

Selective comminution and grinding mechanisms of spent carbon anode from aluminum electrolysis using ball and rod mills

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Abstract: The recovery of spent carbon anode (SCA) materials plays important roles in environment protection and resources recycling, while this cannot be efficiently achieved without liberation. In this study, grinding characteristics of spent carbon anode from aluminum electrolysis in both ball mill and rod mill were analyzed, and compared based on the utilization of the selective Fuerstenau upgrading curves. In addition, the different grinding mechanisms of ball and rod milling were evaluated by analyzing the shape factor and surface roughness of the ground samples. Results of mineralogical characterizations indicated that carbon particles ($d_{50} = 46.86 \mu\text{m}$) presented in the SCA were closely associated with cryolite particles. At 5 min grinding time, the maximum selective comminution factor (β) values of ball milling and rod milling were 2.00 and 1.63, respectively, indicating a higher degree of selective comminution of SCA was achieved from ball milling. Comparisons of the valuable component content (c_v , cum) of $-125 \mu\text{m}$ ground particles and the shape characterizations of $74\text{--}125 \mu\text{m}$ ground particles generated from ball and rod milling manifest that a direct relationship exists between the degree of selective comminution and the grinding mechanism.

Keywords: selective comminution, ball mill, rod mill, spent carbon anode, grinding mechanism

1. Introduction

The International Aluminum Institute (Institute, 2020) reported that the global production of primary aluminum tapped from electrolytic cells (or pots) during the electrolytic reduction of metallurgical alumina (aluminum oxide) was 63,697 thousand metric tons (t) in 2019. Further, China is the largest producer of aluminum (35,795 thousand t in 2019), accounting for 56.20% of global production. Spent carbon cathode (SCC) is the carbon cathode of the Hall-Héroult cell, eroded by high-temperature molten salt and metallic sodium (Grjotheim et al., 1982). Approximately 24–30 kg of SCC is generated from 1 t of aluminum production (Holywell and Breault, 2013). In addition, the carbon anode of aluminum reduction cell is damaged and dropped in the reduction cell due to the selective oxidation of anode, erosion of electrolytes, secondary reaction in reduction process and mechanical damage during operation (Grjotheim et al., 1982; Li et al., 2015). This process results in the presence of the floating carbon residues on the surface of the aluminum reduction cell, which is separated by artificial fishing (Xu et al., 2009). Approximately 5–15 kg spent carbon anode (SCA) is continuously generated from 1 t of aluminum production (Chen et al., 2009). Thus, it has been estimated that the world's aluminum smelters generated 1,528–1,910 thousand t of SCC and 318–955 thousand t of SCA in 2019. It is commonplace for SCC and SCA to be disposed of in landfills, posing a risk of soil contamination due to

the leaching of soluble fluoride and highly toxic cyanide (Sleap et al., 2015). SCC and SCA have been listed in the national list of hazardous wastes by China in 2016 (Yang et al., 2020b). Moreover, SCC and SCA also contain carbon, fluoride, alumina, cryolite, aluminosilicate, and a trace of cyanide, which is a recoverable resource (Flores et al., 2017). Accordingly, from the perspective of resource recycling and environmental protection, it is important to achieve effective green treatment and highly efficient recoveries of SCC and SCA.

The importance of comprehensive use of solid waste (SCC and SCA) in aluminum smelters is in the recovery of valuable materials (carbon and cryolite) and the harmless treatment of fluoride and cyanide. There are two major categories of SCC treatment methods: pyrometallurgical and hydrometallurgical (Bishoyi, 2015; Tropenauer et al., 2019). Pyrometallurgical processes is used for the industrial recycling of solid waste because they facilitate the harmless treatment of solid waste to remove toxic fluoride and cyanide (Bruno, 2003; Chai et al., 2016; Personnet, 2013; Xie et al., 2020; Zhou, 2015; Zou et al., 2018). However, valuable components such as carbon and cryolite are usually lost in slag during the process (Flores et al., 2017; Shi et al., 2012; Yang et al., 2020a; Zou et al., 2018). Moreover, high energy consumption, high equipment investment, and environmental pollution also hinder the wide application of pyrometallurgical processes. Consequently, various hydrometallurgical processes have been developed to treat and recycle SCC and SCA, including flotation and chemical leaching (Tropenauer et al., 2019; Yang et al., 2020b).

Flotation is an effective method of selectively separating hydrophobic carbon from hydrophilic cryolite following the comminution and grinding processes of SCC and SCA. As shown in Table 1, a concentrate with 96.10% carbon content and tailings with 4.10% carbon content can be obtained using a four-stage flotation circuit; however, further upgrading by flotation is challenging. SinghMaitra (1986) introduced a method for recovering carbon and fluoride (while also reducing pollution) by recycling flotation tailings of aluminum smelters. Tropenauer et al. (2019) developed a technique for combing flotation and carrying out chemical treatment to produce purified carbon and silicate fractions from industrial waste (primary aluminum production). It is worth noting that while chemical leaching (acid, soluble aluminum salt solution, and caustic) is more effective than flotation techniques for purifying SCC and SCA (Xiao et al., 2018; Yang et al., 2020b), it can cause severe environmental problems.

Table 1. Summary of flotation studies of SCC and SCA

Solid waste	Carbon content of raw ore (%)	Grinding Method	Number of flotation stages	Carbon content (%)		Ref.
				Concentrate	Tailings	
SCC	71.20	Ball mill	4	96.10	4.10	(Li et al., 2013)
	49.10			85.30	19.80	(Li et al., 2013)
	36.10			73.10	26.60	(Li et al., 2013)
	51.33	Ball mill	3	91.20	5.50	(Lu and Qiu, 1997)
SCA	30.50	Rod mill	3	94.59	1.01	(Mei et al., 2016)

Prior to flotation, SCC and SCA are commonly comminuted and ground to disassociate carbon from cryolite (see Table 1). While grinding-flotation circuits have been widely used for SCC and SCA treatment, differences in selective comminution between ball and rod mills have not yet been analyzed systematically. Selective comminution is a promising method for reducing specific energy consumption. Moreover, appropriate selective comminution preconcentrates the ore and removes the coarse liberated gangue before entering the subsequent milling circuit, which is a potential strategy for substantial energy saving (Hesse et al., 2017; Lowes et al., 2018). Selective comminution is the result of a comminution system that comprises appropriate comminution parameters (such as operational parameters and the design of the comminution machine) and inherently suitable material properties (Hesse, 2014; Stepanov et al., 1991). The different distributions of valuable and waste minerals (as a function of particle size) are commonly employed to characterize selective comminution qualitatively (Hesse et al., 2017). Hesse et al. (2017) defined the parameters to quantify selectivity (S) and η_{ore} , where SZ quantified the change in selectivity between the feed material and product. They confirmed that selective comminution requires a systemic approach – comminution behavior of the ore under a certain load, type of comminution machine, and number of operational parameters of the system. A Fuerstenau

upgrading diagram with recovery plots is another method for estimating selective comminution, which has been used to study the selective comminution of quartz-topaz glimmer Greisen ore (Leißner et al., 2014), iron ore (Reichert et al., 2015), and spent lithium-ion batteries (Widijatmoko et al., 2020).

Some researchers have paid attention to the differences in the grinding mechanisms of ball and rod mills, which commonly results in the following variations: the shape properties of ground particles (Hicyilmaz et al., 2004; Ulusoy, 2008b; Ulusoy and Kursun, 2011), the residence time distribution (RTD) of particles (Gupta and Patel, 2015), the size distribution of ground products (Foszcz et al., 2018), and the flow characteristics of the tracer particles (dolomite, magnetite, and galena) (Abouzeid and Fuerstenau, 2012). AbouzeidFuerstenau (2012) reported that rod mill products have a narrow size distribution compared to those from a ball mill, which is partly attributable to the plug flow mode of the material transport in the rod mill and partly to the well-known bridging action of the rods. Foszcz et al. (2018) found that ball mills are more efficient in terms of incrementally small-sized fractions compared to rod mills. Moreover, based on probability theory, Malyshev et al. (2017) revealed that rod mills have superior capacity for comminuting large particles compared to ball mills due to the combined effect of steric and activation factors. FoszczGawenda (2012) demonstrated that ball mills were optimum for fine grinding (<0.1 mm) while rod mills were efficient for thicker and harder particles in the feed. It is believed that the line load on particles in rod mills helps to break hard materials (Gupta and Yan, 2016). By comparison, the point load on the particles in ball mills is conducive to the attrition process (Bu et al., 2019b; Bu et al., 2020a). Abrasion and attrition result in rounder (more spherical) ball-milled particles with smoother surfaces than rod-milled particles (Bu et al., 2019a; Hicyilmaz et al., 2004; Ulusoy, 2008b; Ulusoy and Kursun, 2011). However, it is found that there is no related study on the difference of selective comminution of SCA between ball and rod mills.

Accordingly, the objective of this study was to compare the selective comminution of SCA using ball and rod mills. The degree of selective comminution was estimated based on Fuerstenau upgrading curves. In particular, the different degrees of selective comminution were associated with the different breakage mechanisms of ball and rod milling. In addition, it is notable that the comparison between ball and rod milling is only a preliminary study because of the lack of the optimization of the operating conditions for ball and rod milling, such as wet and dry grinding, rotational speed of grinding devices, and the filling fractions of grinding medium and powder samples. The effects of those factors on the selective comminution of spent carbon anode from aluminum electrolysis will be presented in the future research.

2. Materials and methods

2.1. Sample preparation

A spent carbon anode (SCA) sample was obtained from an aluminum and power company (Shandong Weiqiao Aluminum & Electricity Co., Ltd., Binzhou, Shandong Province, China) in the form of -1 mm powder. The SCA sample was mixed by thorough coning and quartering, and representative sub-samples were obtained for sieving, X-ray diffraction (XRD), X-ray fluorescence (XRF), and grinding tests.

2.2. Sieving test

The sample was sieved using a set of 1000, 500, 250, 125, 74, and 45 μm sieves (ASTM E11). The industry standard YS/T 273.2-2006 (for the chemical analysis and physical properties of cryolite, part 2: determination of ignition loss) was used to evaluate the approximate loss on ignition (LOI, %). This value of LOI was then used to represent the approximate carbon content of the products obtained from the sieve analysis.

2.3. Grinding test

The grinding tests were carried out in dry conditions. Ball milling tests were performed using a QM-5 laboratory-scale ball mill (Changsha Tianchuang Powder Technology Co., Ltd., Changsha, China). The diameter and the length of the drum for the ball mill were 12.5 and 16.0 cm, respectively. A 2.0 L stainless steel cylinder was used as the drum in the ball milling tests, in which 2.22 kg of stainless steel balls with

diameters of 1.20, 0.90, 0.62, and 0.55 cm were used. The weight percentages of the stainless steel balls were 33.75%, 31.40%, 18.37%, and 16.48%, respectively. The operational speed was 115 rpm, and a 200 g mass of -1 mm feed sample material was used for the ball milling tests.

The critical rotational speed for centrifuging in ball and rod milling can be calculated as follows (Abouzeid and Fuerstenau, 2012):

$$N_c = \frac{30}{\pi} \sqrt{\frac{2g}{D-d_r}} \quad (1)$$

where d_r is the diameter of the rod. For this study, the critical rotational speed for the rod mill was 138–140 rpm (dependent on rod diameter) with a feed size of -1 mm and operating speed of 50 rpm, respectively.

Compared to the commonly used Eq. 1, the following Eq., related to a variety of parameters (including the ball diameter d , the volume fill fraction ϕ of material within the tumbler, and the material angle β_s of repose), was applied to calculate the critical rotational speed for ball milling (Bu et al., 2019b; Juarez et al., 2011; Kallon et al., 2011; Walton and Braun, 1993):

$$N_c = \frac{30}{\pi} \sqrt{\frac{2g}{(D-d_b)\sin\beta_s\sqrt{1-\phi}}} \quad (2)$$

here, D and d_b are the diameters of the drum and ball, respectively. The critical rotational speed for the ball mill was 183–188 rpm, with the variation caused by different ball diameters.

Rod milling tests were conducted using a KJXMB III laboratory-scale rod mill (Yancheng Kejie Test Instrument Factory, Yancheng, Jiangsu Province, China) equipped with a 1.2 L stainless steel cylinder ($D \times L$: 10.5 cm \times 14.0 cm). The grinding tests were carried out using rods of 1.1, 1.3, and 1.5 cm diameters and the weight percentages of the stainless steel rods (length = 14 cm) were 30.37%, 34.96%, and 34.67%, respectively. During the rod milling tests, the operational speed of the rod mill, the weight of the stainless steel rods, and the weight of the feed sample (-1 mm) were 50 rpm, 3.16 kg, and 200 g, respectively.

2.4. XRD and XRF tests

The sample was ground to a fine powder (<200 mesh size) using an XPM- ϕ 120 \times 3 three-headed grinding machine (Nanchang Source of Mining and Metallurgy Equipment Co., Ltd., Nanchang, China). The XRD and XRF analyses were performed using a D8 Advance X-ray diffractometer (Bruker, Germany) and an S8 TIGER X-ray fluorescence spectrometer (Bruker, Germany), respectively. The detailed operating procedure for the XRD measurements has already been described in the literature (Bu et al., 2017; Chang et al., 2020).

2.5. Polarized light microscopy measurement

Thin sections of the SCA sample were prepared by Beijing Riyueshi Mining Technology Development Co., Ltd. (Beijing, China), and photographs of them were taken using a CX40P-series polarizing microscope (Sunny Optical Technology Co. Ltd).

2.6. Shape factor measurement

A 3D dynamic image analysis (Micromeritics® Instrument Corp., Norcross, USA) was conducted to characterize the circularity of the ground products produced by the ball and rod mills. Detailed working procedures and measurement principles for this apparatus can be found in the literature (Ulusoy and Igathinathane, 2014). Circularity (C) is computed from the projected area and bounding circle diameter, which is a dimensionless ratio. Round particles is valid for $C=1$ and elongated particles is valid for $C<1$ especially when C approaches to zero. Accordingly, this can be considered a fraction of the bounding circle's area covered by the actual projected area of the particle (Ulusoy and Igathinathane, 2014).

2.7. Surface roughness measurement

Scanning electron microscopy (SEM) analysis was carried out using an FEI Quanta 250 SEM system (FEI Company, Hillsboro, Oregon, USA). Detailed information on the SEM analysis can be found in the

literature (Bu et al., 2017). The SEM measurement results can only offer a qualitative comparison of roughness of ground particles produced by ball and rod milling. It is notable that the perimeter and area of the particles by 2 D can be calculated based on the axis measurement of the SEM images (Ulusoy et al., 2003; Ulusoy, 2008a). In addition, BET (Brunauer, Emmett and Teller) gas adsorption techniques can be used to calculate a surface roughness factor (Hicyilmaz et al., 2005; Hicyilmaz et al., 2006; Yekeler and Ulusoy, 2004). In this study, the surface roughness was also calculated quantitatively based on the BET results from the following relationship given in Eq. (3) (Gungoren et al., 2019; Guven et al., 2015):

$$\Lambda = \frac{\rho \cdot d_{AVG} \cdot A_{BET}}{6} \quad (3)$$

here, ρ is the density of the solid, d_{AVG} is the mean size of the size fraction, and A_{BET} is the specific surface area calculated using the BET isotherm. In this study, the specific surface areas (area per unit mass or volume) of the ground products obtained from ball and rod milling were determined using a Quantachrome™ Autosorp-1 MP device with N₂ gas using optimized protocols.

3. Results and discussion

3.1. Characterization of SCA

The LOI of the SCA sample was 12.29%. However, according to China Standard GB/T 4291-2017 (for synthetic cryolite), the LOI of the cryolite product should be <2.5%. This means the SCA sample contained approximately 10% carbon, which originated from the carbon anode during aluminum electrolysis. Purified carbon can be used to synthesize silicon carbide (SiC) (Yuan et al., 2018), be recycled as a low-cost material for the anodes of Li-ion batteries (Yang et al., 2020a), and present an accepted energy potential for various ironmaking applications (Flores et al., 2017). The chemical composition of the raw sample is given in Table 2. The SCA sample contained more in fluorine (F, 53.90%), sodium (Na, 26.70%), and aluminum (Al, 16.56%), although it was lacking in calcium (Ca, 2.67%). The XRD patterns of the SCA sample are shown in Fig. 1. The mineral components of the SCA sample were cryolite and corundum, which accorded well with the content of major elements obtained from the XRF analysis shown in Table 2. Cryolite, rather than corundum, was considered the main mineral.

Table 2. Major element constituents from the XRF analysis

Element	F	Na	Mg	Al	Si	S	K	Ca
Wt. (%)	53.90	26.70	0.28	16.56	0.11	0.38	0.19	2.67

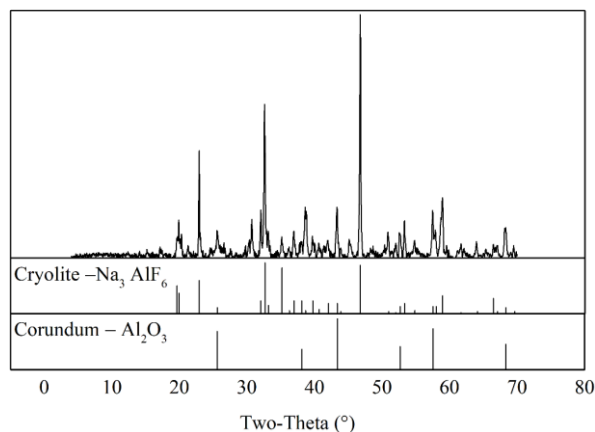


Fig. 1. XRD patterns of the SCA sample with respect to mineral identification

Fig. 2 presents the size distribution of the SCA sample. The LOI value was used to represent the approximate content (c_w) of the waste component. The content (c_v) of the valuable component was calculated as $c_v = 100 - \text{LOI}$. As shown in Fig. 2, c_v increased significantly from 9.17% to 64.13% with increasing particle size (from <45 μm to 500–1,000 μm), indicating that most of carbon material consisted of larger sized fractions. Observations of thin sections of the SCA sample are presented in Fig. 3. Fig. 3

(a) and (b) indicate that there were many carbon particles $>100\ \mu\text{m}$ in a state of dissociation. Fig. 3(d) indicates that large cryolite particles were associated with carbon materials. These phenomena resulted in a high waste component content with size classes coarser than $0.5\ \text{mm}$ (see Fig. 2 (b)). It can also be observed from Fig. 3 that a larger number of locked particles (marked by blue squares) existed in the SCA sample where the carbon material was strongly embedded in cryolite particles. Fig. 4 presents the size distribution of carbon particles in the SCA sample. The sizes of carbon particles of microscope photos were estimated using ImageJ software. The calculation method of the cumulative size distribution of carbon particles referred to the literature (Bu et al., 2020a; Dunbar and Hickey, 2000). It was observed that the size distribution of the carbon particles in the SCA sample had an average size (d_{50}) of $46.86\ \mu\text{m}$, while the d_{50} of the SCA sample was approximately $125\ \mu\text{m}$. Therefore, it was necessary to liberate the valuable component from the waste component by efficient breakage.

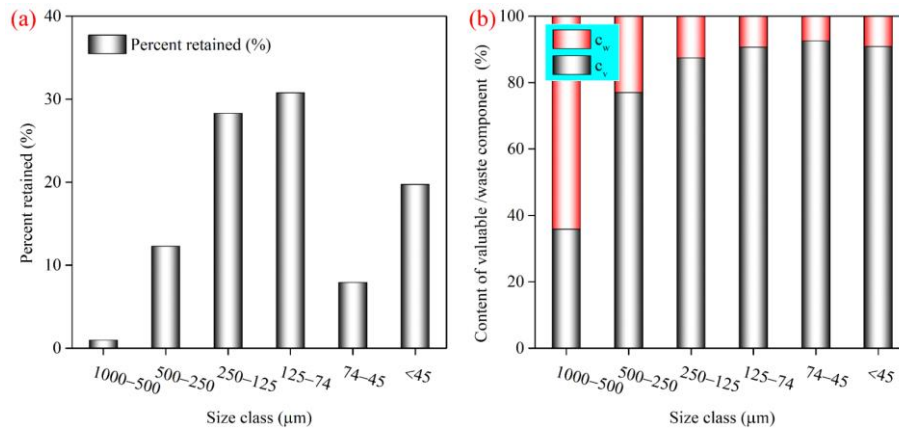


Fig. 2. Size distribution of the SCA sample: (a)-% retained; (b)-content of valuable/waste component

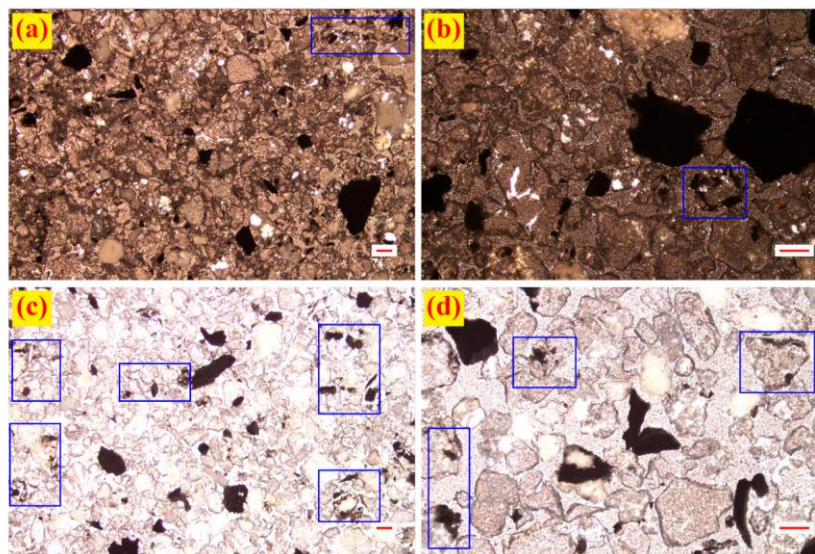


Fig. 3. Microscope photographs of thin sections of the SCA sample. Images (a) and (b) were taken from thin sections of the raw ore, while (c) and (d) were taken from thin sections of a $+45\ \mu\text{m}$ size fraction. The length of the red line represents $100\ \mu\text{m}$

3.2. Comparison of selective comminution using ball and rod mills

For a given t_d (a certain separation cut size), the ground product could be separated into two fractions: fine and coarse. Moreover, the calculated recoveries of valuable (R_v) and gangue (R_w) materials for various t_d values could be plotted diagrammatically. The calculated recovery of valuable, R_v , material is the cumulative recovery of this component with size classes finer than the selected sieve size (undersized product):

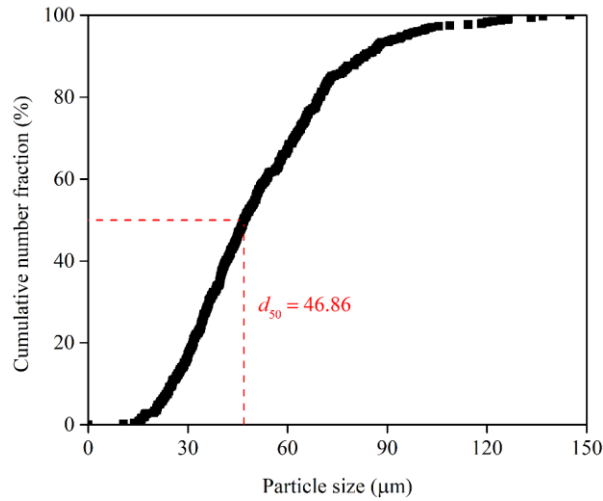


Fig. 4. Size distribution of carbon particles in the SCA sample.

$$R_V = \frac{\sum_{i=1}^N \omega_{V,i} \cdot c_{V,i}}{\sum_{i=1}^M \omega_{V,i} \cdot c_{V,i}} \quad (4)$$

here, N is the number of the size class smaller than the selected sieve size, M is the total number of the size class obtained from sieving analysis, $\omega_{V,i}$ is the weight percentage (valuable material) of the size class i , and $c_{V,i}$ is the content of the valuable material in size class i .

The calculated recovery of gangue, R_W , material is :

$$R_W = \frac{\sum_{i=1}^N \omega_{W,i} \cdot c_{W,i}}{\sum_{i=1}^M \omega_{W,i} \cdot c_{W,i}} \quad (5)$$

here, N is the number of the size class smaller than the selected sieve size, M is the total number of the size class obtained from sieving analysis, $\omega_{W,i}$ is the weight percentage (gangue material) of the size class i , and $c_{W,i}$ is the content of the component (gangue material) in size class i .

The recovery plots for ball and rod milling are presented in Fig. 5 (applied to the coarse fraction). Drzymala and Ahmed (2005) used various mathematical formulae (including formulas with one, two, or more adjustable parameters) to approximate the separation results in a Fuerstenau upgrading graph. According to the plot shape shown in Fig. 5, the two exponential asymmetric Eqs. were modified to approximate the effectiveness of selective comminution for the recovery plots (applied to the coarse fraction) as follows:

$$R_{cum,V} = \frac{100^\alpha - (100 - R_{cum,W})^\alpha}{100^{\alpha-1}} \quad (6)$$

$$R_{cum,V} = \frac{R_{cum,W}^\beta}{100^{\beta-1}} \quad (7)$$

here, a and β are adjustable parameters for Eqs. 6 and 7, respectively, and they are the selective comminution factors. When a (or β) = 1 (dashed line in Fig. 5), there was no selective comminution between the valuable and waste components. When $a > 1$ (or $0 < \beta < 1$), selective comminution existed for the coarse fraction. Finally, when $0 < a < 1$ (or $\beta > 1$), selective comminution existed for the fine fraction.

Comparisons of the fitting performance of Eqs. 6 and 7 using $RMSE$ (root mean square error) and $Adj. R^2$ are displayed Table 3. The parameters of Eqs. 6 and 7 were evaluated via the nonlinear least squares function of MATLAB software. A smaller $RMSE$ (or a greater $Adj. R^2$) indicates a better fitting performance of a model. It is observed that most of $RMSE$ values were smaller than 3 and most of $Adj. R^2$ were greater than 0.97. The magnitudes of $RMSE$ and $Adj. R^2$ were consistent with the literature (Bu et al., 2016; Bu et al., 2020b; Yang et al., 2021). Both Eqs. 6 and 7 provided an acceptable fitting performance from the viewpoint of statistical analysis. However, the $RMSE$ for the fitted data using Eq. 6 was relatively less than that using Eq. 7 for both ball and rod milling. Moreover, the $adj. R^2$ obtained with Eq. 7 was relatively greater than for Eq. 6, and a comparison of the Fuerstenau upgrading curves was fitted by Eqs. 6 and 7 at a grinding time of 5 min. Furthermore, comparisons of the Fuerstenau

upgrading curves fitted by Eqs. 6 and 7 (Fig. 6) indicated that Eq. 7 had a superior ability to describe Fuerstenau upgrading curve compared to Eq. 6. Thus, it is concluded that the assessment results indicated that Eq. 7 exhibited superior fitting performance compared to Eq. 6.

Table 3. Comparisons of the fitting performance (*RMSE* and *Adj. R²*) of Eqs. 6 and 7

Grinding time (min)	Ball milling				Rod milling			
	Eq. 6		Eq. 7		Eq. 6		Eq. 7	
	<i>RMSE</i>	<i>Adj. R²</i>	<i>RMSE</i>	<i>Adj. R²</i>	<i>RMSE</i>	<i>Adj. R²</i>	<i>RMSE</i>	<i>Adj. R²</i>
0	5.83	0.9798	1.68	0.9983	5.83	0.9798	1.68	0.9983
5	7.1	0.9688	1.83	0.9979	5.33	0.9831	1.29	0.999
10	7.4	0.9683	3.47	0.9992	5.53	0.9815	1.47	0.9987
20	3.98	0.9882	1.49	0.9983	5.22	0.9825	2.95	0.9944
30	3.8	0.9893	1.75	0.9977	5.38	0.981	3.3	0.9929
60	1.28	0.999	0.33	0.9999	5.31	0.9842	3.62	0.9926

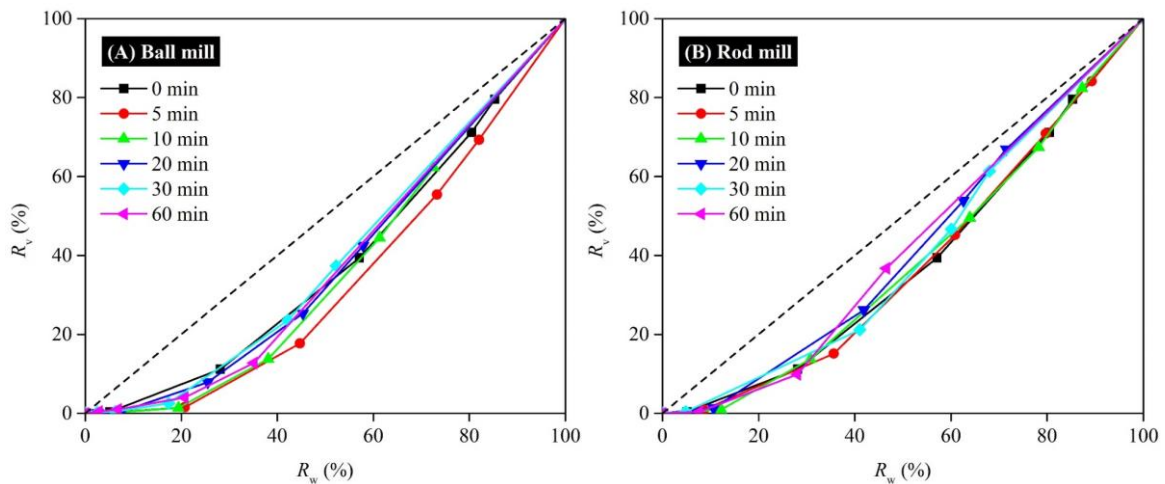


Fig. 5. Recovery plots for the Fuerstenau upgrading curves for ball and rod milling (applied to the coarse fraction). The original data including the separation cut size were attached in the appendix

Eq. 6 was used to calculate the selective comminution factor (β) for ball and rod milling. Fig. 7 presents a comparison of the calculated β for ball and rod milling at various grinding times. The β values for the ball milling at various grinding times were greater than those of the feed material ($\beta = 1.6$). By comparison, the β value for the rod milling was less than that for the feed material when the grinding time was >10.47 min (approximately). At 5 min grinding time, the ball and rod milling had maximum β values of 2.00 and 1.63, respectively. These results indicated that ball milling performed more effectively than rod milling for selective comminution.

3.3. Relationship between grinding mechanisms and selective comminution

The valuable component content ($c_{v, cum}$) of $-125 \mu\text{m}$ ground particles produced for various ball and rod milling times are presented in Fig. 8. Cryolite (2.5–3) has a similar level of Mohs hardness to gypsum (Mohs hardness: ~ 2.2) (Pauly, 1985); thus, the compression strength of cryolite is approximated to that of gypsum, which ranges from 1.5 to 6.7 MPa. According to the difference in Mohs hardness, the cryolite and carbon present in the SCA were considered the weaker material and the stronger mineral, respectively. Moreover, the XRF and XRD analyses indicated that cryolite was the major mineral for this SCA. In this study, the magnitude of $c_{v, cum}$ approximately equalled the cryolite content. When the grinding time was < 10 min, the $c_{v, cum}$ of the $-125 \mu\text{m}$ ground particles from rod milling were smaller than from ball milling. This revealed that rod milling had superior breakage ability for stronger minerals (carbon) compared to ball milling. Gupta and Yan (2016) concluded that the line load was more conducive to the breakage of hard materials than the point load on the particles produced by ball mills,

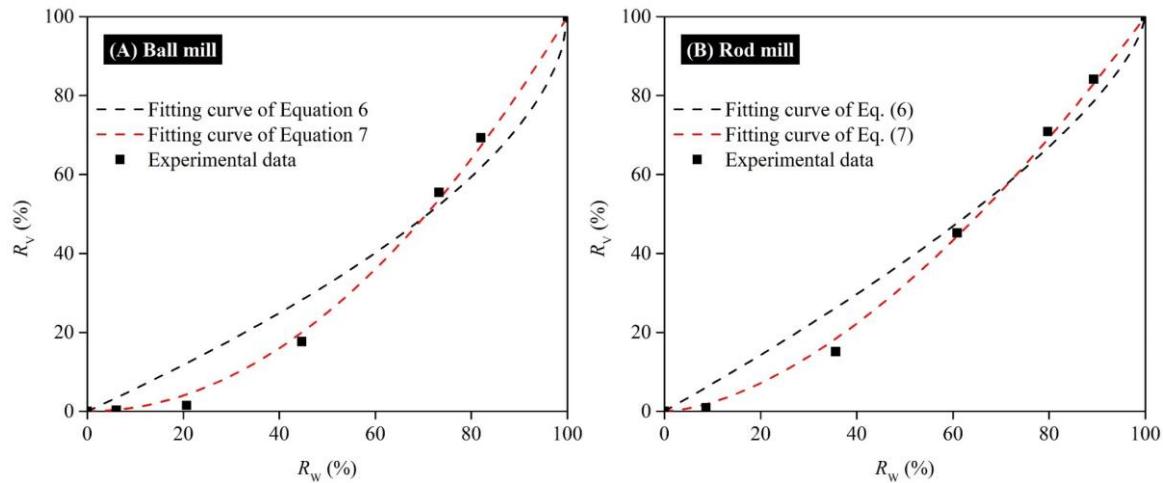


Fig. 6. Comparisons of the Fuerstenau upgrading curves fitted by Eqs. 6 and 7: (A) ball milling; (B) rod milling. The experimental data were the recovery plots for ball and rod milling at 5 min grinding time

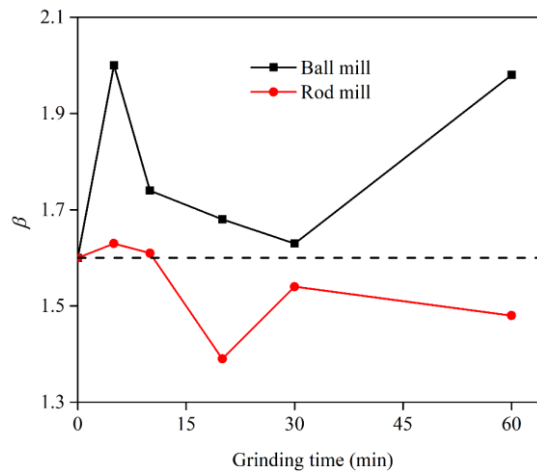


Fig. 7. Comparison of the selective comminution factor β for ball and rod milling at various grinding times

which was consistent with the findings of this study. When the grinding time was >10 min, the $c_{v, cum}$ of the $-125 \mu\text{m}$ ground particles for both ball and rod milling decreased constantly as the grinding time further increased. With this increased grinding time, increasing numbers of carbon and cryolite particles were fractured into smaller particles, which increased the difficulty of selective separation and the consumption of grinding energy. Compared to the raw sample (0 min grinding time, $c_{v, cum} = 90.98\%$), the $c_{v, cum}$ of the $-125 \mu\text{m}$ ground particles produced by ball milling increased and reached a peak of 91.39 % at a grinding time of 5 min, which was consistent with the estimated results for the selective comminution for ball milling (see Section 3.2). Meanwhile, the $-125 \mu\text{m}$ ground product of the rod mill had a smaller $c_{v, cum}$ than the raw sample, indicating that the stronger mineral was fractured by rod milling in larger quantities. This resulted in a deterioration of the degree of selective comminution for rod milling.

As shown in Fig. 9, ground samples produced by ball milling had smoother particle surfaces compared to those produce by rod milling, which exhibited relatively rough particle surfaces. In addition, the surface roughness factors (Λ) of 74–125 μm ground particles produced by ball and rod milling were 18.16 and 15.98, respectively. Rahimi et al. (2012) reported a similar result—the Λ value of ground particles from rod milling is greater than that of ball milling. Compared to ball milling, the increased grinding energy produced during rod milling increased the frequency of the fracturing of the feed material, promoting the formation of rough surfaces (Ahmed, 2010; Feng and Aldrich, 2000). By comparison, the presence of large numbers of fine particles increased the frequency of abrasion for ball

milling, which was conducive to the generation of smoother and cleaner ground particle surfaces (Bu et al., 2019a, 2020a).

Fig. 10 displays a comparison of the shape characterizations of 74–125 μm ground particles produced

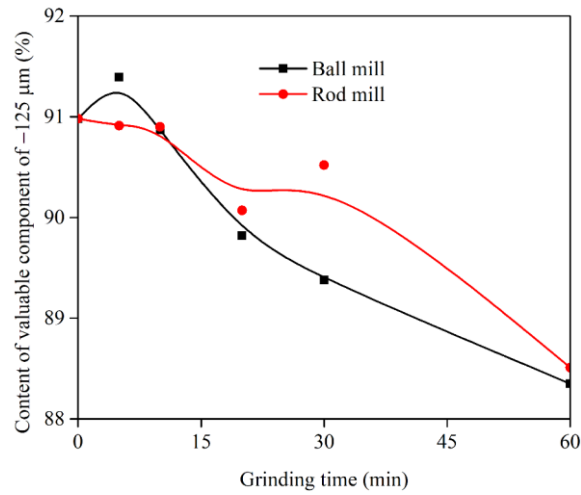


Fig. 8. Valuable component content of -125 μm ground particles produced by ball and rod milling

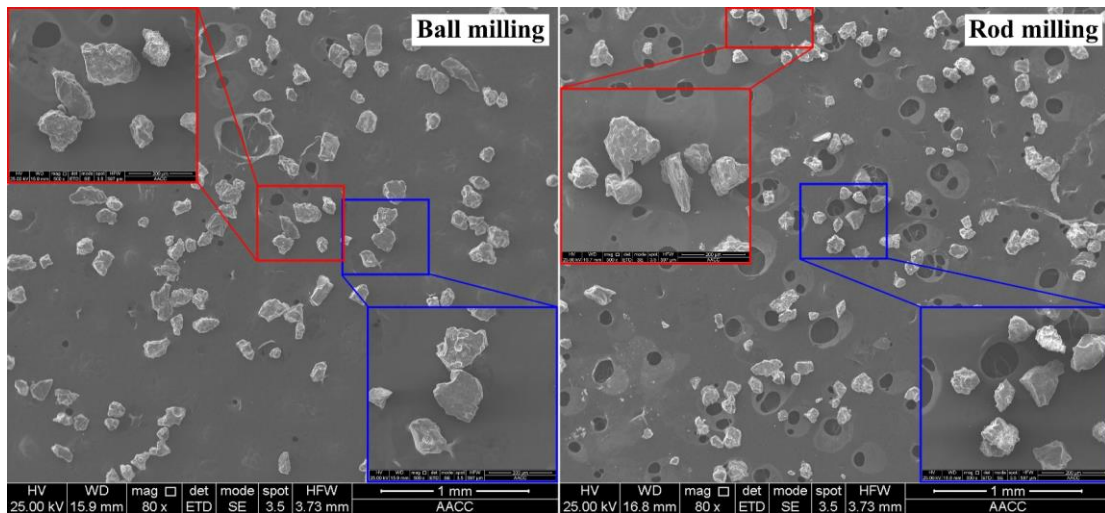


Fig. 9. SEM images of 74–125 μm ground particles produced by ball and rod milling at a grinding time of 10 min

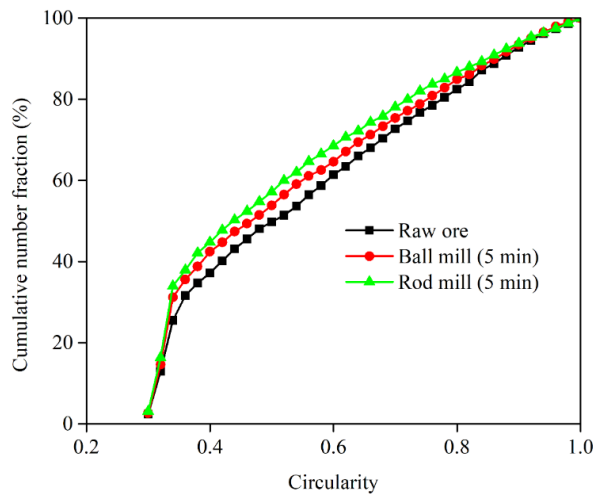


Fig. 10. Comparison of the circularity of the 74–125 μm ground product produced by ball and rod milling at a grinding time of 5 min

Table 2s. Original data for the Fuerstenau upgrading curves for rod milling (applied to the coarse fraction)

Cut size (mm)	Grinding time											
	0 min		5 min		10 min		20 min		30 min		60 min	
	R _w	R _v	R _w	R _v	R _w	R _v	R _w	R _v	R _w	R _v	R _w	R _v
1	0	0	0	0	0	0	0	0	0	0	0	0
0.5	5.17	0.40	8.66	0.93	12.23	0.92	0.00	0.00	0.00	0.00	0.00	0.00
0.25	28.16	11.18	35.61	15.11	30.80	13.74	10.68	1.34	4.88	0.39	0.00	0.00
0.125	57.11	39.38	60.88	45.20	63.94	49.53	41.90	26.18	41.06	21.18	7.70	0.41
0.074	80.46	71.18	79.70	70.88	78.23	67.41	62.59	53.88	60.09	46.66	28.09	9.86
0.045	85.27	79.56	89.25	84.14	87.25	82.39	71.50	66.89	68.04	61.37	46.56	36.72
0	100.0 0	100.0 0	100.0 0	100.0 0	100.0 0	100.0 0	100.0 0	100.0 0	100.0 0	100.0 0	100.0 0	100.0 0

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