Variation in Pore Structure and Associated Fractal Dimensions of Barakar and Barren Measures Carbon-Rich Gas Shales of Jharia Basin, India

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ABSTRACT: The carbon-rich Barakar and Barren Measures shale beds of the Jharia basin were evaluated for variation in pore size, pore structure, and fractal dimensions. The shale core samples were obtained from exploratory boreholes drilled at the Jharia basin. The shale samples were analyzed for organo-inorganic composition by FTIR, pore size, and pore structure using BET low-pressure N₂ adsorption and pore geometry through FE-SEM photographs. The shale samples have significant carbon-rich content and are intercalated-banded in nature. The pore structures were evaluated through N₂ isotherms and validated by SEM images, revealing the mixed contribution of organo-inorganic matter in pore formations controlled by geochemical alteration, diagenesis, and mineral interaction. The rough internal surfaces of the pore were evaluated by categorizing them into fractals D₁, D₂, and D₃. It is observed that the D₂ type of fractal is in abundance associated with mesopores. The positive trend of fractals with pore size, pore structure, depth, fixed carbon, and TOC suggests the influence of different parameters on the formation of pore internal rugged surfaces in shale beds. The FE-SEM images indicate shallow to deep pores with different pore structures with fair to good pore connectivity. In summary, the shale beds of Jharia have heterogeneous complex pore structures, a rough surface, and sorption mechanisms controlled by weathering/alteration, depositional conditions, and organo-inorganic content. In shale beds, gas storage and transport phenomena are directly related to pore size distribution, pore structure, and associated fractal dimensions. The calculated values using the proposed empirical models for porosity (EPOf) and permeability (EPEf) showed excellent linear correlation with the measured porosity (MPOc, R² = 0.8577) and permeability (MPEc, R² = 0.8577), which are close to measured values. The curve matching of EPOf with MPOc and EPEf with MPEc follows a similar path, validating the results and suitability of the models. Hence, the proposed models may be considered to estimate the porosity and permeability of shale and coal beds.

1. INTRODUCTION
Shale gas is a natural gas, which dominantly contains methane derived from shale deposits.¹⁻⁵ The world energy scenario has rapidly changed in the past few decades due to the shale gas revolution. New technologies of horizontal drilling and hydraulic fracturing have paved the way for the United States to become an oil and gas exporter.⁶⁻¹⁰ It has also changed the energy and fossil-fuel-governing geopolitics of the world.¹⁻¹⁰ The shale matrix has complex systems of pores of different sizes from micro-, meso-, and macropores attributed to a heterogeneous mix of organo-inorganic content.⁴⁻¹⁶,¹⁴⁻¹⁹,⁹⁰ The gas storage and release mechanism in shale depends on pore type, pore structure, pore size, and pore fractals.²⁻¹⁹,¹⁸,⁸⁸ The shale matrix pore types and pore structures are mainly influenced by depositional conditions, clay type, moisture, and volatile matter content.⁵,⁸⁸ Likewise, pore surface formation can be contingent on the cracking of organic compounds, pore evolution, stages of
Fourier transform infrared spectroscopy (FTIR) provides crucial information about the functional groups present in shale. The absorption peaks also show the clay and mineral matters\textsuperscript{24−31} and the kerogen transformation level.\textsuperscript{32−37} It also provides information on the transformation of the aliphatic to aromatic components due to the organic content and thermal maturation.\textsuperscript{2,22,26,30}

In India, shale gas exploration and its commercial recovery are emerging as a clean energy resource development. The government is looking for improvements in shale gas development to reduce its petroleum import and dependence and fulfill the growing energy needs of 1.25 billion people.\textsuperscript{38,39} The Proterozoic, Gondwana, and Tertiary sedimentary basins in India have been assessed for shale gas potential and estimated 63 tcf of

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**Figure 1.** Geological map of the Jharia coalfield showing the study area and shale core sampling borehole locations\textsuperscript{53} (reprinted in part with permission from Elsevier-Publisher; Verma et al., 1979). [Verma, R. K.; Bhuin, N. C.; Mukhopadhyay, M. Geology, Structure and tectonics of Jharia CoalField, India-A 3-D Model. Geoexploration 1979, 17, 305−324].

**Figure 2.** Profile and cross sections based on gravity and borehole data along the section lines in the geological map of the Jharia basin\textsuperscript{63} (reprinted in part with permission from Elsevier-Publisher; Verma et al., 1979). [Verma, R. K.; Bhuin, N. C.; Mukhopadhyay, M. Geology, Structure and tectonics of Jharia CoalField, India-A 3-D Model. Geoexploration 1979, 17, 305−324].
recoverable resources. The exploration and exploitation of Gondwana coal in India have continued since 1774. However, associated shales were considered seal (cap-rock) and not a resource; hence, scientific documentation has never been done. Presently, gathering information on fundamental shale gas reservoir characteristics is in full swing in India. The coal seams of the Jharia basin are placed in Category-I, known for maximum future commercial methane potential among Indian CBM blocks. There are thick shale horizons in Barren Measures and the strata between coal seams of the Barakar Formation. The micro properties of these shales are yet to be understood. In this study, the carbon-rich Barakar and Barren Measures shale beds of the Jharia basin have been evaluated for variation in pore size, pore structure, and fractal dimensions with significance to gas storage. This study attempted an analysis of the pore structure, associated fractal dimensions, and sorption mechanism variation in thick shale sequences of Barakar and Barren Measures shale beds of the Jharia basin have been evaluated for variation in pore size, pore structure, and fractal dimensions with significance to gas storage.

This study attempted an analysis of the pore structure, associated fractal dimensions, and sorption mechanism variation in thick shale sequences of Barakar and Barren Measures Formation of the Jharia basin. Clay-associated pores significantly store methane and play an important role in the adsorption and desorption phenomena in shale. The gas flow in shale rocks is governed mainly by clay containing a pore network by the non-Darcy effects. Therefore, the porosity and permeability of shale depend on clay type and mineral content. Low-pressure N₂ adsorption/desorption isotherms and their hysteresis patterns provide helpful information about the physisorption mechanism, surface area, pore volume, pore size, and pore structure characteristics of the shale/coals. The fractal dimensions derived following FHH models indicated a complex fractal system associated with a matrix containing micro-, meso-, and macropores. The hysteresis patterns corresponding to the shale sample correlate with the pore structure identified with scanning electron microscopy (SEM). The study provides insights into the variation of organo-inorganic formed pores, pore structures, surface area, and fractal dimensions regarding shale gas storage and the recovery mechanism.

2. GEOLOGY OF THE JHARIA BASIN

Jharia is one of the most prospective coking coal-bearing basins of the Damodar valley, Jharkhand, and stretches over 450 km². It lies between latitudes 23°37′ and 23°52′ and longitudes 86°06′ and 86°30′. It occurs in a “half-graben” structure, trending in an east–west direction and plunging in the west curtailed by a key fault at the southern boundary having over 1500 m throw. The geological map of the Jharia basin showing the study area and shale core sampling boreholes is given in Figure 1. The profile and cross sections based on gravity and borehole data along the section lines in the geological map of the Jharia basin have been modified after others have considered it postdepositional faults.

### Table 1. Stratigraphic Succession of the Jharia Basin (Modified after Mukhopadhyay et al., 2010)

| Age (Ma) | System/Period | Series/Epoch | Stage/Age | Formation/Thickness | Lithology |
|----------|---------------|--------------|-----------|---------------------|-----------|
| 66 110   | Cretaceous    |              |           | Igneous Intrusive   | Basaltic dykes and sills |
| 252      | Upper Permian | Lopingian    | Canghsingian | Ranigunj (800 m)    | Fine to medium grained greenish sandstone with shale and coal seams |
| 260      | Middle Permian| Gasadalpurian| Wordian   | Barren Measures (750m) | Clay-rich and carbon-rich shale dominated hetero-lithic sequence. Massive sandstone |
| 271 290  | Lower Permian | Cisuralian   | Kungurian | Barakar (1250 m)    | Dirty white coarse and medium grained felspathic sandstones, grits, grey shale, carbon-rich shale with coal seams |
| 299 302  | Upper Carboniferous | Ghelhian | Talchir (245 m) | White coarse grained sandstone, green shales and fine grained sandstone, conglomerate lenses and basal tillite |

Unconformity

| Age (Ma) | System/Period | Series/Epoch | Stage/Age | Formation/Thickness | Lithology |
|----------|---------------|--------------|-----------|---------------------|-----------|
| Pre-cambrian |               |              |           | Chotanagpur Complex | Granite gneiss, hornblende schist/gneiss, metabasic rocks, pegmatite and quartz veins etc. |
According to Varma et al., internal basin tectonics played a significant role in folding the formations. The gradual decrease in thickness of the Barakar formation can be observed from the east to the west. The maximum thickness of the Barren Measures has been recorded in the central part of the basin, decreasing westward and eastward. The stratigraphic succession of the Jharia basin, modified after, showing the lower Gondwana sequence comprising shale and coal horizons, is presented in Table 1.

### 3. RESULTS AND DISCUSSION

#### 3.1. Organo-Inorganic Composition by FTIR Spectroscopy

The results of FTIR analysis of shale core samples are offered in the tabular form showing the presence of O–H stretching, C–C stretch, C=O, carbonates, kaolinite, Si–O stretching, etc. (Table 2). Organic matter in shale is a complex mix of components of numerous physico-chemical assets shown by FTIR spectra, which replicate features inbred from the original organic matter and also assimilated during subsequent diagenesis.

The facets of the microscopic organic constituents (macerals) of shale/coal are fundamental in defining economic utility, oil and gas-generating potential, and a congregation of other properties. The small peaks between 3610 and 3640 cm\(^{-1}\) indicate the hydrous kaolinite intermixed with organic compound hydroxyl (O–H) groups. The O–H stretch of hydrogen bond (3300 cm\(^{-1}\)) with aliphatic to aromatic chain signifies the organic matter thermal gas genesis pattern in shale. The minor peak of aromatic stretch (CH\(_x\)) at 3100–3000 cm\(^{-1}\) indicates initiation of aromatization at an early stage.

### Table 2. Results of FTIR Analysis of Shale Core Samples

| sample no. | formation   | OHstretching/phyllosilicate | C–C stretch/kerogen | aromatic C=O | carbonates | kaolinite | Si–O stretching/quartz | A-factor | C-factor | IAL | IAR |
|------------|-------------|----------------------------|---------------------|--------------|------------|-----------|------------------------|--------|--------|----|----|
| JHK-1      | Barren Measures | √− | − | √ | √ | √ | 0.42 | 0.54 | 0.35 | 0.53 |
| JHK-2      | Barren Measures | √− | − | √ | √ | √ | 0.38 | 0.41 | 0.24 | 0.62 |
| JHK-3      | Barren Measures | √− | − | √ | √ | √ | 0.41 | 0.17 | 0.20 | 0.45 |
| JHK-4      | Barren Measures | √− | − | √ | √ | √ | 0.35 | 0.07 | 0.03 | 0.61 |
| JHK-5      | Barren Measures | √− | − | √ | √ | √ | 0.52 | 0.38 | 0.21 | 0.55 |
| JHK-6      | Barren Measures | √− | − | √ | √ | √ | 0.38 | 0.24 | 0.13 | 0.48 |
| JHK-7      | Barren Measures | √− | − | √ | √ | √ | 0.34 | 0.32 | 0.18 | 0.62 |
| JHK-8      | Barren Measures | √− | − | √ | √ | √ | 0.18 | 0.26 | 0.14 | 0.51 |
| JHK-9      | Barren Measures | √− | − | √ | √ | √ | 0.32 | 0.29 | 0.16 | 0.48 |
| JHK-10     | Barren Measures | √− | − | √ | √ | √ | 0.29 | 0.29 | 0.16 | 0.45 |
| JHK-11     | Barren Measures | √− | − | √ | √ | √ | 0.30 | 0.27 | 0.15 | 0.38 |
| JHK-12     | Barren Measures | √− | − | √ | √ | √ | 0.44 | 0.38 | 0.15 | 0.64 |
| JHK-13     | Barren Measures | √− | − | √ | √ | √ | 0.36 | 0.19 | 0.10 | 0.47 |
| JHK-14     | Barren Measures | √− | − | √ | √ | √ | 0.42 | 0.18 | 0.08 | 0.52 |
| JHK-15     | Barren Measures | √− | − | √ | √ | √ | 0.49 | 0.41 | 0.22 | 0.56 |
| JHK-16     | Barren Measures | √− | − | √ | √ | √ | 0.34 | 0.32 | 0.18 | 0.62 |
| JHK-17     | Barren Measures | √− | − | √ | √ | √ | 0.18 | 0.26 | 0.14 | 0.51 |
| JHK-18     | Barren Measures | √− | − | √ | √ | √ | 0.32 | 0.29 | 0.16 | 0.48 |
| JHK-19     | Barren Measures | √− | − | √ | √ | √ | 0.29 | 0.29 | 0.16 | 0.45 |
| JHK-20     | Barren Measures | √− | − | √ | √ | √ | 0.30 | 0.27 | 0.15 | 0.38 |
| JHK-21     | Barren Measures | √− | − | √ | √ | √ | 0.44 | 0.38 | 0.15 | 0.64 |
| JHK-22     | Barren Measures | √− | − | √ | √ | √ | 0.36 | 0.19 | 0.10 | 0.47 |
| JHK-23     | Barren Measures | √− | − | √ | √ | √ | 0.42 | 0.18 | 0.08 | 0.52 |
| JHK-24     | Barren Measures | √− | − | √ | √ | √ | 0.49 | 0.41 | 0.22 | 0.56 |

*Explanations: A-factor = \(\frac{(2940 + 2850) cm^{-1}}{(2940 + 2850 + 1604) cm^{-1}}\); C-factor = \(\frac{1740 cm^{-1}}{(1740 + 1604) cm^{-1}}\); IAL, aliphaticity index; IAR, aromaticity index.

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Figure 3. FTIR spectra showing different stretchings of aliphatic and aromatic chains in shale samples of the Jharia basin. (1) C–Br stretch, alkyl halides. (2) Aromatic CH\(_x\) out-of-plane deformation. (3) C–O stretch. (4) Aromatic carbon C=C. (5) Oxygenated groups. (6) Aliphatic CH\(_x\) stretching. (7) Aromatic CH\(_x\) stretching. (8) O–H stretch, hydrogen-bonded. (9) O–H stretch, free hydroxyl.

Figure 4. Relation of IAL and IAR with TOC content in shale, (a) Barren Measures and (b) Barakar Formations. IAL, aliphaticity index; IAR, aromaticity index; TOC, total organic carbon.
stage under reducing conditions. The annihilation of the aliphatic chain can be attributed to the aromatic constituents’ enrichment at the peak of 2800 cm\(^{-1}\). The CO stretch of carbonates associated with oxygenated groups is shown at the peak of 1800 cm\(^{-1}\). It accentuates the aggregation of secondary mineral carbonates due to the increase in salinity approaching dry conditions in macropores and fractures of shale. The well-developed aromatic chain with trivial absorption (C=C) recorded between 1600 and 1550 cm\(^{-1}\) specifies that the studied shale passed through the moderate stage of thermal maturation (Figure 3).

The distinct peak of C-O of organic compounds, mainly the aromatic chain, indicates the significant conversion of aliphatic components to aromatic (oil/gas) accentuated by the liberation of volatiles and dry gases through organo-inorganic pores. Also, the more or less uniform symmetric peaks observed between 1250 and 1000 cm\(^{-1}\) denote the deformation of the –CH\(_3\) group during alteration of aliphatic to aromatic components.\(^{59-62}\) The peaks of aromatic regions between 900 and 750 cm\(^{-1}\) illustrate the removal of aromatic substitutes while attaining the thermal maturity >1.20% (bituminous stage) resulting upsurge in the aromatic condensation. Further, it is concluded that the substantial aromatization show a very good gas genesis pattern in shale, and it might have been stored in the pore-associated matrix system.\(^{58,63,64}\)

The FTIR spectra have been used to draw the indications on thermal maturity and kerogen type in the studied shale. The comparative ratio of aliphatic and aromatic bands is defined as the A-factor. A part of a carbonyl or carboxyl and the aromatic bands are calculated as the C-factor following the equation suggested by ref 69.

Figure 5. Relation of A-factor and C-factor derived from FTIR analysis of shale showing the presence of mainly type III/IV kerogen\(^{32}\) (after Ganz and Kalkreuth, 1987). [Ganz, H.; Kalkreuth, W. Application of infrared spectroscopy to the classification of kerogen types and the evolution of source rock and oil shale potentials. Fuel 1987, 66, 708–711].

Figure 6. Adsorption and desorption curve showing variation in the hysteresis pattern due to different pore structures in Barren Measures shale samples.
A-factor (aliphatic/aromatic groups) = \((2940 + 2850) \text{ cm}^{-1} / (2940 + 2850 + 1604) \text{ cm}^{-1}\).

C-factor (carboxyl and carbonyl/aromatic groups) = \(1740 \text{ cm}^{-1} / (1740 + 1604) \text{ cm}^{-1}\).

Similarly, the aliphaticity index (IAL) and the aromaticity index (IAR) are calculated using the distinct CH\(_2\) and CH\(_3\) stretching peaks as given in eqs 1 and 2.

\[
\text{IAL} = \frac{(2950 + 2940 + 2850) \text{ cm}^{-1}}{(3050 + 2950 + 2940 + 2850 + 1604) \text{ cm}^{-1}}
\]

\[
\text{IAR} = \frac{(3100 + 1600) \text{ cm}^{-1}}{(3100 + 3000 + 2850 + 2800 + 1600) \text{ cm}^{-1}}
\]

The values of the A-factor, C-factor, IAL, and IAR indicate a very narrow range due to uniform transformation of the organic content in the shales of Barakar and Barren Measures Formations (Table 2). According to ref 70, the absorption peaks of different functional groups have changed uniformly between 2800 and 3300 cm\(^{-1}\), revealing continuous reduction in aliphatic compounds (during transformation of kerogen I and II to III and IV), a general phenomenon obtained in matured carbon-rich shale or coal. Thus, the IAL and IAR values of the studied shales can ascertain the relative abundance of aliphaticity and aromaticity over the total aliphatics and aromatics.\(^{71,72}\) The relation of IAL and IAR with the TOC content indicates the hydrocarbon generation trend in the studied shale (Figure 4a,b). There is a very minute difference in the Barakar and Barren Measures shale hydrocarbon genesis patterns when converting aliphatic to aromatic rings. However, the slightly higher expulsion of the aliphatic in the Barakar shale can be attributed to greater thermal maturity, which improved the relative aromaticity. The relation of A-factor and B-factor derived from the FTIR analysis of shale shows mainly type III/IV kerogen (Figure 5). It is interpreted that the organic matter of both shale formations significantly transformed and achieved moderate to matured thermal maturity.

3.2. Sorption Pattern and Pore Structures. Low-pressure N\(_2\) adsorption and desorption curves give information on the free pore surfaces, fractals, pore openings, pore types, and pore structures present in the shale matrix.\(^{2,5,11,20,73}\) The gas is adsorbed and stored mainly in micro- (<2 nm), meso- (2–50 nm),...
and macropores (>50 nm). However, some of the macropores
>50 nm and fracture interconnections provide a path for the gas
to flow and seep into the shale. The adsorption pattern indicates
the type II isotherm with desorption hysteresis of H2 and H3
(Figures 6 and 7). The likely cause for the hysteresis pattern
was that capillary condensation happened in the major pores,
signifying that the shale samples have abundant mesopores of slit
and cylindrical structures.3,4,74 The larger loop of the adsorption
and desorption hysteresis pattern indicates a cylindrical pore
with a slit type of opening. The unusual sorption pattern of the
sample JHK-2 has mesopores mainly >10 nm, as shown in BJH
plots in Figure 8. The negative adsorption in shale is due to the
narrow and blind opening micropores that are difficult to admit
N2 gas, bearing low kinetic energy of adsorption between relative
pressure (P/Po) range 0.2 and 1.0. In JHK-12, JHK-13, and
JHK-15, open curves of adsorption and desorption indicate that
the hysteresis phenomenon in low-temperature N2 adsorption
isomers is usually associated with capillary condensation in
mesopore structures. Generally, different shapes of hysteresis
loops are caused by different types of adsorbents. Moreover,
swelling/shrinking of the sample due to interaction with N2 can
result in low-pressure hysteresis. The pore types are classified
into four categories taking into account the pore structures and
openings, such as (i) cylindrical, (ii) slit, (iii) combined, and (iv)
condensate (Table 3). The shale samples of Barren Measures
having cylindrical (samples JHK-1, JHK-3, JKH-4, JHK-5, and
JHK-6) and condensate (sample JHK-2) pores are suitable for
gas adsorption and release. However, the shale belonging to
the Barakar Formation has cylindrical (samples JHK-7, JHK-8,
JHK-9, and JHK-10), slit (samples JHK-13 and JHK-15), combined
(samples JHK-11), and condensate (sample JHK-15) pores due
to the comparatively complex geochemical properties. Barakar
shale has these properties because it is older and has passed
through a higher degree of thermal maturity than the Barren
Measures shale, as discussed in section 3.1. Further, the desorption
pattern presents a downward modulation point between the
relative pressure (P/Po) range of 0.4 and 0.6, suggesting that a
significant quantity of adsorbed gas may be released into the free
wide space condensate pore structure (Figures 6 and 7). The
combined pores replicate the interconnectivity of pores attributed
to the organo-inorganic content (maceral and clay/mineral pore
interaction) due to geochemical alterations.75 The minute variation
in the surface area indicated by single-point BET and multipoint
BET ranges from 1.38 to 13.38 and 1.38 to 13.95 m2/g, respectively,
for Barren Measures and from 3.38 to 10.91 and 3.75 to 11.26 m2/g
for the Barakar shale, respectively, demonstrating the similar influence of volatile and moisture
contents on rugged (fractal) surfaces of pores (Table 3). The
comparatively low surface area in Barakar shale pores speci-
fies the abundance of aromatic ring numbers associated with the
aromatic series comprising the narrow pore openings and con-
densate structures due to the higher degree of thermal maturaton.
Likewise, the surface area was also determined using methods and
processes such as Langmuir, BJH, t-method, DR, and DFT
(Table 3). The DFT method is always considered a reliable
method for determining the surface area, pore size, and pore

Figure 8. BJH plot for determination of pore size distribution and pore volume of Barren Measures shale samples.
The average surface area determined by DFT varies from 0.67 to 13.05 and 3.40 to 9.48 m²/g for Barren Measures and Barakar shale, respectively. Hence, it is concluded that shale beds of Jharia have heterogeneous complex pore structures, a rough surface, and a sorption mechanism controlled by weathering/alteration, depositional conditions, and organo-inorganic content. 

3.3. Evaluation of Pore Size and Its Distribution. The pore size distribution and pore volume were determined from N₂ adsorption data following BJH and DFT methods (Table 4) according to the pore classification suggested by ref 76. The BJH curves show pore size distribution and cumulative pore volume, mainly the presence of mesopores in Barren Measures and Barakar shales, the nontraceable micropores, and a small

| sample no. | single-point BET | multipoint BET | Langmuir | BJH | DH | t-method external | t-method micropore | DR | DFT |
|---|---|---|---|---|---|---|---|---|---|
| JHK-1 | 13.38 | 13.95 | 22.6 | 7.32 | 7.40 | 9.95 | 4.00 | 38.99 | 10.19 |
| JHK-2 | 7.97 | 8.24 | 13.24 | 4.81 | 4.86 | 8.24 | 25.72 | 6.66 |
| JHK-3 | 12.97 | 13.3 | 20.11 | 3.67 | 3.71 | 6.53 | 6.77 | 23.30 | 13.05 |
| JHK-4 | 1.38 | 1.38 | 2.75 | 0.34 | 0.38 | 0.57 | 0.81 | 2.61 | 0.67 |
| JHK-5 | 1.53 | 2.60 | 2.10 | 0.48 | 0.51 | 1.40 | 1.20 | 0.08 | 3.46 |
| JHK-6 | 2.88 | 3.07 | 4.87 | 2.03 | 2.06 | 3.07 | 6.30 | 3.96 |
| JHK-7 | 8.93 | 9.15 | 14.49 | 4.36 | 4.40 | 3.73 | 1.42 | 25.64 | 8.20 |
| JHK-8 | 6.29 | 6.35 | 10.83 | 3.10 | 3.13 | 5.64 | 0.88 | 6.46 | 5.22 |
| JHK-9 | 8.92 | 9.35 | 16.14 | 5.18 | 5.23 | 7.10 | 2.25 | 8.96 | 8.22 |
| JHK-10 | 5.09 | 9.11 | 47.41 | 4.89 | 4.95 | 5.13 | 3.98 | 19.02 | 3.61 |
| JHK-11 | 7.25 | 8.59 | 2.04 | 6.02 | 6.09 | 8.18 | 0.41 | 2.64 | 5.22 |
| JHK-12 | 8.17 | 9.29 | 19.54 | 6.04 | 6.10 | 8.14 | 1.15 | 27.94 | 5.90 |
| JHK-13 | 3.38 | 3.75 | 6.69 | 2.94 | 2.97 | 3.75 | 12.52 | 3.40 |
| JHK-14 | 5.38 | 7.98 | 24.35 | 6.59 | 6.66 | 5.88 | 2.10 | 22.68 | 3.84 |
| JHK-15 | 10.91 | 11.26 | 18.05 | 5.98 | 6.04 | 8.53 | 2.73 | 21.98 | 9.48 |

| sample no. | avg. BJH | DH | DR | DA | DFT |
|---|---|---|---|---|---|
| JHK-1 | 8.69 | 2.98 | 2.98 | 11.68 | 2.90 | 3.97 |
| JHK-2 | 9.46 | 2.99 | 2.99 | 12.88 | 3.14 | 3.97 |
| JHK-3 | 7.38 | 5.09 | 5.09 | 1.82 | 1.68 | 3.97 |
| JHK-4 | 4.27 | 3.16 | 3.16 | 6.21 | 2.10 | 1.41 |
| JHK-5 | 9.46 | 2.99 | 2.99 | 3.14 | 1.38 | 0.78 |
| JHK-6 | 8.97 | 3.53 | 3.53 | 2.36 | 1.98 | 1.41 |
| JHK-7 | 8.17 | 2.99 | 2.99 | 11.89 | 2.98 | 3.97 |
| JHK-8 | 8.13 | 3.16 | 3.16 | 1.66 | 1.70 | 3.97 |
| JHK-9 | 6.77 | 2.99 | 2.99 | 1.70 | 1.68 | 3.97 |
| JHK-10 | 4.52 | 2.98 | 2.98 | 12.56 | 3.00 | 3.17 |
| JHK-11 | 8.12 | 3.33 | 3.33 | 13.12 | 3.12 | 3.97 |
| JHK-12 | 7.26 | 2.99 | 2.99 | 12.76 | 3.06 | 3.97 |
| JHK-13 | 5.46 | 3.34 | 3.34 | 13.29 | 3.16 | 3.80 |
| JHK-14 | 5.26 | 3.32 | 3.32 | 13.37 | 3.08 | 2.77 |
| JHK-15 | 6.74 | 2.99 | 2.99 | 2.09 | 1.84 | 3.97 |

| pore size (nm) | pore volume (cc/g) |
|---|---|
| BJH | DH | t-method | DR | DFT |
|---|---|---|---|---|
| 0.022 | 0.017 | 0.002 | 0.014 | 0.019 |
| 0.019 | 0.017 | 0.007 | 0.009 | 0.016 |
| 0.018 | 0.013 | 0.012 | 0.003 | 0.008 | 0.016 |
| 0.002 | 0.002 | 0.002 | 0.001 | 0.004 |
| 0.017 | 0.006 | 0.006 | 0.002 | 0.006 |
| 0.007 | 0.006 | 0.006 | 0.007 | 0.016 |
| 0.019 | 0.016 | 0.016 | 0.009 | 0.016 |
| 0.008 | 0.011 | 0.011 | 0.002 | 0.011 |
| 0.016 | 0.013 | 0.013 | 0.001 | 0.003 | 0.014 |
| 0.006 | 0.009 | 0.009 | 0.007 | 0.009 |
| 0.017 | 0.016 | 0.015 | 0.009 | 0.014 |
| 0.017 | 0.015 | 0.015 | 0.009 | 0.014 |
| 0.008 | 0.004 | 0.004 | 0.007 | 0.007 |
| 0.010 | 0.010 | 0.010 | 0.008 | 0.009 |
| 0.019 | 0.016 | 0.015 | 0.001 | 0.007 | 0.016 |

Table 3. Low-Pressure N₂ Adsorption Isotherm Type, Hysteresis Pattern, Pore Structures, and Results of Surface Area Derived from Different Methods

Table 4. Results of Pore Size and Pore Volume Determined Using Different Methods from Low-Pressure N₂ Adsorption Isotherms

volume of the shale samples. The average surface area determined by DFT varies from 0.67 to 13.05 and 3.40 to 9.48 m²/g for Barren Measures and Barakar shale, respectively. Hence, it is concluded that shale beds of Jharia have heterogeneous complex pore structures, a rough surface, and a sorption mechanism controlled by weathering/alteration, depositional conditions, and organo-inorganic content. 

3.3. Evaluation of Pore Size and Its Distribution. The pore size distribution and pore volume were determined from N₂ adsorption data following BJH and DFT methods (Table 4) according to the pore classification suggested by ref 76. The BJH curves show pore size distribution and cumulative pore volume, mainly the presence of mesopores in Barren Measures and Barakar shales, the nontraceable micropores, and a small
quantity of macropores (Figures 8 and 9). The average pore sizes are determined to be $5.38 - 9.46$ and $4.52 - 8.17$ nm for Barren Measures and Barakar Formation shales, respectively. In DFT plots, the major part of the pore volume in Barren Measures shale is attributed to mesopores having a diameter between 2.00 and 6.00 nm. However, the Barakar shale also has a comparatively larger size mesopore distribution between 2.00 and 8.00 nm, except for shale sample JHK-13, which has a relatively more significant variation in pore distribution due to its sandy characteristics (Figures 10 and 11). According to pore abundance, the studied shale ensures that the successive trend is mesopores > macropores > micropores. Also, the DFT curves’ trimodular pattern and the pore surface area concerning pore size contributed similarly to the pore volume. Moreover, the variation in pore size indicates the increase in pore volume with a decrease in pore size distribution in the shale, which suggests that the adsorption capacity in shale is highly related to the pore size difference, openings, and their structures. In general, it is recorded that pores of small size have a large pore volume with a greater adsorption capacity.

The pores observed from SEM photographs compared with the low-pressure N$_2$ sorption pattern validate the presence of cylindrical, combined, pore-pipes, and slit pores in shale (Figure 12) such as JHK-1: large open cylindrical pores, JHK-10: combined fractured pores, JHK-11: deep rounded pore-pipes, and JHK-13: wide spacing slit pores. These pores associated with organic matter, clay, and silt are formed from the framework of organo-inorganic content and microfractures. Hence, the microfractures have a higher complexity due to the heterogeneity attributed to the banded nature of the shale. Therefore, it is summarized that Barren Measures and
Barakar shale have a complex pore morphology and a narrow variation in pore size that vary from nanometers to micrometers and may significantly influence the gas storage and fluid flow mechanism in the shale.

3.4. Types of Fractal Dimensions. Identifying the type of fractal pore surfaces in the shale matrix system helps characterize the complex structures and gas storage capacity. Different researchers propose several models to determine pore fractal dimensions based on gas adsorption and desorption patterns like the BET model, the Langmuir model, the FHH model, and the thermodynamic model. In this study, the FHH model has been used to determine the fractal characteristics of shale pores, which can be expressed as

$$\ln \frac{V}{V_0} = A \left[ \ln \left( \ln \left( \frac{P}{P_0} \right) \right) \right] + \text{constant}$$

where $V$ is the adsorbed gas volume, $V_0$ is the monolayer adsorbed gas volume, $P$ is the equilibrium pressure, $P_0$ is the saturated vapor pressure of gas, and $A$ is the slope depending on the adsorption pattern and pore fractals. Thus, the values of fractal dimensions ($D$) can be calculated using eqs 4 or 5

$$D = A + 3$$  \hspace{1cm} (4)

or

$$D = 3A + 3$$  \hspace{1cm} (5)

The equation $D = A + 3$ was applied for the hollow portion in shale pores shown by hysteresis loops initiating capillary condensation; however, the equation $D = 3A + 3$ applies to the van der Waals force.$^{5,79,88}$ The $\ln (V)$ vs $\ln [\ln (P_0/P)]$ plots of Barren Measures and Barakar shale samples display three distinct fractal dimension surfaces corresponding to micro-, meso-, and macro-pores are shown in Figures 13 and 14. The fractal dimension $D_1$ was calculated from linear sections at $P/P_0$ between 0.0002 and 0.0090, $D_2$ was calculated from linear sections at $P/P_0$ between 0.0090 and 0.3000, and $D_3$ was estimated from the linear sections at $P/P_0$ between 0.3000 and 1.0000. The fractal fitting equations and fractal dimension values for the studied shale samples are given in Tables 4 and 5, respectively. The values of $D_1$, $D_2$, and $D_3$ of the Barren Measures shale are in the range of 1.25–2.17, 2.09–2.55, and 2.57–2.76, respectively, while the Barakar shale fractal values varying in the range of 1.45–2.09, 1.11–2.53, and 2.69–2.76, respectively (Table 6). The correlation coefficient of fractal fitting equations ($R^2_1$ and $R^2_3$) of $D_2$ and $D_3$ is greater than 0.95, indicating Barren Measures and Barakar shale pore systems with noticeable fractal surfaces mostly from meso- and macropores. However, it is interpreted that both the formation shales have composite pore internal structures attributed to three different kinds of fractals. The Barakar shale with a comparatively higher thermal maturity developed more irregular pore surfaces during dehydration and devolatilization. It results in the amalgamation of complex macropores to mesopores (Figure 14). Also, the fractal surfaces amplified with increasing $P/P_0$ specifying that compared with the bigger pores (macropores), the smaller pores (micro- and mesopores) are more complex and composite. The $D_2$ and $D_3$
represent the meso- and macropores with a fractal value near 3. Mainly, the intergranular pores that occur between grains of organic and inorganic contents dominate the meso- and macro pores. However, the organic pores, kaolinite fissility openings along the bedding planes, and the subrounded to rounded spacing in grains of quartz, feldspar, and carbonate minerals also contribute to the meso- and macropores. Numerous researchers have stated that the clay mineral content is an important factor

Figure 11. DFT plot for determination of pore size distribution of Barakar shale samples.
influencing the shale pore structure; hence, the clay mineral also contributes to the fractal dimension.\(^5,15,80,88\)

### 3.5. Relationship of Fractal Dimensions with Shale Properties.

The different properties of shale directly influence fractal pore surfaces.\(^3,11,79,85\) The fractal dimensions \(D_1\), \(D_2\), and \(D_3\) show successive transitions due to change in thermal maturity (Figure 15a,c). The release of volatiles due to the cracking of hydrocarbon compound of shale matrix leads to the formation of rugged pore surfaces. A similar trend shown by the relation of fractals with fixed carbon specify succeeding carbon enrichment due to thermal transformation of organic matter contributed to the rough pore surfaces (Figure 15b,e). The TOC content influences the advancement of pores in shales. More pores evolved in higher TOC content shales, and the complex pore surfaces and structures led to greater fractal dimensions. Earlier investigations had shown that the clay and mineral content are a vital aspect controlling the pore types and pore structure of shale.\(^3,15,80,81\) The increasing trend of fractal dimension with the ash content of shale demonstrates that inorganic content also contributed to the formation of pore surfaces (Figure 15d). The rounded to subrounded grains of minerals (e.g., quartz, feldspar, etc.) associated with clays shaped pore surfaces. The variation in fractal dimensions \((D_1, D_2,\) and \(D_3\)) with depth indicates that heterogeneity in lithotype, organo-inorganic content, and thermal maturity controls the construction anomaly of fractal surfaces (Figure 15f).

### 3.6. Empirical Model for Estimation of Porosity and Permeability.

The measurement of porosity and permeability of shale is time-consuming and expensive. Therefore, an empirical method is formulated and proposed to estimate porosity and permeability considering the fractal dimensions, pore size, and pore volume considering their constructive and adverse influences. The relations of measured porosity (MPOc) and measured permeability (MPeC) were evaluated with fractal dimensions, pore size, and pore volume. The moderate linear relationship of MPOc with MPeC \((R^2 = 0.4165)\) indicates pore

Figure 12. Low-pressure N\(_2\) sorption isotherms showing the presence of different types of pores validated by SEM photographs.
connectivity suitable for gas flow in a shale (Figure 16a). The $D_1$ showed a negligible relationship with MPOc, emphasizing that plane surfaces seldom contribute to porosity in shales (Figure 16b). However, the very good linear relation of $D_2$ ($R^2 = 0.8442$) and $D_3$ ($R^2 = 0.6809$) with MPOc (Figure 16c,d) specifies that these fractals are mainly remunerating the porosity in shale through the active rough surfaces of pores formed by the heterogeneous content of organo-inorganic matter.

Similarly, fractals $D_2$ and $D_3$ also showed a direct moderate correlation with MPEc, indicating the combined effects of rugged surfaces on the gas flow mechanism in the pore-associated matrix system of the shale (Figure 16e). The moderate positive relationship of average pore size and MPEc illustrates that the larger pore opening supports the gas flow because of low turbulence and low capillary pressure (Figure 16f). The MOPc and MPEc showed a linear moderate to very good relation with the BET pore volume ($R^2 = 0.5213$ and 0.8422), signifying that matrix containing pores are mainly interconnected (Figure 16g) and interdependent. However, the maceral, mineral, and clays have different porous features due to intricacy in their chemical composition; pore size variation led to fractal dimensions appropriate for adsorption, storage, and flow of gas in shales.

Considering the significance of the above-explained parameters, the following empirical equation is drawn to estimate porosity and permeability in shale:

estimated porosity ($EPO_f$) = \[
\left(\frac{0.2 \times PV_{\text{BET}} + D_2 + D_3}{(4.0 - 0.2 \times D_1 - 0.7 \times D_2)}\right)\times 100\] \[= 1.3\]  

(6)

where $EPO_f$ is the estimated porosity by an empirical method using pore fractals, $PV_{\text{BET}}$ is the pore volume determined by the BET method, and $D_2$ and $D_3$ are fractal dimensions.

estimated permeability ($EPE_f$) = \[
\frac{(0.3 \times PV_{\text{BET}})}{0.2 \times (PS_A + 0.5 \times D_3 + 5.8 \times D_2 + 0.5 \times D_1)}\] \n
(7)

where $EPE_f$ is the estimated permeability by the empirical method using pore fractals, $PV_{\text{BET}}$ is the pore volume determined by the BET method, $D_1, D_2$, and $D_3$ are fractal dimensions, and $PS_A$ is the average pore size.

The estimated porosity ($EPO_f$) and permeability ($EPE_f$) derived from the suggested empirical formula (Table 7) vary
from 0.98 to 1.58% and 0.35 to 0.63 mD for Barren Measures; however, for Barakar Formations, it is estimated to be 0.75–1.65% and 0.41–0.62 mD, respectively. The estimated values of porosity (EPOf) and permeability (EPEf) showed an excellent linear correlation with the measured porosity (MPOc, $R^2 = 0.8577$) and permeability (MPEc, $R^2 = 0.8577$), which are close

**Figure 14.** Presence of three distinct fractal dimensions derived by the FHH method for Barakar shale samples region I ($P/P_0 = 0.0002–0.0090; D_1$), region II ($P/P_0 = 0.0090–0.3000; D_2$), and region III ($P/P_0 = 0.3000–1.0$).
to the measured values. It is concluded that the proposed empirical formula for EPO\text{I} and EPE\text{I} may be a suitable indirect method for the estimation of porosity and permeability of shale samples (Figure 17a,c). Further, the validation of EPO\text{I} and EPE\text{I} calculated values has been done through a curve match with measured porosity and permeability and are presented in Figure 17b,d. The excellent curve matching demonstrates that the proposed empirical models can be used for the estimation of porosity and permeability.

4. SUMMARY AND CONCLUSIONS

The carbon-rich Barakar and Barren Measures shale beds of the Jharia basin were evaluated for organo-inorganic composition by FTIR, pore size, and pore structure using BET low-pressure N\textsubscript{2} adsorption and pore geometry through FE-SEM photographs. The study provides insights into the variation of organo-inorganic formed pores, pore structures, surface area, and fractal dimensions regarding the gas storage and recovery mechanism from shale. The following conclusions have been drawn from the study.

i. The shale samples have significant carbon-rich content and are intercalated-banded in nature, indicating fluvial-lacustrine facies of deposition.

ii. The fractal dimensions \(D_1\), \(D_2\), and \(D_3\) showed successive transitions due to change in thermal maturity.

iii. The values of A-factor, C-factor, IAL, and IAR indicate a very narrow range due to the uniform transformation of organic content in the shales of Barakar and Barren Measures Formations.

iv. The FE-SEM images indicated shallow to deep pores with different pore structures having fair to good pore connectivity.

v. The shale beds of Jharia have heterogeneous complex pore structures, rough surface, and sorption mechanism controlled by weathering/alteration, depositional conditions, and organo-inorganic content.

vi. The variation in pore size indicates the increase in pore volume with a decrease in pore size distribution in the shale, which suggests that the adsorption capacity in the shale is highly related to the pore size difference, openings, and their structures.

vii. The slightly higher expulsion of the aliphatic in Barakar beds of Jharia have heterogeneous complex pore structures, surface area, and sorption mechanism controlled by weathering/alteration, depositional conditions, and organo-inorganic content.

viii. The calculated values using proposed empirical models for porosity (EPO\text{I}) and permeability (EPE\text{I}) showed an

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Table 5. Fractal Fitting Equations for the Studied Shale Core Samples

| sample no. | \(A_1\) | \(D_1 = 3 + A_1\) \(D_2 = 3 + 3A_1\) \(R_1^2\) | \(A_2\) | \(D_2 = 3 + A_2\) \(D_3 = 3 + 3A_2\) \(R_2^2\) | \(A_3\) | \(D_3 = 3 + A_3\) \(D_3 = 3 + 3A_3\) \(R_3^2\) |
|-----------|--------|-----------------|--------|-----------------|--------|-----------------|
| JHK-1     | -1.35  | 1.85            | -1.05  | 0.99            | -0.50  | 2.50            | 1.49   | 1.00            | -0.31  | 2.69            | 2.06   | 0.98            |
| JHK-2     | -1.51  | 1.38            | -1.22  | 0.98            | -0.61  | 2.47            | 1.60   | 0.99            | -0.35  | 2.72            | 2.14   | 0.97            |
| JHK-3     | -1.75  | 1.25            | -2.24  | 0.99            | -0.57  | 2.43            | 1.29   | 1.00            | -0.24  | 2.76            | 2.29   | 0.93            |
| JHK-4     | -0.99  | 1.11            | 0.03   | 1.00            | -0.91  | 1.85            | 0.28   | 1.00            | -0.43  | 2.27            | 1.71   | 0.93            |
| JHK-5     | -1.14  | 1.86            | -0.42  | 1.00            | -0.45  | 2.55            | 1.66   | 1.00            | -0.28  | 2.72            | 2.18   | 0.99            |
| JHK-6     | -0.83  | 2.07            | 0.52   | 1.00            | -0.79  | 2.21            | 0.64   | 1.00            | -0.28  | 2.72            | 2.15   | 0.92            |
| JHK-7     | -1.37  | 1.63            | -1.12  | 1.00            | -0.47  | 2.53            | 1.60   | 1.00            | -0.28  | 2.72            | 2.16   | 0.99            |
| JHK-8     | -1.42  | 1.58            | -1.27  | 0.98            | -1.89  | 1.61            | -2.67  | 0.99            | -0.25  | 2.18            | 2.25   | 0.83            |
| JHK-9     | -1.02  | 1.98            | -0.05  | 1.00            | -0.49  | 2.51            | 1.53   | 1.00            | -0.25  | 2.75            | 2.24   | 0.95            |
| JHK-10    | -1.20  | 1.80            | -0.60  | 0.98            | -1.77  | 1.63            | -2.30  | 0.99            | -0.26  | 2.44            | 2.22   | 0.74            |
| JHK-11    | -1.55  | 1.45            | -1.64  | 0.99            | -0.53  | 2.47            | 1.41   | 0.99            | -0.31  | 2.69            | 2.07   | 0.97            |
| JHK-12    | -1.41  | 1.59            | -1.24  | 0.99            | -0.82  | 2.18            | 0.54   | 1.00            | -0.29  | 2.71            | 2.13   | 0.92            |
| JHK-13    | -0.91  | 2.09            | 0.28   | 0.99            | -1.01  | 1.99            | -0.04  | 0.99            | -0.27  | 2.73            | 2.18   | 0.72            |
| JHK-14    | -1.17  | 1.83            | -0.50  | 0.95            | -0.89  | 2.11            | 0.33   | 0.99            | -0.24  | 2.76            | 2.30   | 0.85            |
| JHK-15    | -1.51  | 1.49            | -1.53  | 1.00            | -0.49  | 2.51            | 1.53   | 0.99            | -0.25  | 2.75            | 2.25   | 0.94            |
excellent linear correlation with the measured porosity (MPOc) and permeability, which are close to measured values. The proposed models may be considered to estimate the porosity and permeability of shale and coal beds.

5. MATERIALS AND METHODS

5.1. Shale Core Sampling and Preparation. A total of 14 shale core samples were obtained from four boreholes (BH-1, BH-2, BH-3, and BH-4) drilled in the Jharia basin with laterally varying depth and thickness of the beds. The shale samples were manually crushed and sieved for further analysis like TOC, low-pressure N2 sorption, and scanning electron microscopy (SEM) and in the sizes of 72 mesh (212 μm), 0.8−1.0 mm, and 1.0−2.0 mm, respectively.

5.2. Total Organic Carbon (TOC) Analysis. The prepared shale samples of size 212 μm were analyzed for TOC using the Vinci Technologies “Rock-Eval 6 Plus TOC module” system. The TOC content of the shale was measured through oxidation under air. The contents of the pyrolyzable carbon and mineral-carbon were determined by thermally disintegrating the sample using a pyrolysis oven. More details about TOC determination using the Rock-Eval 6 system have been explained by different researchers.

5.3. Low-Pressure N2 Adsorption Isotherm. The Quantachrome Autosorb iQ was used to measure the low-pressure N2 adsorption and desorption isotherm following the Brunauer−Emmett−Teller (BET) method. The surface area, pore size, pore volume, and fractal dimensions of micro-, meso-, and macropores were determined following the procedures and models like multipoint BET, density functional theory (DFT), Langmuir, Barrett−Joyner−Halenda (BJH), t-test, and Frenkel−Halsey−Hill (FHH).

5.4. FTIR Spectroscopy. The shale samples were crushed to −75 μm, and 1 mg was mixed with 100 mg of KBr, followed by grounding the mixtures in an agate mortar pestle. The homogenized mixture was pressed in an exiled die to form pellets.
Figure 16. Relations of porosity and permeability with fractal dimensions, (a) porosity vs permeability, (b) fractal dimension ($D_1$) vs porosity, (c) fractal dimension ($D_2$) vs porosity, (d) fractal dimension ($D_3$) vs porosity, (e) fractal dimension ($D_1$) vs permeability, (f) fractal dimension ($D_2$) vs permeability, (g) fractal dimension ($D_3$) vs permeability, (h) average pore size vs permeability, (i) BET pore volume vs porosity, and (j) BET pore volume vs permeability.
following the procedure described in ref 25. Pellets were dried in a vacuum oven for 48 h to remove extra moisture and minimize their influence on FTIR spectra. FTIR analysis was conducted under a Bruker, 3000 Hyperion Microscope with a Vertex 80 FTIR system at IIT Mumbai in the wavelength range of 4000–400 cm\(^{-1}\) in the absorbance mode.

5.5. FE-SEM Analysis. SEM is an advanced tool to study the microfeatures of shale/coal. SEM photographs were used to find out the pore morphology, pore structures, secondary mineral infilings in fracture-pores, organo-inorganic homogeneity, and cleat spacing. The Carl Zeiss-make "FE-SEM Supra 55" with an EDX attachment under magnifications ranging from 4k× to 40k× was used. We took out small chips of shale samples from the core with the help of a chisel. The chips were mounted on brass stubs using silver glue paste and coated with platinum. The photographs were taken from a focused ion beam of different magnitudes. The SEM photographs were marked with 2D surface features (e.g., pore types, openings, structures, aperture, spacings, organo-inorganic allied pores, and fractures).

5.6. Measurement of Porosity and Permeability under Reservoir-Simulated Confining Pressure. The porosity and permeability determination of core samples under confining pressure with accuracy is very vital to design and plan gas recovery from a shale reservoir. The whole core permeameter

Table 7. Results of Measured Permeability and Porosity under Reservoir-Simulated Confining Pressure and Proposed Empirical Model of Shale Core Samples

| sample no. | depth (m) | inlet pressure (psi) | confining pressure (psi) | permeability (MPEC) (mD) | equilibrium pressure (psi) | measured permeability (MPEc) (mD) | measured porosity (MPOc) (%) | estimated permeability (EPEf) (mD) | estimated porosity and permeability (%)
|------------|-----------|----------------------|--------------------------|--------------------------|-----------------------------|--------------------------------|-------------------------------|--------------------------------|--------------------------------|
| JHK-1      | 197       | 260.11               | 282.74                   | 0.63                     | 123.45                      | 285.33                         | 1.48                          | 1.73                           | 0.84                           |
| JHK-2      | 289       | 396.45               | 412.18                   | 0.58                     | 285.02                      | 415.23                         | 1.28                          | 1.71                           | 0.74                           |
| JHK-3      | 329       | 448.69               | 468.75                   | 0.55                     | 310.56                      | 458.69                         | 1.32                          | 1.67                           | 0.63                           |
| JHK-4      | 331       | 458.05               | 472.23                   | 0.35                     | 328.96                      | 475.02                         | 0.98                          | 0.53                           | 0.05                           |
| JHK-5      | 343       | 462.85               | 486.32                   | 0.49                     | 368.42                      | 483.79                         | 1.58                          | 1.86                           | 0.68                           |
| JHK-6      | 351       | 480.69               | 500.21                   | 0.48                     | 394.08                      | 502.14                         | 1.28                          | 1.28                           | 0.25                           |
| JHK-7      | 408       | 565.23               | 582.76                   | 0.62                     | 412.26                      | 585.36                         | 1.48                          | 1.82                           | 0.71                           |
| JHK-8      | 416       | 575.26               | 592.63                   | 0.44                     | 436.85                      | 594.65                         | 0.76                          | 0.26                           | 0.23                           |
| JHK-9      | 431       | 595.18               | 615.49                   | 0.52                     | 456.78                      | 614.96                         | 1.65                          | 1.81                           | 0.57                           |
| JHK-10     | 450       | 605.14               | 635.29                   | 0.41                     | 485.12                      | 636.32                         | 0.75                          | 0.42                           | 0.14                           |
| JHK-11     | 476       | 660.47               | 680.12                   | 0.59                     | 512.20                      | 682.45                         | 1.32                          | 1.68                           | 0.63                           |
| JHK-12     | 496       | 686.69               | 706.58                   | 0.52                     | 525.87                      | 708.29                         | 1.28                          | 1.23                           | 0.56                           |
| JHK-13     | 594       | 827.30               | 845.36                   | 0.48                     | 645.29                      | 848.25                         | 1.22                          | 0.99                           | 0.23                           |
| JHK-14     | 646       | 894.55               | 919.05                   | 0.45                     | 746.95                      | 922.85                         | 1.37                          | 1.17                           | 0.30                           |
| JHK-15     | 904       | 1270.08              | 1286.36                  | 0.56                     | 1052.52                     | 1288.20                        | 1.62                          | 1.81                           | 0.67                           |

Figure 17. Validation of estimated porosity and permeability, (a) estimated porosity (EPOf) vs measured porosity (MPOc), (b) matching curve of measured and estimated porosity, (c) estimated permeability (MPEf) vs measured permeability (MPEc), and (d) matching curve of measured and estimated permeability.
and pycnometer systems used in this study were installed at CSIR-CIMFR, Dhanbad. It is a very fast, precise, and non-destructive determination of permeability and porosity of core samples under confining pressure.84,85 Shale core samples having diameter 52 mm and length 150 mm were taken and polished on both ends with the help of a rotator machine. The core was fixed into the core holder, and it was allowed to maintain the desired confining pressure. An extremely delicate pulse-decay permeameter devoted to the setup was used to measure the trivial gas flow through the core. It is an unsteady-state permeameter intended to measure permeability in the range of 1 milli Darcy (mD) to 10 nano Darcies (10 nD).84,86

The details of the procedure have been described by ref 4, 87.

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Notes
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