<sup>39</sup> Samples were measured using a Siemens D-500 X-ray diffractometer using CuK $\alpha$  radiation (Bruker AXS, Madison, WI, USA) and quantification was accomplished using alumina powder (Al<sub>2</sub>O<sub>3</sub>) as an internal standard. Peak identification used JADE software's search-match function (Materials Data, Inc., Livermore, CA, USA). The measured patterns were compared to a library of XRD patterns of known minerals, and the relative abundance of individual minerals was estimated based on peak intensities, providing quantitative mineralogy information.

The two sandstones are rich in quartz, ranging in abundance from 85.4 wt% to 91.5 wt% (Table 3). K-feldspar (Microcline) and Clay (Kaolinite) are also identified in varying proportions. The limestone predominantly consists of calcite with a value around 99 wt% and a minor constituent of quartz of 1 wt%.

Table 3 Mineral weight percentages for the studied rock samples

| Sample                 | Calcite | Quartz | Microcline | Kaolinite |
|------------------------|---------|--------|------------|-----------|
| Amherst Gray Sandstone | —       | 85.4%  | 10.8%      | 3.7%      |
| Birmingham Sandstone   | —       | 91.5%  | 5.6%       | 2.8%      |
| Indiana Limestone      | 98.8%   | 1.2%   | —          | —         |

### 3.3 Diffusion theory

The two-bulb method has been extensively used to measure gaseous diffusion over wide temperature ranges from 65 to 400 K. Due to the relative compactness and convenient opening mechanisms (without moving parts), the two-bulb method facilitates thermal equilibrium, which makes it easier to measure diffusion at different temperatures.<sup>40</sup> Because the bulb volume is substantially larger than the bridge volume, the following assumptions are made:<sup>40–42</sup>

- Diffusion through the rock sample is in quasi-steady-state – constant flow flux along the connecting tube.
- A linear variation in concentration is established in the tube bounded by the concentration in the bulbs.
- Concentration gradients occur only in the connecting tube, and uniform concentration is surmised in the bulbs.

The first two assumptions are derived from the quasi-steady state that neglects the transient period required to establish a constant gradient across the entire length of the tube. Departure from a quasi-steady state often arises due to design artifacts of the apparatus when the bulbs are not sufficiently large enough compared to the connecting tube. Because our system has bulb volume exceeding the connecting tube volume by more than tenfold, the tube and the rock sample inside it are surmised to be in a quasi-steady state. The last assumption is made on the basis of much smaller diffusivity in the connecting tube than in the bulbs. This occurs because the rock samples inside the connecting tube occupy the available void space for diffusion, which leads to a significant decrease in the diffusion rate.

To determine the effective diffusion coefficient, the time-series concentration of the bulb is given by<sup>40–42</sup>

$$\frac{c(t)}{c_0} = \left(1 - \frac{c_\infty}{c_0}\right) e^{-\frac{DA(V_1+V_2)}{LV_1V_2}t} + \frac{c_\infty}{c_0} \quad (1)$$

where  $c(t)$ ,  $c_0$ ,  $c_\infty$  correspond to H<sub>2</sub> concentration at current, initial and equilibrium condition.  $V_1$ ,  $V_2$  correspond to cell volume of top and bottom bulb, respectively.  $A$  and  $L$  correspond to the cross-sectional area and length of the rock sample.

In the above formula,  $c(t) \rightarrow c_\infty$  for sufficiently large diffusion time (*i.e.*, time to reach equilibrium conditions). A linear relationship can be found by log-transformation of the concentration solution, *i.e.*,

$$\ln(c_\infty - c(t)) = -\frac{DA(V_1 + V_2)}{LV_1V_2}t + \ln(c_\infty - c_0) \quad (2)$$

The effective diffusion coefficient of the rock samples can be determined from the slope of the semi-log plot of  $(c_\infty - c)$  against diffusion time.  $(c_\infty - c)$  is the concentration change in the spiked or breakthrough bulb, monitored through continuous sampling





and mass spectrometry analysis. The detailed derivation can be found in ESI.†

## 4. Results and discussion

### 4.1 Experimental quantification of diffusion coefficients for different rock samples

Fig. 2 shows the measurements of  $H_2$  concentration in the top and bottom bulbs as the system approaches equilibrium, which is found to follow an exponential series. All experimental trials have the same initial  $C/C_0$  of  $H_2$  of 1. The equilibrium  $C/C_0$  varies slightly among rock samples, ranging from 0.43 to 0.47. Since the top and bottom bulbs have the same volume, the concentration, in theory, should approach  $0.5 \text{ mol cm}^{-3}$  at equilibrium. The difference between measured and theoretical equilibrium concentration implies the existence of gas retention within the rock samples, likely caused by bulk phase gas trapped in dead-end pores and/or gas condensed at the pore surface (adsorption).

Because the bulbs have substantially larger volumes than the volume of the connecting tube, initially the transient period

can be neglected, and a quasi-steady state is surmised. As a result, a linear relationship is found in the semi-log plot of  $H_2$  concentration *versus* diffusion time for all rock samples studied during the initial  $H_2$  breakthrough period (first 20–60 minutes) (see Fig. 3). The slope of these linear lines is used to determine the effective diffusion coefficient for the studied rock samples following eqn (2). Table 4 summarizes the measured diffusion coefficient for Birmingham Sandstone, Amherst Gray Sandstone and Indiana Limestone at investigated temperatures, where Birmingham Sandstone has the highest  $H_2$  diffusion coefficient ( $0.045\text{--}0.07 \text{ cm}^2 \text{ s}^{-1}$ ) and limestone possesses the lowest  $H_2$  diffusion coefficient ( $0.026\text{--}0.033 \text{ cm}^2 \text{ s}^{-1}$ ). With reference to  $H_2$  diffusion coefficients in air at comparable temperatures, the measured diffusion coefficients for all tested rocks are approximately two orders of magnitudes smaller. This is caused by solid grains present in the diffusive pathway, which reduces the available void volume for flow and elongates the streamlines. Thus, sandstones with higher porosity have higher diffusion coefficients than limestone with lower porosity.

Previous work<sup>43</sup> indicates that the methane diffusion coefficient in sandstone with porosity of 10–20% typically falls within the range of  $0.002$  to  $0.01 \text{ cm}^2 \text{ s}^{-1}$  at ambient temperature.

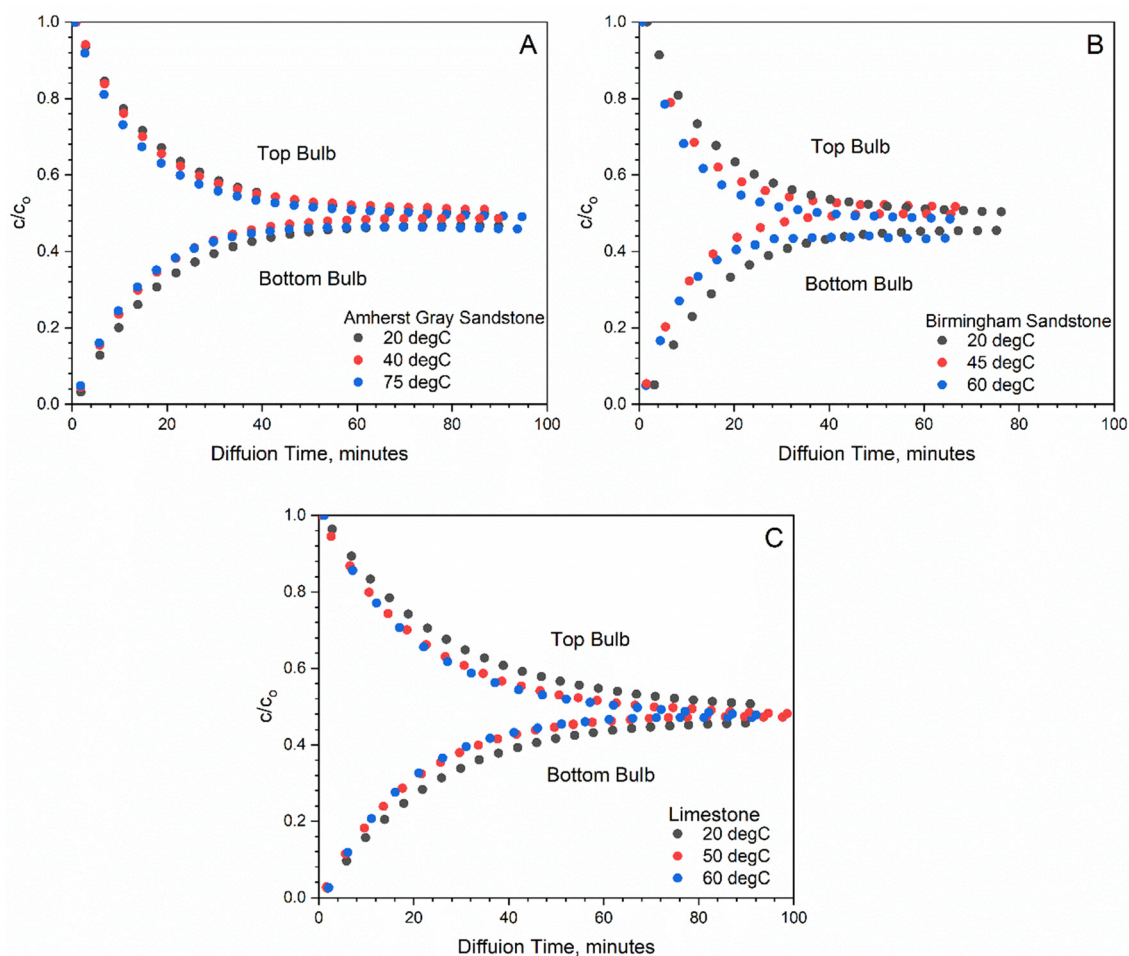


Fig. 2 Changes in concentration of hydrogen in the top and bottom bulb for two sandstone (A, B) and one limestone (C) samples at different temperatures.



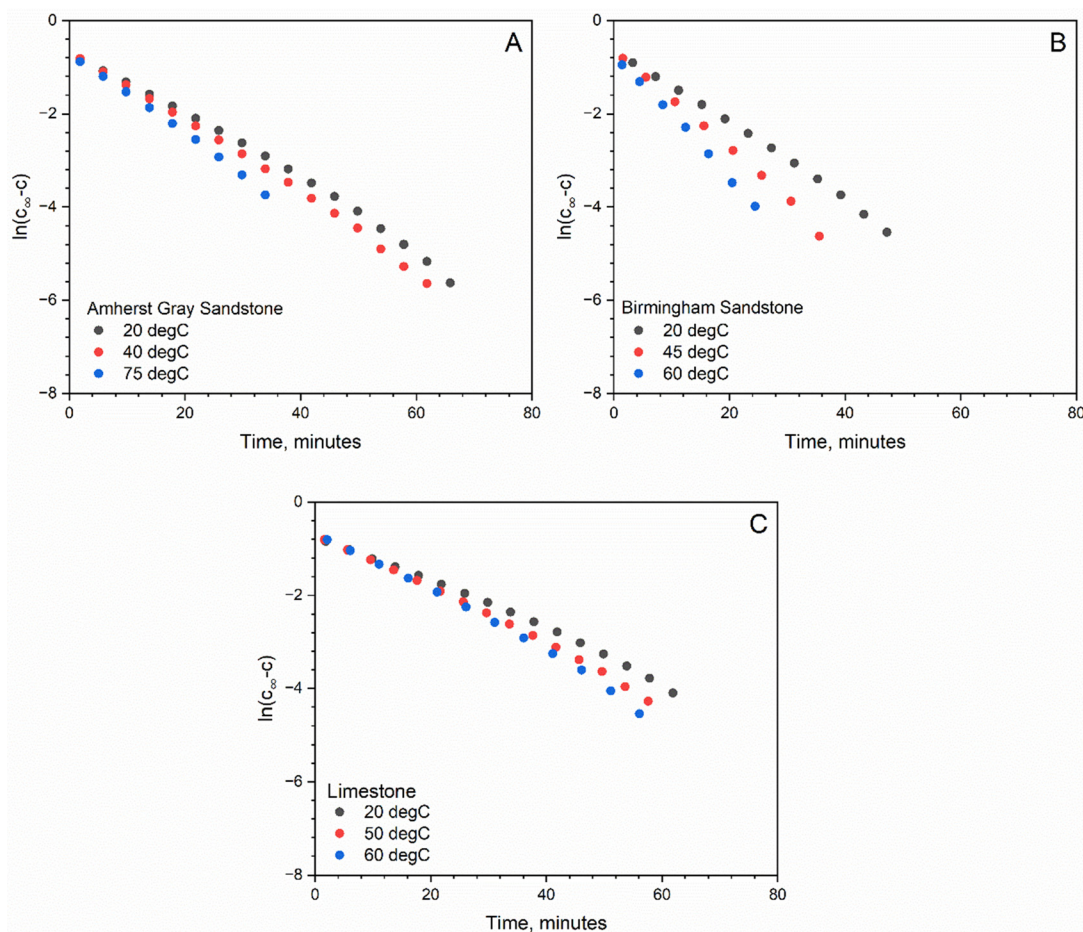


Fig. 3 log transformation of time series concentration to calculate diffusion coefficient according to eqn (2) for two sandstone samples (A, B) and one limestone sample (C).

**Table 4** Measured diffusion coefficients for  $H_2$  diffusion into air-filled rock samples at different temperatures in  $10^{-6} m^2 s^{-1}$ , activation energy ( $E_A$ ) in  $J mol^{-1}$ , and associated MIP-derived porosity. Permeability ( $K$ ) are in units of mD to air, which can be found from the vendor's website (Core Sample|Quarry Source (<https://petroleumcores.com>)). ( $T_{ambient}$ : 20 °C;  $T_{middle}$ : 40–50 °C;  $T_{high}$ : 60–75 °C)

| Sample                 | $D_{ambient}$ | $D_{middle}$ | $D_{high}$ | $E_A$ | $\phi$ | $K$     |
|------------------------|---------------|--------------|------------|-------|--------|---------|
| Amherst Gray Sandstone | 3.6           | 4            | 4.7        | 4174  | 0.185  | 15–150  |
| Birmingham Sandstone   | 4.5           | 5.9          | 7          | 9108  | 0.228  | 250–500 |
| Indiana Limestone      | 2.6           | 3.1          | 3.3        | 4650  | 0.053  | 2–20    |
| In ordinary air        | 71            | 87           | 95         |       | 1      |         |

In this work, we expect the methane diffusion coefficients of our sandstone samples, which have similar porosities (see Table 4), to be relatively close in value. Because of the relatively small molecular size and lower viscosity of  $H_2$  molecules, the obtained  $H_2$  diffusion coefficient is notably greater than that of methane, reaching values up to 20 times higher. Most reservoir engineering tools used in conventional hydrocarbon gas production are often limited to passive transport (advective flow equations) due to the comparatively small flux transported through Brownian motion (molecular diffusion), which is no longer a valid assumption for subsurface  $H_2$  flow in reservoir

rocks. Although Fick's diffusion law is considered in some unconventional reservoir engineering tools, simplifications are made in diffusion modeling to reduce computational cost.<sup>44,45</sup> For example, diffusion coefficient is often treated as a constant throughout reservoir and over time,<sup>46,47</sup> which conflicts with the reality that diffusion varies with fluid composition and reservoir conditions. Therefore, an advection–diffusion system must be coupled with the governing equations for incompressible flow within porous media to develop hydrodynamic modeling tools for the geologic storage of  $H_2$ .

## 4.2 Relationship between temperature and effective diffusion coefficients of $H_2$

The temperature dependence of the diffusion coefficient is useful to correlate diffusion behaviors of reservoirs at different depths. For all rocks studied, the  $H_2$  diffusion coefficient has a monotonically increasing trend with increasing temperatures (see Fig. 4). This is because higher temperatures lead to increased gas molecular kinetic energy, facilitating faster breakthrough of gas molecules into the receiving bulb. In particular, Birmingham Sandstone has diffusion behavior that is most sensitive to temperature: its diffusion coefficient increases by 57% as temperature increases from 20 °C to 60 °C. When subjected to a



similar temperature increase, the measured diffusion coefficients of the other two rocks only increase by less than 30%. Overall,  $H_2$  diffusion in reservoir rocks is strongly dependent on temperature. Storage reservoirs at higher temperatures are thus expected to have more diffusive flow. On average, the growth rate of  $H_2$  diffusion coefficient per  $1^\circ\text{C}$  increase in temperature is estimated to range from 0.5% to 1.4%. Because the rate of diffusion is temperature-dependent, the Arrhenius equation is applied to describe how  $H_2$  diffusion through rock samples varies with temperature. The linear relationship between the natural log of the effective diffusion coefficient and the inverse temperature was also used to calculate activation energy,  $E_A$ , according to the Arrhenius equation (Table 4). Knowing the value for  $E_A$  is critical to quantify how sensitive the diffusion rate is to temperature changes and for predicting diffusion rate at temperatures where experimental measurements are not available.

Despite the inherent heterogeneity present in reservoir rocks, a single value of diffusion coefficient is often used to represent the transport capacity for the entire rock sample. This is valid as long as effective medium approximation is applicable, where the average domain size (microscopic length scale) is substantially larger than the molecular dimension but much smaller than the characteristic length of the macroscopic sample.<sup>48,49</sup> To evaluate this assumption, mercury intrusion porosimetry (MIP) was conducted to characterize pore volume and its distribution for the studied rock samples. MIP operates on the basis of using high injection pressure to force mercury into pore space in a porous medium. Since smaller pores have higher resistance to mercury penetration, the externally applied pressure builds up as mercury accesses smaller pores. Based on the correlation between capillary (entrance) pressure and pore size, pore size distribution can be determined from the recorded mercury intrusion volume at specific pressures.<sup>50</sup>

Assuming a cylindrical pore model, the pore size distribution is determined using the Washburn law:<sup>51</sup>  $P = (-4\gamma \cos \theta)/d$ , where  $P$  is the mercury injection pressure,  $\gamma$  is the surface tension of mercury ( $\sim 484 \text{ mN m}^{-1}$ ),  $\theta$  is the solid/mercury

contact angle ( $\sim 130^\circ$ ),  $d$  is the penetrating pore throat size. The obtained porosity values for the three rocks studied are listed in Table 4 with Birmingham Sandstone being the most porous rock (hence having the largest diffusion coefficient) and the corresponding pore size distributions (PSDs) are shown in Fig. 5. The two sandstone samples have similar bimodal pore size distributions between 20 nm and 300  $\mu\text{m}$  with two clearly separated maxima above 1  $\mu\text{m}$ . In comparison, a lower proportion of large pores is found in the Limestone sample, but with comparable pore sizes ranging between 30 nm and 300  $\mu\text{m}$ . The microscopic length scale is approximated as the average pore size, ranging from 0.66  $\mu\text{m}$  (Limestone) to 1.2  $\mu\text{m}$  (Birmingham Sandstone), which is substantially smaller than the macroscopic length scale involved in the testing system ( $\sim 2.54 \text{ cm}$ ). As the average domain size is significantly smaller than the characteristic length of the sample, it fulfills one of the requirements for using the effective medium approximation.

In order to apply the effective medium approximation, the other requirement is that the pore scale involved in diffusion is much greater than the molecular dimension of diffusing species. To satisfy this requirement, bulk diffusion must be the dominant diffusion regime. The mean free path of  $H_2$  under normal conditions (1 atm and  $25^\circ\text{C}$ ) is 130 nm<sup>52</sup> and is expected to be larger as kinetic energy increases with increasing temperature. Since the lower limit of the MIP-derived PSDs is less than 130 nm, it is possible for gas molecules to collide with the pore wall in addition to intermolecular collisions. In other words, both Knudsen and bulk diffusion are likely to occur in bulk-phase  $H_2$  diffusion for the studied reservoir rocks when a concentration gradient exists. Theoretically, Knudsen diffusion is driven by collisions between gas molecules and the pore wall, whereas intermolecular diffusion causes bulk diffusion.<sup>53,54</sup> The contribution of individual diffusion regimes to overall diffusive flow flux depends on the relative magnitude of the mean free path to pore size (Knudsen number).<sup>55,56</sup> In general, if Knudsen number is greater than 0.1, then Knudsen diffusion should be considered. At the investigated temperatures, the

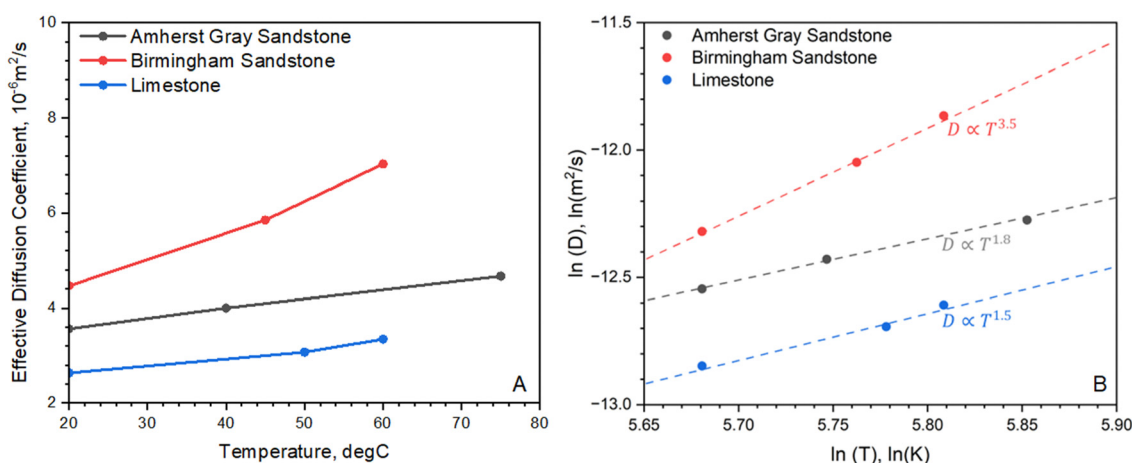


Fig. 4 Temperature dependence of diffusion coefficient in (A) normal scale and (B) log–log scale for Amherst Gray Sandstone, Birmingham Sandstone and Indiana Limestone.





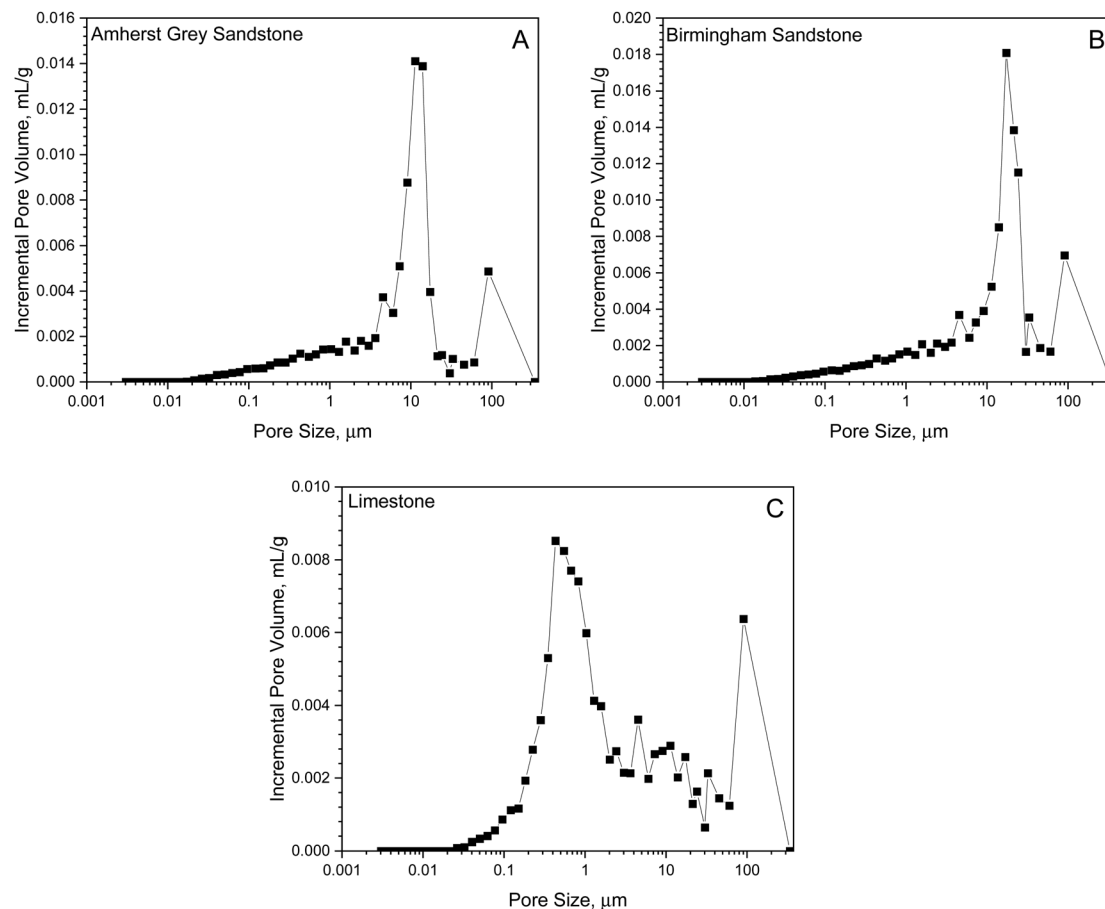


Fig. 5 Pore size distribution on (A) Amherst Gray Sandstone, (B) Birmingham Sandstone, and (C) Indiana Limestone obtained from MIP with  $\theta = 130^\circ$ .

mean free path of  $H_2$  molecules range from 111 nm to 118 nm and therefore, Knudsen diffusion manifests at pores with pore size less than 1  $\mu\text{m}$ , which contributes to  $\sim 10\%$  of the total pore volume of studied sandstone samples. As the mean free path increases with temperature, the frequency of  $H_2$  molecules colliding with the pore wall also increases at higher temperatures, adding additional momentum to diffusive flow. The contribution of Knudsen diffusion to overall diffusive flux, therefore, varies with temperature and the microscopic length scale involved.

If molecular collisions are of cardinal importance in the overall diffusive transport, the resulting derivative of diffusion coefficient with temperature  $\left(\frac{\partial \ln D}{\partial \ln T}\right)_p$  lies between  $3/2$  and  $2$  for binary molecular diffusion.<sup>40</sup> The temperature dependence of the binary gaseous diffusion coefficient,  $D$  under bulk diffusion is inferred from the Chapman–Enskog kinetic theory of gas,<sup>57</sup>

$$D = \left(\frac{M_1 + M_2}{2M_1M_2}\right)^{\frac{1}{2}} T^{\frac{3}{2}} \frac{1}{P \omega \Omega_0} \quad (3)$$

where  $M_1M_2/(M_1 + M_2)$  is the reduced mass of a pair of molecules.  $\omega$  is a measure of molecular size.  $\Omega_0$  is the collision

integral that contains the contribution of the intermolecular forces and, as such, exhibits some degrees of temperature dependence. The derivative of  $\Omega_0$  with temperature is found to vary between  $-1/2$  and  $0$  (dependent upon the magnitude of repulsive or attractive forces), and thereby  $D$  varies as  $T^{\frac{3}{2}}$  to  $T^2$ .

Fig. 4B shows the log transformation of temperature–diffusion coefficient data. The resulting derivative  $\left(\frac{\partial \ln D}{\partial \ln T}\right)_p$  for Amherst Sandstone is  $1.8$ , and for limestone is  $1.5$ , within the range of theoretical prediction between  $T^{\frac{3}{2}}$  and  $T^2$ . However, the derivative  $\left(\frac{\partial \ln D}{\partial \ln T}\right)_p$  for Birmingham Sandstone exceeds the limit of bulk diffusion. It is likely to be caused by Knudsen diffusion processes, which provides additional momentum through pore wall collisions, resulting in a higher temperature dependence of the diffusion coefficient.

Although most pore volume in the studied reservoir rocks is provided by large pores ( $> 1 \mu\text{m}$ ) and nanopores only contribute less than  $20\%$  of the total porosity, the outcome of this work suggests that the confinement effect on  $H_2$  diffusion flow in Birmingham Sandstone due to collisions with pore walls manifests as reservoir temperature increases. Based on Fig. 5, the limestone sample appears to have a more uniform pore size





distribution than the two sandstone samples. Therefore, less temperature dependence is found in  $H_2$  diffusion through limestone compared to sandstone.

### 4.3 Geometric factor of natural rocks

Compared with the molecular diffusivity of  $H_2$  in ordinary air, the measured diffusion coefficients in the studied rock samples are one order of magnitude smaller than the diffusivity in the absence of porous media. This is caused by  $H_2$  interaction with the surrounding pore network, which increases diffusional resistance and reduces diffusion rate. In any porous system, the presence of solid grains decreases the void volume for transport and simultaneously, causes the diffusive path of gas molecules to deviate from straight lines. Consequently, the diffusion coefficients must be scaled with tortuosity and porosity. As noted earlier, the effective medium approximation is reasonable for the system studied herein, because the microscopic length scale is much smaller than the macroscopic length scale. Based on effective medium approximation, the relationship between the effective diffusion coefficient of porous media,  $D$  and the self-diffusion coefficient (*i.e.*, gaseous diffusion coefficient in ordinary air),  $D_p$  is given by<sup>58</sup> (Millington 1959),

$$D = D_p \phi / \tau. \quad (4)$$

The above formula only considers transport through available pore spaces and neglects mass flux through the solid grain. The values for  $D_p$  can be obtained from Table 4, which lists  $H_2$  diffusion coefficient in ordinary air at various temperatures. From a geometrical point of view,  $\phi$  accounts for the reduced area for diffusion, and  $\tau$  accounts for the elongated diffusion path between two points in a porous medium relative to the direct distance between points. There are no consistent observations between  $\phi$  and  $\tau$ , but in general, a decrease in  $\phi$  causes an increase in  $\tau$  as more solid grains are present in the diffusion path.<sup>59</sup> Unlike  $\phi$ ,  $\tau$  is not directly measurable for natural rocks. The magnitude of  $\tau$  depends on various factors, including grain geometry and dimension, pore size distribution and connectivity, and diffusing species.<sup>60</sup> Thus, the effective diffusion coefficient of any given system is unique to that system and requires experimental quantification.

In this work, molecular diffusion measurements are applied to determine the tortuosity of pore channels. The porosity and tortuosity of reservoir rock samples are directly related to the ratio of the effective diffusion coefficient to the molecular diffusion coefficient in the absence of a porous medium. Using eqn (4), the tortuosity values for the studied rock samples was estimated from the measured effective diffusion coefficients and the results are shown in Fig. 6. The estimated tortuosity from diffusion measurements varies between 3 and 4 for the two sandstones. These values resemble results from Donaldson *et al.* (1976),<sup>36</sup> with reported tortuosity ranging from 3 to 10 for  $H_2$ -air diffusion in sandstone. The limestone has a slightly lower tortuosity that ranges from 1 to 2. Because the limestone has a more homogenous pore size distribution with lesser variations in pore volume than the two sandstones (see Fig. 5), the elongation

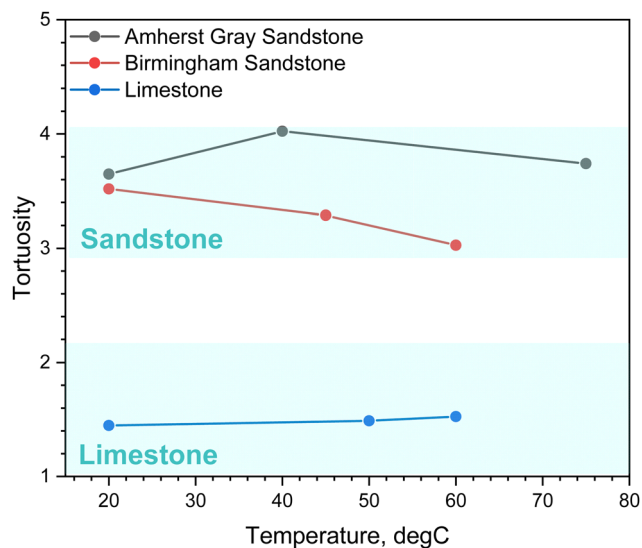


Fig. 6 Effect of temperature on tortuosity of  $H_2$  diffusion through reservoir rocks.

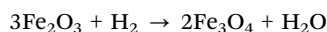
of the diffusive path caused by constrictions of changing pore size is less severe in limestone, resulting in a lower tortuosity. Limestone is generally fine-grained and tightly packed; thereby, its tortuosity value closely aligns with the literature-reported tortuosity value for a system consisting of closely packed uniform spheres ( $\sim 1.34$ ).<sup>61</sup>

The tortuosity exhibits slight temperature dependence for Indiana Limestone and Amherst Gray Sandstone, whereas Birmingham Sandstone appears to have less sinuous diffusive path at higher temperatures. Despite the remarkable similarity in the pore size distributions of the two sandstones, hydraulic tortuosity derived from diffusion measurements reacts differently in response to temperature changes. The two sandstones also share a similar mineral composition, mainly made up of quartz, which comprises over 85% of their mineralogy. Considering that the  $H_2$  diffusion coefficient in pure quartz is on the order of magnitude of  $10^{-6} \text{ cm}^2 \text{ s}^{-1}$ ,<sup>62</sup>  $H_2$  molecules should be primarily transported through available pore space, rather than within the solid grains. It is evident that there exist additional mechanisms that influence the flow of  $H_2$  through the porous network of Birmingham Sandstone. One limitation with MIP characterization technique is that it provides information on pore size distribution but fails to characterize pore connectivity and tortuosity. Even though the two sandstones have comparable pore size distribution, the actual arrangement of pores may differ. The current characterization technique therefore may not provide direct information into these properties.

Nevertheless, it is worth noting that a noticeable gas retention effect is observed in Birmingham Sandstone when examining the stabilized concentration of spiked and breakthrough chambers (see Fig. 2). Additionally, the extent of gas retention in Birmingham Sandstone diminishes as temperature increases, as evidenced by the reduced difference in concentration between the two chambers at high temperatures. It is highly likely that Birmingham Sandstone contains some dead-end pores, which



could contribute to the observed gas retention effect. As temperature increases, the intensified Brownian motion of  $H_2$  molecules is likely to release the gas trapped in these dead-end pores, explaining the reduction in gas retention. The decreased gas retention effect at higher temperatures results in an overall decrease in the distance that diffusing molecules need to travel, leading to the observed reduced tortuosity with temperature. Birmingham sandstone is also distinguishable from other Berea sandstones because it contains iron cement spots,<sup>63</sup> though not in quantities sufficient to be measured using our QXRD. Spots in Birmingham samples were visible as reddish pots, which were revealed using scanning electron microscopy (SEM) and energy dispersive X-ray spectroscopy (EDS) to be iron oxides, while iron present in Amherst samples was primarily pyrite and iron carbonates (Fig. 7). These iron oxide inclusions may interact differently with  $H_2$  gas. The dominant reaction between iron oxide and  $H_2$  is a redox reaction, as described by Yekta *et al.*, (2018):<sup>64</sup>



However, under geological conditions, pyrite reduction reactions with hydrogen have not been observed in previous study.<sup>65</sup> Consequently, Birmingham Sandstone has the potential to react with  $H_2$  at higher temperatures. Due to the consumption of  $H_2$ , a

higher concentration gradient occurs at elevated temperatures, which facilitates the diffusion process and leads to stronger temperature-dependent behavior. Thus, in our diffusion experiments, we observed that the  $H_2$  diffusion coefficient in Birmingham Sandstone is more sensitive to temperature changes compared to Amherst Sandstone (see Fig. 4).

In light of findings from this and previous work,<sup>36</sup> most reservoir rocks tend to have relatively low tortuosity (less than 5) in  $H_2$  diffusion. In practical scenarios where experimental diffusion measurements is unavailable, the effective diffusion coefficient,  $D$  for the reservoir rock can be estimated as the product of porosity and the self-diffusion coefficient, serving as a reasonable first-order approximation. Nevertheless, achieving an accurate estimation of  $D$  necessitates a comprehensive analysis of the underlying diffusion mechanisms involved. As noted earlier, molecular diffusion occurs by bulk diffusion and/or Knudsen diffusion. Bulk diffusion dominates in pores with radii greater than  $1\ \mu\text{m}$  and Knudsen diffusion dominates in pores with radii less than  $0.01\ \mu\text{m}$ . The pore size distributions measured for the two sandstones and one limestone studied in this work reveal that most pores possess radii exceeding  $0.01\ \mu\text{m}$ , with a noticeable portion of pores lying between  $0.01\ \mu\text{m}$  and  $1\ \mu\text{m}$ . Herein, Knudsen and bulk diffusion occur simultaneously for  $H_2$  diffusion in the two sandstones and one limestone studied.

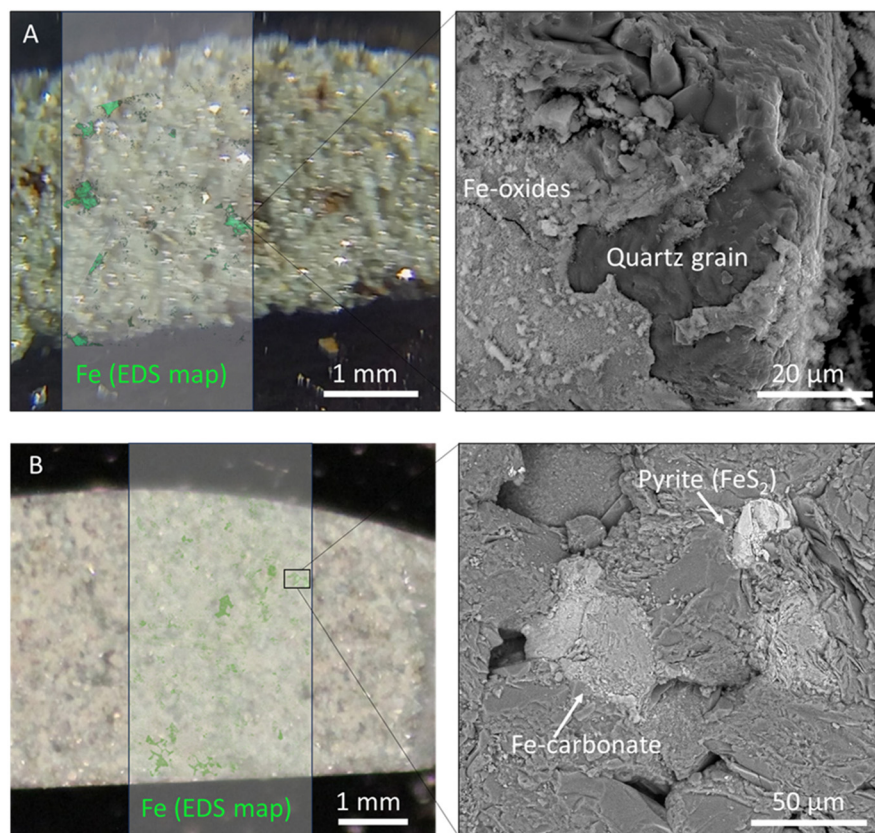
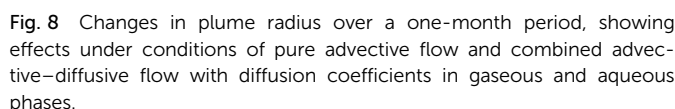


Fig. 7 Optical photography images, left, and SEM images, right, for (A) Birmingham Buff sandstone and (B) Amherst Grey sandstone. The rectangle is the selected analyzed area. Green is pixels with high proportion of Fe relative to all other identified elements. EDS spectra and SEM images were collected with a ThermoFisher Apreo 2 S SEM operating at 1.6 nA, 20 kV.









**Fig. 8** Changes in plume radius over a one-month period, showing effects under conditions of pure advective flow and combined advective-diffusive flow with diffusion coefficients in gaseous and aqueous phases.

In both sandstone and limestone reservoirs, plume sizes slightly expand when diffusion is considered; however, as water saturation increases, this expansion diminished as a result of reduced diffusive flow. If cushion gas is injected into the storage reservoir during the site initialization stage, the simulated plume size due to diffusion sees an increase of up to 7% at the lowest water saturation investigated; at higher water levels, the impact of  $H_2$  diffusion on plume size becomes negligible. In water-saturated reservoirs, the effective  $H_2$  diffusion coefficient is determined by the diffusion coefficients in both the aqueous and gaseous phases. Since the  $H_2$  diffusion rate in water is significantly lower than in gas, reservoirs with higher water saturations result in smaller plume areas. This smaller plume area aids in more efficient  $H_2$  recovery during withdrawal periods. Overall, with a gaseous  $H_2$  diffusion coefficient in reservoir rocks on the order of  $1 \times 10^{-6} \text{ m}^2 \text{ s}^{-1}$ , diffusion flow has a minimal impact on plume migration. Advection predominantly governs  $H_2$  transport in these layers.<sup>3,13</sup>

## 6. Conclusions

### 6.1 Measured H<sub>2</sub> diffusion coefficient in reservoir rocks

This work provides a benchmark dataset for temperature-dependent  $H_2$  diffusion in various reservoir rock types (including

As methane diffusion coefficients typically ranging from 0.002 to 0.01 cm<sup>2</sup> s<sup>-1</sup>, we surmise that concentration-driven diffusion will be more predominant during H<sub>2</sub> storage compared to methane. Therefore, hydrodynamic modeling of the subsurface flow of H<sub>2</sub> during UHS requires an accurate H<sub>2</sub> diffusion coefficient.

## 6.2 Temperature dependence of H<sub>2</sub> diffusion coefficient

For all reservoir rocks studied,  $H_2$  diffusion rate shows a noticeably increasing trend with increasing temperature, where the growth rate of  $H_2$  diffusion coefficient per 1 °C of temperature ranges between 0.5% and 1.4%. Furthermore, the measured  $H_2$  diffusion coefficients are found to be directly proportional to temperature in logarithmic scale, consistent with Chapman–Enskog gas kinetic theory. Consequently, the dominant diffusion regime for  $H_2$  diffusion in reservoir rocks is bulk diffusion due to intermolecular collisions. The presence of Knudsen diffusion, though it only contributes minorly to total mass flux, amplifies the temperature dependence of  $H_2$  diffusion.

### 6.3 Effect of H<sub>2</sub> diffusion on forming plume size

When storing  $H_2$  in depleted hydrocarbon reservoirs,  $H_2$  diffusion results in an up to 7% increase in  $H_2$  plume size for storage reservoirs, and this expansion effect diminishes as initial water saturation of the reservoir rocks increases. While this work only provides a preliminary analysis of the diffusion effect on geologic  $H_2$  storage, it is suggested that conventional reservoir simulation tools built upon pressure-driven advection (Darcy) flow can be improved for diffusion flow to predict the fate of injected  $H_2$ , which is the key to design and optimize field operations.

## Data availability

The data supporting this article have been included as part of the ESI† and the Results and discussion section of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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