Comparative Studies on Flotation Performance of Saturated Fatty Acids and Unsaturated Fatty Acids Separated from Hogwash Oil

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Abstract: Low flotation recovery, high pulp temperature, and large dosage of reagents are the typical disadvantages when using mixed fatty acids (MFA) prepared from hogwash oil for flotation directly. To determine the type of fatty acid that yields poor flotation performance, flotation performance and adsorption characteristics of saturated fatty acids (SFA) and unsaturated fatty acids (UFA) separated from the MFA were studied in our work. GC-MS, FT-IR, iodine value detection and melting point measurement showed that UFA contained \(-\text{(CH}=\text{CH}-\text{CH}_2\text{)}_n\)- groups and had much lower melting point. Quartz flotation tests were used to compare the flotation performance of UFA and SFA, which showed that UFA had excellent low-temperature floatability, and the flotation recovery of UFA was 35 percentage points higher than that of SFA at 20 °C and pH = 11.5. Zeta potential, FT-IR and XPS analysis indicated that UFA and SFA could adsorb onto the surface of activated quartz through chemisorption and hydrogen bonding. However, the adsorption of UFA was much stronger and more favorable; thus, the reason MFA have poor flotation performance was the presence of SFA.

Keywords: hogwash oil; fatty acids; separation; quartz flotation; adsorption mechanism

1. Introduction

Quartz is a type of widely distributed minerals on the surface of the earth, mainly existing in the forms of crystal, quartzite, quartz sandstone, vein quartz, and so on. In addition, it is one of the most common gangue minerals, associated with oxides, sulfides, silicates, and phosphates [1]. Direct flotation with cationic amine collectors is a typical method to select quartz. Another typical method is activated by the polyvalent metal ions, and then using anionic collectors for flotation [2]. The most extensively used anionic collectors are long carbon chain length fatty acids and their salts [3]. For example, RA series collectors (RA-315, RA-515, RA-715, and RA-915) [4,5] are fatty acids collectors which have been proved to be efficient, economical, and nontoxic, and have been widely used in quartz flotation in China.

According to estimates, there are approximately 8 to 15 million tons of hogwash oil annually produced from the catering industry in China [6]. Only 40–60% of the hogwash is able to be collected for recycling [7], and the rest is mostly disposed of as rubbish. The main chemical composition of hogwash is the high fatty acid glycosides [8], which are reliable sources of fatty acids and can be used as collectors for mineral flotation. More than 800 thousand tons fatty acid collector are consumed in China’s ore flotation ore. The traditional process of preparing fatty acids is mainly through chemical synthesis [9]. As the cost of flotation reagents increases, the use of fatty acids refined from hogwash oil as flotation collectors is increasingly economical and environmentally friendly. Sun et al. [10] refined the mixed fatty acids from hogwash oil, which was used to separate diaspore from kaolinite. An Al\textsubscript{2}O\textsubscript{3} recovery of 81.80% could be obtained. Liu et al. [11] used mixed fatty...
acids refined from hogwash oil as the collector of phosphate ore flotation and they got a concentrate with $P_2O_5$ 28.46% and $P_2O_5$ recovery of 87.2%.

The flotation recovery is relatively low, and high pulp temperatures and large quantities are necessary when using the mixed fatty acids refined from hogwash oil as flotation collectors directly [12,13]. Studies have attempted to enhance flotation performance of mixed fatty acids by mixing with other surfactants. Xu et al. [14] found that the preparation of 10% dodecyl trimethyl ammonium chloride in mixed fatty acids could improve its collectability and reduce consumption by approximately 60% for the flotation of lithium pegmatite ore. Cao et al. [15] found that the recovery of apatite flotation increased by 10% when fatty acids collector was mixed with 10% sodium dodecyl sulfonate. Jong et al. [16] confirmed that fatty acids collector mixed with some oleic acid amide exhibited a better collecting capacity on apatite. The mixed fatty acids prepared from hogwash oil contained saturated fatty acids (SFA), such as stearic acid and palmitic acid, and unsaturated fatty acids (UFA), such as oleic acid, linoleic acid, and linolenic acid [10,17–19]. Yu et al. [18] compared the flotation performance of four different mixed fatty acids prepared from hogwash oil which are from different regions in China. They found that mixed fatty acids containing more unsaturated fatty acids could yield better flotation results. Thus, in our study, we attempted to improve the flotation performance of original fatty acids prepared from hogwash oil by separating the original fatty acids into its constituent components to determine what types of fatty acids were responsible for poor flotation performance.

In this work, the mixed fatty acids (MFA) prepared from hogwash oil were separated into SFA and UFA via a chemical method. Gas chromatography-mass spectrometry (GC-MS), FT-IR, iodine value detection and melting point measurements were used to confirm the separation. Quartz flotation tests were used to compare the flotation performance of SFA and UFA. The adsorption properties and mechanisms of SFA and UFA on quartz surface were compared by zeta potential, FT-IR and XPS analysis.

2. Materials and Methods

2.1. Mineral Sample

Hand-picked pure quartz samples were obtained from Sijiaying Iron Mine, Hebei Province, China. The samples were crushed with a hammer. In addition, then the crushed samples were ground with ceramic ball mill and subsequently wet-sieved to obtain size fractions $-0.150 \text{ mm}$. The chemical compositions of the obtained quartz sample were presented in Table 1. The quartz sample was super pure comprising 99.20% SiO$_2$.

| Sample     | SiO$_2$ | Al$_2$O$_3$ | K  | S   | CaO  |
|------------|---------|-------------|----|-----|------|
| Pure quartz| 99.20   | 0.67        | 0.054 | 0.0046 | <0.001 |

2.2. Reagents

The hogwash oil was provided by Wuhan Zeyu Waterproof Building Materials, Ltd. (Wuhan, China). Calcium chloride (CaCl$_2$) with analytical purity was used as activator in the quartz flotation, which was supplied by Tianjin Kemiu Chemical Reagent Co., Ltd., Tianjin, China. The 0.10 mol/L HCl and 0.10 mol/L NaOH solutions were used to adjust the pH of the flotation pulp. The MFA, SFA and UFA prepared from hogwash oil were used as flotation collectors for quartz flotation. In particular, all collectors were saponified by 20% NaOH solution in a mass ratio of 1:1 and diluted to 0.2% solution before use.

2.3. FT-IR Analysis

In the measurement of the FT-IR spectrum (Nicolet 380 FT-IR spectrometer, Thermo Fisher Scientific, Waltham, MA, USA), the spectral pure KBr and quartz were further ground to $-2 \mu\text{m}$ in an agate mortar. Five grams of the ground quartz was mixed with 50 mL aqueous solution with 50 mg/L CaCl$_2$ and then treated with 100 mg/L collectors.
(UFA or SFA) at pH 11.50, and 25 °C. After being stirred for 0.5 h, the pulp was filtered and then washed with distilled water three times and finally dried in a vacuum oven at 50 °C. Finally, a mixture of the quartz sample and KBr at a mass ratio of 1:100 was pressed into a thin disk for FT-IR spectroscopy.

2.4. Flotation Tests

Micro-flotation tests were conducted using an XFGII50 with a 50-mL cell (Changchun Prospecting Machinery Factory, Changchun, China). A 5 g quartz sample was placed in the cell which are filled with ultra-pure water. A pH regulator (0.10 mol/L HCl or NaOH), CaCl$_2$ activator, and collector (MFA, SFA or UFA) were sequentially added to the cell. The conditioning time of each reagent was 2 min at a rotation speed of 1992 rpm, and flotation was conducted for 4 min. Finally, the Equation (1) is used to calculate the recovery $\varepsilon$, where $a$ and $b$ are the dry weights (g) of froth products and tailings, respectively.

$$\varepsilon = \frac{a}{a + b} \times 100\% \quad (1)$$

2.5. Zeta Potential Measurements

Before the measurement of zeta potential (Malvern Instruments Nano-ZS90, Malvern Panalytical, Malvern, UK), the quartz sample was further ground to below 0.005 mm and prepared as a 0.06% suspension. Zeta potential was measured in the absence or presence of CaCl$_2$ and collectors at 25 °C. After agitating for 10 min, 0.10 mol/mL HCl and NaOH were used to regulate the pulp pH. Each measurement was repeated three times, and the average was adopted.

2.6. X-Ray Photoelectron Spectroscopy

XPS analysis of quartz and quartz treated with CaCl$_2$ and collector (UFA or SFA) at pH 11.50 were measured with an American Thermo VG ESCALAB 250 spectrometer (Thermo Fisher Scientific, Waltham, MA, USA). The X-ray source was monochromatized Al Kα (1486.7 eV) with a power of 150 W (15,000 V × 10 mA). The pressure in the sample chamber was 5.0 × 10$^{-8}$ Pa. A value of 284.8 eV was chosen as the standard C(1s) to calibrate the binding energy.

3. Results and Discussion

3.1. Preparation of Collectors

Hogwash oil was first water washed and treated by 40% NaOH solution and 40% H$_2$SO$_4$ solution to obtain the mixed fatty acids (MFA). GC–MS was used to identify the main components of the MFA [20] and listed them in Table 2. The MFA is comprised of unsaturated fatty acids (29.6% oleic acid, 32.1% linoleic acid and 4.5% linolenic acid) and saturated fatty acids (7.9% stearic acid and 24.7% palmitic acid).

| Component     | Structure                                      | Content/% |
|---------------|-----------------------------------------------|-----------|
| Palmitic acid | CH$_3$(CH$_2$)$_{14}$COOH (C16:0)              | 24.7      |
| Stearic acid  | CH$_3$(CH$_2$)$_{16}$COOH (C18:0)              | 7.9       |
| Oleic acid    | CH$_3$(CH$_2$)$_{17}$-CH=CH-(CH$_2$)$_2$-COOH (C18:1) | 29.6      |
| Linoleic acid | CH$_3$(CH$_2$)$_{16}$-[CH=CH-(CH$_2$)$_2$]-COOH (C18:2) | 32.1      |
| Linolenic acid| CH$_3$CH$_2$-[CH=CH-(CH$_2$)$_2$]-COOH (C18:3) | 4.5       |
| others        | /                                             | 1.2       |

"/" means no specific structure.

The SFA and UFA collectors were prepared from MFA using urea inclusion method [21–23] as described in Equation (2). As reported in our previous study [24], this separation mechanism was investigated by molecular simulation, and the SFA and UFA were characterized by iodine value, melting point measurement and Fourier transform infrared (FT-IR) spectroscopy. The results indicated that UFA comprised fatty acids containing -(CH=CH-CH$_2$)$_n$- with an iodine
value of 162.9 and a melting point as low as −17.3 °C, which means that the UFA were easy to disperse and dissolve in flotation pulp at low temperatures. In contrast, no -(CH=CH-CH\_2)_n- group was detected in the SFA, and the SFA comprised saturated fatty acids. The iodine value and melting point of the SFA were 49.0 and 38.8 °C, respectively. Thus, a high pulp temperature greater than 38 °C was necessary to ensure the dissolution and dispersion of SFA, which was consistent with the results of the flotation temperature test (Figure 4).

\[ \text{MFA} + m\text{CO(NH}_2\text{)}_2 \xrightarrow{\text{Ethanol}} \text{SFA}[\text{CO(NH}_2\text{)}_2]_m \downarrow + \text{UFA} + \Delta H \]  

(2)

3.2. Flotation Tests

3.2.1. Conditional Tests of pH

Quartz flotation tests were used to compare the flotation performance of SFA, UFA, and MFA. Firstly, quartz was floated at various pH with collector (MFA, UFA or SFA) concentration of 120 mg/L and activator CaCl\_2 concentration of 50 mg/L at 20 °C, and the results was shown in Figure 1. Pulp pH of quartz flotation is typically greater than 8, and the optimum pH is approximately 11.5. As the pulp pH increased from 8 to 11.5, quartz recovery by collector UFA, MFA and SFA improved, with maxima of 99.01%, 89.45% and 64.10%, respectively. Additionally, the UFA collector had better collectability than those of SFA and MFA at pH 8 to 12.5.

![Flotation recoveries of quartz as a function of pH](image)

**Figure 1.** Flotation recoveries of quartz as a function of pH (collector MFA/UFA/SFA 120 mg/L, activator CaCl\_2 50 mg/L, at 20 °C).

3.2.2. Conditional Tests of Concentration of Collector

Figure 2 shows the effects of the collector concentration (MFA, SFA and UFA) on quartz flotation at 20 °C and pH 11.50. As the UFA concentration of pulp was more than 80 mg/L, the quartz recovery was almost 100%. For SFA, quartz recovery was greatest (approximately 64%) at concentrations between 100 mg/L to 120 mg/L, which was comparatively achieved with only 30 mg/L UFA. For MFA, 89.45% quartz recovery was obtained at 80 mg/L with negligible improvement, but at higher concentrations. UFA collector separated from MFA showed better quartz flotation performance.
3.2.2. Conditional Tests of Concentration of Collector

Figure 2 shows the effects of the collector concentration (MFA, SFA and UFA) on flotation recoveries of quartz as a function of the concentration of collector MFA, SFA or UFA (activator CaCl$_2$ 50 mg/L, pulp pH = 11.5, at 20 °C).

3.2.3. Conditional Tests of Concentration of CaCl$_2$

Figure 3 shows the effects of CaCl$_2$ activator on quartz flotation. For the UFA collector, when the concentration of CaCl$_2$ was only 25 mg/L, a high flotation recovery of 99.50% was obtained. Even at a low 10 mg/L CaCl$_2$ concentration, quartz recovery by UFA collector remained above 90%, whereas the quartz recoveries were 75% and 30% for MFA and SFA respectively. CaCl$_2$ activator concentration should be more than 50 mg/L for when using MFA and SFA collectors to ensure sufficient activation. The UFA collector required lower CaCl$_2$ concentrations to activate quartz.

3.2.4. Conditional Tests of Temperature

The flotation properties of collector UFA, MFA and SFA with different pulp temperatures are presented in Figure 4. The UFA collector had excellent adaptability to pulp temperature. Under flotation temperatures between 10 °C and 50 °C, the recoveries of quartz flotation by UFA were higher than 98%. However, similar quartz recoveries by MFA and SFA flotation required pulp temperatures greater than 30 °C and 35 °C, respectively. The quartz flotation results demonstrated that the flotation performance of SFA separated from MFA was very unfavorable, whereas UFA was an excellent collector with superior low-temperature adaptability for quartz flotation.
Zeta potential analyses were used to investigate the differences in electrostatic interactions between quartz and the two separate components (SFA and UFA) from MFA. The zeta potential measurements of pure at various pH values are shown in Figure 5. The zeta potential of quartz surface was negative within a pulp pH range from 2 to 14; and as the pH value increased, the zeta potential continued to decrease. After treated with 50 mg/L CaCl$_2$ (Figure 5), the zeta potential significantly increased compared with Figure 5. This was due to the adsorption of calcium ions on the negative quartz surface, and the electrostatic force in the adsorption process. The CaCl$_2$ activated quartz surface remained negative throughout the studied pH range.

Figure 5 respectively show the changes in zeta potential when the SFA or UFA collectors was added at 100 mg/L to 50 mg/L activator CaCl$_2$ activated quartz system. The SFA and UFA collectors significantly reduced the negative zeta potential of activated quartz when the pulp pH was above 5, which was reflected in the flotation results of the pH test (Figure 1). To reveal the effects of SFA and UFA on the surface potential of activated quartz, the 100 mg/L SFA and UFA were analyzed. Based the compositions of fatty acids shown in Table 2, palmitic acid (C16:0) and linoleic acid (C18:2) were selected to represent the parameters of SFA and UFA respectively. The solution equilibria of SFA and UFA were as follows. (3) to (6) [25].

Where $S$, $k_1$, $k_2$ and $k_3$ are dissolution, acid dissociation, ions association and ion-molecular complexation constants, respectively having values of $S = 1.6 \times 10^{-5}$ mol/L,
$k_1 = 10^{-10.0}, k_2 = 10^{4.0}, k_3 = 10^{4.7}$ for SFA and $S = 5.7 \times 10^{-6}$ mol/L, $k_1 = 10^{-9.0}, k_2 = 10^{4.5}, k_3 = 10^{5.2}$ for UFA [25–27]. The $\log_c p\text{H}$ diagrams of the SFA and UFA solution were calculated and are shown in Figure 6A,B. UFA primarily existed as fatty acid molecules from pH 2 to 10. Above pH 10, fatty acid anions and anion associations were the main compositions. The dissociation pattern of SFA was similar to that of UFA, but the significant change in composition was observed at approximately pH 12. These results were consistent with the results shown in Figure 1, i.e., the suitable pH values for the SFA and UFA slurries were approximately 10 to 13 and 12 to 13, respectively. In particular, the concentration of $RCOO^-$ anions in the UFA solution at suitable pH was 1.6 times that for SFA, this accounted for the higher UFA, which was the reason the quartz recovery of UFA was higher. When the pulp pH was approximately 12, both SFA and UFA were acid dissociated, thereby preventing electrostatic attraction between SFA/UFA anions and the negatively activated quartz surface. It was obvious that there was no electrostatic attraction between SFA/UFA anions and the negative activated quartz surface. The zeta potential decrease observed when activated quartz was treated with SFA or UFA may be due to chemisorption between the collectors and the activated quartz surface.

![Figure 6](image_url). The species distribution diagrams of 100 mg/L (3.8 $\times$ 10$^{-4}$ mol/L) SFA (A) and 100 mg/L (3.6 $\times$ 10$^{-4}$ mol/L) UFA (B) solution as a function of pH.

### 3.4. FT-IR Analyses

Figure 7 showed the FT-IR spectra of quartz, the FT-IR spectra of quartz treated with 50 mg/L CaCl$_2$ and 100 mg/L UFA and the FT-IR spectra of quartz treated with 50 mg/L CaCl$_2$ and 100 mg/L SFA. The vibrations approximately 3128 cm$^{-1}$ and 1400 cm$^{-1}$ were caused by H$_2$O in air or adsorbed H$_2$O on quartz surface [28]. The vibrations near 1089 cm$^{-1}$, 796 cm$^{-1}$ and 691 cm$^{-1}$ belonged to the symmetric stretching and bending vibrations of Si-O-Si [29].

After treatment with CaCl$_2$ and collector UFA/SFA, as shown in Figure 7, in both spectra, stretching vibration absorption vibrations of the $\text{-CH}_3$, $\text{-CH}_2$ and C=O groups could be seen at positions near 2928 cm$^{-1}$, 2852 cm$^{-1}$ and 1793 cm$^{-1}$, respectively [30,31], suggesting that both the collector UFA and SFA had adsorbed on the activated quartz surface. Additionally, the stretching vibration absorption vibrations of the C=O group shifted approximately 84 cm$^{-1}$ compared to those of pure UFA and SFA (1709 cm$^{-1}$), indicating that C=O group strengthened and chemisorption occurred [24]. The vibrations located at 3420 cm$^{-1}$ (both in Figure 7) illustrated the presence of hydrogen bond adsorption [29]. Thus, FT-IR revealed that both UFA and SFA could be adsorbed onto the activated quartz surface by chemical and hydrogen bonding adsorption.
Figure 7. FT-IR spectra of pure quartz (a), quartz treated with 50 mg/L CaCl$_2$ and 100 mg/L SFA (b), and quartz treated with 50 mg/L CaCl$_2$ and 100 mg/L UFA (c).

3.5. XPS Analysis

Figure 8 showed the XPS analysis of pure quartz, the XPS analysis of quartz treated with 50 mg/L CaCl$_2$ and 100 mg/L UFA at pH 11.50, and 25 °C and the XPS analysis of quartz treated with 50 mg/L CaCl$_2$ and 100 mg/L SFA at pH 11.50, and 25 °C. The results were summarized in Table 3. In the XPS spectra of quartz (Figure 8), an extremely weak C(1s) vibration at 284.8 eV was due to C-C and C-H bonds of hydrocarbon contamination from the environment [32]. In addition, the vibrations around 532 eV, 153 eV and 103 eV were related to the presence of oxygen and silicon [29]. There was no other obvious vibrations, which indicated that the quartz was relatively pure and clear.

Table 3. XPS characterization of reference compounds.

| Sample                        | Element at.% (BE, eV)     |
|-------------------------------|---------------------------|
|                               | C           | O            | Si           | Ca           |
| Quartz                        | 5.1 (284.8) | 64.0 (532.4) | 31.0 (102.9) |              |
| Quartz + CaCl$_2$ + SFA       | 13.8 (284.7) | 58.4 (532.3) | 25.5 (102.9) | 2.3 (346.5)  |
| Quartz + CaCl$_2$ + UFA       | 23.1 (284.7) | 51.1 (532.3) | 24.2 (102.9) | 0.3 (346.8)  |

After treatment with CaCl$_2$ and SFA (Figure 8), Ca adsorption (Ca 2p$_{3/2}$ at 346.5 eV, concentration of 2.3 at.%) was observed on the quartz surface. Furthermore, O(1s) and
Si(2p) contents decreased by 5.6 at.% and 5.5 at.%, respectively, whereas the contents of C(1s) increased by 8.8%, compared with that of the initial quartz surface. Similar XPS results were observed for quartz treated with CaCl₂ UFA (Figure 8) wherein a new Ca vibration (Ca 2p₃/₂ at 346.8 eV concentration of 0.4 at.%) appeared and the contents of O(1s), Si(2p) and C(1s) changed significantly. However, the contents of O(1s), Si(2p) and Ca(2p) were 7.3 at.%, 1.3 at.% and 1.9 at.% lower, respectively, and the C(1s) content was 9.3 at.% higher, compared with the activated quartz surface treated by SFA. Namely, more UFA molecules were adsorbed and covered the O, Si and Ca atoms on the activated quartz surface. These results revealed that both SFA and UFA could adsorb onto quartz surfaces activated by calcium ions, but the adsorption of UFA was much stronger.

To further compare the adsorption of SFA and UFA on quartz, the curve fitting of C(1s) vibrations of different samples are shown in Figure 9A–C. Figure 9A shows that the C(1s) vibration of the pure quartz surface was the only component at 284.8 eV assigned to the C(1s) of C-C and C-H bonds from the hydrocarbon contamination. Figure 9B,C show that the activated quartz surface treated with SFA and UFA included three components at approximately 284.8 eV, 286.6 eV and 288.9 eV due to the C(1s) bonds from C-C/C-H bonds, O=C-OH bonds and O=C-O⁻ bonds, respectively [29,32]. Obviously, both SFA and UFA had been adsorbed onto the surface of activated quartz. However, the adsorption of UFA was much stronger and more favorable, because the C(1s) content in C-C/C-H bonds from the sample treated by UFA was significantly higher and the binding energy of the C(1s) vibration of component O=C-O⁻ for the sample treated by UFA was 0.4 eV lower, than that of the sample treated by SFA. In addition, the components of O=C-OH bonds were found in both samples treated with SFA and UFA and their binding energies were almost same. This indicated that the adsorptions of both SFA and UFA may involve hydrogen bonding, which confirmed the results of the FT-IR analysis.

![Figure 9. XPS C(1s) spectra of quartz (A), quartz treated with 50 mg/L CaCl₂ and 100 mg/L SFA (B), and quartz treated with 50 mg/L CaCl₂ and 100 mg/L UFA (C).](image-url)

By combining the analyses of flotation test, FT-IR, XPS, and zeta potential measurements, it is evident that the adsorption mechanisms of UFA and SFA on quartz surface are the same as summarized in Figure 10. Both UFA and SFA can adsorb onto quartz surfaces activated by Ca²⁺ via chemical and hydrogen bonding adsorption. UFA is an extremely excellent collector for quartz even at low flotation temperatures. Comparably, the SFA had
poor flotation performance requiring high flotation temperatures and pulp pH to ensure dissolution and dispersion.

Figure 9. XPS C(1s) spectra of quartz (A), quartz treated with 50 mg/L CaCl$_2$ and 100 mg/L SFA (B), and quartz treated with 50 mg/L CaCl$_2$ and 100 mg/L UFA (C).

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Figure 10. The schematic view on the adsorption mechanism of UFA and SFA on quartz surface.

4. Conclusions

Mixed fatty acids (MFA) prepared from hogwash oil consisted of 7.9% stearic acid, 24.7% palmitic acid, 29.6% oleic acid, 32.1% linoleic acid, and 4.5% linolenic acid. MFA can be separated into saturated fatty acids (SFA) and unsaturated fatty acids (UFA) through chemical methods. The quartz flotation results demonstrated that the flotation performance of SFA separated from MFA was very unfavorable, whereas UFA showed strong collectability and superior low-temperature adaptability for quartz flotation. At 20 °C and pH 11.50, the highest quartz recoveries of by UFA, MFA and SFA at were 99.01%, 89.45% and 64.10%, respectively. Finally, the zeta potential, FT-IR, and XPS analyses revealed the adsorption mechanism by which both UFA and SFA could be adsorbed onto activated quartz surfaces via chemical and hydrogen bonding, but the adsorption of UFA was much stronger and more favorable. Thus, the unfavorable flotation performance of MFA was due to large amounts of SFA.

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