Research Article
The Size Effect and Microstructure Changes of Granite after Heat Treatment

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High temperature can change the mechanical properties of granite, with significant nonlinear characteristics, and at the same time change its microstructure. Therefore, two kinds of granites are used in this paper: one is normal temperature granite and the other is granite treated at 600°C, and a detailed comparative study is made. The fracture toughness of two kinds of rocks was tested by fracture tests, and the results were analyzed by a nonlinear fracture mechanics model (SEL). At the same time, in order to understand the influence of high temperature on the mineral composition and microstructure of granite, XRD, optical microscope, and SEM were used to observe the mineral composition, microcracks, and fracture morphology of granite. The results show the following: (1) high temperature significantly changes the fracture mechanics parameters of granite. The fracture toughness of granite treated at 600°C is significantly lower than that of untreated granite, which is reduced by more than 60%. (2) No obvious size effect was found in the untreated granite, while the size effect of the granite after treatment at 600°C was significant. (3) The granite after high-temperature treatment showed strong nonlinear characteristics, and the SEL can reasonably describe and explain its nonlinear fracture characteristics. (4) The brittleness of the granite treated at 600°C decreased and the ductility increased. The microscopic morphology of the fracture was rough, with obvious steps and rivers. The microcracks and porosity had increased significantly, but the main components did not change significantly.

1. Introduction

Due to the needs of nuclear waste storage, geothermal resource development, and many other engineering constructions [1–8], the mechanical properties of rocks after high temperature have caused extensive research by scholars.

So far, there are many reports on the fracture toughness of rocks after high temperature, including granite, marble, basalt, and other rocks [9]. In 2007, Nasseri et al. [10] tested the fracture toughness of Westerly granite after high temperature and found that the fracture toughness of granite continued to decrease with the increase of temperature, and the fracture toughness of granite treated at 800°C decreased by more than 90%.

In 2018, Talukdar et al. [11] heated three types of crystalline rocks in India and tested their fracture toughness. The results show that when the treatment temperature reaches 600°C, the fracture toughness values of the three types of rocks are greatly reduced, exceeding 50%. A large number of similar test results have similar findings; that is, the fracture toughness of the rock is significantly reduced [2, 12–17], as shown in Figure 1.

It should be noted that most of the above test reports are the apparent fracture toughness of the rock, rather than its true value. This is due to the influence of the fracture process zone (FPZ) [18]. As Funatsu et al. [19] pointed out, the FPZ of the rock is related to the shape of the sample and the loading conditions, and the impact of FPZ should be fully considered when testing.
The influence of FPZ on fracture toughness of thermally damaged rock should be considered. Miao et al. [20] carried out a fracture test of granite from room temperature to 800°C and used DIC technology to observe the strain on the rock surface. The results showed that a significant FPZ appeared at the crack tip of the rock specimen after high-temperature treatment; and as the processing temperature increases, the size of the FPZ becomes larger and larger. Yin et al. [21] and Zuo et al. [22] also conducted similar experiments and reported the parameters of the double-k model for heat-damaged rocks [23]. However, the application of nonlinear fracture mechanics in the study of rocks after heat treatment is not sufficient, especially regarding the size effect [24]. Generally, the size of the specimen used in the laboratory is usually smaller than the actual structural size. Due to the size effect, the mechanical properties measured in the laboratory are directly used to estimate the actual structural strength, which is inaccurate. This may lead to serious deviations, especially in the case of using small-sized samples and a large FPZ. The size effect law is the key to reasonably applying the laboratory results to the actual structure [25, 26].

Therefore, in this paper, two temperature levels and three different sizes of granite specimens are used for fracture tests. The size effect of heat-damaged granite has been paid more attention. At the same time, the characteristics of mineral composition and microcrack of heat-damaged granite were observed.

2. Specimen Preparation and Investigation Methods

2.1. Specimen Preparation. A medium-fine-grained granite produced in Zhangqiu, China, was used for this test [24, 27–29]. The diameters of the three types of semicircular bend (SCB) specimens were 50 mm, 100 mm, and 200 mm, the thickness was 40 mm, and the initial crack length to radius ratio was 0.3. At the same time, 6 standard Brazilian split disc samples were produced [30–32]. The detailed geometry and dimensions of the test piece are shown in Figure 2 and Table 1.

The specimens in this test are divided into two groups equally: one group (high temperature group) is processed at 600°C and the other group (room temperature group) is not processed. Use the intelligent muffle furnace to slowly heat the granite specimens of the high-temperature group at a heating rate of 5°C/min. When it reaches 600°C, maintain the specimens at this temperature for 1 h [33] and then stop heating and let the specimens cool slowly [10, 34].

2.2. External Characteristics and Mineral Composition. High temperature usually changes the color of rocks; thus, the surfaces of the granite specimens were first observed, and the surface change phenomenon was observed in detail using an industrial microscope (Shenzhen Jingtuo Youcheng Technology Co. Ltd, JT-H6). Standard X-ray powder diffraction analysis (XRD, PANalytical X’Pert PRO) is used to determine the mineral composition of granite [14]. XRD patterns were recorded in the range 2–50°2θ with a speed of 6°/min, operating voltage of 40 kV, and current of 40 mA.

2.3. Rock Mechanical Properties Test

2.3.1. Fracture Toughness Testing. The three-point bending test was carried out on this batch of Zhangqiu granite specimens using the hydraulic servo-controlled testing machine model MTS 381.10. According to ISRM’s recommendations [28, 35], the SCB specimens were loaded in displacement control mode (0.05 mm/min), as shown in Figure 3(a). In order to avoid confusion, the fracture toughness value obtained by directly applying linear fracture mechanics is called the apparent fracture toughness $K_{ic}$. 

![Figure 1: Relationship between fracture toughness and temperature.](image-url)
According to the method recommended by ISRM [28], the specific formula is as follows.

The apparent fracture toughness \( K_{II} \) is

\[
K_{II} = Y \frac{P_{\text{max}} \sqrt{\pi a_0}}{2RB},
\]

\[
Y = -1.297 + 9.516 \frac{S}{2R} \left( 0.47 + 16.457 \frac{S}{2R} \right)^{\beta}
\]

\[
+ \left( 1.071 + 34.40 \frac{S}{2R} \right) \beta^2,
\]

\[
\beta = \frac{a_0}{R},
\]

where \( P_{\text{max}} \) is the peak load; \( a_0 \) is the initial crack length; \( Y \) is the dimensionless stress intensity factor; \( S \) is the pivot point spacing; \( R \) is the radius; and \( B \) is the thickness.

All SCB specimens are tested using the same span to diameter ratio \( (2R) \) of \( S/2R = 0.8 \).

### Table 1: Detailed geometric dimensions of the specimen.

| Specimen type | Diameter (mm) | Thickness (mm) | Crack length, \( a_0 \) (mm) | Crack length ratio \( (a_0/R) \) | Number of specimens |
|---------------|---------------|----------------|-----------------------------|---------------------------------|-------------------|
| SCB-S         | 50            | 40             | 7.5                         | 0.3                             | 6                 |
| SCB-M         | 100           | 40             | 15                          | 0.3                             | 6                 |
| SCB-L         | 200           | 40             | 30                          | 0.3                             | 6                 |
| BD            | 50            | 25             | —                           | —                               | 6                 |

According to the method recommended by ISRM [28], the specific formula is as follows.

2.3.2. Tensile Strength. Brazilian disc splitting tests were performed on two types of granite (RT and 600) to obtain their tensile strength [30] (Figure 3(b)).

\[
\sigma_t = \frac{2P_{\text{max}}}{\pi DB}
\]

where \( \sigma_t \) is the tensile strength; \( P_{\text{max}} \) is the peak load; \( D \) is the diameter; and \( B \) is the thickness.
The same hydraulic servo-controlled MTS (MTS 381.10) testing machine as the three-point bending test was used to perform the Brazilian splitting test. The maximum load of specimen fracture was recorded by displacement controlled loading mode.

2.4. Optical Microscope. The two types of granite samples after the test were made into thin slices to observe the changes in the mineral composition and internal structure of the granite; and these thin slices were observed under a petrographic microscope (Zeiss Axioskop 40). Because ordinary thin slices are not easy to identify microcracks and other structures, in order to find these structures clearly and intuitively, the granite samples were impregnated with epoxy resin in a vacuum box; and these epoxy resins are added with blue dye to make them more eye-catching. In this way, the pore and crack structure could be analyzed very clearly and intuitively, and the porosity and other characteristics of the rock could be evaluated [36, 37].

2.5. SEM. The SEM analyses were performed on the sections of granite samples using the field emission gun scanning electron microscope (Hitachi SU8010), equipped with the energy-dispersive spectroscopy (HORABA Ltd, X-MaxN50011). The system operated at 20 kV accelerating voltage in a high-vacuum mode. The sections of granite samples were coated with gold [38]. The microstructure of granite fracture can be observed directly and conveniently by SEM [39]. EDS is used to observe changes in the composition of minerals in rocks.

3. Experimental Results

3.1. External Characteristics. The color of granite samples after heat treatment is the most intuitive indicator of the impact of the treatment. Comparing the specimens, it was found that the specimen surface color changed from dark to light after 600°C, with brownish-red spots, as shown in Figures 4(a) and 4(b). This phenomenon could be observed more clearly after magnified observation through an industrial microscope (Figures 4(c) and 4(d)). In many experiments, a color change of rocks after high-temperature treatment has been reported [40–42], which may be due to the dehydration and oxidation of the iron-magnesium minerals. The micro area EDS results (Figure 5) of this experiment also indicate the presence of Fe element.

3.2. XRD Analysis Results. According to XRD diffraction results (Table 2), it can be seen that the main constituent minerals of granite are quartz, biotite, plagioclase, and hornblende, among which quartz, biotite, and plagioclase make up the highest proportions. By comparing the XRD results of the unheated and 600°C high-temperature treatments, it can be found that the main minerals constituting the granite have not changed significantly (Table 2 and Figure 6). This is consistent with the high-temperature exposure of granite reported by Chen et al. [41] and Shang et al. [43].

3.3. Rock Mechanical Properties

3.3.1. Fracture Toughness Test Results. According to the test results of SCB, the maximum load was brought into the fracture toughness expression (1) and the fracture toughness value of granite could be calculated as listed in Table 3. In order to display the results more intuitively, a box plot is shown in Figure 7.

The fracture test results show that the fracture toughness of granite specimen decreases significantly after 600°C, as shown in Table 3 and Figure 7. For example, the average fracture toughness of SCB specimens with a radius of 50 mm measured at room temperature was 1.71 MPa·m^{0.5}. The average value of fracture toughness after 600°C is only 0.59 MPa·m^{0.5}, a decrease of 65.6%. This is consistent with many test results of granite. For example, Nasseri et al. [10] tested granite at 800°C and found that its fracture toughness was only 10% of the original granite. Talukdar et al. [11] reported that the fracture toughness values of the three types of crystalline rocks after 600°C were greatly reduced, exceeding 50%. At the same time, the fracture toughness results
of SCB specimens with different radii after untreated and high-temperature treatment were carefully compared (see Figure 7). It can be observed that as the diameter of the specimen increases, the fracture toughness value of untreated granite does not have a significant size effect. However, the fracture toughness value of granite after 600°C treatment increases obviously with the radii of specimens. Thus, the granite after high-temperature treatment shows a significant size effect, while the untreated granite shows no such phenomenon. Di Luzio et al. [25] also reported that concrete after heat loss also showed a significant size effect.

3.3.2. Tensile Strength Results. According to expression (2), the indirect tensile strength values of granite are calculated as shown in Table 4. The results of tensile strength agree with the fracture test. The average tensile strength of the granite treated at 600°C was significantly reduced from $\sigma_{t}^{RT} = 11.90$ MPa to $\sigma_{t}^{600^\circ C} = 4.19$ MPa at room temperature, a 64.8% decrease. This is very similar to the 65.6% reduction in the fracture test.

Table 2: XRD analysis results.

| Mineral composition (%) | Quartz | Biotite | Plagioclase | Hornblende | Orthoclase | Magnetite | Other | Total |
|-------------------------|--------|---------|-------------|------------|------------|-----------|-------|-------|
| Granite at RT           | 24.3   | 26.7    | 33.4        | 4.4        | 9          | 1.2       | 1     | 100   |
| Granite at 600°C        | 22     | 24.4    | 30.5        | 4          | 7.8        | 1.3       | 10    | 100   |

Figure 4: The surface color of granite (a, c) RT; (b, d) 600°C; (c, d) USB Industrial Microscope (Shenzhen Jingtuo Youcheng Technology Co. Ltd, JT-H6).

Figure 5: The result of energy-dispersive spectroscopy (EDS).

3.4. Microstructure

3.4.1. Results of Optical Microscope. After the fracture test was completed, the two types of granite (RT and 600°C) were placed in a vacuum chamber and impregnated with epoxy resin and then processed into slices which are 30 microns thick. In this way, the microcracks will be very easy to identify, as shown in Figure 8. After being impregnated with epoxy resin, the microcracks of the granite can be clearly observed. The mineral grains of the untreated granite (Figure 8(a)) are in close contact with only a few microcracks. The granite treated at 600°C is filled with a large amount of blue resin around the particles (Figure 8(b)), which means that many intercrystalline cracks have appeared. Obviously, the number and width of microcracks in the granite treated at 600°C are significantly increased compared with RT. To quantitatively analyze microcracks, ImageJ was used to identify microcracks filled with blue resin [44]. The software can accurately identify microcracks and highlight them in red, as shown in Figures 8(c) and 8(d).
In order to quantify the changes in the microstructure of granite, ImageJ software was used to identify and calculate the microcracks on the thin slices. The ImageJ analysis results show that the porosity (surface porosity) of the granite sample at RT is only 0.6%, and this value becomes 4.7% after 600°C, and the porosity increases significantly.

3.4.2. Results of SEM. There is a wealth of information on the fracture of the rock, and there are important traces of the fracture process of the rock, such as steps and river patterns. These important microscopic features can help us analyze the causes of rock failure during the fracture process [38]. SEM is an ideal tool for observing these characteristics, so we used SEM to observe and record the microscopic morphologies of the two types of granite fractures in detail, as shown in Figure 9. The particles of the original granite are flat and smooth, the steps and river patterns on the surface of the particles are not obvious, and there are no cracks between and within the particles, as shown in Figures 9(a) and 9(c). After treatment at

| Heat treatment temperature | Specimen no. | R (mm) | B (mm) | $\alpha_0$ (mm) | Effective structural dimension, $D$ (mm) | Load, $P_{\text{max}}$ (N) | $K_f$ (MPa·m$^{0.5}$) |
|----------------------------|--------------|--------|--------|----------------|----------------------------------------|--------------------------|-------------------|
| RT                        | S01          | 24.97  | 40.09  | 7.6           | 4.99                                   | 4472.4                  | 1.65              |
|                           | S02          | 24.96  | 40.15  | 7.56          | 4.99                                   | 5088.4                  | 1.87              |
|                           | S03          | 25.03  | 40.06  | 7.87          | 5.01                                   | 4389.9                  | 1.62              |
|                           | S04          | 48.29  | 40.13  | 15.03         | 9.66                                   | 5917.0                  | 1.54              |
|                           | S05          | 50.01  | 40.12  | 15.78         | 10.00                                  | 6291.6                  | 1.64              |
|                           | S06          | 49.96  | 40.02  | 15.07         | 9.99                                   | 6498.8                  | 1.69              |
|                           | S07          | 99.13  | 39.82  | 31.43         | 19.83                                  | 9992.3                  | 1.84              |
|                           | S08          | 99.15  | 40.04  | 30.73         | 19.83                                  | 9801.6                  | 1.80              |
|                           | S09          | 99.24  | 40.36  | 30.97         | 19.85                                  | 9658.3                  | 1.78              |
| 600°C                     | S10          | 25.13  | 40.23  | 7.63          | 5.03                                   | 1156.9                  | 0.43              |
|                           | S11          | 25.14  | 40.33  | 7.67          | 5.03                                   | 1148.5                  | 0.42              |
|                           | S12          | 24.74  | 40.33  | 7.59          | 4.95                                   | 1708.7                  | 0.63              |
|                           | S13          | 49.36  | 40.21  | 15.43         | 9.87                                   | 1995.8                  | 0.52              |
|                           | S14          | 49.25  | 40.35  | 15.71         | 9.85                                   | 2090.1                  | 0.54              |
|                           | S15          | 50.07  | 40.23  | 15.92         | 10.01                                  | 2100.2                  | 0.55              |
|                           | S16          | 99.01  | 39.57  | 30.8          | 19.80                                  | 3210.6                  | 0.59              |
|                           | S17          | 99.55  | 40.52  | 30.44         | 19.91                                  | 3741.5                  | 0.69              |
|                           | S18          | 99.39  | 40.38  | 30.24         | 19.88                                  | 5086.9                  | 0.94              |

Note. The effective structural dimension $D$ is one of the parameters of the size effect law (SEL); see expression (7) for details (S01–S09, RT; S10–S18, 600°C).
Table 4: The Brazilian splitting test results.

| Specimen no. | R (mm) | B (mm) | Heat treatment temperature (°C) | Peak strain (%) | Load, $P_{\text{max}}$ (N) | $\sigma_t$ (MPa) |
|--------------|--------|--------|---------------------------------|-----------------|--------------------------|----------------|
| B01          | 50.31  | 24.81  | RT                             | 0.80            | 22538.4                  | 11.50          |
| B02          | 50.3   | 24.74  | RT                             | 0.71            | 25042.7                  | 12.80          |
| B03          | 50.3   | 24.83  | RT                             | 0.85            | 22606.7                  | 11.50          |
| Average      |        |        |                                 | 0.79            |                          | 11.90          |
| B04          | 50.54  | 24.94  | 600                            | 0.79            | 10777.5                  | 5.44           |
| B05          | 50.58  | 24.8   | 600                            | 0.79            | 8977.9                   | 4.56           |
| B06          | 50.56  | 25.02  | 600                            | 0.78            | 5086.9                   | 2.56           |
| Average      |        |        |                                 | 0.79            |                          | 4.19           |

Note. B represents the Brazilian splitting test piece and RT indicates room temperature, without heat treatment.

Figure 7: Box diagram of apparent fracture toughness for granite samples untreated and after high-temperature treatment.

Figure 8: Continued.
600°C, the surface of granite mineral particles becomes rough, with obvious steps, river patterns, and cracks. This is similar to the experimental results of Talukdar et al. [11] on crystalline rocks. Zuo et al. [38] pointed out that temperature is an important factor affecting the fracture morphology of rocks. After high temperature, the rock failure mechanism changes from brittleness to brittleness and ductility.

4. Discussion

4.1. Size Effect Analysis. In order to solve the limitations of linear elastic mechanics, Båzant [24] proposed the famous size effect law (SEL) to study quasi-brittle materials. The specific formula is as follows:

$$\sigma_n = c_n \left[ \frac{EG_f}{g(\alpha_c)c_f + d \gamma(\alpha_c)} \right]$$

$$\sigma_n = \frac{P}{t \cdot d}$$

$$K = \frac{P}{b \sqrt{d \gamma(\alpha)}}$$

where $\sigma_n$ is the nominal strength, $c_n$ is a coefficient, $t$ is the thickness, $d$ is the height, $P$ is the maximum load, $E$ is the elasticity modulus, $G_f$ is the fracture energy, $c_f$ is the...
effective fracture process zone length, and $\sqrt{g(a)}$ is a form function associated with the specimen geometry. For some commonly used geometric forms, the specific expressions can be obtained by referring to the stress intensity factor manual. $K$ is fracture toughness, which can be derived from expressions (3)–(5):

$$K_{if} = \frac{K_{IC}}{\sqrt{1 + \left(g'\left(a_0\right)\frac{\alpha_f}{g\left(a_0\right)}d\right)}}$$  

(6)

where $K_{if}$ is the fracture toughness calculated by LEFM and $K_{IC}$ is the fracture toughness of an infinitely large specimen.

Expression (6) can be rewritten as

$$\begin{cases} \frac{1}{K_{if}^2} = \frac{1}{K_{IC}^2} \frac{1}{D} + \frac{1}{K_{IC}^2}, \\ D = \frac{g\left(a_0\right)}{g'\left(a_0\right)}d, \end{cases}$$

(7)

where $D$ is the effective structural dimension. It is independent of the geometry of specimens.

The SCB experiment results (Table 3) are brought into expression (7) for linear fitting, and the results are shown in Figure 10. The linear fitting equations for untreated and high-temperature treatment are as follows:

$$\begin{cases} \frac{1}{K_{if}^2} = 1.79765E^{-16} \frac{1}{D} + 3.23138E^{-13} \quad \text{RT}, \\ \frac{1}{K_{if}^2} = 1.57411E^{-14} \frac{1}{D} + 1.50799E^{-12} \quad \text{600°C}. \end{cases}$$

(8)

According to the results of linear regression analysis, the size effect parameters $K_{IC}$ and $c_f$ are obtained, which are listed in Table 5.

According to the results in Figure 10 and Table 5, it can be clearly seen that, for the untreated granite, $K_{IC}^{RT} = 1.76$ MPa-m$^{0.5}$ and $c_f^{RT} = 0.56$ mm. $c_f^{RT}$ is very small, so the size effect of untreated granite is not significant, according to SEL. However, for the granite treated at 600°C, $K_{IC}^{600°C} = 0.81$ MPa-m$^{0.5}$ and $c_f^{600°C} = 10.4$ mm. Obviously, after 600°C treatment, the effective fracture process zone length $c_f$ of granite has increased significantly, so the size effect of thermal damaged granite is significant.

To more intuitively show the influence of $c_f$, the apparent fracture toughness values are normalized, as shown in Figure 11. $c_f$ is a key parameter to control the size effect as can be seen from the figure. The more significant the size effect is, the larger $c_f$ is. The results of this experiment also show that the size effect law of Bažant can well describe the size effect of granite after high-temperature treatment.

4.2. Effect of Temperature on $c_f$ According to the size effect law of Bažant, the size effect parameters of untreated and high-temperature treated granite samples are calculated. For untreated granite, $c_f^{RT} = 0.56$ mm, while for granite treated at 600°C, $c_f^{600°C} = 10.4$ mm. The $c_f$ value increases significantly, which indicates that 600°C has a significant effect on the FPZ length of granite. The fracture process zone is closely related to the microcracks at the crack tip. High-temperature treatment
conditions will increase the microcracks of the rock, meaning that the FPZ of the rock increases after high temperature. These microcracks may be the main reason for the increase in FPZ.

After high temperature, there is more intuitive evidence for the increase of FPZ length of granite. Miao et al. [20] used DIC technology to observe the full-field displacement and strain of the granite surface and measured the FPZ length of the granite at different temperatures. The results show that the length of the FPZ increases along with processing temperature. Similarly, Yin et al. [21] used acoustic emission to monitor the three-point bending fracture process of granite treated at different temperatures. The results show that acoustic emission events occur only when the load is close to the peak for granite at room temperature (25°C); this is a brittle fracture. However, the granite treated at high temperature (above 500°C) exhibited a large number of acoustic emission events at lower loads (one-third of the maximum load), indicating that crack propagation occurred before reaching the maximum load.

These are consistent with the results of this test and prove that high-temperature conditions increase the length of the fracture process zone ($c_f$) on rock. Both $c_f$ and $D$ have a significant effect on the size effect. The $K_d/K_{IC}$ values of the effective structure size $D$ 0.005–0.2 m and $c_f$ 0.01–0.2 m are calculated according to expression (6). The $K_d/K_{IC}$ ratio decreases with $c_f$ and increases with $D$, as shown in Figure 12.

Therefore, for small samples and a large FPZ, the apparent fracture toughness value $K_{IF}$ will be significantly underestimated. For example, for the heat-damaged granite samples in this test, the $K_d$ values of SCB specimens with $R = 25$ mm are significantly smaller than the $K_{IC}$ value ($K_d = 59 = 0.493$ MPa$\cdot$m$^{0.5}$; $K_{IC} = 0.81$ MPa$\cdot$m$^{0.5}$). In actual engineering, the core is often obtained through small boreholes to produce the specimens, and linear fracture mechanics are used to calculate the apparent fracture toughness parameters of the rock. However, the estimation of the fracture toughness of the actual structure often causes a large deviation. This will adversely affect the design and safety assessment of actual engineering structures, such as the exposure of rocks to high temperatures.

4.3. Estimate $K_{IC}$ The size effect law is inconvenient for the preparation of specimens of different sizes, which hinders the application of the size effect law. If only one size of specimens can be used, it is best that a single specimen can be used to estimate the size effect parameters, which will be very convenient and useful. Cedolin and Cusatis [45] established an equation for the relationship between tensile strength and size effect parameters, which can be used to easily estimate the size effect parameters. The equation is as follows:

$$f_t = \sqrt{\frac{0.39 E G_f}{c_f}}.$$  \hspace{1cm} (9)

The above equation can be rewritten as

$$c_f = 0.39 \left(\frac{K_{IC}}{f_t}\right)^2.$$  \hspace{1cm} (10)

Combined with the size effect law (expression (6)), the following expression can be obtained:

$$K_{IC} = \frac{1}{\sqrt{\left(1/K_{IC}^2\right) - \left(0.39/f_t^2\right)^4}}.$$  \hspace{1cm} (11)

To verify expression (11), the results of this experiment are brought into the expression as shown in Figure 13 and Table 6. The results show that expression (11) can accurately predict the $K_{IC}$ value of heat-damaged granite, but it shows a large deviation for untreated granite. As Cedolin and Cusatis [45] pointed out, expression (11) can only be applied when the ratio of the FPZ length to the size of the specimen reaches a certain range. Since the fracture process zone of untreated granite is very small, expression (11) is no longer applicable. For the heat-damaged rock, if its tensile strength is known, you can try to use expression (11) to estimate its $K_{IC}$ value. If the tensile strength of the rock is unknown, the Brazilian splitting test can be used to estimate its tensile strength. Undoubtedly, more tests are needed to verify the above methods. It should also be noted that expression (11) estimates only one of the parameters of the size effect law, $K_{IC}$, while the other parameter, $c_f$, is not recommended due to its large dispersion.

4.4. From Brittleness to Ductility. Bažant’s brittleness number [46] is

$$\beta = \frac{g(a_0)}{g(a_0) - c_f} \frac{d}{\sigma_f} \frac{D}{c_f}.$$  \hspace{1cm} (12)
where \( D \) is the effective structural dimension.

Bážant’s brittleness number of three sizes of specimens was calculated using (12), and the results are shown in Figure 14. Obviously, the brittleness of thermally damaged granite is significantly reduced. According to Bážant’s standard [47], when \( \beta > 10 \), the structure can be analyzed by elastic fracture mechanics; when \( 0.1 < \beta < 10 \), the structure needs to be analyzed according to nonlinear fracture mechanics. Therefore, for 600°C treated granite samples, nonlinear fracture mechanics should be used for small specimens.

High temperatures usually reduce the brittleness of rock fractures and increase ductility. The \( c_f \) value can also measure the degree of brittleness or ductility. Materials with larger process zones have more ductility [25]. The size effect parameter \( c_f \) increases significantly after the granite samples are treated at 600°C, indicating that the ductility of the granite increases. The experiments of Yin et al. [21] and Miao et al. [20] also showed that the ductility of thermally damaged granite increased.

At the same time, changes in microscopic features also confirm the increase in ductility. The scanning electron microscopy (SEM) results of fracture morphology shown in Figure 9 show that, in the thermally damaged granite samples, the cleavage becomes rough and the river pattern and step pattern are obvious, similar to the test results observed by Zuo et al. [39]. This change in fracture micromorphology indicates that the fracture behavior of granite changed from brittleness to ductility.

### Table 6: Comparison of \( K_{IC} \) estimated according to expression (11) and \( K_{if} \)

| Effective structural dimension \( D \) (mm) | \( K_{if} \) (MPa·m\(^{0.5}\)) | \( K_{IC} \) estimated according to expression (11) (MPa·m\(^{0.5}\)) |
|------------------------------------------|-----------------------------|-----------------------------------------|
| S01                                      | 4.99                        | 1.65                                   |
| S02                                      | 4.99                        | 1.87                                   |
| S03                                      | 5.01                        | 1.62                                   |
| S04                                      | 9.66                        | 1.34                                   |
| S05                                      | 10.00                       | 1.64                                   |
| S06                                      | 9.99                        | 1.69                                   |
| S07                                      | 19.83                       | 1.84                                   |
| S08                                      | 19.83                       | 1.80                                   |
| S09                                      | 19.85                       | 1.78                                   |
| S10                                      | 5.03                        | 0.29                                   |
| S11                                      | 5.03                        | 0.29                                   |
| S12                                      | 4.95                        | 0.43                                   |
| S13                                      | 9.87                        | 0.36                                   |
| S14                                      | 9.85                        | 0.37                                   |
| S15                                      | 10.01                       | 0.37                                   |
| S16                                      | 19.80                       | 0.41                                   |
| S17                                      | 19.91                       | 0.46                                   |
| S18                                      | 19.88                       | 0.63                                   |

Figure 13: \( K_{IC} \) estimated according to expression (11) (the dotted line represents the size effect parameter \( K_{IC} \), while the blue- and red-dotted lines represent the untreated granite and 600°C treated granite, respectively).
4.5. The Failure Mechanisms. The damage of granite due to high temperature is significant. The granite after the test is made into thin slices, and the optical microscope can be used to intuitively observe that the granite after high temperature has obvious microcracks. The number and width of microcracks are significantly increased compared to the original granite, as shown in Figure 8. This phenomenon can also be observed in the scanning electron microscope image (Figure 9).

During the heating process, the expansion coefficient and thermal conductivity of different grains of granite are different. For example, quartz has a higher thermal expansion coefficient than other minerals. At the same time, 600°C can cause the conversion of low quartz to high quartz (α-β) [46], which further aggravates the expansion of quartz particles. Therefore, during heating, due to inconsistent thermal strain, microcracks are likely to occur between the crystal grains. This is consistent with the observation in Figure 8. At the same time, in the cooling stage after the heating is completed, the crystal grains begin to shrink. This process is just the opposite of heating. Due to the different expansion coefficients, the deformation of different crystal grains during cooling is also different. Therefore, cooling also causes microcracks. This is the main failure mechanism of granite thermal damage.

The failure of the rock is closely related to the initiation and propagation of microcracks. There are a lot of microcracks in the thermally damaged rock. It is precisely because of their effects that an FPZ is formed at the crack tip. Therefore, the fracture process zone size of thermally damaged granite is significantly larger than that of original granite. The nonlinear characteristics of heat-damaged granite, such as size effect, become very obvious due to the influence of FPZ.

5. Conclusions

The obtained results of the research lead to the following conclusions:

(1) High temperature significantly weakens the fracture toughness value of granite.

The fracture toughness value of thermally damaged granite is significantly lower than that of untreated granite. For example, the average fracture toughness of SCB specimens ($R = 50\text{ mm}$) measured at room temperature was 1.71 MPa·m$^{0.5}$, while the average fracture toughness after a temperature of 600°C was only 0.59 MPa·m$^{0.5}$, a decrease of 65.6%.

(2) The granite has a significant size effect after treatment at 600°C.

No obvious size effect was found in untreated granite; however, the size effect of granite after 600°C treatment increased significantly. The fracture toughness value of thermally damaged granite may be underestimated if linear elastic mechanics and small-sized specimens are applied directly.

(3) High-temperature conditions have a significant effect on the size effect parameters of granite.

The granite treated at 600°C had a nonnegligible fracture process area compared with untreated granite. For untreated granite, $c_{RT} = 0.56\text{ mm}$, while for granite treated at 600°C, $c_{600°C} = 10.4\text{ mm}$. The value of $c_f$ increased significantly. The size effect law of Bažant was able to reasonably describe the fracture characteristics of thermally damaged granite.

(4) The microcracks of thermally damaged granite increase and the ductility increases.

After 600°C, the microcracks of granite increased, the fracture morphology became rough, and ductility increased. The composition of thermally damaged granite did not change significantly, but the microcracks increased significantly, which may be directly related to the increase of $c_f$.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

The authors declare that there are no conflicts of interest.

Authors’ Contributions

Zhongping Guo contributed to the conception of the study. Jian Li contributed significantly to design and wrote the manuscript. Yongqi Song performed the data analyses.
Chengqian He and Fuyu Zhang helped perform the experiment.

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