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## **Influence of geochemical features on the mechanical properties of organic matter in shale**

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### **Key Points:**

- Young's modulus of inertinite and vitrinite is much higher than that of bitumen
- Modulus of organic matter increases with thermal maturity
- Intensity ratio of D peak to G peak of inertinite's Raman spectrum exhibits a decreasing trend with maturity

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## Abstract

Organic matter is an important constituent in organic-rich shale, which influences the hydrocarbon generation, as well as the mechanical behavior, of shale reservoirs. The physical, chemical, and mechanical properties of organic matter depend on the source material and the thermal evolution process. Previous works attempted to investigate the impact of thermal maturation on the mechanical properties of organic matter. However, owing to the lack of maceral classification and the limitation of data volume during the mechanical measurement, no consistent trend has been identified. In this work, vitrinite reflectance test, scanning electron microscope observation, nanoindentation, and micro-Raman analysis were combined for geochemical and mechanical characterization. A total of 114 test areas were selected for testing, enhancing reliability of the test results. The Young's moduli of organic matter are from 3.57 GPa to 8.32 GPa. With the same thermal maturity, inertinite has the highest Young's modulus, while the modulus of bitumen is the lowest. The Young's moduli of different organic types all increase with vitrinite reflectance. When vitrinite reflectance increases from 0.62% to 1.13%, the modulus of inertinite and vitrinite is increased by 57% and 78%, respectively. The indentation creep of different organic matter suggests that the stiffer organic type demonstrates less creep influence. In addition, with the increase of thermal maturity, the micro-Raman test results show a decrease of intensity ratio of D peak to G peak, indicating an increase of the ordered structure in organic matter. From the characterization results, it was found that organic type and thermal maturity reflect the diversity of the source material and the chemical structure change during the thermal evolution process, and together they influence the mechanical properties of organic matter.

**Keywords:** organic type; thermal maturity; organic matter; Young's modulus; shale.

## 1. Introduction

Organic-rich shale is not only the source rock of conventional reservoirs, but also provides storage space for shale oil/gas reservoirs. Organic matter is an important constituent in organic-rich shale, which distinguishes organic-rich shale from other mud rocks with low organic content, and significantly influences the exploration and development of shale oil/gas. On the one hand, kerogen is the source material of hydrocarbon, and total organic carbon (TOC) content is an essential index for the evaluation of geological sweet spots (Cudjoe et al., 2016; Naizhen and Guoyong, 2016). On the other hand, the mechanical properties that significantly affect the fracturing behavior of shale are markedly different for organic matter and inorganic minerals in shale (Zhao and Zhang, 2020). Consequently, many indices for the evaluation of engineering sweet spots, for example, the brittleness index (Rybacki et al., 2016; Wang and Carr, 2012; Zhao et al., 2018), are also related to organic matter.

Organic matter in shale comes from deposits of the dead bodies of ancient organisms, and it can be classified on the basis of the diversity of source organisms and sedimentary environments. According to solubility, organic matter can be divided into kerogen and bitumen (Hunt, 1979). Regarding kerogen, two classification schemes exist: optical classification and chemical classification. The optical classification scheme borrows methods and concepts from coal petrology (Hutton et al., 1994; Stach et al., 1982). Through reflected light analysis, kerogen can be classified into maceral groups, such as inertinite, vitrinite, and exinite. Chemical classification focuses on the content of C, H, and O elements in kerogen. By major chemical elements analysis, kerogen can be classified as type I, type II, and type III (Tissot and Welte, 2013). Optical classification and chemical classification schemes are related to each other (Hunt, 1979). However, optical classification usually measures organic matter in certain micro regions, while chemical classification refers to characterization of the whole rock. In addition to organic type, thermal maturity constitutes another important geochemical property

of organic matter. Thermal maturity indicates the thermal evolution and hydrocarbon generation stage of the source rock. Vitrinite reflectance is a key indicator of thermal maturity. Vitrinite mainly originates from lignin and cellulose in higher plant debris, and the reflectance of vitrinite changes with maturation during thermal evolution (Suárez-Ruiz and Crelling, 2008; Taylor et al., 1998). Owing to the absence of higher plants, vitrinite is rare in rocks prior to the Devonian period. Consequently, researchers usually measure the reflectance of bitumen or graptolite as an alternative (Bertrand, 1990; Bertrand and Heroux, 1987; Jacob, 1989). Organic type and thermal maturity together influence the chemical structures of organic matter in shale.

Due to difficulties in obtaining sufficiently large samples, the elastic property of organic matter is difficult to characterize using conventional mechanical test methods. However, micromechanical test techniques, such as nanoindentation and modulus mapping based on nanoindenter and PeakForce quantitative nanomechanical mapping (QNM) by atomic force microscope (AFM), offer convenience of investigation at micro scale (Balooch et al., 2004; Dayal et al., 2009; Dokukin and Sokolov, 2012). Organic matter can be treated as a part of the main constituents in shale, and can be directly characterized through application of micromechanical test techniques (Alstadt et al., 2015; Bennett et al., 2015; Eliyahu et al., 2015; Kumar et al., 2012; Wilkinson et al., 2015; Zargari et al., 2013; Zeszotarski et al., 2004). The reported Young's modulus of organic matter was generally between 0 GPa to 25 GPa, which was much lower than that of inorganic minerals (Zhao et al., 2018). These researches revealed strong micromechanical heterogeneity within different constituents in organic-rich shales.

With the basic understanding of elastic properties from direct characterization, geochemical features of organic matter were taken into consideration in recent works (Emmanuel et al., 2016; Li et al., 2018; Zargari et al., 2016). Kerogen and bitumen with different thermal maturities were usually separated during the measurements. Previous works in the second type suggested that the elastic modulus of organic matter is associated with

organic type and thermal maturity. Due to the lack of classification of kerogen and the limitation of data volume, however, no consistent trend has yet been discerned. Yang et al. (2017) measured the mechanical properties of different macerals in shale using AFM-IR, which combines AFM and IR spectroscopy and collects the topography, IR adsorption, and stiffness map simultaneously with nanometer resolution. Inertinite, bitumen, and tasmanite were separated in the work, but thermal maturity was not considered for mechanical characterization.

Similar to characterization of organic matter in shale, the mechanical properties of macerals in coal and the influence of coal rank have also been investigated (Borodich et al., 2015; Epshtein et al., 2015; Vranjes et al., 2018; Zhang et al., 2019). However, elucidation of the exact influence requires further studies. In addition, molecular simulation has recently been utilized to examine the structures and mechanical properties of organic matter (Bousige et al., 2016; Tong et al., 2016; Ungerer et al., 2014). The chemical structure of organic matter is very complex, and to precisely construct a molecular model, and predict the physical and mechanical properties, remain a challenge. Experimental works at micro and nano scales can provide a useful reference and validation for simulations.

In this study, four shale samples with different thermal maturities from the Bonan Sub-sag in the Bohai Bay Basin, China are prepared. A number of geochemical and micromechanical tests are combined to characterize the properties of organic matter in each sample. Vitrinite reflectance is measured using a photomicrometer. Inertinite, vitrinite, and bitumen are identified based on an optical classification scheme. Nanoindentation is performed to characterize the mechanical properties of organic matter in the selected test areas. Moreover, to reveal the evolution of chemical structures of organic matter, micro-Raman test is carried out. Micromechanical test results are compared with the reported data. The influence of organic type and thermal maturity is discussed on the basis of the characterization results.

## 2. Materials and methods

### 2.1. Sample information and preparation

Samples used in this study are shale cores of the third member of the Shahejie Formation from the Bonan Sub-sag. The Bonan Sub-sag is a third-order tectonic unit in the central part of the Zhanhua Depression in the Bohai Bay Basin with an area of 600 km<sup>2</sup>. During the Paleogene period, the Bohai Bay Basin experienced initial, developmental, prosperous and atrophic periods of the fault depression, and successively developed the Kongdian Formation, the Shahejie Formation, and the Dongying Formation (Jiu et al., 2013; Wang et al., 2015). The Shahejie Formation can be divided into four sub-sections, and the third member of the Shahejie Formation from the Bonan Sub-sag comprises a large number of formations containing high total organic carbon.

Shale pieces with thicknesses from 2 mm to 5 mm were cut from the shale cores. Then, mechanical polishing was performed to keep the upper and lower surface of the shale pieces parallel to each other, and to primarily decrease surface roughness. Finally, the upper surface was subjected to ion polishing to further reduce roughness. The roughness of a 20 μm × 20 μm area in the sample surface is below 100 nm.

### 2.2. Characterization methods

#### 2.2.1. *Vitrinite reflectance test*

Vitrinite reflectance has been universally accepted as an index of coal rank and as a thermal maturation indicator of organic matter (Suárez-Ruiz and Crelling, 2008). The device for the vitrinite reflectance test, the photomicrometer, is a combination of an optical microscope and a spectrophotometer. The optical microscope with an oil immersion objective is utilized for observation and test area selection. The spectrophotometer is used to measure reflectance in a certain area. In addition to the vitrinite reflectance test, a photomicrometer can also assist with optical classification of organic types according to image contrast and reflectance data. The

reflectance test and optical microscope observation, which determine thermal maturity and organic type, constitute the foundation of the following tests.

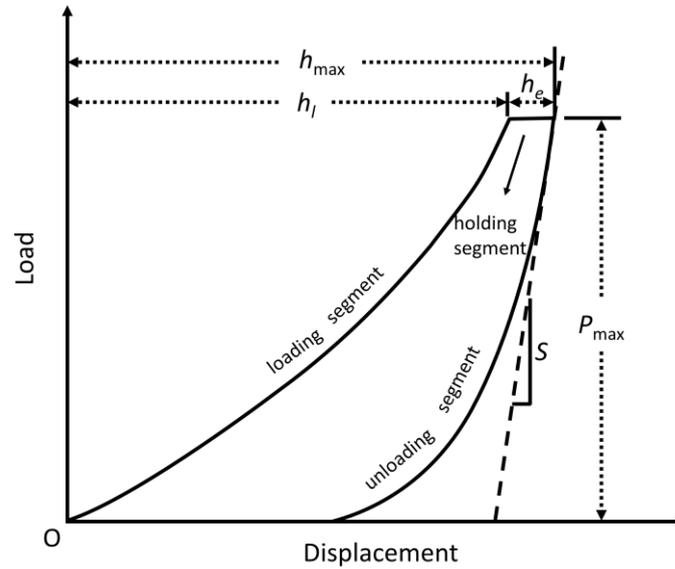
### *2.2.2. Micro-Raman test*

Raman spectrum analysis is widely employed for characterization of carbon materials, such as carbon nanotube and graphene (Dresselhaus et al., 2005; Ferrari et al., 2006; Pimenta et al., 2007). Raman spectrum is highly sensitive to the structure and molecular arrangement of carbon materials. During thermal evolution, the chemical structure of organic matter is gradually altered, and Raman spectrum analysis is a useful tool to characterize the structure change. Numerous researchers have also performed Raman spectrum tests on organic matter in source rocks, and attempted to correlate the spectrum features with the results from conventional vitrinite reflectance tests, to develop novel methods for the determination of thermal maturity (Kelemen and Fang, 2001; Wilkins et al., 2014). Two ways of Raman spectrum analysis exist for organic matter in shale. The first one needs to extract organic matter from shale rocks by mechanical and chemical purification methods. However, this approach necessitates destruction of the shale rock during purification, which makes it unusable for mechanical tests. The second way is via a micro-Raman test. Similar to the photometric test, micro-Raman combines optical observation and Raman spectrum analysis (Lünsdorf, 2016; Tselev et al., 2014), which can locate the micro areas and perform the measurements. Due to its advantages, the second way is chosen for this study.

### *2.2.3. Nanoindentation*

Mainstream commercial nanoindenters are developed on the basis of the method proposed by Oliver and Pharr (Oliver and Pharr, 1992; Oliver and Pharr, 2004). During the nanoindentation measurement, an indenter with a certain geometry penetrates into the sample surface until the targeted force or depth, holds for a set period of time, and then it is pulled out. The load  $P$  on

the sample and the displacement  $h$  of the indenter tip are recorded in the loading, holding, and unloading process (Figure 1).



**Figure 1.** A schematic representation of the load-displacement curve from nanoindentation (modified from Oliver and Pharr (1992)).  $P_{\max}$  is the peak load;  $h_{\max}$  is the maximum indenter displacement;  $h_l$  is the displacement at the end of the loading segment;  $h_e$  is the displacement increase during the holding process; and  $S$  is the initial slope of the unloading segment.

During the loading process, the increasing total displacement contains the elastic displacement and the unrecoverable displacement; whereas, during unloading only, the elastic displacement is gradually recovered. The unloading segment of the  $P$ - $h$  curve reflects the elastic properties of the sample material. According to the contact mechanics (Pharr et al., 1992), the basic model for nanoindentation is as follows:

$$E_r = \frac{\sqrt{\pi}}{2} \frac{S}{\sqrt{A}} \quad (1)$$

where  $E_r$  is the reduced modulus;  $S$  is the slope of the initial unloading segment; and  $A$  is the contact area of the indenter tip on the sample surface, which can be determined by the corrected area function (Oliver and Pharr, 1992):

$$A = 24.5h_c^2 + \sum_{i=0}^7 C_i h_c^{1/2^i} \quad (2)$$

where  $C_i$  are fitting parameters; and  $h_c$  is the contact depth of the indenter, which is related to the maximum displacement  $h_{\max}$  (Oliver and Pharr, 1992):

$$h_c = h_{\max} - \beta \frac{P_{\max}}{S} \quad (3)$$

where  $\beta$  is the parameter corresponding to the geometry of the indenter; and  $P_{\max}$  is the peak load. The reduced modulus is determined by both the sample and the indenter material. The following equation can be used to eliminate the influence from the indenter and calculate the Young's modulus  $E$  of the sample (Johnson, 1987):

$$\frac{1-\gamma^2}{E} = \frac{1}{E_r} - \frac{1-\gamma_{tip}^2}{E_{tip}} \quad (4)$$

where  $\gamma$  is the Poisson's ratio of the sample; and  $E_{tip}$  and  $\gamma_{tip}$  are the Young's modulus and Poisson's ratio of the indenter tip, respectively. For the elastic characterization, organic matter is assumed to be isotropic and linearly elastic.

#### 2.2.4. Experimental procedures

Geochemical test and mechanical characterization are combined in this study (Figure 2). The experimental procedures were as follows:

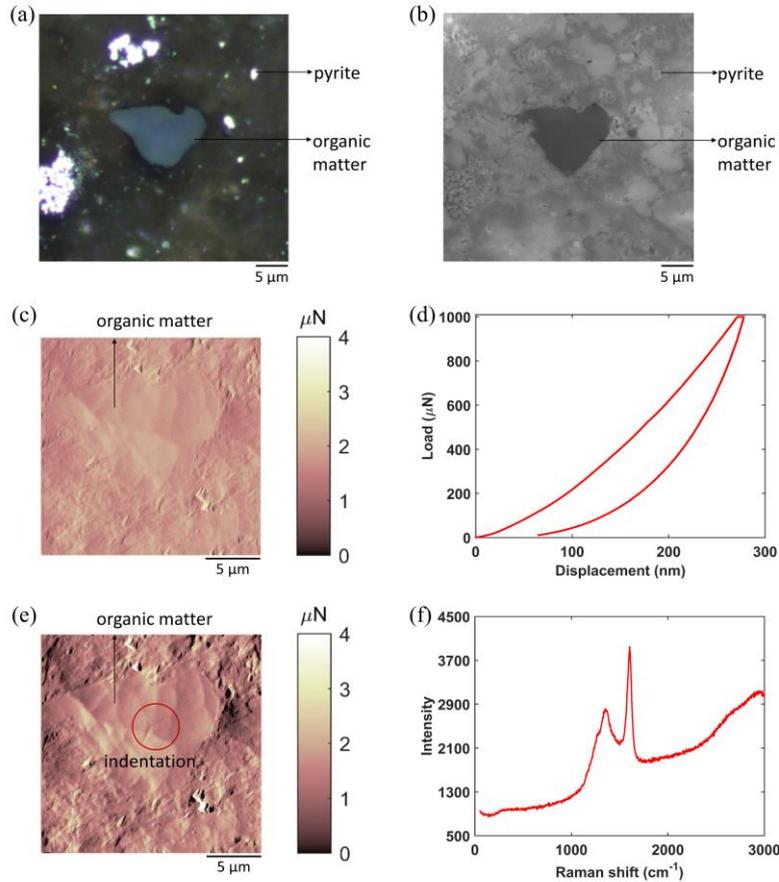
(1) Ion-polished samples were placed into a photomicrometer (QDI 302, CRAIC; DM4500P, Leica). Test areas containing organic matters were observed and selected under an oil immersion objective (Figure 2a). The spectrophotometer was used to measure the reflectance of organic matter. The illumination spot size for reflectance measurement was approximately 3  $\mu\text{m}$ . Organic type was identified through optical image and reflectance data. Vitrinite reflectance constitutes the measurement of the thermal maturity of each sample.

(2) SEM observation (Figure 2b) was then carried out to observe the selected test areas at a low vacuum condition at an accelerating voltage of 15 kV (Quanta 200F, FEI). For convenience of

positioning in the following procedures, the precise coordinate of each test area was recorded, and SEM images of organic matters were captured at various magnifications.

(3) Subsequently, a nanoindenter (Tribo Indenter 950, Hysitron) was utilized for scanning probe microscopy (SPM) and nanoindentation measurement. A Berkovich diamond indenter with a nominal tip radius of 100 nm was chosen for the test. SPM was performed prior to (Figure 2c) and after (Figure 2e) nanoindentation, to determine the indentation position and to observe the residual indentation, respectively. During SPM measurement, the indenter was driven by an oscillating force and scanned the sample at a certain frequency, and  $256 \times 256$  data points of contact force between the indenter tip and the sample surface were obtained. The set point of the driven force was  $2 \mu\text{N}$ , and the dynamic force was  $1 \mu\text{N}$ . Elastic characterization was performed at  $18^\circ\text{C}$ . The indentation was usually placed at the center position of organic matter to avoid interference from surrounding minerals. Under load controlling mode, the maximum load was set to be  $1000 \mu\text{N}$  (Figure 2d), which was uniform for each indentation. The loading, holding, and unloading time were set to be 10 s, 2 s, and 10 s, respectively. The contact area of the indentation was on the order of  $10^6 \text{ nm}^2$  with the experimental setting.

(4) Finally, the micro-Raman test (Figure 2f) was performed to elucidate the structural variation within the organic matter with different thermal maturities (DXRxi, ThermoFisher). The immersion source was a 532 nm laser, and the exposure time was 0.05 s. The spot size of the laser beam was approximately  $1 \mu\text{m}$ . To decrease damage on the sample surface, the laser power was set to be 0.5 mW.



**Figure 2.** Experimental procedures: (a) Vitrinite reflectance test, macerals classification, and test area selection; (b) SEM observation and test area location; (c) SPM (scanning frequency = 0.2 Hz) observation prior to nanoindentation; (d) nanoindentation; (e) SPM (scanning frequency = 0.5 Hz) observation after nanoindentation; and (f) micro-Raman analysis.

### 3. Results and discussion

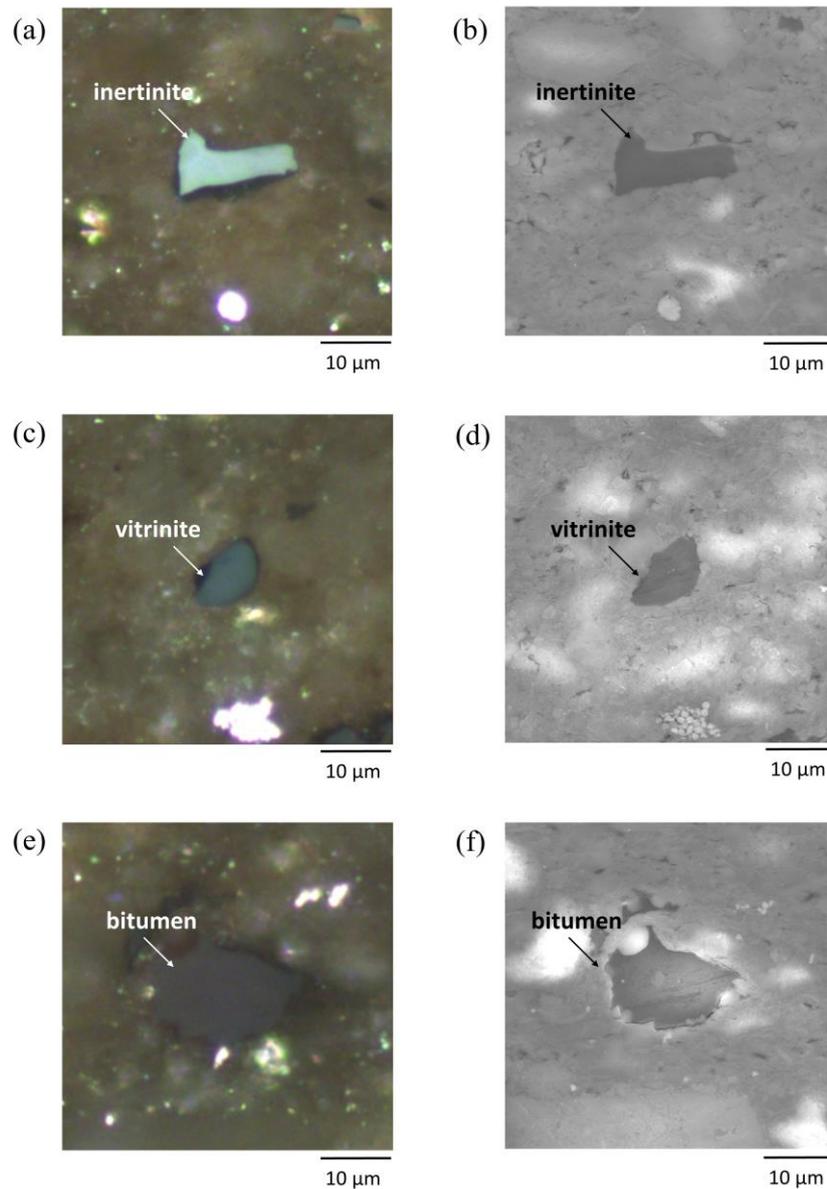
#### 3.1. Organic type and thermal maturity

Table 1 shows the organic type classification and thermal maturity test results. The vitrinite reflectance of shale samples is from 0.62% to 1.13%, which means that all of the samples are in the mature stage. According to the image features under the oil immersion objective, three organic types, including vitrinite, inertinite and bitumen, are identified in the shale samples.

**Table 1.** Vitrinite reflectance and number of areas tested in each sample.

Sample	R <sub>o</sub> (%)	Depth (m)	Number of areas tested		
			Vitrinite	Inertinite	Bitumen
J1	0.62	2966.1	16	11	—
J2	0.69	3049.4	12	9	15
J3	0.94	3005.1	10	11	6
J4	1.13	3101.4	9	12	3

Figure 3 compares the optical images and SEM images of different organic types. Compared with the dry objective, the oil immersion objective can greatly increase the image contrast and the clarity of organic matter. Through the first column of Figure 3, one can see that the image features of the typical test areas are markedly different. Specifically, the organic type in the first test area is inertinite, which is light gray and seems to protrude from the sample surface (Figure 3a). In contrast, vitrinite in the second test area (Figure 3c) and bitumen in the last test area (Figure 3e) are dark gray and brown, respectively. However, it is challenging to differentiate the organic types based only on the image contrast of SEM images (Figure 3b, d, f). In addition to optical images, the reflectance test results can also assist with the classification of organic type. The reflectance of vitrinite is usually higher than the reflectance of bitumen, while lower than that of inertinite. In sample J2, for example, bitumen reflectance is from 0.39% to 0.52%, vitrinite reflectance is from 0.46% to 1.1%, and inertinite reflectance is from 1.1% to 2.2%. A total of 114 test areas that contain different types of organic matter are selected for the nanoindentation test (Table 1).



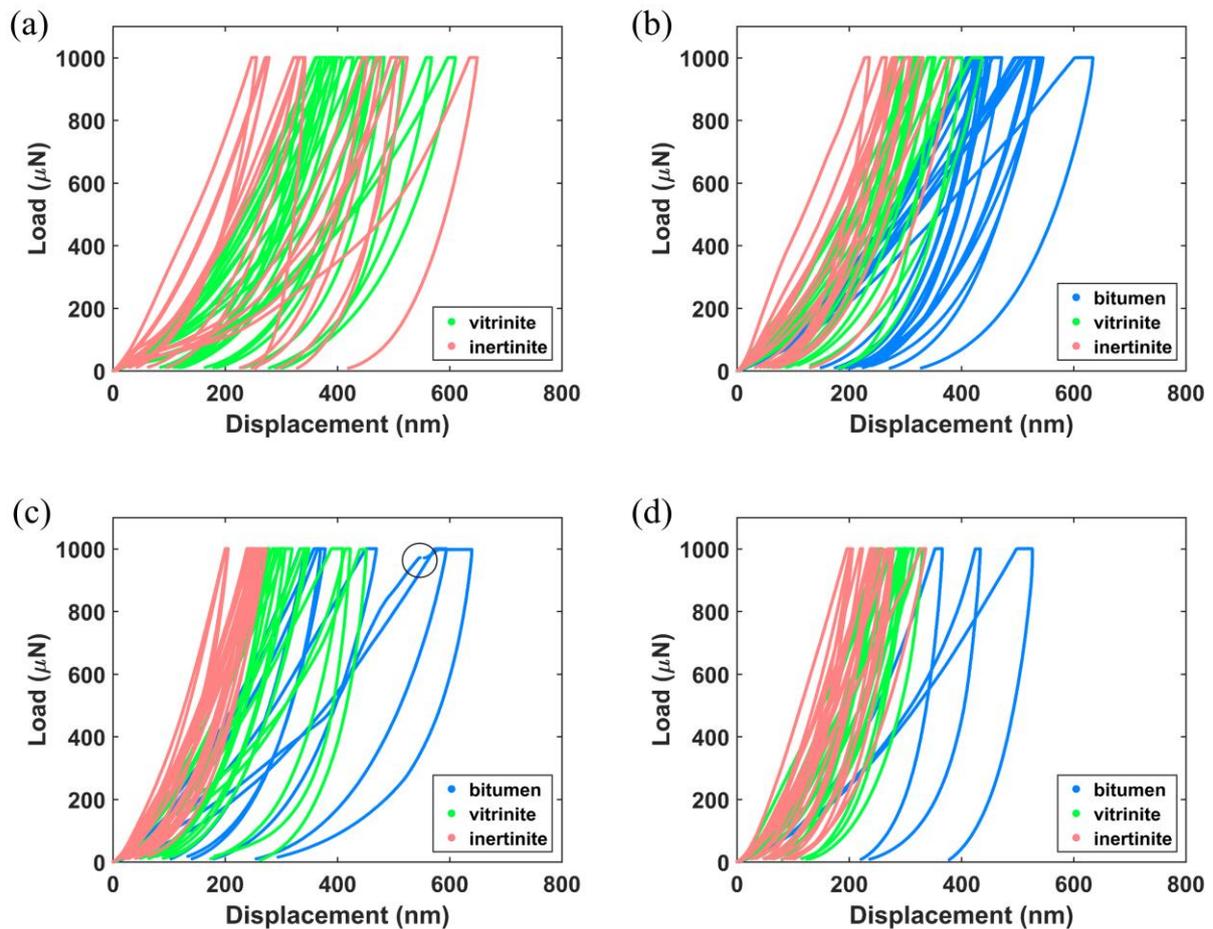
**Figure 3.** Different types of organic matter in sample J3: (a) Optical image and (b) SEM image of inertinite; (c) optical image and (d) SEM image of vitrinite; and (e) optical image and (f) SEM image of bitumen.

### 3.2. Nanoindentation results

The number of areas tested of each kind of organic matter in each sample is listed in Table 1.

Figure 4 assembles the  $P$ - $h$  curves for different kinds of organic matter in each sample. With the same maximum load setting, the maximum displacement of vitrinite is generally lower than

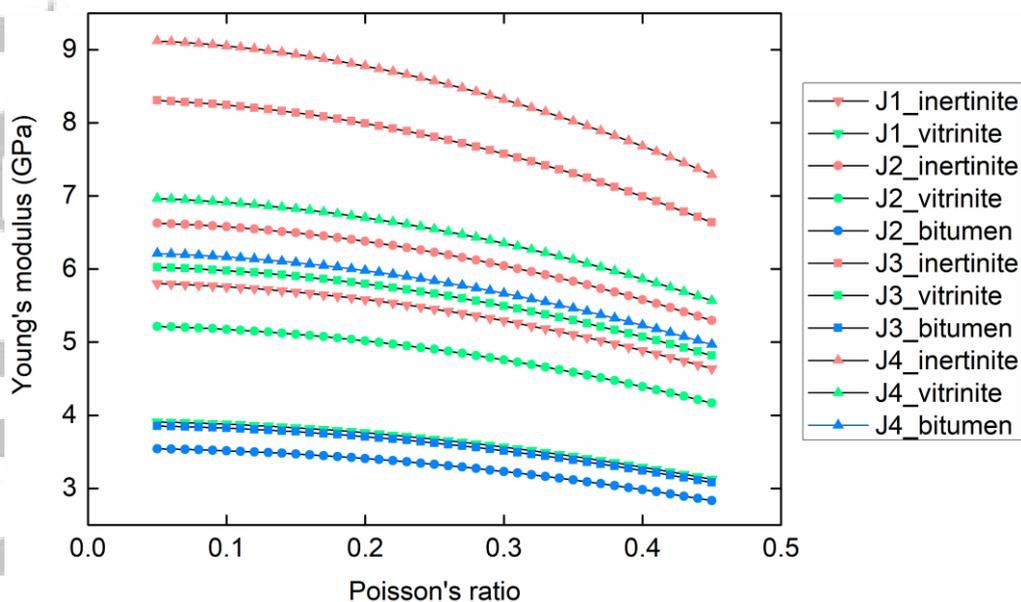
that of inertinite, while higher than that of bitumen. The sudden increase of displacement during the loading process in Figure 4c, which is marked with a black circle, is termed *pop-in*. It suggests the rupture of organic matter. The comparison of the  $P$ - $h$  curves indicates that bitumen is much more compressible than vitrinite and inertinite with the same thermal maturity.



**Figure 4.** Assembly of representative  $P$ - $h$  curves from sample J1 (a), J2 (b), J3 (c), and J4 (d).

The Young's modulus can be calculated using equation (1) to equation (4). The Young's modulus and Poisson's ratio of the diamond indenter tip are 1141 GPa and 0.07 (Field, 1992), respectively. However, to precisely determine the Poisson's ratio of organic matter remains a challenge, and ultrasonic techniques and molecular simulation may be helpful to solve the problem (Rabbani and Schmitt, 2018; Wu and Firoozabadi, 2020). From equation (4), one can know that the estimation of Poisson's ratio influences the calculation results. A wide range of

Poisson's ratio of organic matter, from 0.05 to 0.45, was used in previous micromechanical measurements (Ahmadov et al., 2009; Eliyahu et al., 2015; Emmanuel et al., 2016; Li et al., 2018). Figure 5 illustrates the sensitivity to Poisson's ratio based on the load-displacement data in this work. The calculated Young's modulus of organic matter gradually decreases with the selected Poisson's ratio value. In the following discussions, the Poisson's ratio of organic matter is assumed to be 0.3. It can be found from Figure 5 that the maximum error introduced by the assumption is approximately 12.4%.



**Figure 5.** The sensitivity of calculated Young's modulus to Poisson's ratio of organic matter.

Table 2 summarizes Young's modulus calculation results. It is obvious that inertinite has the highest Young's modulus, while the Young's modulus of bitumen is the lowest. Moreover, the Young's moduli of different organic types all increase with vitrinite reflectance. When vitrinite reflectance increases from 0.62% to 1.13%, the Young's modulus of inertinite increases from 5.29 GPa to 8.32 GPa, by 57%, and the Young's modulus of vitrinite increases from 3.57 GPa to 6.35 GPa, by 78%. From Table 2, it can be concluded that both organic type and thermal maturity affect the mechanical properties of organic matter. Repeated testing on a number of test areas (Table 1) for different organic types in each sample provides a valuable

dataset to examine the variability of the mechanical properties and the reliability of the test results. Except for a couple of test results (i.e., inertinite for sample J1 and bitumen for J3), the standard deviation for each type of testing is moderate (Table 2), indicating either a reliable testing result or moderate variability, or both.

**Table 2.** Young's modulus of different types of organic matter.

Sample	R <sub>o</sub> (%)	Young's modulus (GPa)					
		Vitrinite		Inertinite		Bitumen	
		Mean <sup>a</sup>	90% CI <sup>b</sup>	Mean <sup>a</sup>	90% CI <sup>b</sup>	Mean <sup>a</sup>	90%CI <sup>b</sup>
J1	0.62	3.57±0.83	(3.20, 3.93)	5.29±2.85	(3.73, 6.85)	—	—
J2	0.69	4.76±0.85	(4.32, 5.20)	6.05±1.63	(5.04, 7.06)	3.23±0.80	(2.87, 3.60)
J3	0.94	5.50±0.80	(5.03, 5.96)	7.58±1.38	(6.82, 8.33)	3.52±1.25	(2.49, 4.54)
J4	1.13	6.35±0.97	(5.75, 6.96)	8.32±1.53	(7.53, 9.11)	5.67±0.84	(4.26, 7.08)

<sup>a</sup>± indicates the standard deviation.

<sup>b</sup>CI means confidence interval.

An increase in displacement in the holding segment (Figure 4) indicates the viscoelasticity of organic matter, which was also investigated in previous works (Epshtein et al., 2015; Liu et al., 2019; Rabbani and Schmitt, 2019; Zeszotarski et al., 2004). To approach the problem, indentation creep  $C_{IT}$ , which has been utilized for the characterization of polymers' creep behavior (Chagnon et al., 2013; Koch et al., 2007; Shahdad et al., 2007), is calculated by using the following equation (ISO14577-1, 2002):

$$C_{IT} = \frac{h_e}{h_l} \times 100 \quad (5)$$

where  $h_e$  and  $h_l$  are the displacement increase during the holding and loading process, respectively (please refer to Figure 1). The  $C_{IT}$  values of different types are summarized in Table 3. With the same maximum load and holding time setting, the indentation creep of bitumen is much higher than that of vitrinite and inertinite. The calculation results suggest that

the stiffer organic type demonstrates less creep influence. No clear relation, however, is identified between indentation creep and thermal maturity. While this work focuses on the elastic properties of organic matter, a longer holding time and more suitable test method, for example, dynamic mechanical analysis, would be required for rigorous study of the creep behavior (Chagnon et al., 2013; Koch et al., 2007).

**Table 3.** Indentation creep calculation results.

Sample	R <sub>o</sub> (%)	Indentation creep (%)					
		Vitrinite		Inertinite		Bitumen	
		Mean <sup>a</sup>	90% CI <sup>b</sup>	Mean <sup>a</sup>	90% CI <sup>b</sup>	Mean <sup>a</sup>	90%CI <sup>b</sup>
J1	0.62	2.40±0.31	(2.27, 2.54)	2.21±0.62	(1.87, 2.55)	—	—
J2	0.69	3.09±0.73	(2.71, 3.46)	2.65±0.81	(2.15, 3.15)	4.61±1.11	(4.10, 5.11)
J3	0.94	2.97±1.00	(2.39, 3.55)	2.18±0.51	(1.90, 2.46)	4.66±2.87	(2.30, 7.03)
J4	1.13	2.38±0.75	(1.91, 2.84)	2.18±0.59	(1.88, 2.48)	3.77±1.76	(0.81, 6.73)

<sup>a</sup>± indicates the standard deviation.

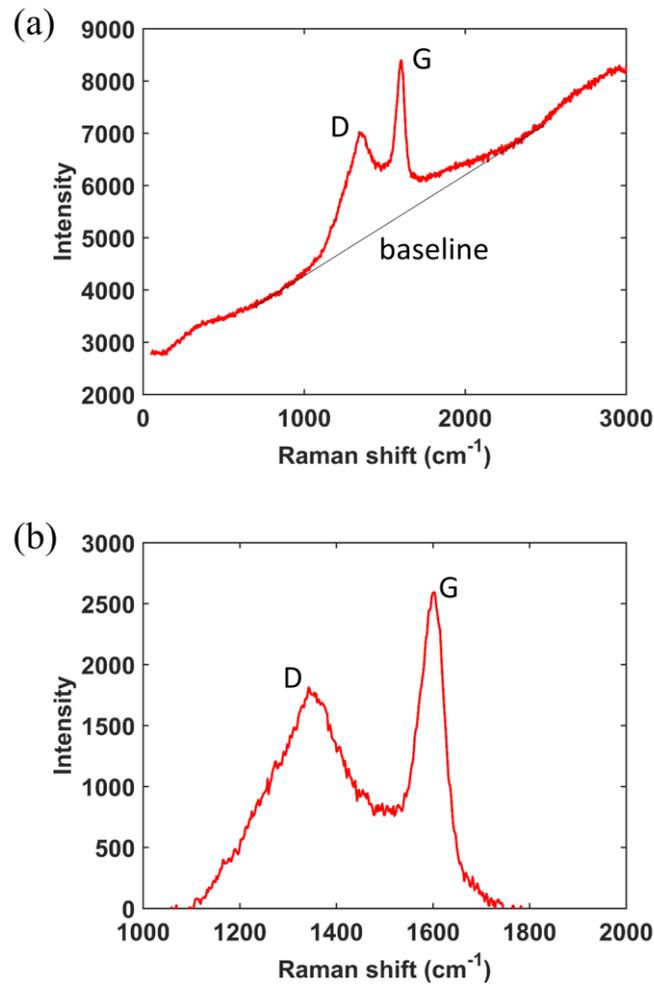
<sup>b</sup>CI means confidence interval.

### 3.3. Micro-Raman analysis

Due to the influence from fluorescence, the Raman spectrum of bitumen and vitrinite has a poor signal-to-noise ratio, and we can only obtain reliable data about inertinite. Through processing of the original spectrum, the micro-Raman test results are analyzed. Figure 6 illustrates a typical Raman spectrum of inertinite. There is a strong fluorescence background in the original Raman data (Figure 6a). Similar to the carbon materials, two characteristic peaks in the spectrum are observed. To remove the background, the baseline is calculated by using the data points around the characteristic peaks. In Figure 6b, the peak around 1350 cm<sup>-1</sup> is the defect peak (D peak), which indicates the defects and the disordered structure in the material.

The peak around 1600 cm<sup>-1</sup> is the graphene peak (G peak), which originates from the plane structure of carbon materials. The intensity ratio of D peak to G peak (IR (D/G)) can be the

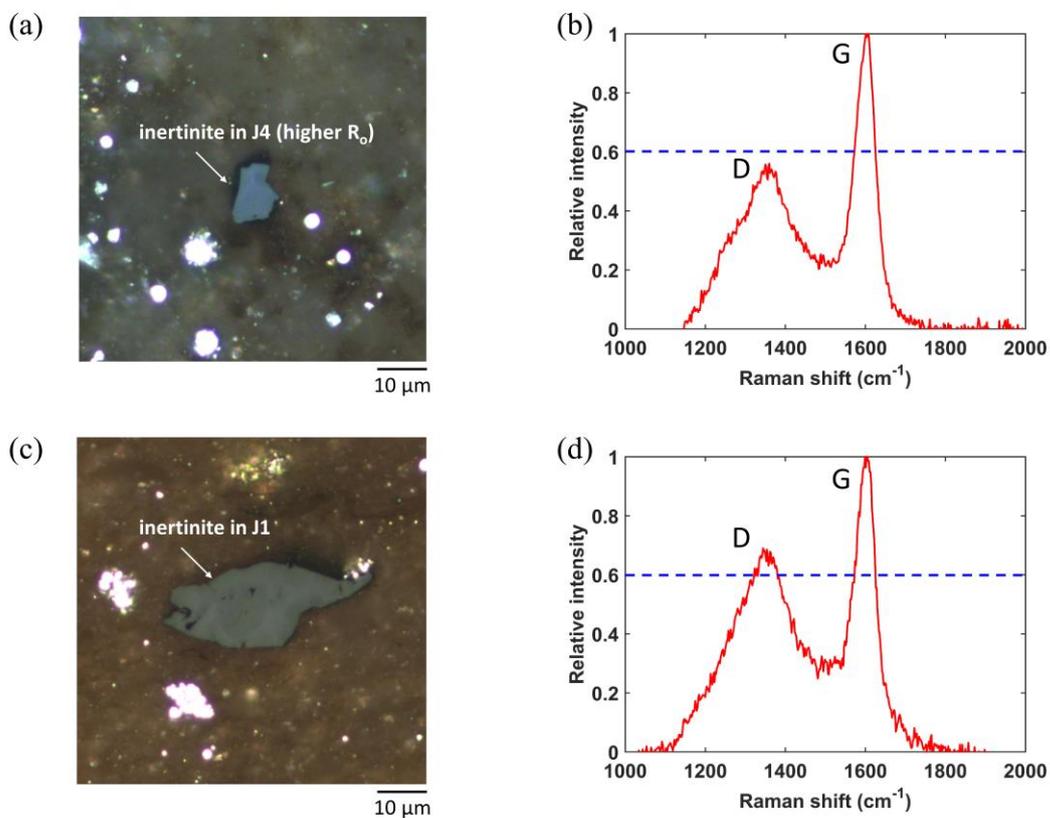
indicator of the degree of graphitization. A higher IR (D/G) indicates fewer defects, a smaller disordered structure, and a higher degree of graphitization.



**Figure 6.** Data processing of Raman spectrum: (a) Original Raman spectrum of inertinite with strong fluorescence background; and (b) processed spectrum in the region of interest. D and G indicate characteristic peaks in the Raman spectrum of carbon material. The baseline is calculated by using the data points around the characteristic peaks. The intensity ratio of D peak to G peak is calculated through the division of maximum intensity value of the D band to that of the G band.

Figure 7 compares two representative Raman spectrums from sample J1 and J4, respectively. The IR (D/G) of the inertinite in sample J1 is higher than that of the inertinite in

sample J4, which suggests that the organic matter in sample J4 has a higher degree of graphitization. The IR (D/G) of each sample is averaged (Table 4) in consideration of the heterogeneity within different inertinites in a certain shale sample. When vitrinite reflectance increases, the IR (D/G) exhibits a general decreasing trend. The chemical structure change results from thermal evolution. With the increase of thermal maturity, the aliphatic structure is gradually decomposed, and the aromaticity of kerogen increases (Craddock et al., 2018; Duan et al., 2018; Tissot and Welte, 2013). Therefore, the calculation results reveal that the organic matter with a higher thermal maturity may contain fewer defects and a more ordered structure.



**Figure 7.** Representative inertinite area and corresponding Raman spectrum from sample J4 (a, b) and J1 (c, d).

**Table 4.** Intensity ratio of D peak to G peak of different samples.

Sample	R <sub>o</sub> (%)	IR (D/G)		Number of areas tested
		Mean <sup>a</sup>	90% CI <sup>b</sup>	
J1	0.62	0.648±0.024	(0.635, 0.661)	11
J2	0.69	0.667±0.022	(0.653, 0.680)	9
J3	0.94	0.620±0.038	(0.599, 0.641)	11
J4	1.13	0.600±0.034	(0.582, 0.617)	12

<sup>a</sup>± indicates the standard deviation.

<sup>b</sup>CI means confidence interval.

### 3.4. Comparison with extant literature

Table 5 presents a comparison of the test results in this study and the reported data. In previous works, utilization of dynamic test techniques is preferred for mechanical characterization, including modulus mapping by nanoindentation system and QNM by AFM. Kerogen and bitumen are distinguished in previous research, but the macerals are not classified. Although Zargari et al. (2016) and Li et al. (2018) both investigate organic matter in Bakken shale, the test results show different trends. Zargari et al. (2016) find the modulus reduction of kerogen from immature to mature stage, while Li et al. (2018) observe that organic matter becomes stiffer as thermal maturity increases. The dissimilar findings may be attributable to the lack of classification of organic type and the limited data volume. The test results provided by Zargari et al. (2016) are storage moduli, which are measured by using modulus mapping with a diamond indenter. Storage modulus is comparable to reduced modulus, and is also influenced by both the sample and the indenter tip (Zhao et al., 2018). The relation between storage modulus and Young's modulus is in the same form as equation (4) (Syed Asif et al., 2001), and the Young's modulus of diamond is much higher than that of organic matter. Therefore, storage modulus and converted Young's modulus should have the same trend from immature to mature stage. The conclusion by Emmanuel et al. (2016) is that the modulus of kerogen significantly increases when vitrinite reflectance increases from 0.40% to 0.82%, while the modulus hardly

changes when vitrinite reflectance increases from 0.82% to 1.25%. In this paper, however, we find that the elastic modulus of vitrinite, inertinite, and bitumen all increase with thermal maturity (Figure 8).

Different from the previous investigations listed in Table 5, this study considers the influence of not only thermal maturity, but also organic type. Vitrinite and inertinite are identified based on optical microscope images and reflectance data. The difference of elastic properties between vitrinite and inertinite are found by nanoindentation measurements. As a consequence, without consideration of maceral classification, conclusions about the influence from thermal maturity may be unreliable, especially when the data volume is limited. In addition, most of the previous studies divide the shale samples into two parts, one of which is for geochemical characterization, while the other one is for the mechanical test. Sample-to-sample variation is also not adequately considered. In this work, a set of geochemical-mechanical experiments is performed on the same selected test areas in a certain sample. Sample-to-sample variation no longer presents a problem, and the experimental data of repeated tests on the same type of test areas provide meaningful statistics. Finally, micromechanical tests on shale are very time consuming, due to the complexity of sample preparation and locating the test area. The number of areas tested in previous works is also relatively small. Here, the data volume is significantly increased, which makes the test results in this work more reliable.

**Table 5.** Comparison with the test results in previous works.

Reference	R <sub>o</sub> (%)	Modulus (GPa)		Test technique
		Kerogen	Bitumen	

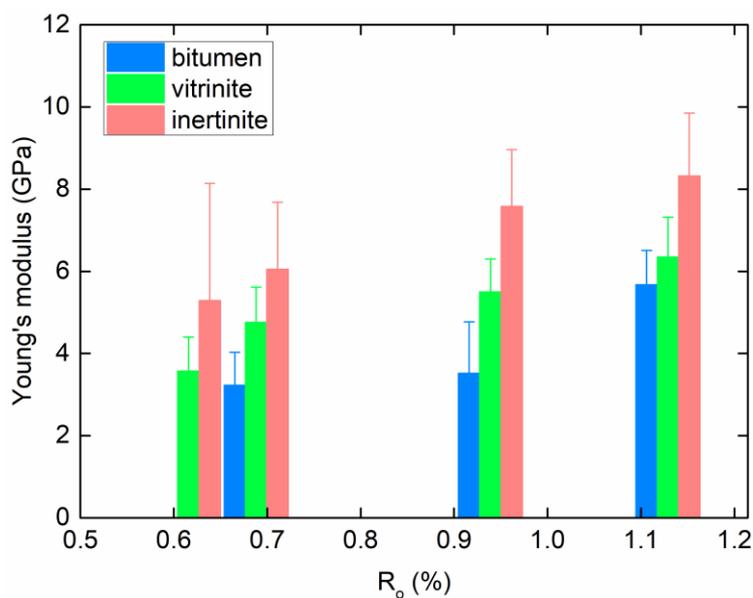
		Vitrinite	Inertinite		Number of areas tested	
Zargari et al. (2016) <sup>a</sup>	Immature	15~20	—	Modulus mapping	12	
	Mature	7~10	—			
	Mature	7~10	—			
	Overmature	7~12	—			
Emmanuel et al. (2016) <sup>b</sup>	0.40	6.10	—	AFM	26	
	0.82	16.0	7.50			
	1.25	15.80	8.50			
Li et al. (2018) <sup>c</sup>	0.35	2.91	—	AFM	3	
	0.64	—	3.33			
	1.04	—	11.77			
This work <sup>d</sup>	0.62	3.57	5.29	Nano-indentation	114	
	0.69	4.76	6.05			3.23
	0.94	5.50	7.58			3.52
	1.13	6.35	8.32			5.67

<sup>a</sup>Modulus in this literature is storage modulus;

<sup>b</sup>Modulus in this literature is Young's modulus;

<sup>c</sup>Modulus in this literature is Young's modulus;

<sup>d</sup>Modulus in this work is Young's modulus.



**Figure 8.** Young's modulus of organic matter with different vitrinite reflectance.

#### 4. Conclusions

In the present paper, the influence of geochemical features on the mechanical properties of organic matter is investigated by combining various geochemical and micromechanical characterization methods. The main findings are as follows:

(1) Inertinite, vitrinite, and bitumen show different image features under the oil immersion objective and have a different reflectance range. It is difficult to distinguish organic type only according to the image contrast of SEM images.

(2) The vitrinite reflectance of the four shale samples is from 0.62% to 1.13%. With the same thermal maturity, mechanical heterogeneity exists within different types of organic matter. Inertinite has the highest Young's modulus, while the modulus of bitumen is the lowest. Taking sample J3 as an example, the Young's modulus of inertinite is more than two times that of the modulus of bitumen.

(3) The Young's moduli of all types of organic matter increase with thermal maturity. When vitrinite reflectance increases from 0.62% to 1.13%, the Young's modulus of inertinite increases from 5.29 GPa to 8.32 GPa, by 57%, and the Young's modulus of vitrinite increases from 3.57 GPa to 6.35 GPa, by 78%. The indentation creep calculation results suggest that the stiffer organic type indicates less creep influence.

(4) Characteristic peaks of carbon material can be found in the Raman spectrum of inertinite. The intensity ratio of D peak to G peak exhibits a decreasing trend with thermal maturity, indicating the increase of graphitization degree of organic matter.

(5) The geochemical and micromechanical characterization results reveal that the mechanical properties of organic matter are influenced by both organic type and thermal maturity. Compared with bitumen, vitrinite and inertinite have more polycyclic aromatic hydrocarbons, fewer saturated aliphatic hydrocarbons, and higher carbon content. Therefore, with the same

thermal maturity, the Young's modulus of vitrinite and inertinite is higher than that of bitumen. With the increase of thermal maturity, the aliphatic structure is gradually decomposed and consumed, and the aromaticity of kerogen increases. Consequently, IR (D/G) decreases, and the Young's modulus increases.

(6) The mechanical behavior of organic matter plays an important role in the predictions of macroscopic mechanical properties at the core scale and even the well scale, especially for organic-rich shales. This can be achieved based on upscaling methods, such as the Mori-Tanaka model and the self-consistent method, with the mechanical characterization results of the main constituents in shale (Abedi et al., 2016; Goodarzi et al., 2017; Zhao et al., 2018). The characterization in this work can serve as a supplement to micro constituent input databases for more reliable overall calculation results. At micro scale, the experimental research can also provide validation for related molecular simulations (Bousige et al., 2016; Wu and Firoozabadi, 2020). In addition, mechanical variation within organic matters has the potential to assist with the identification of organic type and the determination of thermal maturity.

Considering sample-to-sample variation, this work proposes a workflow focusing on fixed micro areas for different characterization. The data volume is also significantly increased compared with that in the extant literature. However, the range of vitrinite reflectance in this study is limited, and all of the four samples are in the mature stage. In the future, shale samples in other thermal maturation stages need to be investigated.

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<http://dx.doi.org/10.17632/hh22sgnnv4.1>.

## References

- Abedi, S., Slim, M., & Ulm, F.-J. (2016). Nanomechanics of organic-rich shales: the role of thermal maturity and organic matter content on texture. *Acta Geotechnica*, *11*(4), 775-787. <https://doi.org/10.1007/s11440-016-0476-2>
- Ahmadov, R., Vanorio, T., & Mavko, G. (2009). Confocal laser scanning and atomic-force microscopy in estimation of elastic properties of the organic-rich Bazhenov Formation. *The Leading Edge*, *28*(1), 18-23. <https://doi.org/10.1190/1.3064141>
- Alstadt, K. N., Katti, K. S., & Katti, D. R. (2015). Nanoscale morphology of kerogen and in situ nanomechanical properties of green river oil shale. *Journal of Nanomechanics and Micromechanics*, *6*(1), 04015003. [https://doi.org/10.1061/\(ASCE\)NM.2153-5477.0000103](https://doi.org/10.1061/(ASCE)NM.2153-5477.0000103)
- Balooch, G., Marshall, G., Marshall, S., Warren, O., Asif, S. S., & Balooch, M. (2004). Evaluation of a new modulus mapping technique to investigate microstructural features of human teeth. *Journal of Biomechanics*, *37*(8), 1223-1232. <https://doi.org/10.1016/j.jbiomech.2003.12.012>
- Bennett, K. C., Berla, L. A., Nix, W. D., & Borja, R. I. (2015). Instrumented nanoindentation and 3D mechanistic modeling of a shale at multiple scales. *Acta Geotechnica*, *10*(1), 1-14. <https://doi.org/10.1007/s11440-014-0363-7>
- Bertrand, R. (1990). Correlations among the reflectances of vitrinite, chitinozoans, graptolites and scolecodonts. *Organic Geochemistry*, *15*(6), 565-574. [https://doi.org/10.1016/0146-6380\(90\)90102-6](https://doi.org/10.1016/0146-6380(90)90102-6)
- Bertrand, R., & Heroux, Y. (1987). Chitinozoan, graptolite, and scolecodont reflectance as an alternative to vitrinite and pyrobitumen reflectance in Ordovician and Silurian strata, Anticosti

Island, Quebec, Canada. *AAPG Bulletin*, 71(8), 951-957. <https://doi.org/10.1306/948878f7-1704-11d7-8645000102c1865d>

Borodich, F. M., Bull, S., & Epshtein, S. (2015). Nanoindentation in studying mechanical properties of heterogeneous materials. *Journal of Mining Science*, 51(3), 470-476. <https://doi.org/10.1134/s1062739115030072>

Bousige, C., Ghimbeu, C. M., Vix-Guterl, C., Pomerantz, A. E., Suleimenova, A., Vaughan, G., Garbarino, G., Feygensohn, M., Wildgruber, C., & Ulm, F.-J. (2016). Realistic molecular model of kerogen's nanostructure. *Nature Materials*, 15(5), 576-582. <https://doi.org/10.1038/nmat4541>

Chagnon, L., Arnold, G., Giljean, S., & Brogly, M. (2013). Elastic recovery and creep properties of waterborne two-component polyurethanes investigated by micro-indentation. *Progress in Organic Coatings*, 76(10), 1337-1345. <https://doi.org/10.1016/j.porgcoat.2013.04.003>

Craddock, P. R., Bake, K. D., & Pomerantz, A. E. (2018). Chemical, molecular, and microstructural evolution of kerogen during thermal maturation: case study from the Woodford Shale of Oklahoma. *Energy & Fuels*, 32(4), 4859-4872. <https://doi.org/10.1021/acs.energyfuels.8b00189>

Cudjoe, S., Vinassa, M., Gomes, J. H. B., & Barati, R. G. (2016). A comprehensive approach to sweet-spot mapping for hydraulic fracturing and CO<sub>2</sub> huff-n-puff injection in Chattanooga shale formation. *Journal of Natural Gas Science and Engineering*, 33, 1201-1218. <https://doi.org/10.1016/j.jngse.2016.03.042>

Dayal, P., Savvides, N., & Hoffman, M. (2009). Characterisation of nanolayered aluminium/palladium thin films using nanoindentation. *Thin Solid Films*, 517(13), 3698-3703. <https://doi.org/10.1016/j.tsf.2009.01.174>

Dokukin, M. E., & Sokolov, I. (2012). Quantitative mapping of the elastic modulus of soft

materials with HarmoniX and PeakForce QNM AFM modes. *Langmuir*, 28(46), 16060-16071.  
<https://doi.org/10.1021/la302706b>

Dresselhaus, M. S., Dresselhaus, G., Saito, R., & Jorio, A. (2005). Raman spectroscopy of carbon nanotubes. *Physics Reports*, 409(2), 47-99.  
<https://doi.org/10.1016/j.physrep.2004.10.006>

Duan, D., Zhang, D., Ma, X., Yang, Y., Ran, Y., & Mao, J. (2018). Chemical and structural characterization of thermally simulated kerogen and its relationship with microporosity. *Marine and Petroleum Geology*, 89, 4-13. <https://doi.org/10.1016/j.marpetgeo.2016.12.016>

Eliyahu, M., Emmanuel, S., Day-Stirrat, R. J., & Macaulay, C. I. (2015). Mechanical properties of organic matter in shales mapped at the nanometer scale. *Marine and Petroleum Geology*, 59, 294-304. <https://doi.org/10.1016/j.marpetgeo.2014.09.007>

Emmanuel, S., Eliyahu, M., Day-Stirrat, R. J., Hofmann, R., & Macaulay, C. I. (2016). Impact of thermal maturation on nano-scale elastic properties of organic matter in shales. *Marine and Petroleum Geology*, 70, 175-184. <https://doi.org/10.1016/j.marpetgeo.2015.12.001>

Epshtein, S. A., Borodich, F. M., & Bull, S. J. (2015). Evaluation of elastic modulus and hardness of highly inhomogeneous materials by nanoindentation. *Applied Physics A*, 119(1), 325-335. <https://doi.org/10.1007/s00339-014-8971-5>

Ferrari, A. C., Meyer, J., Scardaci, V., Casiraghi, C., Lazzeri, M., Mauri, F., Piscanec, S., Jiang, D., Novoselov, K., & Roth, S. (2006). Raman spectrum of graphene and graphene layers. *Physical Review Letters*, 97(18), 187401. <https://doi.org/10.1103/PhysRevLett.97.187401>

Field, J. E. (1992). *The properties of natural and synthetic diamond*. Academic Press, London.

Goodarzi, M., Rouainia, M., Aplin, A., Cubillas, P., & de Block, M. (2017). Predicting the elastic response of organic-rich shale using nanoscale measurements and homogenisation methods. *Geophysical Prospecting*, 65(6), 1597-1614. <https://doi.org/10.1111/1365-2478.12475>

Hunt, J. M. (1979). *Petroleum geochemistry and geology*. W H Freeman and Company, San Francisco.

Hutton, A., Bharati, S., & Robl, T. (1994). Chemical and petrographic classification of kerogen/macerals. *Energy & Fuels*, 8(6), 1478-1488. <https://doi.org/10.1021/ef00048a038>

ISO14577-1 (2002). *Metallic materials—instrumented indentation test for hardness and materials parameters*. International Organization for Standardization, Geneva.

Jacob, H. (1989). Classification, structure, genesis and practical importance of natural solid oil bitumen (“migrabitumen”). *International Journal of Coal Geology*, 11(1), 65-79. [https://doi.org/10.1016/0166-5162\(89\)90113-4](https://doi.org/10.1016/0166-5162(89)90113-4)

Jiu, K., Ding, W., Huang, W., You, S., Zhang, Y., & Zeng, W. (2013). Simulation of paleotectonic stress fields within Paleogene shale reservoirs and prediction of favorable zones for fracture development within the Zhanhua Depression, Bohai Bay Basin, east China. *Journal of Petroleum Science and Engineering*, 110, 119-131. <https://doi.org/10.1016/j.petrol.2013.09.002>

Johnson, K. L. (1987). *Contact mechanics*. Cambridge University Press.

Kelemen, S., & Fang, H. (2001). Maturity trends in Raman spectra from kerogen and coal. *Energy & Fuels*, 15(3), 653-658. <https://doi.org/10.1021/ef0002039>

Koch, T., Kogler, F. R., Schubert, U., & Seidler, S. (2007). Mechanical properties of organic-inorganic hybrid materials determined by indentation techniques. *Monatshefte für Chemie-Chemical Monthly*, 138(4), 293-299. <https://doi.org/10.1007/s00706-007-0612-7>

Kumar, V., Sondergeld, C. H., & Rai, C. S. (2012). *Nano to macro mechanical characterization of shale*. Paper presented at SPE Annual Technical Conference and Exhibition, Society of Petroleum Engineers. <https://doi.org/10.2118/159804-ms>

Li, C., Ostadhassan, M., Gentzis, T., Kong, L., Carvajal-Ortiz, H., & Bubach, B. (2018). Nanomechanical characterization of organic matter in the Bakken formation by microscopy-

based method. *Marine and Petroleum Geology*, 96, 128-138.  
<https://doi.org/10.1016/j.marpetgeo.2018.05.019>

Liu, Y., Xiong, Y., Liu, K., Yang, C., & Peng, P. (2019). Indentation size and loading rate sensitivities on mechanical properties and creep behavior of solid bitumen. *International Journal of Coal Geology*, 216, 103295. <https://doi.org/10.1016/j.coal.2019.103295>

Lünsdorf, N. K. (2016). Raman spectroscopy of dispersed vitrinite—Methodical aspects and correlation with reflectance. *International Journal of Coal Geology*, 153, 75-86.  
<https://doi.org/10.1016/j.coal.2015.11.010>

Naizhen, L., & Guoyong, W. (2016). Shale gas sweet spot identification and precise geo-steering drilling in Weiyuan Block of Sichuan Basin, SW China. *Petroleum Exploration and Development*, 43(6), 1067-1075.

Oliver, W. C., & Pharr, G. M. (1992). An improved technique for determining hardness and elastic modulus using load and displacement sensing indentation experiments. *Journal of Materials Research*, 7(06), 1564-1583. <https://doi.org/10.1557/jmr.1992.1564>

Oliver, W. C., & Pharr, G. M. (2004). Measurement of hardness and elastic modulus by instrumented indentation: Advances in understanding and refinements to methodology. *Journal of Materials Research*, 19(1), 3-20. <https://doi.org/10.1557/jmr.2004.19.1.3>

Pharr, G., Oliver, W., & Brotzen, F. (1992). On the generality of the relationship among contact stiffness, contact area, and elastic modulus during indentation. *Journal of Materials Research*, 7(3), 613-617. <https://doi.org/10.1557/JMR.1992.0613>

Pimenta, M., Dresselhaus, G., Dresselhaus, M. S., Cancado, L., Jorio, A., & Saito, R. (2007). Studying disorder in graphite-based systems by Raman spectroscopy. *Physical Chemistry Chemical Physics*, 9(11), 1276-1290. <http://dx.doi.org/10.1039/B613962K>

Rabbani, A., & Schmitt, D. R. (2018). Ultrasonic shear wave reflectometry applied to the determination of the shear moduli and viscosity of a viscoelastic bitumen. *Fuel*, 232, 506-518.

<https://doi.org/10.1016/j.fuel.2018.05.175>

Rabbani, A., & Schmitt, D. R. (2019). The longitudinal modulus of bitumen: Pressure and temperature dependencies. *Geophysics*, 84(4), MR139-MR151.

<https://doi.org/10.1190/geo2018-0344.1>

Rybacki, E., Meier, T., & Dresen, G. (2016). What controls the mechanical properties of shale rocks?—Part II: Brittleness. *Journal of Petroleum Science and Engineering*, 144, 39-58.

<https://doi.org/10.1016/j.petrol.2016.02.022>

Shahdad, S. A., McCabe, J. F., Bull, S., Rusby, S., & Wassell, R. W. (2007). Hardness measured with traditional Vickers and Martens hardness methods. *Dental Materials*, 23(9), 1079-1085.

<https://doi.org/10.1016/j.dental.2006.10.001>

Stach, E., Murchison, D., Taylor, G. H., & Zierke, F. (1982). *Stach's textbook of coal petrology*. Borntraeger Berlin.

Suárez-Ruiz, I., & Crelling, J. C. (2008). *Applied coal petrology: the role of petrology in coal utilization*. Academic Press.

Syed Asif, S., Wahl, K., Colton, R., & Warren, O. (2001). Quantitative imaging of nanoscale mechanical properties using hybrid nanoindentation and force modulation. *Journal of Applied Physics*, 90(3), 1192-1200. <https://doi.org/10.1063/1.1380218>

Taylor, G. H., Teichmüller, M., Davis, A., Diessel, C., Littke, R., & Robert, P. (1998). *Organic petrology*. Gebrüder Borntraeger, Berlin.

Tissot, B. P., & Welte, D. H. (2013). *Petroleum formation and occurrence*. Springer Science & Business Media.

Tong, J., Jiang, X., Han, X., & Wang, X. (2016). Evaluation of the macromolecular structure of Huadian oil shale kerogen using molecular modeling. *Fuel*, 181, 330-339.

<https://doi.org/10.1016/j.fuel.2016.04.139>

Tselev, A., Ivanov, I. N., Lavrik, N. V., Belianinov, A., Jesse, S., Mathews, J. P., Mitchell, G.

D., & Kalinin, S. V. (2014). Mapping internal structure of coal by confocal micro-Raman spectroscopy and scanning microwave microscopy. *Fuel*, 126, 32-37.

<https://doi.org/10.1016/j.fuel.2014.02.029>

Ungerer, P., Collell, J., & Yiannourakou, M. (2014). Molecular modeling of the volumetric and thermodynamic properties of kerogen: Influence of organic type and maturity. *Energy & Fuels*, 29(1), 91-105. <https://doi.org/10.1021/ef502154k>

Vranjes, S., Misch, D., Schöberl, T., Kiener, D., Gross, D., & Sachsenhofer, R. F. (2018). Nanoindentation study of macerals in coals from the Ukrainian Donets Basin. *Advances in Geosciences*, 45. <https://doi.org/10.5194/adgeo-45-73-2018>

Wang, G., & Carr, T. R. (2012). Methodology of organic-rich shale lithofacies identification and prediction: A case study from Marcellus Shale in the Appalachian basin. *Computers & Geosciences*, 49, 151-163. <https://doi.org/10.1016/j.cageo.2012.07.011>

Wang, M., Sherwood, N., Li, Z., Lu, S., Wang, W., Huang, A., Peng, J., & Lu, K. (2015). Shale oil occurring between salt intervals in the Dongpu Depression, Bohai Bay Basin, China. *International Journal of Coal Geology*, 152, 100-112. <https://doi.org/10.1016/j.coal.2015.07.004>

Wilkins, R. W., Boudou, R., Sherwood, N., & Xiao, X. (2014). Thermal maturity evaluation from inertinites by Raman spectroscopy: the 'RaMM' technique. *International Journal of Coal Geology*, 128, 143-152. <https://doi.org/10.1016/j.coal.2014.03.006>

Wilkinson, T. M., Zargari, S., Prasad, M., & Packard, C. E. (2015). Optimizing nano-dynamic mechanical analysis for high-resolution, elastic modulus mapping in organic-rich shales. *Journal of Materials Science*, 50(3), 1041-1049. <https://doi.org/10.1007/s10853-014-8682-5>

Wu, T., & Firoozabadi, A. (2020). Mechanical properties and failure envelope of kerogen matrix by molecular dynamics simulations. *The Journal of Physical Chemistry C*, 124, 2289–2294. <https://doi.org/10.1021/acs.jpcc.9b09639>

Yang, J., Hatcherian, J., Hackley, P. C., & Pomerantz, A. E. (2017). Nanoscale geochemical and geomechanical characterization of organic matter in shale. *Nature Communications*, 8(1), 2179. <https://doi.org/10.1038/s41467-017-02254-0>

Zargari, S., Prasad, M., Mba, K. C., & Mattson, E. D. (2013). Organic maturity, elastic properties, and textural characteristics of self resourcing reservoirs. *Geophysics*, 78(4), D223-D235.

Zargari, S., Wilkinson, T. M., Packard, C. E., & Prasad, M. (2016). Effect of thermal maturity on elastic properties of kerogen. *Geophysics*, 81(2), M1-M6. <https://doi.org/10.1190/geo2015-0194.1>

Zeszotarski, J. C., Chromik, R. R., Vinci, R. P., Messmer, M. C., Michels, R., & Larsen, J. W. (2004). Imaging and mechanical property measurements of kerogen via nanoindentation. *Geochimica et Cosmochimica Acta*, 68(20), 4113-4119. <https://doi.org/10.1016/j.gca.2003.11.031>

Zhang, Y., Lebedev, M., Smith, G., Jing, Y., Busch, A., & Iglauer, S. (2019). Nano-mechanical properties and pore-scale characterization of different rank coals. *Natural Resources Research*, 1-14. <https://doi.org/10.1007/s11053-019-09572-8>

Zhao, J., & Zhang, D. (2020). Dynamic microscale crack propagation in shale. *Engineering Fracture Mechanics*, 228, 106906. <https://doi.org/10.1016/j.engfracmech.2020.106906>

Zhao, J., Zhang, D., Wu, T., Tang, H., Xuan, Q., Jiang, Z., & Dai, C. (2018). Multiscale approach for mechanical characterization of organic-rich shale and its application. *International Journal of Geomechanics*, 19(1), 04018180. [https://doi.org/10.1061/\(asce\)gm.1943-5622.0001281](https://doi.org/10.1061/(asce)gm.1943-5622.0001281)