Evaluation of Frother Types for Improved Flotation Recovery and Selectivity

Qingqing Huang
West Virginia University, qingqing.huang@mail.wvu.edu

Xinbo Yang
University of Kentucky

Rick Q. Honaker
University of Kentucky

Follow this and additional works at: https://researchrepository.wvu.edu/faculty_publications

Part of the Mining Engineering Commons

Digital Commons Citation
Huang, Qingqing; Yang, Xinbo; and Honaker, Rick Q., "Evaluation of Frother Types for Improved Flotation Recovery and Selectivity" (2019). Faculty & Staff Scholarship. 1559.
https://researchrepository.wvu.edu/faculty_publications/1559
Evaluation of Frother Types for Improved Flotation Recovery and Selectivity

Qingqing Huang 1, *, Xinbo Yang 2 and Rick Q. Honaker 2

1 Department of Mining Engineering, West Virginia University, Morgantown, WV 26506, USA
2 Department of Mining Engineering, University of Kentucky, Lexington, KY 40508, USA; xinbo.yang@uky.edu (X.Y.); rick.honaker@uky.edu (R.Q.H.)
* Correspondence: qingqing.huang@mail.wvu.edu; Tel.: +1-304-293-3456

Received: 15 August 2019; Accepted: 25 September 2019; Published: 28 September 2019

Abstract: A laboratory study was conducted to evaluate and compare the effectiveness of nine different frother types when used in a three-phase, continuously operating froth flotation system. The frothers included several that are commonly used in the industry (e.g., MIBC, 2EH, and F-1) as well as unique frother types (e.g., F-3). The tests were conducted in a 5-cm diameter laboratory flotation column that provided near plug-flow mixing conditions due to a length-to-diameter ratio of around 50:1. Test results indicate that F-1, MIBC, and MPC (in order of decreasing effectiveness) provided the weakest performance in terms of combustible recovery while F-2, MAC, and 2EH were the top three generating the highest separation efficiencies. When processing ultrafine coal, the ash content of the flotation concentrate ranged from 10% to 15% while recovering over 80% of the combustible material. F-3, F-4, and DIBC provided over 80% recovery of combustibles at the expense in the amount of hydraulic entrainment. The flotation performances were also closely examined in accordance with the fundamental properties of the nine tested frothers, and their correlations were addressed in detail.

Keywords: frothers; column flotation; recovery; selectivity; entrainment

1. Introduction

The froth flotation process commonly requires the use of a multitude of chemical reagents including collectors, frothers, surface modifiers, and pH-regulators. Frothers play an essential role in determining the bubble size distribution as well as the stability and mobility of the froth phase [1,2]. The fundamental properties of frothers have been extensively investigated over the past several decades. It is widely acknowledged that frothers preferentially accumulate at the water–gas interface and interact with the collectors absorbed on the solids, which facilitates the solid-bubble collection process [3–6]. Furthermore, frothers impose a significant influence on bubble size distribution, which further affects the superficial bubble surface area flux and the ultimate flotation recovery [7–12]. It has been found that the way frothers affect the bubble size is through both surface tension reduction and the prevention of bubble coalescence for concentrations above the critical coalescence concentration [7,13–15]. Mineral particles report to the froth flotation product either because of the selective attachment of hydrophobic particles or water entrainment of non-selective hydrophilic particles [13,16]. The entrained hydrophilic particles that are recovered to the final flotation product are undesired gangue minerals. As such, entrainment greatly deteriorates the selectivity of the froth flotation process [17,18]. The amount of hydrophilic entrainment associated with a flotation unit is believed to be affected by the amount of water carried in the froth [19–24], which is closely related to the frothing properties of the frother used in the process.

Froth stability and mobility are important factors that affect flotation recovery and grade. A froth that is too stable may prevent adequate drainage of the entrained material and is difficult to handle.
while an unstable froth is least desirable due to the negative effect on recovery and selectivity [25]. The frothers that are more efficient in reducing bubble size have been observed to produce more stable foams [6–8,13]. Laskowski [26] was able to characterize frothers based on their ability to reduce bubble size and increase froth stability. The study distinguished the frothers known as being selective from those being identified as powerful. In a review paper authored by Khoshdast and Sam [1] on the properties of frothers, it was concluded that bubble size, and stability and mobility of the froth phase play a significant role in the kinetic viability of the process. The overall flotation recovery and grade are mainly driven by the physical properties of the frothers such as molecular weight, hydrophile–lipophile balance (HLB), dynamic frothability, and critical coalescence concentration [1].

The froth is established not only by frothers but also by the presence of solid particles [13]. The particle size, shape, and surface hydrophobicity have been reported to greatly affect the performance of a froth flotation process [27–32]. Overall, the behavior of solids particles and the interaction with frothers is a critical but poorly understood step in industrial froth flotation, which often leads to recovery losses in the froth phase [33]. Therefore, to better understand the interaction mechanisms in a froth flotation system driven by the characteristics of the frother types, a laboratory flotation column was utilized in the present study to evaluate the properties of nine frothers in a three-phase (liquid–gas–solid) system. Fundamental properties of the frothers including surface tension and frothing abilities were measured and correlated with their molecular weights and HLB values along with the findings of the flotation tests, which were obtained as a function of frother concentration. Optimum flotation performances were identified for each frother type as a function of recovery, concentrate quality, and process selectivity.

2. Experimental

2.1. Coal Sample

Run-of-mine coal samples were collected from an active coal preparation plant located in Grafton, West Virginia. The coal was extracted from the Lower Kittanning coal seam, which ranks as high-volatile bituminous coal. Upon receiving the samples, the coal was crushed using a laboratory jaw crusher followed by a hammer mill to a top size of approximately 6 mm. The entire sample was then homogeneously split using a riffler into separate sample lot sizes of approximately 0.5 kg, placed into storage bags, and stored in a freezer to prevent oxidation until the time needed for the flotation tests. Prior to each test, a 0.5 kg sample was ground using a laboratory pulverizer to achieve a top particle size of 210 micrometers.

2.2. Chemicals

A total of nine alcohol-based frothers including MPC, MIBC, MAC, 2EH, DIBC, F-1, F-2, F-3, and F-4 were evaluated in the present study. Of the nine frothers, MPC, MAC, 2EH, MIBC, and DIBC were purchased from Sigma-Aldrich with reagent purity greater than 98%, 97%, 99.6%, 98%, and 80%, respectively. The remaining frothers were directly provided by the manufacturers. The chemical structures of all nine test frother types are shown in Figure 1. Their physical properties including their compositions, molecular weights, solubility in water, and densities are shown in Table 1. The water solubility values are of significant importance. The data indicates that DIBC, F3, and F4 have exceptionally low values in the range of 66.4 to 321.5 mg/L (ppm). The relative difference between the water solubility values and the frother solution concentration had a significant impact on the results obtained in the present study. Therefore, the frother concentration of the dilute solution that was fed into the flotation unit ranged from 22 ppm to 66 ppm, which was slightly lower than the lowest water solubility among the nine evaluated frothers. In addition to the frothers, fuel oil No. 2 was used as a collector throughout the whole study. Hydrochloric acid and sodium hydroxide of analytical grades were used for the pH adjustment.
2.3. Physical Properties Measurement

In addition to the properties provided by the chemical vendors (Table 1), laboratory experimental tests were further conducted to determine supplemental characteristics of the nine frothers, which included their hydrophilic–lipophilic balance (HLB) values, surface-active properties, and foamability. As known, the HLB value is a measure of the degree to which a surfactant is hydrophilic or lipophilic, was determined according to Davies’ method [34] following the equation given below:

\[
HLB = 7 + \sum_{i=1}^{m} H_i - n \times 0.475
\]  

(1)

where \( m \) is the number of hydrophilic groups in the molecule, and \( H_i \) and \( n \) the corresponding group number of the \( i \)th hydrophilic group and the number of the lipophilic groups in the molecule, respectively.

Surface-active properties of various frothers were determined by measuring the inter-facial surface tension of the solutions, which were prepared by mixing the frothers with distilled water at different concentrations. A surface tensiometer (Fisher Tensiometer Model 20, Fisher Scientific, Hampton, NH,
USA) that employed the ring method was used to conduct the tests. A platinum-iridium ring was utilized during the measurement at a controlled temperature of 25 °C. The tensiometer was calibrated with pure methanol solutions of various concentrations and known surface tensions. The accuracy was further checked using distilled water. The surface-active properties of the tested frothers were expressed as their capability to reduce the air-water interfacial surface tension in mN/m per ppm of the tested frother. Furthermore, the foamability of the frothers were studied in accordance with the method described by Gupta et al. [5] using a graduated glass burette (2.5 cm in diameter and 50 cm in height), which was fitted with a glass frit and connected to a mercury manometer and air compressor. Detailed operating procedures can be referred to the early study by Gupta et al. [5], and the foamability of the tested frothers are expressed as the froth height generated per unit concentration in mm/ppm.

2.4. Column Flotation Apparatus

A continuously operated laboratory column setup was used to perform the flotation tests. The Plexiglas column measured 5 cm in diameter and 2.4 m tall, which provided a length-to-diameter ratio of around 50:1 and ensured nearly plug-flow operating conditions. As shown in Figure 2, feed slurry was injected at a depth of 450 mm below the overflow lip while 400 mL/min of wash water was added in the froth zone at a depth of 100 mm below the overflow lip, which minimized recovery to the froth concentrate by hydrophilic entrainment. The wash water was added via a water spray dispenser with two 1–4 inch (in diameter) hollow stainless tubes perpendicularly intersecting each other, which was designed in a manner for the effective distribution of wash water.

![Figure 2. Schematic of laboratory column flotation setup used for the standard testing procedure.](image)

The flotation column pumped a portion of the tailings stream through a static mixer under a pressure of 138 kPa (20 psi) for bubble generation. Air was injected into the tailings recirculation line at the bottom of the column prior to the static mixer while frother was injected from a separate sump directly into the feed stream to the column. The frother concentration in the frother sump was varied from 22 ppm to 66 ppm depending on the desired frother concentration in the flotation column, which ranged from 10 ppm to 30 ppm.

Two pressure transducers were installed along the vertical length of the column at a distance of 660 mm apart to: (1) Measure the air fraction within the collection zone and (2) control the froth
depth at a constant value of 356 mm. The air fraction of the collection zone was calculated according to Equation (2):

\[ \epsilon = 1 - \frac{\rho_{\text{water}} \times \Delta h}{\rho_{\text{pulp}} \times L} \]  

where \( \epsilon \) is the air fraction, \( \rho_{\text{water}} \) the density of water, \( \rho_{\text{pulp}} \) the density of slurry pulp, \( \Delta h \) the difference in the pressure transducer readings in millimeters of the water head, and \( L \) the physical distance between the two transducers.

The lower transducer, which was located 1420 mm below the column overflow lip, provided a measurement of pressure that reflected the froth zone level in the column. A controller received the signal from the transducer and controlled a pinch valve located in the column tailings line to maintain the froth level at the pre-set value of 356 mm. Measurements were taken of the tailings volumetric flow rate during each experiment to allow a calculation of the percentage of the wash water reporting to the tailings stream (i.e., bias factor, BF) according to Equation (3):

\[ BF = \frac{Q_T - Q_F}{Q_W} \times 100 \]  

where \( Q_T \), \( Q_F \), and \( Q_W \) are the volumetric flow rate of tailings, feed, and wash water, separately. A bias factor of 60% is typically used to minimize the negative impacts of hydraulic entrainment. If the bias factor was too low, an increase in froth depth was required. However, it was desired to keep the froth depth and the collection zone volume constant for the entire test program. By maintaining a constant froth depth of 356 mm, bias factor variations occurred that reflected the characteristics of the frother type due to the influence on the bubble size distribution and froth stability.

2.5. Flotation Release Analysis

A flotation release analysis was conducted on a representative coal sample to establish the ultimate separation performance achievable by froth flotation. For this test, Fuel Oil No. 2 was used as a collector and MIBC was used as the frother. The test was conducted in two phases. The first phase involved floating the coal in a laboratory batch Denver conventional cell until all of the floatable material was recovered. The tailings from the first stage of flotation were stored in a bucket and the concentrate placed back into the flotation cell to repeat the process. Three stages of flotation were used in the first phase to eliminate all of the hydraulically entrainable particles.

In the second phase, the final concentrate obtained from the first phase was floated under starvation conditions using low aeration rates and agitation speeds. The concentrate collected under this condition contained the lowest amount of ash-bearing material. The air rate was slightly increased along with the impeller speed to collect the next fraction, which contained a higher amount of ash material. This process was repeated to produce a sufficient number of data points to generate a recovery versus product ash relationship under the ultimate flotation condition.

The release analysis results for the Lower Kittanning coal are shown in Table 2. The data suggest that the coal is relatively easy-to-float and has good liberation characteristics as indicated by a relatively consistent low ash product over a range of mass yield values. The flotation data indicated that the ultimate performance would yield a clean coal concentrate containing 9.4% ash while recovering 79.2% of the combustible material.
Table 2. Release analysis results for ultrafine Lower Kittanning coal.

| Sample I.D. | Incremental | Cumulative |
|-------------|-------------|------------|
|             | Weight (%)  | Ash (%)    | Weight (%)  | Ash (%)  | Combustible Recovery (%) |
| P1          | 2.48        | 5.57       | 2.48        | 5.57     | 4.54                  |
| P2          | 3.50        | 5.60       | 5.98        | 5.59     | 10.94                 |
| P3          | 6.76        | 6.00       | 12.74       | 5.81     | 23.25                 |
| P4          | 5.04        | 6.02       | 17.79       | 5.87     | 32.43                 |
| P5          | 6.04        | 6.36       | 23.82       | 5.99     | 43.37                 |
| P6          | 5.29        | 7.50       | 29.11       | 6.27     | 52.85                 |
| P7          | 3.23        | 8.73       | 32.35       | 6.51     | 58.57                 |
| P8          | 2.12        | 10.04      | 34.47       | 6.73     | 62.27                 |
| P9          | 3.34        | 12.32      | 37.81       | 7.22     | 67.93                 |
| P10         | 7.32        | 20.45      | 45.13       | 9.37     | 79.22                 |
| P11         | 8.17        | 42.64      | 53.30       | 14.47    | 88.30                 |
| P12         | 0.35        | 76.07      | 53.65       | 14.87    | 88.46                 |
| P13         | 9.44        | 77.74      | 63.09       | 24.28    | 92.53                 |
| Tailing     | 36.91       | 89.55      | 100.00      | 48.37    | 100.00                |

Total 100.00 48.37

2.6. Column Flotation Tests

Prior to each test, the frother solution was prepared by adding the near pure frother into tap water at a quantity that provided a concentration ranging from 22 ppm to 66 ppm (dependent on the final frother concentration target) followed by mixing in a high shear blender. The blended dilute frother solution was then added to the frother sump shown in Figure 2.

Initial froth flotation tests involved identifying the volumetric feed flow rate that resulted in a near optimum separation performance (i.e., maximum coal recovery and ash rejection). The feed flow rate was varied from 300 mL/min to 2000 mL/min at a constant feed solids concentration of 5% by weight. The optimum feed volumetric flow rate was determined to be 1000 mL/min. As such, all of the frother evaluation experiments were conducted at a constant feed volumetric flow rate of 1000 mL/min and solids content of 5% by weight.

The feed coal slurry was conditioned with fuel oil No. 2 at a concentration of 0.125 kg/t for a period of 15 min. The test was initiated by turning on the gas flow to the column to achieve a superficial gas velocity of 1.67 cm/s. As aforementioned, it was desired to maintain a constant froth depth (356 mm) as well as a superficial gas velocity (1.67 cm/s) for the purpose of assessing the effects of various frothers on the froth flotation performance. It is understood that in practice, an optimum separation can be achieved by plant operators by varying the froth depth and superficial gas velocity. Wash water was then added at a rate of 400 mL/min to fill the column. When the column was full, the bubble generation pump was turned on and the speed adjusted to achieve a 138 kPa (20 psi) feed pressure into the static mixer. The controller was set to achieve the desired froth level. Next, the feed slurry and frother pumps were started and adjusted to provide a combined flow rate of 1000 mL/min entering the column through the feed port. Steady-state operation was ensured before taking samples of the concentrate and tailings by allowing a time equal to three volumetric retention times after any change to the column operation. After collecting the samples, the tailings flow rate was measured and the bias factor was calculated and recorded. The pressure transducers’ readings were also recorded to quantify the air fraction within the collection zone. Detailed operational parameters for the laboratory column flotation tests are summarized in Table 3.
Table 3. Operational parameters for laboratory column flotation tests.

| Parameter                  | Value   |
|----------------------------|---------|
| Feed Flow Rate             | 1000 mL/min |
| Wash Water Flowrate        | 400 mL/min  |
| Solids Content             | 5 wt. %   |
| Superficial Air Velocities | 1.67 cm/s |
| Froth Height               | 356 mm    |
| Collector Dosage           | 0.125 kg/t |
| Solution pH                | ~7       |

Samples of the feed, clean coal concentrate, and tailings were collected, filtered, dried, and weighed. The dried samples were analyzed for ash content using a LECO proximate analyzer (St. Joseph, MI, USA). Mass yield was determined using the two-product equation, which was based on mass and component balance equations and the ash content of the three streams. Combustible and ash recovery values were determined from the yield and ash contents while the differences in the recovery values represented the separation efficiency.

3. Results and Discussion

3.1. Physical Properties Evaluation

The supplemental physical properties of all the nine frothers were measured and provided in Table 4. It can be seen that F-3, F-4, and DIBC were the three most surface-active frothers with a corresponding surface tension reduction ability of −0.22, −0.14, and −0.10 mN/m/ppm, respectively. The surface tension reduction ability of the frothers corresponds relatively well with their frothing ability. F-3, F-4, and DIBC were able to generate noticeably higher froth heights at identical concentrations relative to the majority of the remaining frothers (Table 4). The highest froth height was generated by F-3 corresponding to a value of 2.03 mm/ppm frother, which is indicative of its high surface activity. The surface tension reduction ability of the nine test frothers was further plotted versus the water solubility, as shown in Figure 3, which more clearly shows the order of the surface activity. MPC was the least surface-active of all nine frothers. In addition, Figure 3 also shows the correlation between the surface tension reduction ability and water solubility. A more soluble frother in aqueous solution correlates with less surface activity thereby limiting the reduction in surface tension. This fact agrees well with previous findings that alcohols with lower solubility are generally more effective than those with high solubility [1].

Table 4. Supplemental physical properties of all the nine evaluated frothers.

| Code | HLB | Surface Tension in DI Slope, mN/m/ppm | Froth Height Slope, mm/ppm |
|------|-----|----------------------------------------|---------------------------|
| MPC  | 6.53| −0.0078                                | 0.233                     |
| MAC  | 5.58| −0.0434                                | 1.353                     |
| DIBC | 4.63| −0.1000                                | 1.371                     |
| MIBC | 6.05| −0.0186                                | 0.776                     |
| 2EH  | 5.1 | −0.0791                                | 1.435                     |
| F-1  | 5.1 | −0.0583                                | 0.895                     |
| F-2  | 7.97| −0.0294                                | 1.609                     |
| F-3  | 6.25| −0.2157                                | 2.026                     |
| F-4  | 5.78| −0.1400                                | 1.235                     |
Furthermore, the HLB values of the frothers were evaluated as a function of their molecular weights as indicated in Figure 4. Prior studies have indicated that frothers are normally characterized as either “selective” or “powerful” [26]. The former term refers to the capability of frothers to recover fine particles while the latter indicates the frothing capacity of frothers in floating coarse particles [1,26,33]. The “selective” or “powerful” curves in Laskowski’s early study [26] were reproduced here in Figure 4. As seen, the frothers that fall on the left side of the diagram were more selective while those situated far to the right were more powerful and strong frothers [1,26]. Therefore, it can be derived from the diagram that MPC, MIBC, and MAC exhibited the properties of selective frothers whereas F3, F4, and DIBC were more powerful frothers. F1, F2, and 2EH fell between the two lines, which might represent the combination of two properties.

3.2. Bias Factor and Air Fraction Measurements

The bias factor and air fraction as a function of frother concentration were measured and recorded in each flotation test involving the nine frother types. As aforementioned, a bias factor of 60% was targeted to eliminate the amount of material being recovered by hydraulic entrainment. The test results shown in Figure 5 indicate that the bias factor values obtained during the flotation tests fell in the range of 50% to 85%, which is favorable and efficient for removing nearly all of the undesired gangue minerals recovered by entrainment. It is also noted that 2EH was the frother generating the highest bias factors with all three values equal to or exceeding 80% whereas F-3 produced the lowest value of 50% at the a frother concentration of 20 ppm. This finding reflects the stability of the bubble produced by the F-3 frother.
3.2. Bias Factor and Air Fraction Measurements

The bias factor and air fraction as a function of frother concentration were measured and recorded in each flotation test involving the nine frother types. As aforementioned, a bias factor of 60% was targeted to eliminate the amount of material being recovered by hydraulic entrainment. The test results shown in Figure 5 indicate that the bias factor values obtained during the flotation tests fell in the range of 50% to 85%, which is favorable and efficient for removing nearly all of the undesired gangue minerals recovered by entrainment. It is also noted that 2EH was the frother generating the highest bias factors with all three values equal to or exceeding 80% whereas F-3 produced the lowest value of 50% at the frother concentration of 20 ppm. This finding reflects the stability of the bubble produced by the F-3 frother.

In terms of air fraction, it can be clearly seen from Figure 6 that F-3, F-4, and DIBC generally gave higher air fractions with all values exceeding 20%, regardless of the frother concentration. In contrast, F-1, MPC, and MIBC generated relative lower air fractions, which were all below 13.3%. The findings presented here agree well with the known physical properties such as their surface tension reduction and frothing abilities (Table 4). In addition, the air fractions of the more frothing frothers do not change with various frother concentrations while the values increased slightly with frother concentration with the less effective frothers.

Figure 4. The hydrophile–lipophile balance (HLB)–molecular weight diagram for the nine evaluated frothers (after [26]).

Figure 5. Bias factor values measured over a range of frother concentrations at a fixed aeration rate for each of the nine frother types.

In terms of air fraction, it can be clearly seen from Figure 6 that F-3, F-4, and DIBC generally gave higher air fractions with all values exceeding 20%, regardless of the frother concentration. In contrast, F-1, MPC, and MIBC generated relative lower air fractions, which were all below 13.3%. The findings presented here agree well with the known physical properties such as their surface tension reduction and frothing abilities (Table 4). In addition, the air fractions of the more frothing frothers do not change with various frother concentrations while the values increased slightly with frother concentration with the less effective frothers.
3.3. Column Flotation Performance Evaluation

The effects of frother type and frother concentration on the separation performances achieved when treating ultrafine coal are shown in Figures 7–10, the data points of which have a standard deviation less than 3 percent in accordance with the replicate tests being performed. MPC was identified as the weakest among all the nine test frothers based on the data presented in Table 4 and Figure 5. At a 10 ppm concentration, MPC produced relatively large bubbles and was unable to support a moderate froth depth of around 356 mm. As a result, combustible recovery values obtained using MPC was the lowest among the nine frothers with the highest value being 75% at a concentration of 30 ppm. This finding could be explained by the lowest surface tension reduction and froth depth generated per ppm of the frother (Table 4). Frothers decreased the bubble size by adsorbing at the water–air interface and reducing the interfacial surface tension. Lower surface tension values helped to reduce the bubble size, which increased the flotation rate constant (k) and thus combustible recovery. Increasing dispersion between the air bubbles through elevated electrostatic repulsion forces was another mechanism provided by frothers that results in finer bubble size distributions and enhanced flotation performance. On the contrary, the less surface-active MPC frother resulted in a large bubble size distribution and relatively low bubble-particle collision efficiencies, which yielded slow flotation rates and low flotation recovery values.
Figure 3. At a 10 ppm concentration, MPC produced relatively large bubbles and was unable to support a moderate froth depth of around 356 mm. As a result, combustible recovery values obtained using MPC was the lowest among the nine frothers with the highest value being 75% at a concentration of 30 ppm. This finding could be explained by the lowest surface tension reduction and froth depth generated per ppm of the frother (Table 4). Frothers decreased the bubble size by adsorbing at the water–air interface and reducing the interfacial surface tension. Lower surface tension values helped to reduce the bubble size, which increased the flotation rate constant (k) and thus combustible recovery. Increasing dispersion between the air bubbles through elevated electrostatic repulsion forces was another mechanism provided by frothers that results in finer bubble size distributions and enhanced flotation performance. On the contrary, the less surface-active MPC frother resulted in a large bubble size distribution and relatively low bubble-particle collision efficiencies, which yielded slow flotation rates and low flotation recovery values.

The selectivity provided by the MPC frother was excellent as indicated by the proximity of the separation performances to the ideal performance (release analysis) curve in Figure 9. This finding is likely due to low water recovery and low bubble-particle attachment efficiencies, which limited recovery to the highest grade particles. Similarly, MIBC produced relatively high grade concentrates with an ash content of around 8% and low recovery values. The combustible recovery value was below 80% when using MIBC at a 30 ppm frother concentration. Dry, brittle foams were observed during the column tests with MIBC, which were not ideal optimum flotation performances. It should be noted, however, that the performances of both MIBC and MPC were comparable to the ideal performance when producing low ash concentrates. This corresponds well with the fact that both MPC and MIBC were categorized by Laskowski’s classification [26] as “selective” (Figure 3). The results presented were also consistent with the fact that the aliphatic alcohol frothers containing 5–8 carbon atoms (Figure 1) tended to produce less persistent but more selective froth compared to other alcohol mixtures [1]. Lower volumetric feed flow rates would produce recovery values similar to the other seven frother types. However, weak frothers like MPC and MIBC lead to lower recovery values at high feed rates due to kinetic and carrying capacity limiting conditions.

Figure 7. Combustible recovery values obtained as a function of frother concentration for each frother type.

Figure 8. Product ash content as a function of frother concentration for each frother type. Visual observations of the froth generated using 2EH and F-1, both of which are C-8 alcohols, gave a more productive froth that freely overflowed as compared to the stiffer, brittle froth produced by MPC and MIBC. As a result, higher combustible recovery values were produced by 2EH and F-1 at the same frother concentrations. The highest recovery value of 86.2% was obtained at a frother concentration of 10 ppm (Figure 7). This finding was expected considering the relatively strong surface tension reduction ability of $-0.08 \text{ mN/m/ppm}$ and high froth height of 1.44 mm/ppm associated with the 2EH frother (Table 4).

Figure 9. Combustible recovery of evaluated frothers as a function of clean coal ash; the solid line represents the results obtained from release analysis using MIBC as a frother.
Visual observations of the froth generated using 2EH and F-1, both of which are C-8 alcohols, gave a more productive froth that freely overflowed as compared to the stiffer, brittle froth produced by MPC and MIBC. As a result, higher combustible recovery values were produced by 2EH and F-1 at the same frother concentrations. The highest recovery value of 86.2% was obtained at a frother concentration of 10 ppm (Figure 7). This finding was expected considering the relatively strong surface tension reduction ability of \(-0.08 \text{ mN/m/ppm}\) and high froth height of \(1.44 \text{ mm/ppm}\) associated with the 2EH frother (Table 4).

Figure 8. Product ash content as a function of frother concentration for each frother type.

Figure 9. Combustible recovery of evaluated frothers as a function of clean coal ash; the solid line represents the results obtained from release analysis using MIBC as a frother.

Figure 10. Mass yield of evaluated frothers as a function of clean coal ash.

Figure 11. Separation efficiency of evaluated frothers as a function of frother concentration; error bars represent one standard deviation of three replicate tests.

The selectivity provided by the MPC frother was excellent as indicated by the proximity of the separation performances to the ideal performance (release analysis) curve in Figure 9. This finding is likely due to low water recovery and low bubble-particle attachment efficiencies, which limited recovery to the highest grade particles. Similarly, MIBC produced relatively high grade concentrates with an ash content of around 8% and low recovery values. The combustible recovery value was below 80% when using MIBC at a 30 ppm frother concentration. Dry, brittle foams were observed during the column tests with MIBC, which were not ideal optimum flotation performances. It should be noted, however, that the performances of both MIBC and MPC were comparable to the ideal performance when producing low ash concentrates. This corresponds well with the fact that both MPC and MIBC were categorized by Laskowski’s classification [26] as “selective” (Figure 3). The results presented were also consistent with the fact that the aliphatic alcohol frothers containing 5–8 carbon atoms (Figure 1) tended to produce less persistent but more selective froth compared to other alcohol mixtures [1]. Lower volumetric feed flow rates would produce recovery values similar to the other seven frother
types. However, weak frothers like MPC and MIBC lead to lower recovery values at high feed rates due to kinetic and carrying capacity limiting conditions.

Visual observations of the froth generated using 2EH and F-1, both of which are C-8 alcohols, gave a more productive froth that freely overflowed as compared to the stiffer, brittle froth produced by MPC and MIBC. As a result, higher combustible recovery values were produced by 2EH and F-1 at the same frother concentrations. The highest recovery value of 86.2% was obtained at a frother concentration of 10 ppm (Figure 7). This finding was expected considering the relatively strong surface tension reduction ability of $-0.08 \text{ mN/m/ppm}$ and high froth height of $1.44 \text{ mm/ppm}$ associated with the 2EH frother (Table 4).

With one exception, combustible recovery values generally increased with an elevation in frother concentration, which was the anticipated trend (Figure 7). The one exception was 2EH. The tests involving 2EH were repeated multiple times and the same trend was observed in all cases despite mixing the frother in water at concentrations well below the water solubility limit. A possible explanation could be the synergistic effect of the frother–collector depression interaction or the frother–collector–mineral system on the column flotation performance [25,35]. When relatively high dosages are applied, the extra frother molecules may be adsorbed on the coal particle surface, which converts the solid surfaces from hydrophobic to hydrophilic thereby generating a repulsion force between coal particles and similarly charged bubbles [6,36]. If this is the case for 2EH, this may explain the negative impact on recovery observed when increasing frother dosage above 10 ppm. MAC generates a froth height of $1.35 \text{ mm/ppm}$, which was about the same as the froth height generated by 2EH despite the lower surface tension reduction ability of $-0.04 \text{ mN/m/ppm}$ (Table 4). It can be seen from Figure 7 that MAC achieved a combustible recovery of around 83% with a cleaner coal ash content of 10.3% compared to the value of 14.16% achieved by 2EH at a 10 ppm dosage. The similar froth heights generated by 2EH and MAC correlated well with the similar combustible recovery values. The lower surface tension of 2EH generally implies the ability to produce smaller bubbles with larger superficial bubble surface area flux and more hydrophilic entrainment (higher product ash content) compared to MAC at 10 ppm concentration. Given that the bias factor (Figure 5) was sufficiently high to eliminate hydraulic entrainment, the more likely explanation for the higher product ash contents when using 2EH was the larger bubble surface area flux, which allowed for the recovery of more floatable material, including the more weakly, high ash content particles. Increasing the volumetric feed flow rate and/or the feed solids concentration would make the flotation environment more competitive and thus reduce the product ash content to values similar to MAC. If this hypothesis is correct, 2EH would be the superior frother given that higher mass and volume throughput capacities could be realized while achieving separation performances similar to MAC. Additional tests would be needed to confirm the hypothesis, which would involve conducting tests over a range of volumetric feed rates and feed solid concentrations at a given frother concentration.

A similar trend was obtained for F-3, F-4, and DIBC with an increase in the frother dosage. A combustible recovery of around 83% and 84% was achieved by DIBC and F4, respectively, while the highest combustible recovery of around 87% was generated by F-3 (Figure 7). This can be explained by its super strong surface tension reduction ability as indicated in Table 4 by the $-0.22 \text{ mN/m/ppm}$ value, which was the largest among all nine frother types. In addition, F-3, F-4, and DIBC were the three frothers with the longest hydrocarbon chains (C9–C12, Figure 1) and higher molecular weights (Table 1), which determine the power and performance of the frothers. Frothers with a long carbon chain and high molecular weight normally produce more persistent and less selective froth [1], which were confirmed again by the flotation results presented in the study. The results were also consistent with the aforementioned findings that F-3, F-4, and DIBC were identified as more “powerful” frothers (Figure 4). F-2, F-3, F-4, and DIBC generated relative higher recoveries along with MAC, which corresponded well with the relatively high froth heights generated per unit concentration, which varied from 1.24 to 2.03 mm/ppm frother.
It can be seen from Figure 8 that the corresponding clean coal ash of DIBC, F-3, and F-4 was relatively high ranging from 14.3% to 23.2%. This finding was likely due to the strong surface activity associated with these three frother types (Table 4), which greatly reduced the bubble size and increased the superficial bubble surface area flux as explained for the 2EH frother. Visual observations indicated that these frother types produced the smallest bubble distributions, which was confirmed by the highest air fraction values in the collection zone ranging from 20.8% to 24.6% (Figure 6). The relatively lower bias factors varying from 50% to 75% (Figure 5) suggests the stability of the froth produced by these frother types. F-2 and MAC generated relatively cleaner coal products with ash content values ranging from 10.3% to 13.6% while maintaining a high level of combustible recovery between 82% and 87%. The flotation performances achieved when testing F-2 and MAC approached the release curve performance as shown in Figure 9. The mass yield versus clean coal ash curve shown in Figure 10 further indicates that MPC generated the lowest yield among all the tests while the highest yields (>54%) were produced by F-2, F-3, and F-4. It is also noticed that F-2 and MAC produced a higher mass yield at the same clean coal ash content (<15%). In addition, the separation efficiency values achieved when using the nine frother types at different concentrations are shown in Figure 11 with the error bars representing one standard deviation of three replicate tests. The highest separation efficiency of around 72% was realized when using F-2 and MAC. A concentration of 10 ppm with 2EH performed in the same manner; however, a further increase in the dosage depressed both the recovery and separation efficiency.

![Figure 11. Separation efficiency of evaluated frothers as a function of frother concentration; error bars represent one standard deviation of three replicate tests.](image)

The F-3, F-4, and DIBC frother types produced the highest product ash values among the nine frothers tested. Given that the bias factors are well above the value that was considered critical for eliminating hydraulic entrainment (Figure 5), it was likely that the high bubble surface area flux rates generated by using F-3, F-4, and DIBC resulted in the flotation of weakly hydrophobic, high ash particles. To reject these particles, retention time could be reduced by increasing feed volumetric flow rate and higher feed solid concentrations could be introduced to the system to create a more competitive bubble-particle attachment environment. In other words, the use of F-3, F-4, or DIBC
Frother types have the potential to increase the volumetric and mass throughput capacity of a given flotation system while maintaining high recovery values. Further tests are needed to provide evidence of this potential technical and economic benefit. Besides, it should be kept in mind that the flotation differences for various frothers were generated under the constrained experimental conditions (e.g., fixed froth depth and superficial gas velocity) with one particular high-quality bituminous coal. The differences could be overcome by varying the operating conditions to achieve optimum performance in terms of the mass yield and clean coal grade. Additional tests may be needed when switching to a different coal feedstock, which is of considerable significance from the practical standpoint; however, it is not the scope of the presented work.

4. Conclusions

The results of our study showed that MPC and MIBC produced the poorest performance among all the nine frother types evaluated due to their weak froth properties. Test results also indicated F-1 was a relatively weak frother and higher concentrations than those tested were needed to boost its separation performance. F-2 and MAC were proven to be excellent frother types providing optimum separation performances at the mass and volumetric feed flow rates utilized in the test program. 2EH provided a comparable performance as F-2 and MAC at a concentration of 10 ppm. However, higher frother concentrations deteriorated performance as quantified by recovery and separation efficiency values. DIBC, F-3, and F-4 also proved to be excellent frothers as indicated by relatively high air fractions in the collection zone, which resulted in a high level of recovery. However, the three frother types produced the highest clean coal ash content values among the nine frothers. Given that the bias factor measured during each test involving the frother was sufficient to eliminate hydraulic entrainment, it may be concluded that the high product ash contents were due to an excessive bubble surface area flux, which resulted in the ability to float weakly hydrophobic, low grade particles. This finding means that flotation systems will be capable of treating higher volume and mass feed flow rates using DIBC, F-3, and F-4 compared to the other frothers. Furthermore, the separation performances achieved when using F-3 and F-4 frother types are far to the right of the MAC and F-2 performances on the recovery versus product ash curve, which implies that the mass and volume feed flow rates will be significantly higher when using F-3 or F-4 frother types.

Specific findings from the present study are summarized as follows:

1. The frothers that provided the weakest performances in terms of combustible recovery were F-1, MIBC, and MPC in order of decreasing effectiveness.
2. F-2 was consistently the stronger frother among the nine frothers tested over a 10 to 30 ppm concentration range followed by 2EH and F-3.
3. The tests with 2EH were repeated multiples times. In all cases, recovery decreased with an elevation in frother concentration. This response was obtained despite the froth concentration being significantly below the water solubility limit. The performance at 10 ppm, however, yielded a combustible recovery value comparable to F-2.
4. In regards to selectivity, F-2 and MAC provided the highest separation efficiency values, which agreed well with Laskowski’s HLB–molecular weight diagram [26], according to which F-2 and MAC are classified as more selective frothers.
5. DIBC and F-3 frothers provided high recovery values. However, the product ash values obtained when using these two frothers were among the highest of the nine tested. The F-4 frother provided the highest ash content values. In the case of the three frothers, the bias factors were generally above 60%, which should be sufficient to eliminate the hydraulically entrained material. This is supported by the proximity of the separation performances to the ideal performance curve generated by release analyses. As such, we could conclude that the DIBC, F-3, and F-4 frothers might provide separation efficiency performances comparable to F-2 and MAC if the volumetric feed rate was increased above the 1000 mL/min value used throughout the test program. If this hypothesis were true, the DIBC, F-3, and F-4 frothers provide the potential for higher volumetric throughput capacities.
which is important for retention time constrained flotation systems. Based on a relative comparison of
the separation performance with the ideal recovery-product ash (release analysis) curve, the F-3 has
the potential for treating the highest volumetric throughput capacities.

Author Contributions: Conceptualization, Q.H., X.Y. and R.Q.H.; methodology, Q.H., X.Y. and R.Q.H.; formal
analysis, Q.H., X.Y. and R.Q.H.; resources, Q.H., X.Y. and R.Q.H.; writing—original draft preparation, Q.H. and
X.Y.; writing—review and editing, Q.H., X.Y. and R.Q.H.; supervision, R.Q.H.

Funding: This research received no external funding.

Conflicts of Interest: The authors declare no conflict of interest.

References
1. Khoshdast, H.; Sam, A. Flotation frothers: Review of their classifications, properties and preparation. Open
Miner. Process. J. 2011, 4, 25–44. [CrossRef]
2. Tan, Y.H.; Zhang, W.; Finch, J.A. Frother structure-property relationship: Effect of polyethylene glycols on
bubble rise velocity. Miner. Eng. 2018, 116, 56–61. [CrossRef]
3. Leja, J.; Schulman, J.H. Flotation theory: Molecular interactions between frothers and collectors at
solid–liquid–air interfaces. Trans. AIME 1954, 199, 221–228.
4. Leja, J. Mechanisms of Collector Adsorption and Dynamic Attachment of Particles to Air Bubbles as Derived
from Surface-Chemical Studies. Trans. IMM 1956, 66, 425–437.
5. Gupta, A.K.; Banerjee, P.K.; Mishra, A.; Satish, P. Effect of alcohol and polyglycol ether frothers on foam
stability, bubble size and coal flotation. Int. J. Miner. Process. 2007, 82, 126–137. [CrossRef]
6. Drzymala, J.; Kowalczyk, P. Classification of flotation frothers. Minerals 2018, 8, 53. [CrossRef]
7. Cho, Y.S.; Laskowski, J.S. Effect of flotation frothers on bubble size and foam stability. Int. J. Miner. Process.
2002, 64, 69–80. [CrossRef]
8. Cho, Y.S.; Laskowski, J.S. Bubble Coalescence and its effect on dynamic foamability. Can. J. Chem. Eng. 2002,
80, 299–305. [CrossRef]
9. Laskowski, J.S. Fundamental properties of flotation frothers. In Proceedings of the 22nd International
Mineral Processing Congress, Cape Town, South Africa, 29 September 2003; Volume 2, pp. 788–797.
10. Laskowski, J.S.; Cho, Y.S.; Ding, K. Effect of frothers on bubble size and foam stability in potash ore flotation
systems. Can. J. Chem. Eng. 2003, 81, 63–69. [CrossRef]
11. Laskowski, J.S.; Tlhone, T.; Williams, P.; Ding, K. Fundamental properties of the polyoxypropylene alkyl
ether flotation frothers. Int. J. Miner. Process. 2003, 72, 289–299. [CrossRef]
12. Grau, R.A.; Heiskanen, K. Bubble size distribution in laboratory scale flotation cells. Miner. Eng. 2005, 18,
1164–1172. [CrossRef]
13. Melo, F.; Laskowski, J.S. Fundamental properties of flotation frothers and their effect on flotation. Miner. Eng.
2006, 19, 766–773. [CrossRef]
14. Finch, J.A.; Nesset, J.E.; Acuña, C. Role of frother on bubble production and behaviour in flotation. Miner.
Eng. 2008, 21, 949–957. [CrossRef]
15. Szyszka, D. Critical Coalescence Concentration (CCC) for surfactants in aqueous solutions. Minerals 2018, 8,
431. [CrossRef]
16. Savassi, O.N.; Alexander, D.J.; Franzidis, J.P.; Manlapig, E.V. An empirical model for entrainment in industrial
flotation plants. Miner. Eng. 1998, 11, 243–256. [CrossRef]
17. Yianatos, J.B.; Finch, J.A.; Laplante, A.R. Selectivity in column flotation froths. Int. J. Miner. Process. 1988, 23,
279–292. [CrossRef]
18. Wang, L.; Peng, Y.; Runge, K. Entrainment in froth flotation: The degree of entrainment and its contributing
factors. Powder Technol. 2016, 288, 202–211. [CrossRef]
19. Engelbrecht, J.A.; Woodburn, E.T. The Effects of Froth Height, Aeration Rate, and Gas Precipitation on Flotation;
Mintek: Randburg, South Africa, 1975.
20. Lynch, A.J.; Johnson, N.W.; Manlapig, E.V.; Thorne, C.G. Mineral and Coal Flotation Circuits; Elsevier:
Amsterdam, The Netherlands, 1981; Volume 65, p. 305.
21. Maachar, A.; Dobby, G.S. Measurement of feed water recovery and entrainment solids recovery in flotation
columns. Can. Metall. Q. 1992, 31, 167–172. [CrossRef]
22. Rahal, K.; Manlapig, E.; Franzidis, J.P. Effect of frother type and concentration on the water recovery and entrainment recovery relationship. *Miner. Metall. Proc.* 2001, 18, 138–141. [CrossRef]

23. Wang, L.; Peng, Y.; Runge, K. The mechanism responsible for the effect of frothers on the degree of entrainment in laboratory batch flotation. *Miner. Eng.* 2017, 100, 124–131. [CrossRef]

24. Kursun, H. The influence of frother types and concentrations on fine particles’ entrainment using column flotation. *Sep. Sci. Technol.* 2017, 52, 722–731. [CrossRef]

25. Subrahmanyam, T.V.; Forssberg, E. Froth stability, particle entrainment and drainage in flotation—A review. *Int. J. Miner. Process.* 1988, 23, 33–53. [CrossRef]

26. Laskowski, J.S. Testing flotation frothers. *Fizykochem. Probl. Miner./Physicochem. Probl. Mi.* 2004, 38, 13–22.

27. Ata, S.; Ahmed, N.; Jameson, G.J. Collection of hydrophobic particles in the froth phase. *Int. J. Miner. Process.* 2002, 64, 101–122. [CrossRef]

28. Ata, S.; Ahmed, N.; Jameson, G.J. A study of bubble coalescence in flotation froths. *Int. J. Miner. Process.* 2003, 72, 255–266. [CrossRef]

29. Ata, S.; Ahmed, N.; Jameson, G.J. The effect of hydrophobicity on the drainage of gangue minerals in flotation froths. *Miner. Eng.* 2004, 17, 897–901. [CrossRef]

30. Bournival, G.; Souza, L.D.O.; Ata, S.; Wanless, E.J. Effect of alcohol frothing agents on the coalescence of bubbles coated with hydrophobized silica particles. *Chem. Eng. Sci.* 2015, 131, 1–11. [CrossRef]

31. Honaker, R.Q.; Saracoglu, M.; Huang, Q. Application of hydrophobic and magnetic plastic particles for enhanced flotation recovery. *Miner. Eng.* 2016, 98, 223–231. [CrossRef]

32. Honaker, R.; Saracoglu, M.; Huang, Q. Evaluation of a novel coal flotation improvement approach with the addition of hydrophobic magnetic particles. *Int. J. Coal Prep. Util.* 2017, 1–17. [CrossRef]

33. Pugh, R.J. Experimental techniques for studying the structure of foams and froths. *Adv. Colloid Interfac.* 2005, 114, 239–251. [CrossRef]

34. Davies, J.T. A quantitative kinetic theory of emulsion type. I. Physical chemistry of the emulsifying agent. In Proceedings of the International Congress of Surface Activity, London, UK, 8–12 April 1957; Volume 1, pp. 426–438.

35. El-Shall, H.; Abdel-Khalek, N.A.; Svoronos, S. Collector–frother interaction in column flotation of Florida phosphate. *Int. J. Miner. Process.* 2000, 58, 187–199. [CrossRef]

36. Bueno-Tokunaga, A.; Pérez-Garibay, R.; Martínez-Carrillo, D. Zeta potential of air bubbles conditioned with typical froth flotation reagents. *Int. J. Miner. Process.* 2015, 140, 50–57. [CrossRef]