The Pore Structure of Marine to Continental Transitional Shales in the Permian Shanxi Formation on the East Margin of the Ordos Basin, China

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The pore structure is an important factor in determining the storage capacity for shale gas development. Twelve samples were selected from the marine to continental transitional shales in the Shanxi Formations on the eastern margin of the Ordos Basin, and their nanoscale pore structure was investigated by a variety of low-temperature N2 adsorption (LT-N2GA) and low-pressure CO2 adsorption (LP-CO2GA) and high-pressure mercury intrusion (HPMI) experiments. The shale pores are complex and are mainly mesopores, which range mainly from 10 nm to 40 nm and greatly contribute to the pore volume (PV), whereas pores with a diameter less than 0.8 nm greatly contribute to the specific surface area (SSA) of the shale. The pore structure is affected by the TOC content, organic matter maturity, and illite/smectite (I/S) mixed layer content. The total PV increases with the increase in TOC content, organic matter maturity, and I/S mixed layer content. The effects of pores on the occurrence shale gas were determined by high-pressure methane adsorption experiments. The maximum adsorption amount of methane was positively correlated with the SSA of the micropores, indicating that the micropores have a large SSA, which controlled the adsorbed gas content of the shale. The mesopores provide the majority of the PV, which mainly corresponds to the volume of free gas in the shale, and the macropores are mainly micron-sized pores, which can form the main migration channels for shale gas.

1. Introduction

China’s shale gas exploration and production have made considerable strides in marine shales, notably in the southern Sichuan Basin, after the success of shale gas in North America [1–4]. In 2020, China reported a total production of 200 × 108 m3 of shale gas. Marine to continental transitional shales have attracted scant attention than marine shales, which is mostly found in delta facies where rapid sedimentary environment changes have caused the transitional shale to be thinner and less continuous, making it difficult for development [5, 6]. Nonetheless, the transitional shale gas resources are large, approximately 19.8 × 1012 m3 or 25% of China’s shale gas reserve [7–9]. Future research on the exploration and development of shale gas in China will focus on the transition shales [8]. Shale reservoirs are nano- to micrometer-scale complex and heterogeneous porous solid media having low porosity and poor permeability [10, 11]. Shale gas normally exists in three states: free, adsorbed, and dissolved. The adsorption state of shale gas occurs mostly on the surfaces of organic compounds and inorganic minerals, while the free-state state occurs predominantly in pores and microcracks and the dissolved state occurs predominantly in liquid hydrocarbons and water [12]. The IUPAC classification of pores is divided into three main categories: micropores (<2 nm), mesopores (2–50 nm), and macropores (>50 nm) [13]. The pores in shale are further categorized into three types based on their form.
and origin: interparticle (inter P), intraparticle (intra P), and organic material (OM) pores [14, 15]. The irregularity of the pore structure and the random range of the pore size distribution are the major factors controlling gas storage and flow capacity [16, 17]. Numerous factors affect the pore structure of marine shale; however, mineral content is especially important for the generation and maintenance of pore structure [18–20]. In addition, the content of OM is also an important factor affecting the development of shale pores, and the higher the content of OM is, the more favourable the development of micropores [21–24]. Due to the lack of study on marine to continental transitional shales, it is essential to appreciate the pore structure of these shales to assess shale reservoirs and understand how gas is stored and transported.

Table 1: Geochemical characteristics and mineralogical compositions of shale samples.

| Samples | Depth (m) | Strata       | TOC (%) | $R_o$ (%) | Qu | Pl | Si | Ca | Do | Py | T | C | I/S | K | Ch |
|---------|-----------|--------------|---------|-----------|-----|----|----|----|----|----|---|---|----|---|----|
| Sample 1| 2142.4    | Shanxi       | 5.78    | 2.54      | 60.99 | 0.00 | 0.00 | 0.00 | 2.96 | 2.67 | 33.38 | 17.97 | 9.38 | 4.91 | 1.12 |
| Sample 2| 2143.2    | Shanxi       | 6.43    | 2.33      | 59.74 | 0.00 | 0.00 | 0.00 | 3.83 | 2.79 | 33.63 | 15.34 | 11.07 | 5.50 | 1.72 |
| Sample 3| 2144.4    | Shanxi       | 11.1    | 2.37      | 48.79 | 0.00 | 0.00 | 1.68 | 4.90 | 3.88 | 40.75 | 21.73 | 8.32  | 9.25 | 1.44 |
| Sample 4| 2145.2    | Shanxi       | 8.66    | 2.17      | 37.87 | 0.00 | 0.00 | 11.01 | 2.55 | 4.26 | 44.31 | 19.17 | 9.81  | 14.43 | 0.90 |
| Sample 5| 2148.9    | Shanxi       | 7.83    | 2.38      | 59.68 | 0.00 | 0.00 | 0.00 | 1.32 | 0.00 | 39.00 | 13.03 | 6.64  | 17.36 | 1.98 |
| Sample 6| 2150.9    | Shanxi       | 1.82    | 2.08      | 31.12 | 2.48 | 4.71 | 0.00 | 0.00 | 0.34 | 61.35 | 13.03 | 14.98 | 30.18 | 3.16 |
| Sample 7| 2152.6    | Shanxi       | 1.08    | 2.16      | 37.74 | 0.00 | 5.73 | 0.00 | 0.00 | 3.46 | 53.06 | 13.03 | 6.64  | 10.79 | 2.44 |
| Sample 8| 2154.4    | Shanxi       | 1.68    | 2.23      | 45.12 | 2.67 | 1.50 | 0.00 | 0.00 | 1.13 | 49.57 | 14.29 | 20.08 | 13.49 | 1.72 |
| Sample 9| 2158.6    | Shanxi       | 1.84    | 2.19      | 36.03 | 1.71 | 2.56 | 0.00 | 1.27 | 0.63 | 57.80 | 14.54 | 19.99 | 19.48 | 3.79 |
| Sample 10| 2163.1   | Shanxi       | 0.84    | 2.13      | 34.27 | 1.58 | 1.69 | 0.00 | 0.00 | 0.53 | 61.93 | 12.53 | 29.98 | 15.88 | 3.53 |
| Sample 11| 2165.9   | Shanxi       | 2.37    | 2.36      | 38.00 | 3.99 | 1.12 | 0.00 | 0.00 | 1.11 | 55.78 | 13.03 | 20.37 | 20.06 | 2.32 |
| Sample 12| 2167.2   | Shanxi       | 9.85    | 2.64      | 28.27 | 0.00 | 2.70 | 0.00 | 0.00 | 3.28 | 65.75 | 20.03 | 15.05 | 28.32 | 2.35 |

$R_o$ is the vitrinite reflectance; Qu, Pl, Si, Ca, Do, Py, and TC are the quartz, plagioclase, siderite, calcite, dolomite, pyrite, and total clays, respectively; I, I/S, K, and Ch are the illite, illite/smectite, kaolinite, and chlorite, respectively.
Figure 2: Continued.
In addition to conventional reservoir pore research methods, theoretical methods of porous material characterization are also introduced to pore characteristics of shale, which can be roughly summarized as image analysis, fluid injection method, and nonfluid injection method [25–29]. Image analysis techniques such as scanning electron microscopy [25, 30], field emission scanning electron microscopy [31, 32], focused ion beam scanning electron microscopy [33], transmission electron microscopy, and atomic force microscopy can be used to intuitively observe and analyze the morphology and size of microscopic pores in shale [30–35]. Fluid injection techniques, such as high-pressure mercury injection (HPMI) [25], low-temperature N₂ adsorption (LT-N₂GA) [32, 33], low-pressure CO₂ adsorption (LP-CO₂GA) experiments [32, 33], and He-Hg pore analysis experiments, have been used [31–33, 36–38]. A series of nonfluid injection techniques, such as low-field nuclear magnetic resonance [39], small-angle/ultrasmall-angle neutron scattering [40, 41], and micro- and nano-CT scanning, have been used to quantitatively characterize and analyze shale nanopore structures [35, 38–43]. Low-pressure gas adsorption is the simplest and most common of these experimental methods. While LT-N₂GA and HPMI are useful for studying mesopores and macropores, LP-CO₂GA has been demonstrated to be a more effective method for determining the structure of micropores [25, 31–33, 36, 37].

Recent study indicating about 1.9 × 10¹² m³ of saturated gas resources in the Upper Carboniferous demonstrates the tremendous development prospects of shale gas in Carboniferous and Permian transition shale reservoirs on the eastern margin of the Ordos Basin [7–9]. In addition, Li et al. found that the coal-bearing strata on the eastern margin of the Ordos Basin have a significant potential for the coproduction of tight sand gas, shale gas, and coalbed methane [12]. This analysis focuses primarily on twelve shale samples from the Permian Shanxi Formation on the eastern margin of the Ordos Basin. Considering the differences in determination effectiveness of different test methods for shale porosity, we obtained the pore size distribution (PSD), from nanopores to micropores of the shale in the study area through LP-CO₂GA, LT-N₂GA, and HPMI. We investigated the influence of the abundance, organic matter maturity, and mineral composition on shale pore development through TOC and whole-rock mineral X-ray diffraction analysis. These results, combined with high-pressure methane adsorption (HPMA) experiment results, were then used to discuss the influence of the complete pore structure on the adsorption capacity of shale. Consequently, our research serves as a guide and point of reference for the finding and exploitation of shale gas in the Shanxi Formation along the eastern margin of the Ordos Basin and adjoining basins.

2. Geological Setting

The Ordos Basin is around 37 × 10⁴ km² in size and almost rectangular in form. Based on the existing tectonic features [12, 44], according to the structural morphology, the basin can be subdivided into six subtectonic units (Figure 1(a)). The study area is located on the eastern margin of the Ordos Basin, the structure is relatively stable, the stratigraphy is monoclinal to the west, and the outcropping layers are young from east to west [8, 9, 12]. After the Early Ordovician, during the influence of the Caledonian movement, the sedimentary thickness of the inner layer of the basin decreased, and then, the basin was fully uplifted and denuded. Carboniferous and Permian transitional strata were extensively deposited, including the upper Carboniferous Benxi formations, and the lower Permian Taiyuan Formation and Shanxi Formation were deposited (Figure 1(b)) [9]. The sedimentary tectonic evolution of the Ordos Basin and the stratigraphic characteristics of the studied region are intricately interwoven. The basin, as well as the whole
Figure 3: Continued.
Great North China Block, experienced a facies sequence evolution history from a marine environment with sea-land interaction to an inland lake basin in the period from the Benxi Period of the late Palaeozoic to the Shiqianfeng Period of the late Permian [8]. Due to regional tectonic activity, the water body changed periodically, forming multiple stage sedimentary phases of littoral shallow sea-delta front-littoral shallow lake facies and depositing many sets of marine to continental transition organic-rich shales [8, 9].

3. Samples and Methods

3.1. Samples. The experimental samples used in this study were all extracted from shale gas well cores and came from the Shanxi Formation on the eastern margin of the Ordos Basin. Twelve shale samples with different TOC contents were selected to conduct experiments on LP-CO$_2$GA, LT-N$_2$GA, HPMI, and HPMA.

3.2. TOC, Thermal Maturity, and Mineral Component. Shale samples were first treated with HCl to remove carbonate, then rinsed with deionized water to remove any leftover HCl, and then dried at 110°C for 12 hours. The TOC content was measured by a LECO CS230 carbon/sulfur analyser in line with national standard GB/T19145-2003. Cutting samples were made into pellets and then ground and polished. Before measuring the samples, the MPV-II microphotometer was calibrated via the twofold calibration technique in order to measure the vitrinite reflectance. Two reference samples were selected so that the gadget could be repeatedly calibrated until it reached its optimal state [45, 46]. In accordance with the Chinese Oil and Gas Industry Standard SY/T 5124-2012, measuring points were evenly distributed across sample surfaces to ensure measurement accuracy. The mineral compositions of the shale samples were determined by using Bruker D8 Advance X-ray diffractometers [46]. The minerals and the relative mineral percentages for each sample were estimated following the Chinese Oil and Gas Industry Standard SY/T 5163-2018.

3.3. High-Pressure Mercury Intrusion (HPMI) Porosimetry. The HPMI test was carried out by the Autopore-9505 mercury intrusion tester following national standards GB/T 21650.1-2008, which measured apertures between 3.6 nm and 960 μm, with a maximum pressure of 60,000 psi (413 MPa) and a volumetric accuracy of less than 0.01 cm$^3$. Samples were heated in a vacuum oven at 3 Pa and 105°C for 16 h to remove moisture, and then, the samples were evacuated before mercury injection. The pore diameter distribution was obtained according to the Washburn formula [47].

3.4. Low-Pressure Gas Adsorption Experiments. Using the gas adsorption approach, the nanopore distribution in shale may be successfully determined. Micropores are often

| Loop type | Sample                                              | Pore type            |
|-----------|-----------------------------------------------------|----------------------|
| H3        | Sample 1, sample 6, sample 7, sample 8             | Plate-shaped pore    |
|           | Sample 9, sample 10, sample 11, sample 12          | Wedge-shaped tubular pore |
| H4        | Sample 2, sample 3, sample 4, sample 5             | Parallel slit-shaped pore |
|           |                                                     | Plate-shaped pore     |
|           |                                                     | Ink bottle-shaped pore |

**Figure 3:** Typical sample N$_2$ adsorption-desorption isotherms and pore size distribution of samples.

**Table 2:** Pore types obtained from the adsorption isotherms using N$_2$ adsorption-desorption isotherms.
Figure 4: Continued.
examined by LP-CO$_2$GA, while mesopores may be examined by LT-N$_2$GA.

LT-N$_2$GA and LP-CO$_2$GA analysis was carried out using a Quantachrome Autosorb-IQ specific surface area and micropore analyser, in accordance with the national standards GB/T 21650.2-2008 and GB/T 21650.3-2011. Its limit of SSA is 0.0005 m$^2$/g, its range of pore diameter is from 0.35 nm to 500 nm, and its PV limit is 0.0001 cm$^3$. All the samples were subjected to vacuum at 105°C for 12 h to remove the water and volatile substances in the samples. The LT-N$_2$GA experiment was performed at 77 K with a nitrogen purity greater than 99.999%. The SSA was obtained by using the Brunauer-Emmett-Teller (BET) methods, and the PV and PSD were calculated by the density functional theory (DFT) model [48]. The LP-CO$_2$GA experiment was performed at different partial pressures at 273 K. The PV, SSA, and PSD of the micropores (<2 nm) are calculated by a more suitable DFT model.

3.5. High-Pressure Methane Adsorption (HMPA). The HMPA experiments were performed by a Gravimetric Isotherm Rig 3 instrument with a temperature accuracy of 0.5°C and a high-precision accuracy of 0.1%. This equipment was from the CNPC Key Laboratory of Unconventional Oil & Gas. The methane adsorption experiment was conducted under a fixed temperature of 70°C and a pressure ranging from 0 MPa to 25 MPa. Following the shale samples’ crushing to a mesh size of 60-80, around 100 g of each sample was put into the sample cell and put through a seal test and then underwent a methane adsorption test in accordance with the Chinese standard GB/T 19560-2008.

4. Results

4.1. Organic Geochemistry and Petrology. The TOC contents range from 0.84% to 11.1% (mean 4.94%) (Table 1). The shales are generally the stages of late high mature to early overmaturity, and the vitrinite reflectance ($R_o$) ranged from 2.08% to 2.64% (mean 2.30%). The mineral contents of the shale samples varied (Table 1). The clay mineral content is the highest, ranging from 33.38% to 65.75% (mean 49.69%); the quartz content is the second-highest, with an average of 43.14%, and the shales also contain feldspar, calcite, dolomite, pyrite, and other minerals [41]. The clay minerals are mainly illite and I/S mixed layer with a minor amount of kaolinite and chlorite.

4.2. Pore Structure Determined by Mercury Porosimetry (HPMI). HPMI curves can reflect the development and connectivity of pores in different pore-throat segments [37, 49, 50]. Based on the mercury intrusion and extrusion curves, Figure 2 depicts two unique types of pore architecture (A and B). Type A, represented by sample 4, represents fully open pores with good connectivity and can be described as cylinder-like pores. The pore morphology of type B, represented by sample 1, is mainly that of half-open pores with poor connectivity, similar to that of slit- and plate-like pores.

The mercury intrusion and extrusion curves of some samples are shown in Figures 2(a) and 2(c), showing the typical two-stage pattern represented by sample 4. Such curves have high mercury extrusion efficiency, which indicates that there are more open pores and that the pore connectivity is good [49, 50]. The mercury intrusion and extrusion curves of the other shale samples are shown in Figures 2(b) and 2(d), showing a typical three-section pattern, represented by sample 1. However, such curves have a low mercury extrusion efficiency, indicating that the open pores in the mercury injection test aperture range are smaller and the pore connectivity is worse [50].

4.3. Pore Structures by Low-Temperature N$_2$ Adsorption (LT-N$_2$GA). HPMI cannot accurately characterize the mesopores
in shale samples [51]; therefore, a LT-N$_2$GA experiment is used to characterize the mesopores. The nitrogen gas adsorption-desorption isotherms are as follows (Figure 3). According to the IUPAC, four types of hysteresis loops are classified [13]. The nitrogen gas adsorption-desorption isotherms show two types of pore structures, H3 and H4 (Table 2).

From the hysteresis loops formed (Figure 3(a)), represented by sample 4, the corresponding nitrogen gas adsorption-desorption isotherm rises slowly in the medium-pressure section and rapidly in the high-pressure section. There is no obvious saturated adsorption platform in the adsorption branch. The desorption branch decreases slowly at the beginning, decreases rapidly in the medium-pressure section, and finally decreases slowly again. This pattern is generally consistent with the characteristics of H4-type hysteresis loops and indicates that the pore morphology is generally characterized by parallel slit-, plate-, and ink bottle-shaped pores. According to the massive hysteretic loops of these pores (Figure 3(c)), the shale pores consist of a large amount of ink bottle-shaped structures and parallel slit-shaped structures.

![Figure 5](image.png)

**Figure 5:** (a) Isothermal adsorption curves of shale samples and (b) the interrelationship between TOC content and maximum adsorption capacities.
number of micropores and a small number of mesopores. In terms of the hysteretic loops formed by other shale samples (Figure 3(b)), represented by sample 10, the corresponding adsorption isotherm increases slowly in the middle- and low-pressure sections and rapidly in the high-pressure section, and the isotherm is nearly vertical. The desorption branch and the adsorption branch are nearly parallel in the high-pressure section and exhibit a steep near-vertical pattern. The middle- and low-pressure sections have a gentle trend, and the adsorption branch overlaps the desorption
branch. This pattern is similar to that of the H3-type hysteresis loop and indicates that the pore morphology is dominated by plate-shaped pores and wedge-shaped tubular pores. In addition, this pattern demonstrates that the shale pores are well-connected and sizable (Figure 3(d)). The shale of the Shanxi Formation in the study area has a very complex pore structure that is often irregular and consists of plate-shaped pores, ink bottle-shaped pores, and wedge-shaped tubular pores [51].

4.4. Pore Structures by Low-Pressure CO₂ Adsorption (LP-CO₂GA). CO₂ molecules may measure pore widths in the
Figure 8: Continued.
range of 0.3 to 2.0 nm at high temperatures (273 K) owing to their advantageous properties, such as high energy and rapid balancing [50, 51]. Figures 4(a) and 4(b) show that the LP-CO₂GA isotherms are type I at low temperatures. In the low-pressure region \((p/p_0 < 0.01)\), the adsorption quantity is small but the rate is high, and as the relative pressure is increased, the adsorption quantity continuously increases while the rate progressively drops. Overall, the adsorption capacity increased significantly with increasing TOC content in the 12 samples. The adsorption capacity of sample 3 is the largest, greater than 3.5 cm³/g, indicating that it contains many micropores. Sample 10 has the lowest adsorption capacity, less than 1.0 cm³/g, indicating that the sample contains a few micropores. According to the DFT model, the PSD curve of sample 3 comprises three peaks, the bulk of which are scattered between 0.3 and 0.8 nm (Figure 4(c)). The PSD curve for sample 10 displays six peaks (Figure 4(d)), the bulk of which are dispersed between 0.35 and 0.7 nm and 0.75 and 0.9 nm, indicating that the majority of the micropores are distributed between these pore size ranges.

4.5. Adsorption Capacity from High-Pressure Methane Adsorption (HPMA). HPMA experiments can test the maximum CH₄ adsorption capacity of shale under fixed temperature and pressure conditions [52]. As Figure 5(a) shows, under the conditions of 25 MPa and 70°C, the maximum CH₄ adsorption capacity of the Shanxi Formation shale after balanced water treatment is significantly different. The maximum adsorption capacities of shale samples are between 0.62 cm³/g and 5.40 cm³/g (mean of 2.27 cm³/g). The maximum CH₄ adsorption capacities of sample 3 and sample 10 are 5.40 cm³/g and 0.64 cm³/g, and the corresponding TOC contents are 11.1% and 0.84%, respectively. The maximum CH₄ adsorption capacity of shale tends to increase with the increase in total organic carbon content (Figure 5(b)), and the correlation coefficient is as high as 0.81, indicating that organic carbon content is the main factor affecting the development of nanoscale pores in shale. With the increase in TOC content, CH₄ adsorption capacity of shale also increases, which is consistent with previous research results [52–55]. The reason may be that the increase in shale thermal evolution makes kerogen generate hydrocarbon and form more organic pores. In addition, OM pores have a greater internal pore SSA than other types of pores. As a consequence, the increase in TOC content provides a large amount of storage space for shale gas, thus increasing the overall gas content.

5. Discussion

5.1. Complete Pore Structure Characteristics. Due to the complex and variable pore structure of shale and the vast range of pore diameters (from nanometer to micron development to varying degrees), it would be impossible for a single experimental approach to define the pores at all scales entirely and precisely. Based on the experimental results of HPMI, LT-N₂GA, and LP-CO₂GA [50, 51, 54, 56], the apertures of all pores in the Shanxi Formation shale on the eastern margin of the Ordos Basin are investigated. In this study, the best aperture measurement segments of these methods are selected for superposition analysis; among them, the micropores are characterized by LP-CO₂GA experimental data, the mesopores are characterized by LT-N₂GA experimental data, and the macropores are characterized by HPMI experimental data.

Using the aforementioned three procedures, the entire PSD properties of the Shanxi Formation shale samples in the study area were obtained. In terms of the PV distribution characteristics of the full pore size range, the PV distribution characteristics of different shale samples vary greatly, but in general, mesopores provide the majority of the PV, followed by micropores, and macropores provide a small amount of PV (Figure 6). In terms of specific data (Table 3), the total
PV of shale range from 23.23 to 46.54 × 10⁻³ cm³/g (mean 30.22 × 10⁻³ cm³/g). The majority of micropores are formed between 0.3 and 0.8 nm, and the PV ranges from 2.25 to 12.96 × 10⁻³ cm³/g (mean 7.16 × 10⁻³ cm³/g), accounting for 9.69-39.37% of the total PV, with an average of 23.24%. The majority of mesopores are found in the pore size ranges of 3-6 nm and 10-40 nm, and the PV ranges from 12.68 to 27.81 × 10⁻³ cm³/g (mean 18.90 × 10⁻³ cm³/g), accounting for 49.26-71.71% of the total PV, with an average of 62.93%. The macropore volume is mainly contributed by pores above 50-100 nm, whose value ranges from 1.19 to 7.07 × 10⁻³ cm³/g (mean 4.16 × 10⁻³ cm³/g), accounting for 4.75-23.16% of the total PV, with an average of 13.83%.

The analyses above have found that the pore types and structures of marine to continental transitional shale facies in the Shanxi Formation are quite different from those of marine shales in the Longmaxi formation. Jiang et al. and Shi et al. found that the PV of shale pores in Longmaxi Formation, Sichuan Province, is relatively developed in micropores, mesopores, and macropores [51, 57]. Zhu et al. found that micropores, mesopores, and macropores all contribute to the PV of the Longmaxi formation shale in the Upper Yangtze region, and the contributions of micropores and macropores are more prominent [56]. In this paper, the pore volume of the marine to continental transitional shale in the Shanxi Formation is mainly provided by mesopores, while the contributions of micropores and macropores are lower. This difference is caused by the difference in organic matter types between marine shale and marine to continental transitional shale.

According to the distribution characteristics of the Shanxi Formation shale, the SSA area distribution of shale samples also has strong heterogeneity, and great differences exist among different samples. However, on the whole, micropores and mesopores provide most of the SSA, and the contribution of macropores to the SSA is minimal or even negligible (Figure 7). In terms of specific data (Table 4), the total SSA of the shale samples ranges from
11.88 to 55.47 m$^2$/g (mean 31.64 m$^2$/g). The bulk of the surface area specific surface area (SSA) of micropores is given by pores between 0.3 and 0.8 nm with values ranging from 7.05 to 41.97 m$^2$/g (mean 23.58 m$^2$/g), accounting for 59.35-84.86% of the total SSA, with an average of 71.78%. The SSA of mesopores is mainly provided by the mesopores in the range of 3-6 nm. The SSA of mesopores in the range of 10-40 nm in some samples ranges from 4.50 to 13.12 m$^2$/g (mean 7.86 m$^2$/g), accounting for 14.80-37.90% of the total SSA, with an average of 27.45%. The SSA ranges from 0.06 to 0.38 m$^2$/g (mean 0.20 m$^2$/g), accounting for 0.16-2.75% of the total SSA, with an average of 0.77%. This value range is consistent with that of the marine Longmaxi Formation shale [50, 51, 56, 57]. Because micropores consist of organic pores and clay pores, micropores offer a large adsorption space for shale gas.

5.2. Influencing Factors of Pore Structure. According to a number of researches [21–29, 51], the TOC content, the type of organic material, and the mineral content of the clay have significant roles in controlling the PV and PSD of shale. Although different test methods and samples have been used, all the relevant studies show that the TOC content is the main controlling factor of micropores [23, 50–54, 58, 59].

5.2.1. Effect of TOC Content on Pore Structure. Organic material pores were formed predominantly from the heat decomposition of organic matter, which produced hydrocarbons [22, 29]. Numerous studies have shown that a high TOC content is advantageous for pore formation, the higher adsorption capacity, and the production of shale gas [25, 28, 51, 53, 58].

Figure 8(a) illustrates a weak linear relationship between the total PV and TOC content in the examined shale, indicating that the TOC content provides only part of the PV and that organic pores are not dominant. Figure 8(b) indicates that the SSA of shale does indeed have a significant positive correlation with TOC content, mainly because the
SSA of shale is mainly provided by micropores, and TOC content mainly controls the micropores. The TOC content has obvious positive correlations with the PV and SSA of micropores, with correlation values of 0.93 and 0.95, respectively (Figures 8(c) and 8(d)). However, the TOC content has no apparent relationships with the PV and SSA of mesopores and macropores. These results were consistent with those of most scholars [22, 25, 28, 29].

5.2.2. Effect of Thermal Maturity on Pore Structure. Some researchers have proposed that organic matter pores increase with organic matter maturation and that when the organic maturity of shale exceeds some threshold, the porosity of shale begins to decrease [28, 46, 60, 61].

The vitrinite reflectance ($R_o$) of organic matter in the Shanxi Formation in the study area ranges from 2.08% to 2.64% (mean 2.30%). The total PV or the volume of micropores, mesopores, and macropores in the Shanxi Formation shale of the study area show an increasing trend with increasing $R_o$ (Figures 9(a) and 9(b)). With the increase in $R_o$, the increase in the shale pore volume is not caused by the simultaneous increase in the pore volume of all the micropores, mesopores, and macropores. Therefore, the maturity of $R_o$ is an important factor affecting the development of shale pores in the Shanxi Formation.

5.2.3. Influence of Brittle Mineral Content on Pore Structure. In the Shanxi Formation of the research region, quartz predominates the list of brittle shale minerals. The association between the volume of micropores, mesopores, and macropores or the total PV and the presence of brittle minerals is not visible, as shown in Figures 10(a) and 10(b). Brittle minerals host few micropores, and brittle mineral grains, such as quartz and calcite, have a low SSA [21, 45, 51]. Relative to the micropore SSA of the shale matrix, brittle minerals provide a negligible SSA. For mesopores and macropores, although the brittle mineral occurrence in mineral particles of mesopores and macropores has a certain protective effect, mesopores and macropores and a high content of shale matrix brittle minerals are not widely developed [51, 53]. Consequently, the relationships between the brittle mineral content and the total PV and mesoporous and macropore volume are not obvious.
Figure 12: Cross-plots of maximum adsorption capacity with (a) total PV, (b) total SSA, (c) PV, and (d) SSA.
5.3. PV and SSA on Methane Adsorption Ability. In shale samples, the majority of free gas is contained in the pores or fractures of shale, PV and gas saturation essentially control the quantity of free gas. The majority of adsorption gas is adsorbed on the surface of organic matter (OM) and clay minerals, and the quantity of this gas is directly proportional to the SSA of shale [49, 51, 54]. As demonstrated in Figures 12(a) and 12(b), the maximum adsorption capacity of shale is positively correlated with the PV and SSA, showing that the quantity of methane adsorption in shale samples steadily rises with increasing SSA. As demonstrated in Figures 12(c) and 12(d), the maximum adsorption capacity of shale samples is positively correlated with micropores, but the correlation of mesopores is poor, indicating that micropores SSA affected the amount of methane adsorption. The SSA provided by mesopores is limited and cannot play a decisive role in the amount of methane adsorption. The results show that micropores have a large SSA and control the adsorbed gas content of shale. The free gas volume is mainly determined by the PV of shale, the mesopores of shale pores in the Shanxi Formation play a dominant role in the total PV, and more free gas can be accommodated. Since macropores are mostly micron-scale pores, they may serve as the principal channel for shale gas migration, which is consistent with academic research done in China and worldwide on the adsorption capability of shale from various places [51, 54]. Chalmers et al. found that CH₄ adsorption in shales increases with the increase in micropore volume [16]. Zhong et al. found that the micropores of continental shale in Yanchang Formation had the most influence on CH₄ adsorption capacity, and the correlation coefficient between micropores and adsorption capacity was significantly greater than other factors [61]. Lin et al. conducted a large number of studies on marine shale of Longmaxi formation in Sichuan Basin and discovered that micropores are primarily responsible for shale’s adsorption capacity [62].

6. Conclusions

(1) Although shale contains micropores and macropores, mesopores predominate in the marine to continental transition Shanxi shale on the eastern edge of the Ordos Basin. The total SSA is 11.88–55.47 m²/g (mean 31.64 m²/g), and the total PV is 23.23–46.54 × 10⁻³ cm³/g (mean 30.22 × 10⁻³ cm³/g). Pores from 10 nm to 40 nm contribute most of the PV, and the SSA is mainly contributed by micropores < 0.8 nm

(2) The maximum adsorption capacity in the marine to continental transitional Shanxi shale on the eastern margin of the Ordos Basin ranges from 0.62 cm³/g to 5.40 cm³/g (mean 2.72 cm³/g). With the increase in TOC content, the methane adsorption capacity of shale increases correspondingly, indicating that TOC has a strong adsorption capacity because organic pores can provide sufficient sites for gas adsorption

(3) The TOC content, organic matter maturity, and I/S mixed layer contents are the main factors controlling the pore structure of transition shales in the Permian Shanxi Formation on the east margin of the Ordos Basin

(4) The micropores of the Shanxi Formation shale have a large SSA and control the gas adsorption, the mesopores provide the main PV and the space for free gas, and the macropores form the main migration channels for gas

Data Availability

The data are available upon request.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

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