

# The Hydrocarbon Potential of Carboniferous Reservoirs in the Jimsar Sag, Northwest China: Implications for a Giant Volcanic-Petroleum Reserves

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Gong D, Song Y, Peng M, Liu C, Wang R and Wu W (2022) The Hydrocarbon Potential of Carboniferous Reservoirs in the Jimsar Sag, Northwest China: Implications for a Giant Volcanic-Petroleum Reserves. Front. Earth Sci. 10:879712. doi: 10.3389/feart.2022.879712 As an unconventional petroleum reservoir, the volcanic reservoir is essential for finding largescale oil and gas reserves in deep sedimentary basins. Based on basin modeling, organic geochemistry, and organic petrology, this study evaluates the exploration potential of the Carboniferous volcanic-petroleum system in the Jimsar Sag of the southeastern Junggar Basin. The Carboniferous source rocks in the study area were developed in the Lower Carboniferous Songkharsu Formation, a set of marine-terrigenous transitional source rocks, lithologically composed of carbonaceous mudstone, mudstone, and coal. The kerogen is characterized by type II<sub>2</sub>-III, indicating a gas-prone source rock. Carbonaceous mudstones, mudstone, and coal are classified as medium-good, medium, and poor source rocks, respectively. The hydrocarbon-generating potential is close to that of the Kelameili gas field. The Carboniferous-reservoir oil in well J15 is characterized by <sup>13</sup>C-enriched stable carbon isotopes, relatively high  $\Sigma C_{31-35}/C_{30}$  and Pr/Ph ratios, relatively low density, gammacerane index, regular C<sub>29</sub>-sterane content, and undetected β-carotene, indicating a Songkharsu origin. The Carboniferous-reservoired gas from well J3301 has stable carbon isotope composition enriched in <sup>13</sup>C, similar to the gas from the Kelameili gas field, and should be from the Songkharsu source rock. This set of source rocks is widely distributed in the Jimsar Sag with considerable thickness, among which thickness >100 m reaches 580 km<sup>2</sup>, accounting for 38.7% of the whole sag. The Songkharsu source rocks entered the main oilgenerating window at the end of the Early Jurassic. The area of Songkharsu source rocks in the Jimsar Sag with gas-generating intensity greater than  $20 \times 10^8 \text{ m}^3/\text{km}^2$  reaches 1,015 km<sup>2</sup>, whereas that with oil-generating intensity greater than 500  $\times$  10<sup>4</sup> t/km<sup>2</sup> reaches 1,146 km<sup>2</sup>. However, 13 Carboniferous volcanic lithological traps were found in the sag, covering an area of 230 km<sup>2</sup>. The Carboniferous volcanic-petroleum system in the Jimsar Sag has the resource potential to form large and medium oil and gas fields, which could become a critical replacement field for volcanic oil and gas exploration in the Junggar Basin after the Klameili gas field.

Keywords: Junggar basin, Jimsar Sag, Carboniferous, volcanic rocks, hydrocarbon-generating potential, oil and gas sources, exploration domain

# **1 INTRODUCTION**

Volcanic reservoirs are unconventional petroleum reservoirs whose physical properties are not constrained by depth (Zou et al., 2008) and are important for finding large-scale oil and gas reserves in deep sedimentary basins. Since the first discovery of volcanic reservoirs in the San Joaquin Basin, California, United States, in 1887 (Petford and Mccaffrey, 2003), there has been a history of volcanic oil and gas exploration for more than 130 years. Volcanic reservoirs, such as Scott Reef (Australia), Jatibarang (Indonesia), Kudu (Namibia), Lake Kivu (Congo), Ben Khalala (Algeria), Samgori (Georgia), Urucu (Brazil), Yaraktin (Russia), and Ragusa (Italy) have been discovered globally (Petford and Mccaffrey, 2003).

Volcanic rocks are in and around sedimentary basins in China, and the exploration of volcanic reservoirs has been conducted for more than 60 years. Currently, China has made a series of significant breakthroughs in oil and gas exploration for volcanic reservoirs in the Junggar, Songliao, Erlian, Tarim, Santanghu, and Sichuan basins; volcanic oil and gas resources have been developed (Zou et al., 2008; Dai et al., 2021). Among them, the Junggar Basin has the highest degree of exploration for volcanic-petroleum reservoirs, and several large and medium volcanic reservoirs have been discovered in the northwest margin (Cao et al., 2010; Chang et al., 2019), hinterland (Chen et al., 2017; Zhi et al., 2022), and east (Yu et al., 2014; Sun et al., 2016; Guo et al., 2020) of the basin. In 2006, the Kelameili gas field was discovered in the northeastern part of the basin (Figure 1A), with proven reserves of  $1,033 \times 10^8 \text{ m}^3$ , the largest volcanic gas reservoir discovered in China (Gong et al., 2019,2021).

The Jimsar Sag is approximately 120 km southeast of the Kelameili gas field (Figure 1A). Its shale reservoir in the Permian Lucaogou Formation (P<sub>2</sub>l) has a billion-ton resource scale, which has become an important shale oil production base in China (Bai et al., 2017; Hu et al., 2018). The Kelameili gas field and the Jimsar Sag share a similar geotectonic setting during the Carboniferous; they are both parts of the remnant ocean basin formed after the closure of the Paleo-Asian Ocean (Carroll et al., 1995; Xiao W. J. et al., 2008). In 2001, well J15 in the Jimsar Sag obtained an industrial oil flow of 5.1 t/d in the Carboniferous system. In 2019, well J3301 obtained an excellent oil and gas in the Carboniferous system (Figure 1B). In addition, wells JT1 and J20 drilled in the Carboniferous encountered huge thick volcanic reservoirs (Figure 1B). The information shows that the Carboniferous petroleum system of this sag has excellent exploration prospects.

However, in the past 20 years, efforts have focused on exploring and developing  $P_2$ l shale oil in this area. Thus, the source rock quality, oil/gas source, and hydrocarbon accumulation conditions of the Carboniferous petroleum system have not been studied. By analyzing organic geochemistry, organic petrology, and basin modeling, this study evaluates the exploration potential of the Carboniferous volcanic reservoirs in the Jimsar Sag and reveals a potential giant volcanic-petroleum system. The research results have important implications for domestic and international exploration of volcanic reservoirs.

# 2 GEOLOGICAL BACKGROUND

The Junggar Basin, located in northwest China (**Figure 1A**), is an intraland superimposed basin that developed from the Early Carboniferous to the Quaternary (Xiao W. J. et al., 2008). It is at the intersection of Kazakhstan, Siberian, and Tarim Paleoplates (Xiao W. J. et al., 2008; He et al., 2018). Based on the Permian tectonic pattern, the basin can be divided into six first-order tectonic units (Central Depression, Wulungu Depression, Luliang Uplift, Western Uplift, Eastern Uplift, and North Tianshan Piedmont Thrust Belt). These first-order tectonic units can be further subdivided into 44 second-order tectonic units (**Figure 1A**) (He et al., 2018).

The Jimsar Sag is a "dustpan-shaped" sag developed on a Carboniferous folded basement. It is a second-order tectonic unit of the Eastern Uplift covering approximately 1,500 km<sup>2</sup> (Figures 1A,B) (Bai et al., 2017). During the Late Carboniferous, the Shaqi Salient to the north and the Gusi Salient to the east of the sag were uplifted under the influence of the Hercynian movement (Xiao W. J. et al., 2008; He et al., 2018). The water in the Jimsar Sag was connected to that of the Bogda Mountain Front Sag and the Fukang Sag simultaneously (Li et al., 2015a,b; Zhang L. et al., 2020). During the early Middle Permian, the study area subsided dramatically and became a relatively independent sedimentary unit (Li et al., 2015a,b; Zhang L. et al., 2020). At the end of the Triassic, the Shaqi Salient was strongly uplifted by the Indochinese movement, and the Permian and Triassic strata of the east slope of the Jimsar Sag suffered different denudation (Novikov, 2013; Li et al., 2015a,b; Zhang L. et al., 2020). During the Late Jurassic-Cretaceous, the southeast corner of the sag uplifted, and the study area denudated under the influence of Acts II and III of the Yanshan Movement (He et al., 2013; Novikov, 2013). During the Himalayan period, the sag was uplifted from east to west, and the strata were thinned to the east, forming the present-day tectonic pattern (He et al., 2013; Li et al., 2015a,b).

The Jimsar Sag is developed with Carboniferous-Quaternary strata from the bottom up, with five regional unconformities (i.e., top Carboniferous, top Middle Permian, top Triassic, top Jurassic, and top Cretaceous) (Figure 1C). P<sub>2</sub>l is the most crucial source rock formation in the study area (Carroll, 1998; Wang et al., 2013). It was formed in a saline lacustrine environment after the closure of the residual sea, with high organic matter abundance, and is an excellent oil-prone source rock (Bai et al., 2017; Hu et al., 2018). The Carboniferous in the study area contains the Lower Carboniferous Songkharsu Formation  $(C_1s)$  and Upper Carboniferous Bashan Formation  $(C_2b)$  (Du, 2010; Yu et al., 2014) (Figure 1C). These two sets of strata are coastal-offshore deposits, comprising volcanic clastic rocks with a maximum cumulative thickness of more than 4 km (Du, 2010; Yu et al., 2014). Medium and acidic volcanic lavas and tuffaceous volcanic clastic rocks dominate the lower section of the Songkharsu Formation (C<sub>1</sub>s<sup>a</sup>) (Du, 2010; Gong et al., 2019). The upper section  $(C_1s^b)$  is a clastic rock deposited during the lull volcanic activity, locally interbedded with thin coal seams (Figure 1C), which is the primary source-rock-bearing section of the Carboniferous system in the eastern Junggar Basin (Du, 2010; Gong et al., 2019). C<sub>2</sub>b is dominated by medium-basic volcanic



lava interspersed with tuffaceous volcanic clastic rocks (Figure 1C).

Four sets of the reservoir-caprock assemblages are developed in the Jimsar Sag: 1) a reservoir-caprock with volcanic breccia and volcanic clastic rocks at the top of the Carboniferous as reservoirs and mudstones in the lower part of the Middle Permian Jiangjunmiao Formation  $(P_2j)$  as caprocks; 2) a reservoir–caprock assemblage with sandstones in the lower part of the  $P_2l$  and Upper

Well	Formation	Depth (m)	Lithology	Whole Ro	ock Comp	onents (%)	Organic Maceral Content (%)				
				Total Organic Matter	Pyrite	Other Minerals	Sapropelite	Exinite	Vitrinite	Inertinite	
J15	C <sub>1</sub> s <sup>b</sup>	2,860	Coal	60.7	1.5	37.8	4.3	12.6	54.9	28.2	
J3301 J3301	C <sub>1</sub> s <sup>b</sup> C <sub>1</sub> s <sup>b</sup>	4,135 4,441	Carbonaceous mudstone Mudstone	15.3 2.1	1.3 2.9	83.4 92.1	1.2 4.3	3.5 18.0	88.9 71.5	6.4 6.2	

TABLE 1 | The maceral characteristics of C<sub>1</sub>s<sup>b</sup> source rocks in the Jimsar Basin.

Permian Shangwuerhe Formation ( $P_3w$ ) as reservoirs and mudstones in the middle and upper part of  $P_3w$  as caprocks; 3) a reservoir–caprock assemblage with sandstones in the Lower Triassic Jiucaiyuanzi ( $T_1j$ ) and Shaofanggou ( $T_1s$ ) Formations as reservoirs and thick mudstones developed in their upper part as caprocks; 4) reservoir–caprock assemblages formed by the sand and gravels developed at the bottom of Jurassic groups as reservoirs and mudstones developed in the interior or middle and upper parts as caprocks (Bai et al., 2017; Hu et al., 2018; Zhang S. et al., 2020) (**Figure 1C**).

# 3 SAMPLES AND EXPERIMENTAL METHODS

### 3.1 Sampling

In this study, 137 source rock samples (21 from Jimsar Sag and 116 from Kelameili gas field), 36 oil samples (1 from Carboniferous reservoir and 35 from  $P_2$ l reservoir in the Jimsar Sag), and 21 gas samples (1 from Jimsar Sag and 20 from Kelameili gas field) were discussed, with part of the well locations shown in **Figure 1B**. Natural gas was collected at the wellheads and stored in 1-L gas-tight cylinders. Oil and condensates were collected at the wellheads under ambient pressure and stored in 5-ml glass containers with screw-on Teflon-lined caps. The samples were refrigerated to below  $-6^{\circ}C$ .

### **3.2 Analytical Processes**

# 3.2.1 Total Organic Carbon (TOC) and Rock-Eval Analysis

The analyses were conducted at the China University of Petroleum (Beijing). First, the 146 rock samples were crushed to powder for TOC analysis (**Table 1**). In addition, the powdered samples were split into 200-mg subsamples and treated using HCl at  $60^{\circ}$ C to remove the carbonates and then washed using distilled water to remove the HCl. Finally, the washed subsamples were dried overnight at  $50^{\circ}$ C, and their carbon contents were determined using LECO CS–230 analyzers.

For rock-eval pyrolysis, 100 g of each crushed rock sample was placed in the vessel of an OGE-II instrument. These samples were heated from 300 to 600°C in a helium atmosphere at a heating rate of 50°C/min, and their Rock-Eval parameters ( $S_1$ ,  $S_2$ , and  $T_{max}$ ) were measured. In this study,  $S_1$  is the amount of free hydrocarbon that can be volatilized from the rock sample (mg HC/g rock);  $S_2$  is the amount of hydrocarbon produced by the cracking of organic matter (mg HC/g rock).  $T_{max}$  (°C) is the

temperature at which the  $S_2$  yield is maximized, which estimates the thermal maturity of the sediment (Peters, 1986).

# 3.2.1 Organic Petrology Analysis and Measurement of Vitrinite Reflectance

Optical microscopy analyses were conducted on thin rock sections at the China University of Geoscience (Beijing) (**Table 2**). Before being embedded in a homogeneous mixture of Buehler's epoxy resin and hardener (ratio 5:1), 11 core samples from eight wells were sectioned perpendicular to the bedding. The preparations involve drying and polishing, as described by Taylor et al. (1998) and Amijaya and Littke (2006). The thin rock sections were examined at different magnifications and under different light conditions (incident white- and blue-light excitation) to characterize the organic matter features of  $C_{15}^{b}$  source rock. The optical instrument was a Nikon LV 100 microscope.

The vitrinite reflectances ( $R_o$ ) of 46 samples were measured at the China University of Geoscience (Beijing) (**Table 1**), using a Zeiss Scope A1 incident light microscope at a wavelength ( $\lambda$ ) of 546 nm. The reflectances of samples, rich in vitrinite or solid bitumen particles, were measured at least 50 times.

#### 3.2.2 GC and GC-MS Analysis

Mass spectrometry (MS) and gas chromatography MS (GC–MS) analyses were performed at the China University of Petroleum (Beijing). Soxhlet apparatus was used to extract 30 samples using  $CHCl_3$  for 72 h. The resulting extracts were fractionated using open silica gel column chromatography with *n*-hexane. The resulting saturated hydrocarbons were analyzed using GC and GC–MS.

The GC analysis was performed in an Agilent 7890A gas chromatograph fitted with a 60 m  $\times$  0.25 mm  $\times$  0.25 µm capillary column with nitrogen (99.999%) as the carrier gas. The GC oven temperature was at 40°C for 10 min, and then, it ramped from 40 to 70°C at 4°C/min and 300°C at 8°C/min, and finally held at 300°C for 40 min.

The GC-MS analysis was performed in an Agilent 7890–5975C using the same column type as in the GC analysis but with helium (99.999%) as the carrier gas. During the GC-MS analysis, the GC oven temperature was held at 50°C for 1 min before it ramped to  $120^{\circ}$ C at  $20^{\circ}$ C/min, from 120 to  $250^{\circ}$ C at  $4^{\circ}$ C/min, and from 250 to  $310^{\circ}$ C at  $3^{\circ}$ C/min. Finally, it was held at  $310^{\circ}$ C for 30 min.

#### 3.2.3 Geochemical Analysis of Natural Gas

The geochemical analysis of the natural gas was conducted at the Experimental and Testing Institute of PetroChina Xinjiang Oilfield Company and Northwest Institute of Eco-Environment

Lithology	Formation	Location	Statistics	S <sub>1</sub> (mg HC/g Rock)	S₂ (mg HC/g Rock)	T <sub>max</sub> (°C)	S₁+S₂ (mg HC/g Rock)	тос (%)	PI	HI (mg HC/g TOC)	PC (%)	D (%)	Sample Number
Mudstone	C <sub>1</sub> s <sup>b</sup>	Jimusar Sag	min	0.15	0.64	436	0.88	0.57	0.07	32	0.07	3.70	9
		-	max	0.77	9.60	447	10.37	5.89	0.42	216	0.86	22.90	
			mean	0.45	2.41	442	2.87	1.96	0.22	140	0.24	14.60	
	C <sub>1</sub> s <sup>b</sup>	Kelameili gas	min	0.04	0.24	432	0.39	0.50	0.05	38	0.03	3.52	97
		Field	mean	2.06	14.52	480	16.58	5.85	0.49	248	1.38	26.35	
			max	0.54	2.33	459	2.87	2.04	0.21	106	0.24	11.14	
Carbonaceous	C <sub>1</sub> s <sup>b</sup>	Jimusar Sag	min	0.24	22.27	431	23.47	13.60	0.01	97	1.95	8.12	7
mudstone		Ū	max	5.11	53.28	433	55.60	29.32	0.12	214	4.61	18.57	
			mean	2.28	43.03	432	45.31	24.03	0.05	180	3.76	15.76	
	C <sub>1</sub> s <sup>b</sup>	Kelameili gas	min	0.76	20.41	442	24.08	11.35	0.02	129	2.00	12.54	10
		Field	mean	9.75	88.12	457	96.85	32.87	0.22	288	8.04	26.78	
			max	5.18	45.57	450	50.75	20.59	0.10	215	4.21	19.99	
Coal	C <sub>1</sub> s <sup>b</sup>	Jimusar Sag	min	0.67	35.82	431	36.49	39.60	0.02	87	3.03	7.35	4
		Ū	max	15.84	93.36	441	109.20	57.75	0.15	182	9.06	16.60	
			mean	7.87	70.11	437	77.99	45.83	0.09	152	6.47	13.95	
	C <sub>1</sub> s <sup>b</sup>	Kelameili gas	min	1.18	52.54	437	53.93	42.92	0.02	101	4.48	8.63	10
	·	Field	mean	1.95	87.61	447	88.95	58.15	0.03	204	7.38	17.20	
			max	1.57	72.87	441	74.45	48.77	0.02	151	6.18	12.82	

TABLE 2 | The hydrocarbon generating potential of C1s<sup>b</sup> source rocks in the Jimsar Basin.

and Resources, Chinese Academy of Sciences. A Hewlett-Packard 6890 II gas chromatograph (GC) analyzed the natural gas components. The hydrocarbon gas component was separated using capillary columns (Plot  $Al_2O_3$  50 m  $\times$  0.53 mm). The furnace temperature of the GC was first set to be 30°C and held for 10 min. Then, the temperature was ramped up to 180°C at a rate of 10°C/min. Stable carbon isotope analysis of alkane gas  $(C_1-C_4)$ was conducted using a Finnigan Mat Delta S mass spectrometer interfaced with an HP 6890II gas chromatograph. The alkane gas components (C1-C4) and CO2 were separated using a chromatographic column (Plot Q  $30 \text{ m} \times 0.32 \text{ mm}$ ). The column heating process was as follows: the heating rate was 8°C/min at 35-80°C; the temperature was then increased to 260°C at a heating rate of 5°C/min. The final temperature was held for 10 min. Each sample was analyzed three times with an accuracy of  $\pm 0.3\%$ (VPDB).

#### 3.2.4 Basin Modeling

The burial and thermal histories of the source rocks in the study area were reconstructed using PetroMod software. The current heat flow and thermal conductivity values of the source rocks were adopted from previous studies (Wang et al., 2000a,b; Qiu et al., 2000,2001; Qiu, 2002). The  $R_0$  values were calculated using the Easy%Ro model proposed by (Sweeney and Burnham, 1990; Ren et al., 2020). This model can be applied  $R_0$  ranging of 0.3–4.6%.

# **4 RESULTS AND DISCUSSION**

## **4.1 Source Rock Evaluation** 4.1.1 Types of Organic Matter

The type of organic matter determines the hydrocarbongenerating capacity and product type in a source rock (Tissot et al., 1984; Bordenave, 1993). Therefore, the organic macerals of source rocks and their assemblage characteristics can be studied to characterize the types of organic types (Tissot et al., 1984; Bordenave, 1993; Taylor et al., 1998; Suárez et al., 2012). In this study, the whole-rock maceral contents of  $C_1s^b$  source rocks with three different lithologies (coal, carbonaceous mudstone, and mudstone) in the Jimsar Sag were analyzed based on thin section, oil immersion, and fluorescence observations (**Figure 2; Table 1**). Organic macerals can be classified into four major groups: sapropelite, exinite, vitrinite, and inertinite (Tissot et al., 1984; Bordenave, 1993; Taylor et al., 1998; Suárez et al., 2012). The hydrocarbon generation capacity of the four macerals decreases sequentially, with their products transiting from liquid hydrocarbons to natural gas. The inert maceral group has almost no hydrocarbon-generating capacity.

The original matrices of the sapropelite maceral are primarily lower organisms (i.e., algal and bacteria) and zooplankton (Taylor et al., 1998; Suárez et al., 2012). This study is characterized by mineral bituminous exhibiting weak fluorescence (Figure 2). In addition, sporadically distributed alginate was found in the mudstone samples (Figures 2G,H). The primitive matrices of the exinite group are mainly the plant's reproductive organs and the epidermis and secretions of branches, leaves, and roots (Tissot et al., 1984; Bordenave, 1993; Taylor et al., 1998; Suárez et al., 2012). The exinite maceral group in the  $C_1s^b$  source rocks of the Jimsar Sag mainly include microsporinite, liptodetrinite, cutinite, and resinite (Figure 2). Some dark striped vertical fractures are filled with sphaltenite (Figures 2D,E). The parent material of both the vitrinite and inertinite groups is the xylem of higher plants. The former formed in a reducing-semi-oxidizing environment and the latter in a semi-oxidizing-oxidizing environment (Tissot et al., 1984; Bordenave, 1993; Taylor et al., 1998; Suárez et al., 2012). In this study, the vitrinite maceral



FIGURE 2 | Photomicrographs of macerals for C<sub>1</sub>s<sup>b</sup> source rock in the southeastern Junggar; Note:clay minerals (CI); mineral bituminous (MB); microsporinite (Mis); vtrinite (V); liptodetrinite (Ld); cutinite (Cu); resinite (R); oil satins (O); telinite (T); semi-fusinite (SF); fusinite (F); Pyrite (Py); asphaltenite (Pt); alginate (AI). (A–C) demonstrate the coal sample from well J15 (2,860 m, core). (D–F) demonstrate the carbonaceous mudstone sample from well J3301 (4,135 m, cuttings). (G–I) demonstrate the carbonaceous mudstone sample from well J3301 (4441 m, core). (A,B,D,E,G,H) are inspired by blue light (reflected fluorescence). (C,F,I) are inspired by reflected light (oil immersion).

consists of telinite, collinite, semi-fusinite, and fusinite (**Figure 2**). Fragments of broken fusinite were visible in the center of some semifusinite (**Figure 2C**). Most of them preserved the woody cell structure, indicating that they underwent different degrees of cell wall swelling (**Figure 2C**). Various components can be seen in a parallel arrangement under the microscope, reflecting the hydrostatic sedimentary microenvironment (**Figure 2B**), confirmed by the widespread pyrite observed in  $C_1s^b$  source rocks (**Figure 2**).

The maceral components of the  $C_1s^b$  source rocks, with different lithologies in the Jimsar Sag, are dominated by the vitrinite group (>50%), reflecting their gas-prone characteristics (**Figure 2; Table 1**). The oil-prone maceral content (the sum of the sapropelite and exinite groups) of the mudstone and carbonaceous mudstone reached 22.3 and 16.9%, respectively

(Table 1), showing a bit oil-generating potential (Figure 2; Table 1). The "oil stains" identified through fluorescence observation are evidence (Figure 2B).

The Rock-Eval hydrogen index [HI =  $(100 \times S_2)/TOC$ ] is a fast and economical method to distinguish organic matter types of the source rocks (Tissot et al., 1984; Bordenave, 1993). Based on the HI values of 600, 300–600, 200–300, 50–200, and less than 50 mg HC/g TOC, the organic matter can be classified into five types: Type I (extremely oil-prone), Type II (oil-prone), Type II/III (oilprone/gas-prone), Type III (gas-prone) and Type IV (barren source rock) (Tissot et al., 1984; Bordenave, 1993). Among the C<sub>1</sub>s<sup>b</sup> source rocks of the Jimsar Sag, the HI values of mudstone, carbonaceous mudstone, and coal ranged from 32 to 216 mg HC/g TOC (average 140 mg HC/g TOC), 97–214 mg HC/g TOC (average 180 mg HC/g TOC), and 87–183 mg HC/g TOC



(average 152 mg HC/g TOC), respectively (**Figure 3**; **Table 2**). The  $C_1s^b$  source rocks were dominated by kerogen type III. Carbonaceous mudstone is more oil-prone than coal, and mudstone has the poorest oil-generating capacity (**Figure 3**; **Table 2**). Some of the carbonaceous mudstone samples have high HI values (>200 mg HC/g TOC) and show some oil-generating capacities (**Figure 3**; **Table 2**). Among the  $C_1s^b$  source rocks in the Kelameili gas field, the average HI ratios for mudstone, carbonaceous mudstone, and coal are 106, 215, and 151 mg HC/g TOC, respectively, and the overall characteristics are close to those of the Jimsar Sag (**Figure 3**; **Table 2**).

#### 4.1.2 Source Rock Maturity

In this study,  $R_o$  data were obtained for three  $C_1s^b$  source rock samples from the Jimsar Sag. The  $R_o$  value of the sample from well J15 (2860 m) is 0.65%, and the  $R_o$  values of the two samples from well J3301 (4,441 and 4,445 m) are 1.38 and 1.41%, respectively. Well J15 is in the eastern margin of the Jimsar Sag (**Figure 1B**) and has a lower maturity. According to the Ro values of well J3301, the  $C_1s^b$ source rocks have recently entered at least the main oil-generating window in the depression area. The Rock-Eval maturity indicators of  $C_1s^b$  source rocks in the study area and the Kelameili gas field are



similar. Their hydrocarbon production indices  $[PI = S_1/(S_1+S_2)]$  and  $T_{max}$  are distributed between 0.8 and 1.0 and 440 and 460°C, respectively, in the main oil-generating window and samples with low PI and  $T_{max}$  values are in the immature stage (**Figure 4**; **Table 2**). A few samples with a poor correlation between PI and  $T_{max}$  may have suffered from the contamination of nonindigenous hydrocarbons (**Figure 4**). Some of the  $C_1s^b$  mudstone samples from the Kelameili gas field fall within the "inert carbon" (**Figure 4**), suggesting that those mudstone samples with low HI in **Figure 3** have a poor initial hydrocarbon-generating capacity, in addition to the effect of thermal maturation.

Based on the  $R_o$  and  $T_{max}$  data of  $C_1s^b$  source rock samples from the Jimsar Sag and previous studies on the  $R_o$ -depth relationship in the eastern Junggar Basin (Gong et al., 2021), and combined with the tectonic map of the  $C_1s^b$  bottom boundary of the sag, the  $R_o$ contour map of the study area was plotted (**Figure 5A**). The area of the  $C_1s^b$  source rocks in the Jimsar Sag entering the main oilgenerating window ( $R_o > 0.7\%$ ) and condensate-generating stage ( $R_o = 1.3-2.0\%$ ) reaches 980 and 490 km<sup>2</sup>, respectively (**Figure 5A**). Considering that the humic source rocks produce large amounts of natural gas at the main oil-generating window (Dai et al., 1992; Petersen, 2002; Petersen and Nytoft, 2006; Petersen et al., 2011), the Jimsar Sag favor large- and mediumgas field formation.

In this study, a two-dimensional seismic profile, running longitudinally through the Jimsar Sag in a nearly north-south direction, was selected to recover the burial and thermal evolution history of the  $C_1s^b$  source rocks. At the end of the Carboniferous



(~305 Ma), the C<sub>1</sub>s<sup>b</sup> source rocks were in the immature-low maturity stage ( $R_0 < 0.7\%$ ) and had not reached the main oilgenerating window (Figure 6A). After that, the  $C_1s^b$  source rocks slowly subsided, and by the end of the Early Jurassic (~182 Ma), they entered the early stage of the oil-generating window ( $R_0$  = 0.7-1.0%) (Figure 6B). Subsequently, the subsidence rate accelerated, and the C<sub>1</sub>s<sup>b</sup> source rocks entered the late stage of the oil-generating window ( $R_0 = 1.0-1.3\%$ ) and the condensate/wet gas-generating window ( $R_0 = 1.3-2.0\%$ ) stages in the Cretaceous (~100 Ma). The  $R_0$  of the deepest part of the sag reached 1.7% (Figure 6C). In addition, regional tectonic uplift occurred in the study area, and the hydrocarbon-generating process of C1s<sup>b</sup> source rocks was temporarily stalled (Figure 6D). With further deep burial, the maturity of  $C_1s^b$  source rocks continues to increase. Recently, the deep  $C_1s^b$  source rocks in the Jimsar Sag have entered the massive condensate generation stage. In contrast, the maturity of source rocks at the edge of the sag is low, and a few of them enter the main oil-generating window (Figure 6E).

#### 4.1.3 Hydrocarbon-Generating Potential

Humic source rocks form in fresh marsh-phase environments (oxidizing-semi-oxidizing), whereas lacustrine/marine source rocks form in reducing-semi-reducing environments (Tissot et al., 1984; Bordenave, 1993; Gong et al., 2019). Thus, there are differences in hydrocarbon-generating mechanisms and product types between the two. Their evaluation criteria should be more demanding than those for lacustrine/marine source rocks. The coaly source rocks are rich in aromatic functional groups and poor in lipid functional groups (carbonrich and hydrogen-poor) and have high inert carbon content (Chen et al., 1997; Petersen, 2002; Petersen and Nytoft, 2006; Petersen et al., 2011). Based on thousands of Rock-Eval data of terrigenous source rocks of different lithologies in the major proliferous basins in northwest China, Chen et al. (1997) proposed a scheme to evaluate the hydrocarbon-generating potential of coaly (humic) source rocks using lithologies (mudstone, carbonaceous mudstone, and coal). This scheme has been well tested in the practice of evaluating terrigenous



FIGURE 6 | The hydrocarbon generating profile for C<sub>1</sub>s<sup>b</sup> source rocks in the Jimsar Sag. (A): The thermal evolution history in the Late Carboniferous (305 Ma). (B): The thermal evolution history in the Late Jurassic (182 Ma); (C): The thermal evolution history in the Late Cretaceous (100 Ma); (D): The thermal evolution history in the Late Cretaceous (65 Ma); (E): The thermal evolution history in nowadays (0 Ma);



source rocks in China. This study evaluates the hydrocarbongenerating capacity of C1sb source rocks in Jimsar Sag based on this scheme.

The TOC of C<sub>1</sub>s<sup>b</sup> mudstone in the study area ranged from 0.57 to 5.89%, with an average of 1.96%; the hydrocarbon-generating potential  $(S_1+S_2)$  ranged from 0.88 to 10.37 mg HC/g rock, with an average of 2.87 mg HC/g rock. Thus, the  $C_1s^b$  mudstone belonged to medium-good source rocks (Figure 7A; Table 2). Chen et al. (1997) found that for carbonaceous mudstones and coal, the correlation between  $S_1+S_2$  and HI is better than that between S1+S2 and TOC, and it is more suitable as an index for evaluating organic matter abundance. The C<sub>1</sub>s<sup>b</sup> carbonaceous mudstone is dominated by medium source rocks, with their  $S_1+S_2$ and HI ratios being 23.47-55.60 mg HC/g rock (average 45.31 mg HC/g rock) and 97-214 mg HC/g TOC (average 180 mg HC/g TOC), respectively (Figure 7B; Table 2). The average  $S_1+S_2$  and HI values of coal were 77.99 mg HC/g rock and 152 mg HC/g TOC, respectively, indicating a poor source rock (Figure 7C; **Table 2**). The hydrocarbon-generating capacity of  $C_1s^b$  source rocks in the Jimsar Sag and Kelameili gas fields is similar (Figure 7; Table 2).

# 4.2 Oil and Gas Sources

#### 4.2.1 Ga Source Correlation

One gas sample from the Carboniferous reservoir has been obtained in the Jimsar Sag (from well J3301). The methane content of the natural gas is 78.79%, and the drying factor  $(C_1/\sum C_{1-4})$  is 0.89, indicating a wet gas  $(C_1/\sum C_{1-4} < 0.95)$ . The methane content of the natural gas from the Kelameili gas field ranged from 84.90 to 93.56%, with an average of 88.80%. The natural gases have their  $C_1/\sum C_{1-4}$  ratios ranging from 0.89 to 0.96 (average 0.92), and wet gas accounted for 85.9% of the total samples. The  $C_1/\sum C_{1-4}$  ratios of the gas from the Kelameili gas field are higher than those of the Jimsar Sag, reflecting a relatively higher thermal maturity.

The stable carbon isotope compositions of methane  $(\delta^{13}\text{C-CH}_4)$ , ethane  $(\delta^{13}\text{C-C}_2\text{H}_6)$  and propane  $(\delta^{13}\text{C-C}_3\text{H}_8)$  in well J3301 are -33.8‰, -28.6‰, and -26.8‰, respectively (**Figure 8**). The  $\delta^{13}\text{C-CH}_4$ ,  $\delta^{13}\text{C-C}_2\text{H}_6$ , and  $\delta^{13}\text{C-C}_3\text{H}_8$  of natural gas from Kelameili gas field are -32.3‰ ~ -29.2‰

(average -30.5‰), -28.9‰ ~ -25.6‰ (average -27.1‰), and -26.8‰ ~ -23.1‰ (average -24.7‰), respectively, which are more enriched in <sup>13</sup>C than those of the gas from well J3301 (Figure 8). Considering similar carbon isotopic compositions of kerogen ( $\delta^{13}C_{kerogen}$ ) in the C<sub>1</sub>s<sup>b</sup> source rocks of the Jimsar Sag and Kelameili gas field (Gong et al., 2019,2021), the difference in the stable carbon isotopic compositions of the two types of gas reflects the higher maturity of the gas in the Kelameili gas field, consistent with that in  $C_1 / \sum C_{1-4}$  ratios. The humic source rocks are characterized by kerogen types III and II<sub>2</sub>, composed of aromatic structures relatively enriched in <sup>13</sup>C isotopes (Barry and Fang., 2014; Dai et al., 2014; Liu et al., 2008; Liu et al., 2019; Yao et al., 2020). In contrast, the sapropelic source rocks are characterized by kerogen types I and II<sub>1</sub>, composed of aliphatic structures enriched in <sup>12</sup>C isotopes (Barry and Fang., 2014; Dai et al., 2014; Liu et al., 2019; Wu et al., 2020). Therefore, when humic and sapropelic source rocks generate natural gas at similar maturity, the former generates natural gas (coal-type gas) with a more enriched <sup>13</sup>C carbon isotope composition than the latter (oil-type gas).

Previous studies have confirmed that natural gas from the Kelameili gas field is derived from C<sub>1</sub>s<sup>b</sup> source rocks (Dai et al., 2016; Sun et al., 2016), which exhibit the characteristics of coaltype gas in the  $\delta^{13}$ C-CH<sub>4</sub>- $\delta^{13}$ C-C<sub>2</sub>H<sub>6</sub>- $\delta^{13}$ C-C<sub>3</sub>H<sub>8</sub> diagram proposed by (Dai et al., 2014) (Figure 8). The natural gas from well J3301 has  ${}^{13}$ C-CH<sub>4</sub>,  $\delta^{13}$ C-C<sub>2</sub>H<sub>6</sub>, and  $\delta^{13}$ C-C<sub>3</sub>H<sub>8</sub> values similar to those from the Kelameili gas field (Figure 8), demonstrating the gas was derived from C<sub>1</sub>s<sup>b</sup> source rocks. Although influenced by maturity, the  $\delta^{13}$ C-C<sub>2</sub>H<sub>6</sub> values are more reflective of the original parent material and can distinguish coal-type gas from oil-type gas (Dai et al., 2005; Liu et al., 2016; Wu et al., 2021). Previous studies have shown that the limit of  $\delta^{13}$ C-C<sub>2</sub>H<sub>6</sub> value for both is around –29.0‰ (Fu et al., 1990; Gang et al., 1997; Liu et al., 2004; Dai et al., 2005, 2014; Xiao Z. H. et al., 2008; Liang et al., 2013; Liu et al., 2016; Wu et al., 2021). The  $\delta^{13}$ C-C<sub>2</sub>H<sub>6</sub> value of gas from well J3301 is -28.6‰, which shows the characteristics of coal-type gas (Figure 8).

Oil-type gas is drier than coal-type gas at the same thermal evolutionary stages (Bernard et al., 1978; Whiticar, 1999; Liu et al., 2019; Ci et al., 2020). In the  $\delta^{13}$ C-CH<sub>4</sub>-C<sub>1</sub>/C<sub>2+3</sub> gas genetic





identification plate, gases from the J3301 well and Kelameili field gas fall in the evolution trend line of coal-type gas (**Figure 9**), reflecting their origin from  $C_{1s}^{b}$  source rocks. **Figure 9** reflects the lower maturity of gas from well J3301 than gas from the Kelameili gas field.

#### 4.2.2 Oil-Source Correlation

Two sets of effective source rocks, i.e.,  $P_2l$  and  $C_1s^b$ , were developed in the Jimsar Sag (the Middle and Lower Jurassic source rocks are immature). Previous studies showed that the  $P_2l$  source rocks are brackish lacustrine source rocks deposited

in a reducing environment (Bai et al., 2017; Hu et al., 2018). The oil from the Jimsar billion-ton shale oil field is derived from this set of source rocks (Bai et al., 2017; Hu et al., 2018). The  $C_1s^b$  source rocks are a set of marine-terrigenous transitional deposits in an oxidizing environment with a predominantly terrestrial higher plant input (Gong et al., 2019,2021). A tiny amount of oil has been obtained in the Jimsar Sag from the Carboniferous weathered crust reservoir in well J15. This study compared the physical properties, biomarker fingerprints, and carbon-isotope characteristics of oil samples from well J15 with oil from the P<sub>2</sub>l reservoirs in the Jimsar Sag.

As discussed in **Section 4.2.1**, sapropelic source rocks (P<sub>2</sub>I) have more <sup>13</sup>C-depleted  $\delta^{13}C_{kerogen}$  values than the humic source rocks (C<sub>1</sub>s<sup>b</sup>) when their maturity is close. In addition, the carbon isotopic composition of global Carboniferous sediments is enriched in <sup>13</sup>C (Wang et al., 2013). Therefore, the carbon isotopes of oil derived from C<sub>1</sub>s<sup>b</sup> source rock should be more <sup>13</sup>C-enriched than those from P<sub>2</sub>I source rocks. The stable carbon isotopic compositions of saturated ( $\delta^{13}C_{saturated}$ ) and aromatic ( $\delta^{13}C_{aromatic}$ ) fractions of oil from well J15 are -26.7‰ and -25.9‰, respectively. The  $\delta^{13}C_{saturated}$  and  $\delta^{13}C_{aromatic}$  values of P<sub>2</sub>I reservoired oil in the study area are -33.4‰ to -29.8‰ (average -31.9‰) and -31.4‰ to -27.6‰ (average -30.4‰), respectively, which are lower than those in the J15 well. Therefore, the J15 oil should be derived from C<sub>1</sub>s<sup>b</sup> source rock.

The  $\beta$ -carotene is associated with anoxic/saline environments (Jiang and Fowler, 1986; Peters et al., 2005). The prevalence of  $\beta$ -carotene is a characteristic of Permian-sourced oils in the Junggar Basin (Wang et al., 2013; Cao et al., 2020). In this study,  $\beta$ -carotene was undetected in the oil from well J15 (**Figure 11A**), whereas some amount of  $\beta$ -carotene was detected in the P<sub>2</sub>l reservoired oil (**Figure 11D**), reflecting the different genetic sources of the two.



The Pr/Ph vs gammacerane index plot; (C): The Pr/Ph vs gammacerane index plot; (B): The  $\Sigma C_{31-35}/C_{30}$  hopane ratio vs density plot; (C): The  $\alpha\alpha\alpha$ -20R-C<sub>28</sub>/C<sub>29</sub> sterane ratio vs C<sub>29</sub> regular steranes rato plot.

The main precursors of pristane (Pr) and phytane (Ph) are chlorophylls of photosynthetic organisms, and their abundance is related to the sedimentary environment (Powell, 1988; Peters et al., 2005). The high Pr/Ph value of 1.76 for oil from well J15 (**Figure 10B, Figure 11A**) indicates an oxidizing environment with abundant terrigenous organic matter input Powell, corresponding to the  $C_1s^b$  source rock. In contrast, Pr/Ph values for oil from the P<sub>2</sub>l reservoir are lower, ranging from 0.83 to 1.52, with an average of 1.30 (**Figure 10B, Figure 11D**), reflecting that the source rocks were deposited in a reducing–oxidizing transitional environment dominated by algal and bacterial inputs, with some input from the terrigenous higher plant (Powell, 1988; Peters et al., 2005).

Note:1) n-C<sub>17</sub>; 2) Pr; 3) n-C<sub>18</sub>; 4) Ph; 5)  $\beta$ -carotene; 6) C<sub>19</sub> tricyclic terpane; 7) C<sub>20</sub> tricyclic terpane; 8) C<sub>21</sub> tricyclic terpane; 9) C<sub>30</sub> hopane (10) gammacerane 11) C<sub>31</sub> hopane20S 12) pregnanes 13)  $\alpha\alpha\alpha$ -C<sub>28</sub>20S-regular sterane 14)  $\alpha\beta\beta$ -C<sub>28</sub>20R-regular sterane 15)  $\alpha\beta\beta$ -C<sub>28</sub>20S-regular sterane 16)  $\alpha\alpha\alpha$ -C<sub>28</sub>20R-regular sterane 17)  $\alpha\alpha\alpha$ -C<sub>29</sub>20S-regular sterane 18)  $\alpha\beta\beta$ -C<sub>29</sub>20R-regular sterane 19)  $\alpha\beta\beta$ -C<sub>29</sub>20S-regular sterane (20) $\alpha\alpha\alpha$ -C<sub>29</sub>20R-regular sterane.

The gammacerane reflects the stratified water in the depositional environments in which different source rocks were developed (Peters et al., 2005). The stratified water is due to the longitudinal salinity gradients (Sinninghe et al., 1995; Vu et al., 2009). An increase in water salinity increases the gammacerane index [gammacerane/(gammacerane +  $C_{31}$  hopane)] and decreases the Pr/Ph values (Sinninghe et al., 1995; Vu et al., 2009). The P<sub>2</sub>l reservoired oils have a higher gammacerane index of 0.35–0.65 (average 0.46), showing that they were derived from a set of saline and reducing source rock (**Figure 10B**, **Figure 11E**). In contrast, the gammacerane index of oil from well J15 is lower at 0.30 (**Figure 10B**, **Figure 11B**), indicating a set of fresh source rock (Sinninghe et al., 1995; Vu et al., 2009), corresponding to the C<sub>1</sub>s<sup>b</sup> source rocks.

The oil density from well J15 is 0.7898 g/cm<sup>3</sup>, approximately 0.1 g/cm<sup>3</sup> lower than that of the P<sub>2</sub>l reservoired oil (0.8352–0.9732 g/cm<sup>3</sup>, average 0.8809 g/cm<sup>3</sup>), indicating that they may have different genetic sources (**Figure 10C**). A higher abundance of  $C_{31-35}$  hopane is a feature of the marine environment (Peters et al., 2005). The  $\Sigma C_{31-35}/C_{30}$  hopane ratio of the P<sub>2</sub>l reservoired oil is low, ranging from 0.56 to 0.86



and J251–H (P<sub>2</sub>I reservoir); **(B,E)**: m/z 191 chromatograms of oils from wells J15 (Carboniferous reservoir) and J251–H (P<sub>2</sub>I reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir) and J251–H (P<sub>2</sub>I reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir) and J251–H (P<sub>2</sub>I reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 217 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 207 chromatograms of oils from wells J15 (Carboniferous reservoir); **(C,F)**: m/z 208 chromatograms of oils from wells J15 (Carboniferous reservoir); **(D,F)**: m/z 208 chromatograms of oils from wells J15 (Carboniferous reservoir); **(D,F)**: m/z 208 chromatograms of oils from wells J15 (Carboniferous reservoir); **(D,F)**: m/z 208 chromatograms of oils from wells J15 (Carbonifer

(average 0.74) (Figure 10C). In contrast, the  $\Sigma C_{31-35}/C_{30}$  hopane ratio of oil from well J15 is higher, 0.96 (Figure 10C), corresponding to the depositional environment of  $C_1s^b$  source rock whose sediment environment was affected by seawater (Du, 2010).

From the  $\alpha\alpha\alpha$ -20R-C<sub>28</sub>/C<sub>29</sub> sterane ratio, the difference between the oil from well J15 and P<sub>2</sub>l reservoir is negligible (**Figure 10D**, **Figures 11C**,**F**). However, the relative content of C<sub>29</sub> regular steranes in oil from well J15 was low at 45.86% compared to 48.08–52.43% (average 50.12%) for the P<sub>2</sub>l reservoired oil (**Figure 10D**, **Figures 11C**,**F**). High levels of C<sub>29</sub> steranes indicate significant plant biomass input, reflecting a poor organic matter type (Volkman, 1986; Schwark and Empt., 2006), which seems to be the opposite of the situation in the study area. In previous studies, abundance  $C_{29}$ -regular steranes appeared in some algae, such as diatoms and green algae (Orcutt and Patterson, 1975; Grantham, 1986; Volkman, 1986). However, the abundance of  $C_{29}$  steranes in oil does not signify that most organic matter comes from vascular plants. One possible explanation is that the  $C_{29}$ -regular steranes in  $P_2$ l reservoired oil may come from specific zooplankton that feeds on algae and bacteria. Similar situations have appeared in Paleozoic oils in Oman and the Tarim Basin in China (Grantham, 1986; Zhang and Cheng, 2021).

### 4.3 Oil and Gas Exploration Potential

It is confirmed that a set of  $C_1s^b$  marine-terrigenous transitional source rocks was developed in the Jimsar Sag and that oil and natural gas derived from this set of source rocks have been identified. A planar tracing based on 2D and 3D seismic data evaluated the scale of this source rock. The results show that  $C_1s^b$  source rocks are distributed in the Jimsar Sag with considerable thickness, among which the range of thickness >100 m reaches 580 km<sup>2</sup>, accounting for 38.7% of the whole sag; the range of thickness >50 m reaches 220 km<sup>2</sup> (Figure 5B). Three subdepositional centers developed in the sag (Figure 5B).

The lithology of C<sub>1</sub>s<sup>b</sup> source rocks in Jimsar Sag is dominated by carbonaceous mudstone, followed by mudstone, and a few thin coal seams are developed. Therefore, kinetic simulation of hydrocarbon generated from C<sub>1</sub>s<sup>b</sup> carbonaceous mudstone and mudstone was conducted in a closed system (golden tubes), adopting Tang et al. (2000) and Xiong et al. (2004) experimental method (the details will be discussed in another manuscript). The experimental results showed that the maximum gas yield of the mudstone and carbonaceous mudstone is 213.69 and 250.16 mg HC/g TOC, respectively, at a heating rate of 2°C/h. Their maximum oil yield was 26.22 and 162.46 mg HC/g TOC, respectively. Based on the obtained parameters, such as pre-exponential factor and activation energy, according to the Arrhenius formula, and the characteristics of thermal evolution and hydrocarbon generation histories of source rocks (Figure 6), this study sketched the planar contour maps of the gas- (Figure 5C) and oil- (Figure 5D) generating intensities of C<sub>1</sub>s<sup>b</sup> source rocks in the Jimsar Sag.

The relationship between the distribution of large and medium gas fields and the gas-generating intensity of source rocks in China shows that large gas fields (proven geological reserves >300 × 10<sup>8</sup> m<sup>3</sup>) in China are distributed in areas with gas-generating intensity >20 × 10<sup>8</sup> m<sup>3</sup>/km<sup>2</sup> (Dai et al., 1996,1999,2000; Li et al., 2020; Wei et al., 2020). The area of  $C_1s^b$  source rocks in the Jimsar Sag with gas-generating intensity >20 × 10<sup>8</sup> m<sup>3</sup>/km<sup>2</sup> (Figure 5C), accounting for 67.7% of the study area. Approximately one-third of the sag has gas-generating intensity >100 × 10<sup>8</sup> m<sup>3</sup>/km<sup>2</sup> (Figure 5C). The area of  $C_1s^b$  source rocks with the oil-generating intensity >500 × 10<sup>4</sup> and 1,000 × 10<sup>4</sup> t/km<sup>2</sup> are 1,146 and 702 km<sup>2</sup> (Figure 5D), respectively, showing a favorable oil-generating potential.

Seismic, gravitational, and magnetic data identified 13 Carboniferous volcanic lithologic traps (buried hills), covering  $230 \text{ km}^2$ , in the Jimsar Sag (**Figure 5**). These traps are distributed in areas with high gas and oilgenerating intensities, and source rocks have entered the main oil-generating window (**Figure 5**), conducive for forming self-generated and -stored petroleum reservoirs. In summary, the Jimsar Sag has the resource potential to form large and medium oil/gas fields and is an alternative for petroleum exploration in the Junggar Basin after the Kelameili gas field.

# **5 CONCLUSION**

1) In the Jimsar Sag (in the southeastern part of Junggar Basin), a set of marine–terrigenous transitional source rocks were developed in the Lower Carboniferous Songkharsu Formation, comprising carbonaceous mudstone and mudstone. They belong to medium–good source rocks that are characterized by kerogen types II2–III and are primarily gas-prone, with some oil-generating ability. The quality of  $C_1s^b$  source rocks in the Jimsar Sag is close to that in the Kelameili gas field. At the end of the Early Jurassic,  $C_1s^b$ source rocks reached the oil-generating peak in the Jimsar Sag. Recently, most of them have reached the highly-mature stage.

2) The Carboniferous-reservoired oil in well J15 has more  $^{13}C$ -enriched  $\delta^{13}C_{saturated}$  and  $\delta^{13}C_{aromatic}$  values and higher  $\Sigma C_{31-35}/C_{30}$  hopane and Pr/Ph values (0.96 and 1.76) than the P\_2l-reservoired oil. The density, gammacerane index, and  $C_{29}$  regular sterane content of the J15 oil are 0.7898 g/cm<sup>3</sup>, 0.39, and 45.86%, respectively, which were lower than those of the oil from the P\_2l reservoir. The  $\beta$ -carotene was undetected in the J15 oil. Therefore, J15 oil should be derived from  $C_1 s^b$  source rock.

3) The  $\delta^{13}$ C-CH<sub>4</sub>,  $\delta^{13}$ C-C<sub>2</sub>H<sub>6</sub>, and  $\delta^{13}$ C-C<sub>3</sub>H<sub>8</sub> values of gas from well J3301 are –33.8‰, –28.6‰ and –26.8‰, respectively, which are similar to carbon isotope values from the Kelameili gas field and show the characteristics of coal-type gas, indicating a C<sub>1</sub>s<sup>b</sup> origin.

4) The  $C_1s^b$  source rocks are distributed in the Jimsar Sag with considerable thickness, among which the area with thickness >100 m reaches 580 km<sup>2</sup>, accounting for 38.7% of the study area. In addition, 13 Carboniferous volcanic lithological traps have been identified in the Jimsar Sag, covering 230 km<sup>2</sup>. The Carboniferous petroleum system of the Jimsar Sag can form large and medium oil and gas fields, which would be an alternative oil and gas exploration in volcanic reservoirs in the Junggar Basin after the Kelameili gas field.

# DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

# **AUTHOR CONTRIBUTIONS**

DG: Conceptualization, Writing Review; Editing, Supervision; YS: Writing Original Draft, Formal analysis; MP: Formal analysis, Methodology; CL: Investigation, Data Curation; RW: Investigation, Data Curation; WW: Geophysical research.

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Conflict of Interest: YS, MP, and CL were employed by Xinjiang Oil Company.

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

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