



Article Numerical Simulation of the Proppant Settlement in SC-CO₂ Sand-Carrying Fluid in Fracturing Fractures

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Abstract: Supercritical CO₂ fracturing has unique advantages for improving unconventional reservoir recovery. Supercritical CO₂ can penetrate deep into the reservoir and increase reservoir reform volume, and it is less damaging to reservoir and easy to flow back. However, when the supercritical CO₂ flows as the sand-carrying fluid in the fracture, the settlement of the proppant is still worth studying. Based on the study of supercritical CO₂ density and viscosity properties, assuming that the reservoir has been pressed out of the vertical crack by injecting prepad fluid, the proppant characteristics in sand-carrying fluid under different conditions were studied by numerical simulation. After the analysis, the proppant accumulation and backflow will occur at the end of the crack. Large sand diameters, high fluid flow rates, high sand concentrations, high reservoir temperatures, and low reservoir pressures can help to shorten deposition time, and the small particle size, high fluid flow rate, low sand concentration, low reservoir temperature, and high reservoir pressure can help increase the uniformity of sand deposition. Shortening the sand deposition time can help to complete the fracturing efficiently, and increasing the deposition uniformity can improve the fracture conductivity. This article has studied the proppant settling and crack formation characteristics. It is hoped that this study can provide theoretical support for field fracturing and provide theoretical assistance to relevant researchers.

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Copyright: © 2022 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). **Keywords:** reservoir fracturing; supercritical CO₂; sand-carrying fluid; proppant; settlement characteristics

1. Introduction

Unconventional resources are an important area for oil and gas exploration and development in the future, and there are many methods for reservoir modification and stimulation at present [1]. CO_2 fracturing is one of the core technologies for oil-gas industrial development [2,3]. Since the 1980s, North America has used the fracturing fluid system based on liquid CO_2 for reservoir reconstruction. In 1981, hydrated ethanol/ CO_2 emulsions were successfully used for fracturing in the Western Canadian Sedimentary Basin (WCSB), and these fracturing fluids had completed more than 3000 fracturing operations in the WCSB area by 2008 [4]. In 1993, the United States implemented a CO_2 sand fracturing operation in the Big Sandy field for the first time; since then, carbon dioxide fracturing technology has gradually developed. CO_2 fracturing has unique advantages over other fracturing methods [5,6]. Experimental and site construction show that after CO_2 fracturing, the yield is five times higher than that of nitrogen foam fracturing [7,8].

When temperature and pressure conditions reach the critical point of carbon dioxide, that is, the temperature is higher than 31.06 °C and the pressure is higher than 7.38 MPa, the interface between gas and liquid carbon dioxide disappears completely and finally becomes a compressible high-density fluid, i.e., supercritical carbon dioxide [9–11]. Supercritical CO₂ fracturing employs non-aqueous media (CO₂, chemical reagents, etc.) as fracturing fluids to fracture unconventional reservoirs. In theory, supercritical CO₂ fracturing can

produce more microcracks without causing clay expansion and reduce the negative impact on the reservoir. At the same time, it can use unconventional reservoirs to store a large amount of CO_2 gas to achieve greenhouse gas emission reduction [12,13].

At present, domestic and foreign researchers have conducted a large number of SC-CO2 applications in oil and natural gas development. Studies have shown that the proppant suspension and sedimentation in fracturing fluid are crucial to fracture morphology and conductivity capability. At present, the numerical models describing proppant transport laws can be divided into two categories: continuum theory model and sedimentation theory model. The continuum theory model treats the fracturing fluid and proppant particles as a continuous medium to describe the fracturing fluid flow and proppant transport process. Based on the continuum theory, Wasp and Aude (1970) used Froude number correlations to calculate the deposition rate of particles in the mixed phase and compared the solid suspension state under different transport conditions [14]. Wasp and Aude pointed out that the inhomogeneity degree which can be tolerated in pipelines is dependent on the specific application; Clifton and Wang (1988) obtained proppant transport equations by establishing proppant motion equations and fracturing fluid motion equations [15]. Based on the slurry transport and settling experiments, Barree and Conway (1994) obtained a new slurry transport model and applied it to 3D fracture simulation [16]. Studies have shown that the model has good accuracy in simulating the proppant slurry delivery process. Sharma and Gadde (2005) presented a model of the proppant particle velocity in the channel flow, indicating that the proppant has a retarding or accelerating effect relative to the fluid depending on the ratio of the proppant size to the crack width [17]. Wood and Wheeler et al. (2007) defined a Slurry Properties Index for a given proppant condition and fluid composition [18]. Based on this index, the minimum horizontal flow rate required for solid-phase suspension is deduced to guide the fracturing design.

However, for low-viscosity fracturing fluids, the sedimentation theory model is a better choice. Proppants tend to settle in a low-viscosity fluid to form a packed bed. Based on sedimentation theory, Clark and Quadir (1981) pointed out that the sedimentation rate of particles in fracturing fluid is the main factor that determines the proppant distribution in hydraulic fractures and predicts the terminal settling velocities of proppants under different fracturing conditions [19]. Patankar et al. (2002) studied the influence of the lifting force acting on the particle deposition through two-dimensional numerical simulations and laboratory experiments and played a catalytic role in the study of proppant settlement [20]. Gadde et al. (2004) proposed a new model for proppant transport and settlement in hydraulic cracks and studied the influence of different factors on proppant settling velocity [21]. At the same time, the new proppant settlement model was incorporated into the three-dimensional hydraulic fracture simulator (UTFRAC-3D) [22,23]. The above studies provide a good basis for revealing the motion of proppant in fracturing fluids.

In the field of fluid mechanics, the coupled computational fluid dynamics-discrete element method (CFD-DEM) is gradually becoming widely used [24–26]. CFD represents computational fluid dynamics, and DEM represents the discrete element method [27,28]. The fluid and solid are coupled through the exchange of momentum and energy. Because DEM reveals the essence of particle motion, the coupled CFD-DEM has been widely used in many applications related to fluid–particle interaction [29,30]. Considering the typical heat transfer between particles and between fluids and particles, Zhang and Tahmasebi (2019, 2018) established a coupling multiphysical and micromechanical model based on DEM [24,25]. A series of studies have been performed on fluid flow and particle movement in porous media. The CFD-DEM method is evidently very suitable for simulating the flow characteristics of liquid and solid mixtures at the microscale and other flow conditions. The above research provides theoretical support for the development of this study.

In the actual hydraulic fracturing process, because the fracturing conditions can reach the CO₂ critical point, the CO₂ fracturing fluid will flow in a supercritical state in the fracture. The physical parameters of the fracturing fluid will also change significantly with temperature and pressure conditions. In this study, in order to describe the characteristics of proppant suspension and settling in SC-CO₂ fracturing fluid, firstly the changes in density and viscosity of supercritical carbon dioxide under different temperature and pressure conditions were analyzed. Secondly, ICEM CFD and FLUENT software were used to simulate the proppant settlement under different fracturing conditions. In the current research, the application of ICEM CFD and FLUENT software mainly focuses on the flow of liquid or the mixed fluid of liquid and solid and on the flow in the pipeline or microporosity [31–34]. However, there is no research on the sedimentation of mixed fluid in underground high-temperature and high-pressure fractures. Therefore, this study is the first to obtain the settling characteristics of proppant in supercritical CO₂ in reservoir fractures.

2. The Change Characteristics of SC-CO₂ Density and Viscosity

For carrying proppants, the density and viscosity of the fracturing fluid are important factors that affect the ability of proppant to suspend and settle. The density and viscosity of supercritical CO_2 are greatly affected by pressure and temperature, and the flow of CO_2 fracturing fluid in the reservoir can easily reach the critical condition. Therefore, here we analyze the effect of temperature and pressure on the density and viscosity of CO_2 fracturing fluids.

2.1. SC-CO₂ Density Variation

There is a connection between the density, pressure, volume, and amount of gas. For actual gas, there is an interaction force between gas molecules, and the volume of actual gas molecules cannot be ignored, which is quite different from the ideal gas state equation. Table 1 shows the statistics of current actual gas state equations.

Table 1. Actual gas state equation.

Presenter	State Equation
Van der Waals (VdW) (1887)	$P = \frac{RT}{V-b} - \frac{a}{V^2}$
Redlich & Kwong (RK) equation (1949)	$P = \frac{RT}{V-b} - \frac{a/\sqrt{T}}{V(V+b)}$
Soave-Redlich-Kwong (SRK) (1972)	$P = \frac{RT}{V-b} - \frac{a(T)}{V(V+b)}$
Peng-Robinson (PR) (1976)	$P = \frac{RT}{V-b} - \frac{a(T)}{V(V+b) + b(V-b)}$
Patel-teja (PT) (1982)	$P = \frac{RT}{V-b} - \frac{a(T)}{V(V+b)+c(V-b)}$
Span & Wagner (1994)	$rac{M\cdot p(\delta, au)}{ ho RT}=1+\delta \phi^r_{\delta}$

As can be seen from Table 1, researchers have proposed a large number of actual gas state equations such as RK, SRK, PR, and PT equations, etc., but different equations have certain limitations [35,36] and only have better performance under certain temperature and pressure conditions. In contrast, the Span & Wagner equation of state used experimental data to fit the actual gas state equation in the form of Helmholtz free energy [37], which has high accuracy and a wide range of uses overall. This model calculates temperatures up to 1100 K (827 °C) and pressures up to 800 MPa (8000 bar) and can be used to calculate the state parameters near the CO₂ critical point. Meanwhile, in order to simplify the calculation, the empirical formula proposed by Ouyang (2011) [38] was used.

The CO₂ density expression is

$$\rho_{CO_2} = A_0 + A_1 P + A_2 P^2 + A_3 P^3 + A_4 P^4 \tag{1}$$

where ρ_{CO_2} denotes CO₂ density, kg/m³; *P* denotes the pressure, 0.0069 MPa; and $A_0 \sim A_4$ denotes correlation coefficients, which can be obtained by

$$A_i = b_{i0} + b_{i1}T + b_{i2}T^2 + b_{i3}T^3 + b_{i4}T^4 \quad (i = 0, 1, 2, 3, 4)$$
⁽²⁾

where $b_{i0} \sim b_{i4}$ denotes correlation coefficients, which are given in Tables A1 and A2 (Appendix A), and *T* denotes the temperature, °C.

Figure 1 shows the influence of temperature and pressure on CO₂ density.



Figure 1. The relationship between SC-CO₂ density and temperature-pressure.

As shown in Figure 1, the density of SC-CO₂ is between that of gaseous CO₂ and liquid CO₂. At a certain temperature, SC-CO₂ viscosity increases with increasing pressure, and the increase in density is more pronounced at relatively low temperatures (40~60 °C), while density changes gently under high pressure (above 15 MPa). When the pressure is certain, the density of SC-CO₂ is reduced by the increase in temperature and decreases rapidly under low pressure (Around 10 MPa). The greater the density of the sand-carrying fluid, the more favorable the suspension of the proppant. Therefore, in terms of SC-CO₂ density, the sand-carrying fluid has a higher ability to carry proppant under conditions of about 40 °C and high pressure.

2.2. SC-CO₂ Viscosity Variation

In 1990, Vesovic et al. obtained a formula for calculating CO_2 viscosity by analyzing and summarizing a large amount of experimental data [39]. According to the experimental data supplemented in the later period, in 1998 Fenghour revised the fitting formula of the residual viscosity in Vesovic's model, taking into account the effects of temperature and density on the residual viscosity, and improved the model accuracy. As shown in Figure 2, the variation of the viscosity of CO_2 was obtained using the viscosity model proposed by Fenghour (1998) [40].

The fluid viscosity is formed by the internal friction generated by the relative motion between different molecules. According to the mechanism of viscosity, the viscosity can represent the sum of zero-density viscosity, residual viscosity, and singular viscosity [41,42]. Among them, the ratio of singular viscosity to total viscosity is relatively low, generally less than 0.01. The viscosity calculation formula can be expressed as

$$\eta(T,\rho_{\text{CO}_2}) = \eta_0 + \Delta\eta(T,\rho_{\text{CO}_2}) + \Delta\eta_c(T,\rho_{\text{CO}_2})$$
(3)

where η denotes total fluid viscosity, Pa·s; η_0 denotes zero-density viscosity, Pa·s; $\Delta \eta$ denotes residual viscosity, Pa·s; and $\Delta \eta_c$ denotes singular viscosity, Pa·s.

Among them, the zero-density viscosity calculation can be expressed as

$$\eta_0(T) = \frac{1.00697(T+273.15)^{0.5}}{G'_\eta(T')}$$
(4)

where $\eta_0(T)$ denotes the zero-density viscosity, μ Pa·s, and $G'_{\eta}(T')$ denotes the reduced effective cross-section, which can be represented by the empirical equation

$$\ln G'_{\eta}(T') = \sum_{i=0}^{4} \left(a_i \ln(T' + 273.15) \right)^i$$
(5)

$$(T' + 273.15) = k(T + 273.15)/\varepsilon, \quad \varepsilon/k = 251.196K$$
 (6)

where *T'* denotes the reduced temperature, ${}^{\circ}C$; ε/k denotes the energy scaling parameter, K; and $a_0 \sim a_4$ denote empirical coefficients, which are listed in Table A3 (Appendix B). The formula for calculating the residual viscosity is

$$\Delta \eta(\rho_{\rm CO_2}, T) = d_{11}\rho_{\rm CO_2} + d_{21}\rho_{\rm CO_2}^2 + \frac{d_{64}\rho_{\rm CO_2}^6}{\left(T' + 273.15\right)^3} + d_{81}\rho_{\rm CO_2}^8 + \frac{d_{82}\rho_{\rm CO_2}^8}{\left(T' + 273.15\right)}$$
(7)

where $d_{11}, d_{21}, d_{64}, d_{81}, d_{82}$ denote empirical coefficients, which are listed in Table A3 (Appendix B).

From Figure 2, the viscosity of SC-CO₂ is about $0.02 \sim 0.12$ cp. Under certain temperature conditions, the SC-CO₂ viscosity increases with increasing pressure and increases faster at 40~60 °C. Meanwhile, under certain pressure conditions, the viscosity of SC-CO₂ decreases with increasing temperature and decreases rapidly at 10~15 MPa. The higher the viscosity of the sand-carrying fluid, the less likely the proppant is to settle and the more beneficial it is to proppant suspension and migration. At conditions of similar density, high pressure, and 40 °C, the proppant is better carried by the sand-carrying liquid.





(**b**) Pressure greater than 20.69 MPa

Figure 2. The relationship between SC-CO₂ viscosity and temperature-pressure.

3. Numerical Simulation of Proppant Settlement

In this study, a shale gas well in China's Sichuan Basin is analyzed as a case. The fracturing construction and fracture diagrams are shown in Figure 3. The characteristics of the shale gas reservoir are shown in Table 2.

First, the ICEM CFD is used to establish a long and narrow fracture model (90 m-long, 18 m-high, and 5 cm-wide rectangular rectangles), with one end as the fluid inlet and one end as the closed boundary. The supercritical CO_2 sand-carrying liquid entered from the left side at a certain flow velocity (speed inlet boundary), and other boundaries were closed boundaries. When the sand-carrying liquid volume filled the entire crack, the fluid stopped entering ().

Then, the fluid parameters were set, including the selection of sand-carrying fluid (supercritical CO_2) and proppant (Ceramsites), as well as the particle size, density, and

sand ratio. The parameters involved include sand diameter: 0.6 mm; sand concentration: 0.4; sand density: 2650 kg/m^3 .

Then, the fluid flow conditions were set, including temperature, pressure, and flow rate. The flow conditions were temperature: $60 \,^{\circ}$ C; pressure: $30 \,$ MPa; sand-carrying fluid inlet flow rate: $1 \,$ m/s.

Finally, under the above conditions, the settlement characteristics of proppant at the characteristic time points of the flow process (such as 20 s, 40 s, 60 s, 80 s, 100 s, 140 s) after supercritical CO₂ flows into the fracture were simulated.



Figure 3. Fracturing construction and fracture diagrams.

Parameter	Value	Parameter	Value
Reservoir temperature	60 °C	Reservoir pressure	30 MPa
Reservoir depth	3650 m	Sand-carrying fluid	Supercritical CO ₂
Porosity	1.38%	Proppant	Ceramsites
Gas layer thickness	34 m	Sand-carrying fluid inlet flow rate	1 m/s
Gas content	2.48 m ³ /t	Sand diameter	0.6 mm
Organic carbon (TOC) mass fraction	3.5%	Sand concentration	0.4
Vitrinite reflectance	2.2%	Sand density	2650 kg/m^3

Table 2. Basic parameters of shale gas reservoir.

The key parameters involved in the model establishment process are shown in Table 2. Figure 4 shows the sand concentration distribution cloud map at different times (20 s; 40 s; 60 s; 80 s; 100 s; 140 s). The flow conditions were temperature: 60 °C; pressure: 30 MPa; sand diameter: 0.6 mm; sand-carrying fluid inlet flow rate: 1 m/s; sand concentration: 0.4; sand density: 2650 kg/m³.

At a temperature of 60 °C and a pressure of 30 MPa, the supercritical CO₂ has a density of 829.78 kg/m³ and a viscosity of 0.0768 mPa·s. As can be seen from Figure 4, when supercritical CO₂ carries sand into the fracture, the liquid phase and sand will fall under the action of gravity while flowing horizontally. While falling, it produces a forward velocity that accelerates the sand flow. When the gravel reaches the top of the fracture, it will also return due to impact on the wall and show stacking and reflux phenomena at the end of crack. Due to the large differences in liquid- and solid-phase densities and small CO₂ viscosities, sedimentation will continue to occur during the sand flow. After a period of time, when the fluid no longer enters the fracture, it tends to be stable.



Figure 4. Gravel settlement diagrams at different flow moments.

3.1. Solid Particle Diameter

In order to study the sedimentation characteristics of solid particles with different sizes in the sand-carrying liquid, we set five different particle sizes: 0.00025 m; 0.0006 m; 0.001 m; 0.0015 m; and 0.002 m. The flow conditions were temperature: $60 \text{ }^{\circ}\text{C}$; pressure: 30 MPa; inlet flow rate of sand-carrying fluid: 1 m/s; sand concentration: 0.4; sand density: 2650 kg/m^3 . The distribution of sand sedimentation after 80 min of flowing in the five cases is as follows.

From Figures 5 and 6, it can be seen that the smaller the particle size, the faster the particles flow with the liquid phase, and the particles will quickly reach the end of the fracture and flow back. The smaller the particle size, the longer the particles are suspended in the liquid phase and the less likely sedimentation is to occur. On the contrary, the sand particles with a larger particle size finally reach a shorter deposition stability time. However, the sand distribution in the cracks will be more uniform when the sand diameter is small.



Figure 5. Gravel settlement diagram with different particle sizes.



Figure 6. Settling time at different particle diameters.

3.2. Fluid Velocity

In order to study the sedimentation characteristics of solid particles under different sand-carrying and solid-phase particle flow speeds, we set five different inlet velocity sizes: 0.5 m/s; 1 m/s; 1.5 m/s; 2 m/s; and 2.5 m. The flow conditions were temperature: $60 \degree \text{C}$; pressure: 30 MPa; solid particle diameter: 0.0006 m; sand concentration: 0.4; sand density: 2650 kg/m^3 . The sand distribution after 80 min in the five cases is as follows.

It can be seen from Figures 7 and 8 that the fluid velocity has a significant effect on the sedimentation characteristics of solid particles. When the flow rate is low, the flow of sand-carrying fluid and sand is relatively stable, and sand is more prone to sedimentation. A lot of gravel does not reach the end of the crack and is deposited at the bottom. At high flow rates, most of the sand will be deposited by hitting the trailing edge of the fracture, and eventually the sand will be more evenly distributed. As can be seen from Figure 8, it takes a long time for the sand deposition to stabilize at a low flow rate, and the deposition stability time at a high flow rate significantly decreases. Due to the increased turbulence of the fluid at high flow rates, the settling time becomes gentle at 1.25 m/s~2.25 m/s.



Figure 7. Gravel settlement diagram with different fluid velocity.



Figure 8. Settling time at different inlet velocities.

3.3. Sand Concentration

For obtaining the sedimentation characteristics of solid particles under different sand concentrations, we set five different sand concentrations: 0.2; 0.3; 0.4; 0.5; and 0.6. The flow conditions were temperature: 60 °C; pressure: 30 MPa; solid particle diameter: 0.0006 m; inlet flow rate of sand-carrying fluid: 1 m/s; sand density: 2650 kg/m³. The sand distribution after 80 min in the five cases is as follows.

As can be seen from the Figures 9 and 10, the higher the sand concentration, the faster the sand front moves. The higher the concentration of sand, the greater the overall density of the sand-carrying liquid, the faster the settling velocity, and the shorter the settling time. However, at high concentrations, sand tends to accumulate at the tail of the fracture, and sand concentration at high concentrations is not as uniform as at low concentrations.



Figure 9. Gravel settlement diagram with different sand concentrations.



Figure 10. Settling time at different sand concentrations.

3.4. Flow Temperature

In order to obtain the effect of flow temperature on sand deposition in sand-carrying liquid, we set five different flow temperatures: 40 °C; 60 °C; 70 °C; 80 °C; and 100 °C. The flow conditions were pressure: 30 MPa; solid particle diameter: 0.0006 m; inlet flow rate of sand-carrying fluid: 1 m/s; sand concentration: 0.4; sand density: 2650 kg/m³. The sand distribution after 80 min in the five cases is as follows.

In Figures 11 and 12, the flow temperature mainly affects the sedimentation of sand by affecting the density and viscosity of supercritical CO_2 . When the pressure is constant (30 MPa), the temperature increases from 40 °C to 100 °C, and the density and viscosity gradually decrease. Therefore, the sand-carrying capacity of the liquid phase decreases with increasing temperature, and the shorter the final settling time, the more uneven the distribution of sand.



Figure 11. Gravel settlement diagram with different flow temperatures.



Figure 12. SC-CO₂ physical properties and sand settling time at different flow temperatures.

3.5. Average Pressure

In order to obtain the effect of average pressure on sand deposition in sand-carrying liquid, we set five different flow temperatures: 20 MPa; 30 MPa; 40 MPa; 50 MPa; and 60 MPa. The flow conditions were flow temperature: 60 $^{\circ}$ C; solid particle diameter: 0.0006 m; inlet flow rate of sand-carrying fluid: 1 m/s; sand concentration: 0.4; sand density: 2650 kg/m³. The sand distribution after 80 min in the five cases is as follows.

In Figures 13 and 14, pressure affects the density and viscosity of supercritical CO₂. When the flow temperature is 60 °C, the average pressure rises from 20 MPa to 60 MPa, and the density and viscosity of supercritical CO₂ gradually increase. At the same time, the sand front moves faster, the sand-carrying capacity of the liquid phase increases slightly, and the final stable time increases but the sand deposition uniformity increases.

In summary, solid particle diameter, fluid velocity, sand concentration, flow temperature, and average pressure all have an effect on the sedimentation of the proppant in the fracture, and they will ultimately affect the fracturing process and the fracturing effect.



Figure 13. Gravel settlement diagram with different average pressures.



Figure 14. SC-CO₂ physical properties and sand settling time at different average pressures.

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4. Conclusions

This article studies the sedimentation characteristics of proppant in sand-carrying fluid by numerical simulation during supercritical CO₂ fracturing. Several meaningful conclusions can be drawn:

- (1) For cracks with a certain height and length, when the SC-CO₂ sand-carrying liquid enters the crack at a certain speed under the action of gravity and viscous forces, the proppant will continuously deposit and the accumulation and backflow will occur at the end of the crack.
- (2) The large sand diameters, high fluid flow rates, high sand concentrations, high reservoir temperatures, and low reservoir pressures can help to shorten deposition time, and shortening the sand deposition time can help to complete the fracturing efficiently.
- (3) The small particle size, high fluid flow rate, low sand concentration, low reservoir temperature, and high reservoir pressure help increase the uniformity of sand deposition, and increasing the deposition uniformity can improve the fracture conductivity.
- (4) According to the reservoir conditions, the reasonable adjustment of the fracturing parameters can significantly improve the fracturing effect, and we hope that the results of this study can play a guiding role in field construction.

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Appendix A. SC-CO₂ Density Calculation Correlations

Table A1. Value of correlation coefficients $b_{i0} \sim b_{i4}$ (P \leq 20.69 MPa).

Symbol	b _{i0}	b _{i1}	b _{i2}	b _{i3}	b _{i4}
i = 0	$-2.15 imes10^5$	$1.17 imes10^4$	$-2.30 imes 10^{2}$	1.97	$-6.18 imes10^{-3}$
i = 1	$4.76 imes 10^2$	-2.62×10	$5.22 imes 10^{-1}$	$-4.49 imes10^{-3}$	$1.42 imes 10^{-5}$
i = 2	$-3.71 imes10^{-1}$	$2.07 imes10^{-2}$	$-4.17 imes10^{-4}$	$3.62 imes 10^{-6}$	$-1.16 imes10^{-8}$
i = 3	$1.23 imes10^{-4}$	$-6.93 imes10^{-6}$	$1.41 imes10^{-7}$	$-1.23 imes10^{-9}$	$3.95 imes10^{-12}$
i = 4	$-1.47 imes10^{-8}$	$8.34 imes 10^{-10}$	$-1.70 imes 10^{-11}$	$1.50 imes 10^{-13}$	-4.84×10^{-16}

Table A2. Value of correlation coefficients $b_{i0} \sim b_{i4}$ (P > 20.69 MPa).

Symbol	b _{i0}	b _{i1}	b _{i2}	b _{i3}	b _{i4}
i = 0	$6.90 imes 10^2$	2.73	-2.25×10^{-2}	$-4.65 imes10^{-3}$	$3.44 imes 10^{-5}$
i = 1	$2.21 imes10^{-1}$	$-6.55 imes10^{-3}$	$5.98 imes10^{-5}$	$2.27 imes10^{-6}$	$-1.89 imes10^{-8}$
i = 2	$-5.12 imes10^{-5}$	$2.02 imes10^{-6}$	$-2.31 imes10^{-8}$	$-4.08 imes10^{-10}$	$3.89 imes10^{-12}$
i = 3	$5.52 imes10^{-9}$	$-2.42 imes10^{-10}$	$3.12 imes10^{-12}$	$3.17 imes10^{-14}$	$-3.56 imes10^{-16}$
i = 4	-2.18×10^{-13}	$1.01 imes 10^{-14}$	$-1.41 imes10^{-16}$	$-8.96 imes 10^{-19}$	1.22×10^{-20}

Appendix B. SC-CO₂ Viscosity Correlations

Coefficient	Value	Coefficient	Value
a0	0.235156	d11	$4.071119 imes 10^{-3}$
a1	-0.491266	d21	$7.198037 imes 10^{-5}$
a2	0.05211155	d64	$2.411697 imes 10^{-17}$
a3	0.05347906	d81	2.971072×10^{-23}
a4	-0.01537102	d82	$-1.627888 imes 10^{-23}$

Table A3. Viscosity calculation coefficients.

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