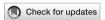
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Influencing factors of the carbon sequestration coefficient in saline aquifers based on multiphase flow displacement experiments

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Introduction: The volumetric method is the primary approach for calculating geological CO_2 storage potential, with its accuracy largely dependent on the pore volume of reservoir rocks and the effective storage coefficient. While the precision of reservoir rock pore volume can be enhanced through more sophisticated geological exploration techniques, the current selection of effective storage coefficients lacks a theoretical foundation. Thus, obtaining a more accurate effective storage coefficient is crucial for improving the evaluation precision of CO_2 geological storage potential.

Methods: To explore the factors influencing the effective carbon sequestration coefficient in saline aquifers and accurately assess their storage potential, nine sets of multiphase flow core displacement experiments were conducted using orthogonal design, with porosity, confining pressure, and pressure difference as variables.

Results: The results indicate that among these three factors, porosity has the most significant impact on maximum residual CO₂ saturation.

Discussion: Qualitative analysis of water migration in cores during displacement was performed using nuclear magnetic resonance (NMR) T_2 curves, revealing a close correlation between water movement and pore structure: water in mesopores and macropores is preferentially displaced, whereas water in nanopores and micropores is more resistant to displacement. Additionally, NMR was employed to analyze the maximum residual CO_2 saturation of artificial cores under different conditions, leading to the establishment of a multiple linear regression equation for maximum residual CO_2 saturation. By incorporating the volume coefficient derived from numerical simulations, the geological CO_2 storage coefficient for actual engineering sites can be estimated.

KEYWORDS

 ${\rm CO_2}$ geological storage, carbon sequestration coefficient, residual ${\rm CO_2}$ saturation, multiphase flow displacement, nuclear magnetic resonance



FIGURE 1Nuclear magnetic resonance analysis system for multiphase flow displacement.

1 Introduction

Geological CO₂ storage refers to the process of injecting CO₂—primarily captured from large industrial emission sources—into suitable geological formations (e.g., deep saline aquifers and depleted oil and gas reservoirs) through engineering techniques to achieve its long-term isolation from the atmosphere (Bert et al., 2005; Jing, 2021; Guo et al., 2014). It is estimated that by 2060, the contribution of CCUS technology to China's carbon neutrality goal could exceed 500 million tons (Ding, 2021; Xiaochun et al., 2006). Among all geological formations suitable for CO₂ storage, saline aquifers account for 95.6% of the total storage potential, far exceeding that of other formation types. Owing to their enormous potential and wide distribution, carbon sequestration in saline aquifers is regarded as a critical

supporting technology for achieving carbon peak and carbon neutrality targets.

Referring to the calculation method for geological CO_2 storage in saline aquifers proposed by the U.S. Department of Energy based on volumetric theory, the formula for estimating the CO_2 storage potential of saline aquifers is as follows (Goodman et al., 2011; USDOE, 2006; Song et al., 2015):

$$P = A \times h \times \phi \times \rho_{CO_2} \times E$$

where P (kg) is geological potential, A (m²) is reservoir area, h (m) is thickness, φ (%) is porosity, and ρ CO₂ (kg/m³) is CO₂ density. E, the carbon sequestration coefficient, is defined as the ratio of the volume of supercritical CO₂ to the total pore space of the formation. And it is a dimensionless geological coefficient.

As indicated by the formula, the accuracy of CO_2 geological storage potential calculations depends largely on the pore volume of reservoir rocks and the effective storage coefficient. While the precision of reservoir rock pore volume can be improved via advanced geological exploration techniques, the current selection of effective storage coefficients lacks a theoretical basis. Thus, obtaining a more accurate effective storage coefficient is crucial for enhancing the precision of CO_2 geological storage potential evaluations.

Most existing studies on effective storage coefficients for geological CO₂ sequestration have been conducted under idealized experimental conditions or via numerical models (Dai et al., 2022; Diao et al., 2023; Fu et al., 2022; Lei et al., 2022; Wei et al., 2013; 2021; 2022; Yang et al., 2019; Zhang et al., 2005; Zhang et al., 2019), failing to reflect the high complexity of real-world on-site scenarios. Currently, the mainstream storage coefficient used in calculating CO₂ storage potential in saline aquifers is 0.02, which involves significant uncertainties (Goodman et al., 2011; Song et al., 2015; Diao et al., 2017; Guo et al., 2014). In practice, saline aquifers exhibit substantial variations in geological characteristics across regions, such as porosity, permeability, formation pressure, temperature,

TABLE 1 Information of artificial cores.

Number	Specification	Photo
A-9	Porosity 5%, φ 25 × 50 mm	caA-9
B-4	Porosity 10%,φ25 × 50 mm	B-4
C-9	Porosity 15%,φ25 × 50 mm	C49

TABLE 2 Experimental Condition Design Table for Core Displacement with Different Porosities, confining pressures and Pressure Differences.

Serial number	Core	Porosity (φ/%)	Confining Pressure pressure difference (P/MPa) $(\Delta P/MPa)$		Temperature (°C)
1	A-9	5	8	0.9	33
2	A-9	5	9	0.3	33
3	A-9	5	10 0.6		33
4	B-4	10	8 0.3		33
5	B-4	10	9	0.6	33
6	B-4	10	10	0.9	33
7	C-9	15	8	0.6	33
8	C-9	15	9	0.9	33
9	C-9	15	10	0.3	33



FIGURE 2
Vacuum pressure saturation device.

and formation water chemistry. These factors are interrelated, and traditional experimental methods struggle to systematically analyze the role of each factor, making it difficult to determine a unified and accurate storage coefficient.

To investigate the carbon sequestration coefficient in saline aquifers, this study adopted an orthogonal experimental design, using artificial cores with significantly different porosities as experimental materials. A multiphase flow displacement NMR analysis system was employed to conduct nine sets of core multiphase flow displacement experiments via orthogonal design, with porosity, confining pressure, and pressure difference as variables (Dongjiang et al., 2021). The maximum residual $\rm CO_2$ saturation of artificial cores under different conditions was obtained, thereby accurately identifying the relative importance and interactions of each factor on the storage coefficient. This effectively addresses the limitations of existing studies in comprehensively analyzing multiple factors. A multiple linear regression equation for maximum residual $\rm CO_2$ saturation was established, which can provide a reference for determining carbon storage coefficients in saline aquifers in future research.

2 Methodology

2.1 Experimental apparatus

The MesoMR12-060H-I multiphase flow displacement NMR system (Figure 1) was used, featuring.

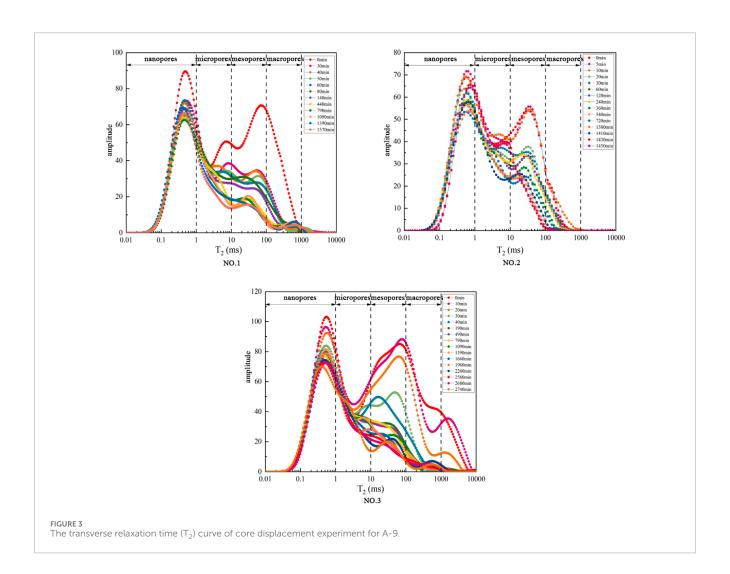
- Relaxation spectrum testing: Analyzes porosity, pore size distribution, saturation, and layered water content.
- High/low temperature and pressure displacement: Supports pressures of 0–15 MPa, temperatures of –30 °C–40 °C, and core lengths of 25–60 mm (1-inch diameter), enabling real-time monitoring of fluid distribution during displacement.

2.2 Principle

NMR measures nuclear magnetic resonance of magnetic nuclei in external magnetic fields, providing rapid, non-destructive analysis of porous media properties (e.g., porosity, permeability, water saturation). In ${\rm CO}_2$ sequestration research, it is used to track the migration of wetting-phase fluids and characterize reservoir properties.

TABLE 3 Nuclear magnetic resonance (NMR) sequence parameters table.

Nuclear magnetic resonance (NMR) sequence	Parameter setting	Acquisition time/min	
CPMG	Waiting time TW = 5000 ms Echo time TE = 0.2 ms Number of accumulations NS = 16	3	



2.3 Experimental materials

Artificial sandstone cores (supplied by Yilai Bo Technology Co., Ltd.) with porosities (5%, 10%, 15%) were used, and their porosity accuracy was pre-tested (Table 1).

2.4 Experimental design

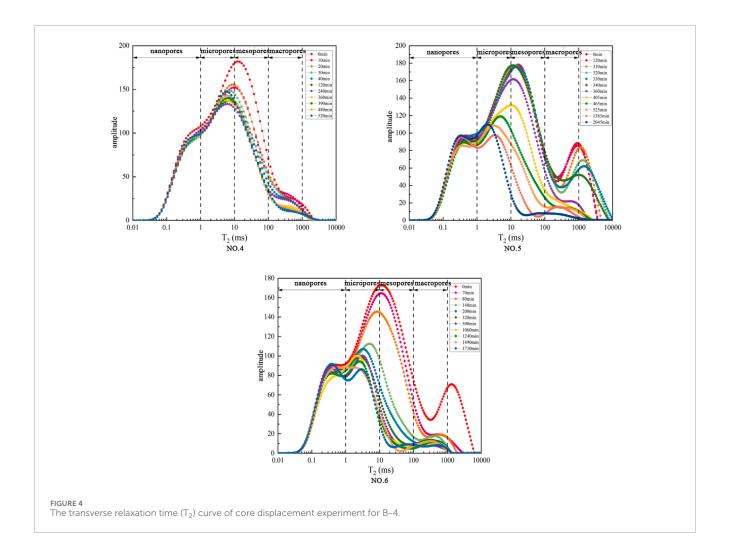
A three-factor, three-level orthogonal design (L9 (3^3)) was adopted, with variables including porosity (5%–15%), confining pressure (8–10 MPa), and pressure difference (0.3–0.9 MPa), while

temperature was maintained constant at 33 °C. Details of the nine experimental groups are provided in Table 2.

2.5 Procedures

2.5.1 Core saturation

Cores were fully saturated with water using a vacuum-pressure saturation device (Figure 2) to ensure uniform wetting. This device is primarily composed of a vacuum system, a liquid storage tank, a sample saturation chamber, a manual pump, a pressure gauge, and valves. When the liquid level in the storage tank ceases to drop, the valve is closed, and the hand pump is used to pressurize to 25 MPa,

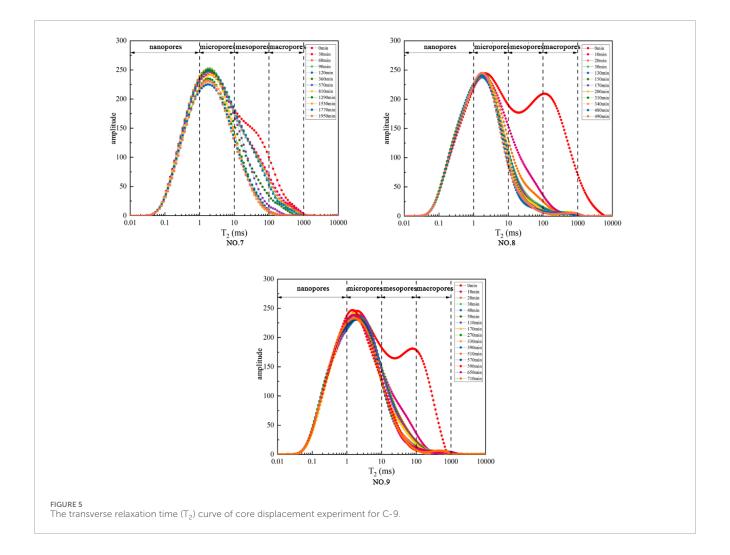


with this pressure maintained for 24 h. At this stage, the core is considered to be fully water-saturated.

2.5.2 Displacement experiments

The core sample to be tested, which was fully saturated with water, was prepared prior to conducting the NMR core displacement experiment, following these steps.

- System self-inspection: The NMR measurement system was powered on according to specifications, a calibration sample was placed into the core holder, and a system self-inspection was performed.
- Sample preparation and loading: The core sample to be tested was retrieved, a heat-shrinkable tube was applied, and a heat gun was used to shrink the tube tightly around the core. The prepared core sample was loaded into the core holder.
- 3. Pipeline connection: The core holder was positioned within the NMR coil, and pipelines for the confining pressure unit, gas injection unit, back pressure unit, and low-temperature circulation unit (which maintains the NMR system temperature at 25 °C to eliminate temperature-induced interference with NMR signals) were connected.
- 4. Pressurization: After purging the confining pressure fluid (fluorinated fluid FC770), the fluid was heated to the required experimental temperature using the heating unit. The industrial control computer was used to synchronously apply confining pressure and back pressure in the following sequence: 3-5-7-9-12 MPa for confining pressure and 1-3-5-7-8 MPa for back pressure. Meanwhile, CO₂ from the gas cylinder was injected into the intermediate container and pressurized to the required experimental pressure via a constant-rate, constant-pressure pump.
- 5. Initial signal acquisition: T_2 curves and images of the sample in its initial saturated state were collected using the Carr-Purcell-Meiboom-Gill (CPMG) sequences. Additionally, the industrial control computer was used to set the sampling sequence and time intervals for the dynamic displacement process.
- 6. Water injection: Since high temperatures from the heat gun during sample preparation might lead to partial water loss in the core, the core was re-saturated with water at the experimentally designed back pressure for 12 h prior to the displacement experiment. This ensured 100% water saturation in the core before gas injection.
- 7. $\rm CO_2$ injection: Once water saturation was complete, $\rm CO_2$ gas (pressurized to the required experimental level) was



injected into the core holder—with the confining pressure maintained 3–5 MPa higher than the inlet pressure throughout the experiment—to initiate the displacement process.

- 8. Data collection: Relevant NMR signals were acquired at predefined time points during displacement using the set sampling sequence, and the dynamic displacement process was monitored in real time.
- 9. Residual weight measurement: Throughout the displacement process, CPMG sequence tests were performed at regular intervals to obtain a series of core T2 curves, one-dimensional water distribution maps, and MRI images after varying displacement durations. As displacement progressed, reduced water content gradually weakened the hydrogen proton signal intensity. When the signal intensity approached the background level, MRI images were simultaneously observed: if fluid distribution boundaries became indistinct (i.e., imaging quality was impaired), it indicated that pore water had been displaced to irreducible saturation. Alternatively, the experiment could be terminated when changes in the core's T₂ curve were negligible and the curves nearly overlapped. The core sample was removed, and its residual weight was measured. All other experimental groups followed the same procedure.

2.5.3 NMR parameters

The CPMG sequence parameters were: waiting time (TW) = 5,000 ms, echo time (TE) = 0.2 ms, 16 accumulations, with an acquisition time of 3 min (Table 3).

3 Results and discussion

3.1 NMR T₂ spectra analysis

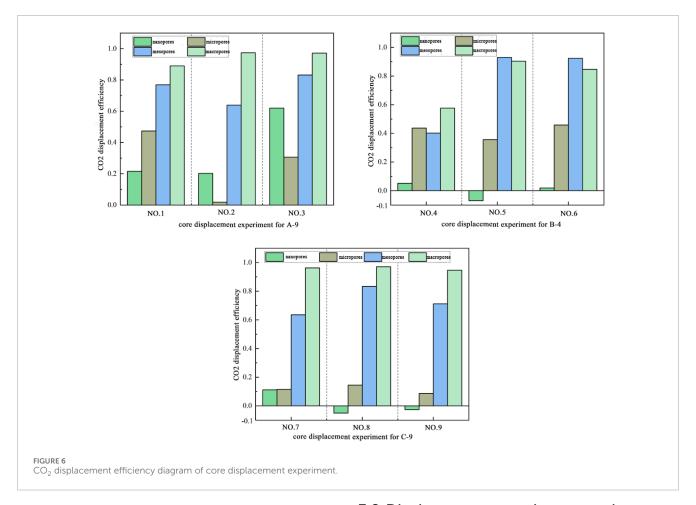
 $\rm T_2$ curves (Figures 3–5) show bimodal distributions for cores A-9 and B-4, with rapid water loss in macropores/mesopores and slow loss in micropores/nanopores. Core C-9 exhibits rapid initial displacement, with most water removed within 10 min.

Using the Laplace equation:

$$P_c = \frac{2\gamma \cdot COS\theta}{R}$$

(Where P_c is capillary entry pressure, γ is interfacial tension, θ is contact angle, and R is pore radius), smaller pores (nanopores/micropores) require higher P_c , explaining why residual water concentrates in these regions.

To conduct an in-depth analysis of displacement processes across different pore size groups, the final ${\rm CO_2}$ displacement



efficiency for each pore size group in each experimental set was calculated using the initial T_2 curve from the displacement experiment and the T_2 curve at the final state upon displacement completion. The formula for calculating ${\rm CO}_2$ displacement efficiency is as follows.

$$R = \frac{A_1 - A_2}{A_1}$$

Where: R is the CO_2 displacement efficiency; is the total nuclear magnetic signal within the range of nanopores, micropores, mesopores, or macropores in the initial T_2 curve; is the total nuclear magnetic signal within the range of nanopores, micropores, mesopores, or macropores in the T_2 curve at the final state after displacement.

Water can only migrate into nanopores if the pressure is sufficient to overcome the capillary entry pressure, which is typically high due to the small pore radii. CO_2 displacement efficiency (Figure 6) shows that mesopores and macropores have displacement efficiencies of >60% and >80%, respectively, while nanopores and micropores have <50%. Negative efficiency in some nanopore calculations (Groups 5, 8, 9) reflects water influx from larger pores. The distribution of residual CO_2 (Figure 7) shows that mesopores hold \geq 40% of residual CO_2 , due to their higher proportion in the pore structure and the limited space in nanopores/micropores occupied by residual water.

3.2 Displacement experiment results

Maximum residual CO_2 saturation (S_R) and residual water saturation (S_W) were calculated using:

$$S_R = \frac{m_1 - m_2}{m_1 - m_0} \times 100\%, S_W = 1 - S_R$$

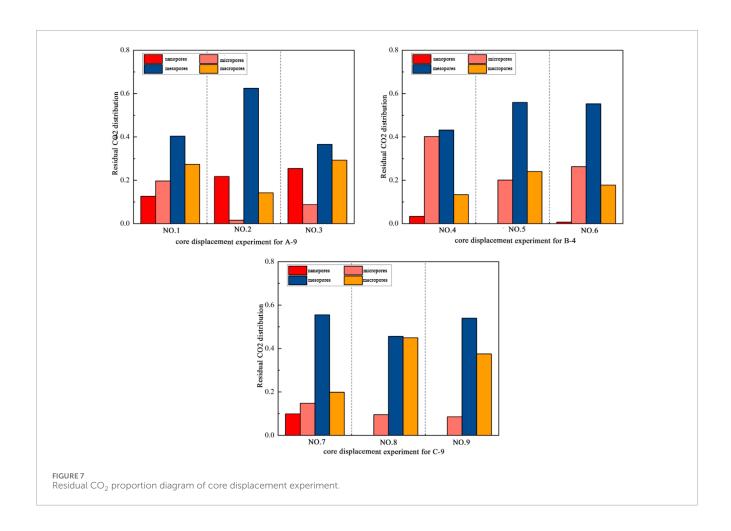
where m_0 , m_1 , m_2 are dry, saturated, and residual core weights, respectively. Results for nine groups are listed in Table 4.

3.3 Data analysis

Assuming no interference from other factors and the generalizability of laboratory results, SPSS was employed to construct a regression model. The model (Table 5) showed an $R^2 = 0.80$, indicating that 80% of the variation in S_R is explained by the three factors. The Durbin-Watson statistic (1.877) confirmed the independence of the data. ANOVA (Table 6) indicated a significant model fit (F = 6.674, p < 0.05).

Regression coefficients (Table 7) indicated.

• Porosity (x_1) had a negative impact on S_R $(\beta = -1.206, p < 0.05)$: a 10-percentage-point increase in porosity reduced S_R by 12.06%. This is consistent with the results of the NMR displacement experiments, primarily due to the influence of capillary forces.



 ${\sf TABLE\ 4\ Maximum\ residual\ CO_2\ saturation\ (S_R)\ and\ residual\ water\ saturation\ (S_W)\ in\ displacement\ experiments\ of\ artificial\ cores. }$

Serial number	Maximum residual CO ₂ saturation (S _R)/%	Residual water saturation (S _W)/%
1	67.01	32.99
2	53.35	46.65
3	66.49	33.51
4	47.83	52.17
5	66.52	33.48
6	69.13	30.87
7	47.59	52.41
8	52.19	47.81
9	50.88	49.12

TABLE 5 Durbin-watson test.

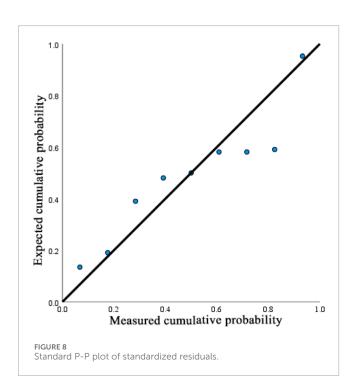
Model	R	R2	Adjusted R2	Standard error of estimate	Durbin-watson
1	0.895a	0.800	0.680	0.5165	1.877

TABLE 6 Analysis of variance (ANOVA) for the model.

Model		Sum of squares	Degree of freedom	Mean square	F	Significance
	Regression	0.053	3	0.018	6.674	0.034
1	Residual	0.013	5	0.003		
	Total	0.067	8			

TABLE 7 Coefficients of the multiple linear regression equation.

Model		Unstandardized coefficients		Standardized coefficients	t	Significance	Collinearity statistics	
		В	Standard error	Beta			Tolerance	VIF
	(Constant)	0.218	0.200		1.090	0.326		
1	Porosity	-1.206	0.422	-0.572	-2.861	0.035	1.000	1.000
	Confining pressure	0.040	0.021	0.380	1.903	0.115	1.000	1.000
	Pressure difference	0.202	0.070	0.573	2.867	0.035	1.000	1.000



- Pressure difference (x_3) had a positive impact on S_R (β = 0.202, p < 0.05): a 1 MPa increase raised S_R by 20.2%. This is mainly because the pressure difference facilitates the formation of "preferential pathways."
- Confining pressure (x_2) exhibited no significant effect (p = 0.115).

Porosity had the strongest influence (largest absolute coefficient). The regression equation is:

$$S_R = 0.218 - 1.206x_1 + 0.04x_2 + 0.202x_3$$

Residual analysis (Figure 8) confirmed a normal distribution, validating the model assumptions.

The maximum residual CO_2 saturation measured in the experiment corresponds to the displacement efficiency of the core under the relevant operating conditions. To derive the CO_2 storage coefficient for actual engineering sites, an additional key parameter—the volume coefficient—is required alongside the displacement coefficient. The CO_2 storage coefficient of saline aquifers can be calculated using the following formula:

$$E = E_V \times S_R$$

In this formula, E denotes the geological ${\rm CO_2}$ storage coefficient; ${\rm S_R}$ represents the maximum residual ${\rm CO_2}$ saturation; and ${\rm E_V}$ stands for the volume coefficient.

The volume coefficient is associated with the area and formation thickness, while the heterogeneity of the formation complicates its calculation. In practical engineering, the volume coefficient is typically derived via three-dimensional geological modeling and numerical simulation methods. By integrating the maximum residual $\rm CO_2$ saturation equation developed in this study, the geological $\rm CO_2$ storage coefficient for actual engineering sites can be calculated, thereby supporting the evaluation of $\rm CO_2$ geological storage potential in saline aquifers.

4 Conclusion

To investigate the factors influencing the effective storage coefficient for CO_2 sequestration in saline aquifers and accurately

assess their storage potential, nine sets of multiphase flow core displacement experiments based on orthogonal design were conducted, with porosity, confining pressure, and pressure difference as variables. The experimental materials used were artificial cores with significantly different porosities. The key findings are as follows.

- 1. 80% of the variation in "maximum residual CO_2 saturation" is influenced by "porosity," "confining pressure," and "pressure difference." Among these three factors, porosity has the most significant impact on maximum residual CO_2 saturation.
 - 2. Qualitative analysis of water migration in cores during displacement via NMR T_2 curves revealed that water migration during displacement is closely related to pore structure: water in mesopores and macropores is preferentially displaced, while water in nanopores and micropores is more resistant to displacement.
 - 3. NMR was used to analyze the maximum residual CO_2 saturation of artificial cores under different conditions, and a multiple linear regression equation relating maximum residual CO_2 saturation to "porosity," "confining pressure," and "pressure difference" was established. By incorporating the volume coefficient derived from numerical simulations, the geological CO_2 storage coefficient for actual engineering sites can be estimated. This serves as a reference for subsequent calculations of geological CO_2 storage potential in saline aquifers.

Data availability statement

The original contributions presented in the study are included in the article/supplementary material, further inquiries can be directed to the corresponding authors.

Author contributions

WZ: Writing – original draft. CZ: Writing – review and editing. TJ: Writing – original draft. JbZ: Writing – original draft. JaZ: Writing – original draft. MW: Writing – original draft. JnZ: Writing – original draft. LF: Writing – original draft.

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Conflict of interest

Author JbZ was employed by Shanxi Institute of Geological Survey CO., LTD.

Author MW was employed by National Energy Group Inner Mongolia Shanghai Temple Power Generation Co., Ltd.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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