Influence of *Bacillus cereus*-Gold Interaction on Bio-flotation of Gold in the Presence of Potassium Butyl Xanthate

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Received: 3.01.2021; Revised: 30.01.2021; Accepted: 1.02.2021; Published: 7.02.2021

Abstract: *Bacillus cereus* is used to enhance the gold floatability in the presence of potassium butyl xanthate (PBX) as a collector and pine oil as a frother. The effect of bacteria interaction on the gold and quartz behavior as the main components of the gold ore was investigated using zeta-potential, contact angle, SEM, and FTIR measurements. The effect of pH, contact time, temperature, and concentration of xanthate, pine oil, and *Bacillus cereus* on the floatability of the gold and quartz was investigated. The zeta-potential of gold is strongly affected by *Bacillus cereus* interaction, while the maximum floatability is achieved at neutral pH. A concentrate of 10612 g/t gold with 95% recovery is obtained from a binary mixture contains 500 g/t gold in the presence of $1 \times 10^8$ *Bacillus cereus* cells, $5 \times 10^{-3}$ M potassium butyl xanthate and $5 \times 10^{-3}$ M pine oil at pH 7 and 35°C for 10 min.

Keywords: gold; *Bacillus cereus*; bio-flotation; potassium butyl xanthate; bio-beneficiation.

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1. Introduction

Gold element has special characteristics such as lustrous color and high resistance. It is used in jewelry, electronic circuits, and medical applications due to its unique physical and chemical properties [1]. Gold has been used in the electronic industries due to its higher electrical conductivity and corrosion resistance. Gold occurs in nature as metal or as alloys with other metals. It is usually associated with quartz or pyrite minerals. It is a rare element with a concentration of 0.005 g/t of the earth’s crust. The low-grade deposits contain 3 and 6 g/t, so it is challenging to meet the commercial grade (3000-4000 g/t). Usually, gold is concentrated in its ores using gravity, flotation, or cyanidation techniques, while refining is carried out by electrolysis [2, 3].

Although the flotation technique is widely applied for the gold enrichment in the gold-bearing mineral ores, there is no enough research for the flotation of gold-bearing ores. Flotation is essential to processing the low-grade gold ore and removing the undesired impurities before the next extraction process. It is also used for treating complex gold ores such as refractory and metallic ores. Except for the gravity separation technique, flotation is a cost-effective method for gold ore processing. However, gold flotation has many problems depending on the ore’s composition [4, 5].
Nowadays, biological processing routes are sought to solve the problems associated with lean grade ores. The traditional methods fail to separate the minerals from complex ores. Bio-beneficiation refers to the selective removal of undesirable mineral constituents from ore by utilizing microorganisms as surface modifiers, depressants, collectors, or dispersing agents to enhance the separation of one mineral from another either by flotation or flocculation [6-8], thus enriching it concerning the desired valuable minerals. Bio-beneficiation processes are relatively new and are under intense investigation in recent years. Most of the studies are so far confined to the laboratory [9-11]. Since the bacteria adhere to a mineral surface within a few minutes and alter the surface properties essential in mineral beneficiation techniques, the microorganisms have numerous applications in the flotation and flocculation processes. The behavior of bacterial cells on the mineral surface is the basis for successfully bio-beneficiation process. The selective bacterial adhesion on the mineral surface is important for selective surface modification, which leads to efficient separation [12].

This work investigates Bacillus cereus's behavior as a surface modifier for gold to enhance its floatability and thus its separation from its synthetic mixture with quartz.

2. Materials and Methods

2.1. Materials

Two pure gold samples were delivered from “Alfa-Easer” Company, UK. The first sample of less than 0.149 mm particle size (99.998%) was employed in flotation experiments, while the second is less than 0.074 mm (spherical, 99.9%) was used for surface behavior investigation. The pure quartz sample of 0.149 mm particle size was supplied by the Egyptian Mineral Resources Authority (EMRA). Analytical grade NaOH, HCl, and HNO₃ were used in this study. Potassium butyl xanthate (PBX), was used as a flotation collector. All chemicals are supplied by Sigma-Aldrich, Germany. Pure Pine oil of chemical grade is supplied by El-Nasr Chemical Company and Guangzhou chemicals Co., Ltd. Freshly prepared nutrient broth/ agar was used for cultivation of bacterial strains.

2.2. Methods

2.2.1. Bacterial isolation

Bacillus cereus is isolated, purified by streaking on nutrient agar plates, then transferred to nutrient agar slopes stored at 4°C and sub-cultured monthly. The nutrient agar was used for the cultivation of bacterial strains. The counting of the bacterial cells was carried out using a hemocytometer [10].

2.2.2. Measurements

A laser Zeta Meter ‘Malvern Instruments Model Zeta Sizer NAno ZS’ was used for zeta potential measurements. A 0.05 g of a solid sample is placed in 50 ml 10⁻² M NaCl solution. A definite amount of the bacteria is added, and then it is conditioned for 60 minutes at definite pH. The contact angle measurements for the pure gold and quartz before and after conditioning with different bacteria concentration were carried out using Contact Angle meter Model Drop-Master DM-701 at room temperature (25°C). All measurements were performed three times and averaged as a confirmed result.
The morphology and structural formation were determined using a scanning electron microscope (SEM) model JEOLJSM-5400, Japan. For FTIR analysis, the spectrum is obtained with KBr pellets prepared with solid sample and analytical grade KBr from Merck. The FTIR spectrum was obtained with a Spectrum 2000 Perkin Elmer spectrometer of 4000 and 400 cm\(^{-1}\) range.

The gold was determined in the binary mixture and concentrates using “Rigaku super Mini 200” X-ray fluorescence. The sample was pressed into a pellet by mixing the sample with Boreox BM-0008 binder, and the mixture was subjected to 20-ton press using “Fluxana PR-25N” for 20 Sec.

2.2.3. Flotation experiments

A bench-scale flotation experiment was carried out using a flotation column with 100 ml capacity. One gram of solid was conditioned using definite conditions of pH, bacterial concentration/PBX, and conditioning time. The flotation was conducted for 5 minutes at an air follow the rate of 0.65 cm\(^3\)/min. Both floated and sank fractions were collected, dried, weighted, and chemically analyzed. All flotation experiments, whether pure gold and quartz or their mixture, were carried out three times and averaged as a confirmed result.

3. Results and Discussion

3.1. Surface characteristic of pure gold and quartz

3.1.1. Zeta potential and contact angle

Fig. 1. shows that the isoelectric point for quartz is located at pH 2.1. The quartz as an oxide mineral forms a hydroxylated surface when it is in contact with water vapor. The adsorption of H\(^+\) onto the hydroxylated quartz surface compresses the diffuse electrical double layer. Therefore, the zeta potential of quartz becomes more positive. On the contrary, OH's adsorption onto the hydroxylated surface would increase the thickness of the diffuse electrical double layer [13]. The gold particles have a negative charge at all pH range. The Zeta potential of the gold depends on pH, indicating that both H\(^+\) and OH\(^-\) are potentially determining ions. The results showed that the electronegativity of zeta potential increases gradually with increasing the pH. The IEP of the gold was not determined. It was reported that the isoelectric point of gold in chloride media is around pH 2 and suggested that the OH\(^-\) ions are adsorbed at the inert surface [14].

_Bacillus cereus_ is positively charged at pH 2 with an isoelectric point at pH 2.25, which agrees with the reported IEP of 2.5. The zeta potential of bacterial isolates alone was performed. It explained that the bacterial cell charge originates from dissociation or protonation of carboxyl and amino groups and consequently depends on pH. At higher pH, it becomes progressively negatively charged due to proton dissociation [15].

The gold surface is strongly affected due to bacterial interaction, and its IEP became 2.1 rather than 1.5. On the other hand, there is no significant change in quartz minerals' zeta potential after treatment. Although both bacteria and the mineral surfaces are negatively charged, the adhesion is based on the bacteria's surface heterogeneity with a polysaccharide envelope. It also includes hydroxyl, hydrophobic and ionic moieties. The flexible fimbriae regulate the surface charge. The dissolving ions from the mineral alter the bacteria's surface
charge to reduce repulsion [6]. The hydrogen bonding and chemical interaction also play significant roles in bacterial interaction with single minerals [16].

The contact angle characterizes the surface properties related to the flotation. The greater the contact angle, the greater is the work of adhesion between the particle and bubble, and the more resilient the system is against disruptive forces [17]. Fig. 2 shows the contact angle of the gold and quartz before and after treatment with Bacillus cereus. The contact angle of quartz surface is decreased from 28.9° to 9.6° with increasing bacteria concentration up to 5×10⁶ cells/ml. The contact angle of gold is slightly decreased with increasing bacteria concentration.

![Figure 1. Zeta potential of pure gold and quartz and their treated forms with Bacillus cereus.](image1)

![Figure 2. Contact angle of pure gold and quartz and their treated forms with Bacillus cereus.](image2)

3.1.2. FTIR of gold, quartz, and Bacillus cereus

The FTIR of bacteria usually includes the O−H, C−C, CH₂, C=O, C−N, and C=O bands for polysaccharides and lipids (protein) [18]. The main characteristic peaks for Bacillus cereus are located in the range from 425-874 cm⁻¹ and 1637 cm⁻¹ which belongs to C=O of amide group and of O−C=O carboxylic groups, 2075 cm⁻¹ for stretch C−H, C−H₂ and C−H₃ of alkyl groups, with stretching vibration bands of O−H and N−H at 2823-3719 cm⁻¹, Fig.3 [19].

FTIR of the pure quartz exhibited the characteristic bands associated with quartz crystals in 400-1200 cm⁻¹, Fig. 3A. It includes the peak at 695 cm⁻¹, which is due to the octahedral site bending symmetry of Si–O vibration group. It is characteristic of crystalline quartz. The tetrahedral silicate ion vibrations band is located at around 784 cm⁻¹. The bands around 1080 cm⁻¹ are due to the silicon-oxygen stretching vibrations. The absorption bands...
located at 440-650 cm\(^{-1}\) and 1885 cm\(^{-1}\) involve atomic motions within the SiO\(_4\) tetrahedra. As a result of *Bacillus cereus* interaction, there is no significant change in the quartz spectrum. It is believed that there is no interaction between *Bacillus cereus* and the quartz surface [20].

The FTIR spectrum of the pure gold includes a broad band at 3486-3668 cm\(^{-1}\) due to stretching of O–H which may be adsorbed on the gold surface. The spectrum of the treated gold with *Bacillus cereus* showed new functional groups, Fig. 3B. The bands are located at 1076, 1247, 1383, 1532 and 1648 cm\(^{-1}\) which belongs to C=O of the amide group and O–C=O of the carboxylic group. Also, stretch vibration band of O–H and N–H at 3417-3441 cm\(^{-1}\) and 3794 cm\(^{-1}\). The stretching peak of C–H, C–H\(_2\) and C–H\(_3\) of alkyl groups are shifted to 2853-2922 cm\(^{-1}\) [21]. It indicated the role of hydrogen bonding and chemical interaction. Result in significant surface chemical changes, not only on the cell surfaces but also on the gold [22]. It also proved that the type of adsorption that occurred is mainly chemical adsorption, making the gold surface more hydrophobic than that of the quartz.

![FTIR spectra of pure gold and quartz and their treated forms with Bacillus cereus.](image)

**Figure 3.** FTIR spectra of pure gold and quartz and their treated forms with *Bacillus cereus*.

### 3.1.3. SEM images of gold, quartz, and *Bacillus cereus*

Fig. 4 shows the Scanning Electron Microscope images for the pure quartz (A) and its treated form with *Bacillus cereus* (B). It is clear that there is no detected *Bacillus cereus* on the quartz surface. However, there is no significant change in the quartz surface after interaction with *Bacillus cereus*. On the other hand, Fig. 5 shows significant adsorption of *Bacillus cereus* on the gold surface. The gold surface is covered with condensed bio-film of both bacterial cells and biological metabolite. The results also showed the presence of nitrogen, oxygen, and carbon elements on the gold surface. This is due to the biofilm formation of bacteria onto the gold surface. It is proved the significant change in gold surface behavior, such as in the zeta-potential measurements and thus its floatability [23].
3.2. Floatability of pure gold and quartz

3.2.1. Effect of pine oil

It was reported that the gold exhibited 53% floatability while quartz cannot be float without any flotation reagents [24]. Pine oil was used as a frother in this study. Fig. 5 shows the floatability of pure gold metal and quartz minerals as a pine oil concentration function at pH 7. The pine oil increases the floatability of both gold and quartz due to its oleophilic character. Although the gold is naturally floating, pine oil enhanced its floatability to reach complete flotation at $10^{-2}$ M of pine oil. Increasing gold recovery as the concentration of pine oil increased due to the increase of bubble production. On the other hand, quartz is not floating. Increasing the pine oil leads the excess bubble formation; hence hydrophilic quartz particles are lifted with a gas bubble [24]. The quartz is floated to reach 89.5% at $8 \times 10^{-2}$ M pine oil. The maximum floatability difference ($\approx 84\%$) is achieved at $5 \times 10^{-3}$ M pine oil.

3.2.2. Effect of potassium butyl xanthate

Fig. 6. shows the floatability of pure gold and quartz as a function of potassium butyl xanthate (PBX) concentration as a selective collector in the presence of $1 \times 10^{-3}$ M pine oil. It is stated that the xanthates are absorbed on the surface of the minerals due to chemical reactions between the polar group and the surface, strongly hydrophobic insoluble metal xanthates being formed [25]. The floatability of quartz is slightly increased up to 11.6% at $8 \times 10^{-2}$ M then it is decreased to 7.8%. The maximum floatability difference ($\approx 85\%$) is achieved at $5 \times 10^{-3}$ M PBX.

Floatability of gold metal increases as PBX concentration increases to 95.9% at 0.01 M PBX then decreased to 84.5% at 0.05 M PBX. The flotation recovery is increased as the PBX
concentration increases because the PBX acts as a surfactant that increases the hydrophobic characteristic of the solid particles. The xanthate is chemisorbed on the gold surface at a potential which the xanthate can oxidize to dixanthogen as follow: [26]

\[
\text{ROCS}_2^- \rightleftharpoons (\text{ROCS}_2) \text{ads} + e^- \\
2 \text{ROCS}_2 \text{ads} \rightleftharpoons (\text{ROCS}_2)_2
\]

Where R denotes an alkyl chain.

![Graph](https://biointerfaceresearch.com/)

**Figure 5.** Effect of pine oil concentration on the floatability of pure gold and quartz particles.

**Figure 6.** Effect of potassium butyl xanthate concentration on the floatability of pure gold and quartz particles.

3.2.3. Effect of *Bacillus cereus*

Fig. 7 shows the effect of *Bacillus cereus* concentration (cell count) on the floatability of pure gold and quartz in the presence of 1×10⁻³ M pine oil at pH 7. *Bacillus cereus* increases the floatability of both gold and quartz. The gold presented its higher floatability (≈ 99%) in the presence of low bacterial concentration (1.5×10⁸ cells), while quartz showed its maximum floatability (≈ 99%) at a higher bacteria concentration (3.75×10⁸ cells). The maximum floatability difference (≈ 84%) is achieved in the presence of 1×10⁸ bacteria cells. At which the floatability of gold is 94%, while it is only 9.7% for quartz. These results agree with zeta potential, contact angle, FTIR, and SEM measurements, which showed the adsorption of
bacteria on the gold surface and the strong effect of bacterial interaction with gold. It is suggested that the attachment and agglomeration of bacterial cells to solid surfaces provide a stable growth environment for the cells and enhances catalytic functions through the localization of cells into biofilms. The excretion of an extracellular polymeric substance composed of macromolecules as polysaccharides, proteins, and lipids promotes biofilms' development on mineral surfaces. It is suggested that a Hydrogen bond is formed between bacteria and mineral surface as a result of the presence of \((\text{OH}^-)\) groups of polysaccharide of metabolite. The charge and spacing between them control the biomolecules' interaction with the solid surfaces [27].

![Figure 7. Effect of Bacillus cereus concentration on the floatability of pure gold and quartz particles.](https://biointerfaceresearch.com/)  

3.2.4. The dual effect of *Bacillus cereus* and potassium butyl xanthate  

The floatability of gold and quartz in the presence of \(5 \times 10^{-3}\) M potassium butyl xanthate and \(1 \times 10^{-3}\) M pine oil as a function of *Bacillus cereus* concentration is presented in Fig. 8. For conditioning with xanthate followed by *Bacillus cereus*, the floatability of gold and quartz increases with increasing bacteria concentration. The maximum floatability difference (≈ 85 %) is achieved in the presence of \(1 \times 10^8\) of bacteria cells. At which the floatability of gold is about 91% while it is only 6% for quartz.

Although the floatability of gold and quartz by conditioning with xanthate first is more than that of conditioning with *Bacillus cereus* first, the separation efficiency is better. This is due to the higher amount of floating quartz. Thus, the maximum floatability difference (≈ 89 %) is achieved in the case of conditioning with *Bacillus cereus* first in the presence of \(1 \times 10^8\) of bacteria cells. The floatability of gold is about 94%, while it is only 4% for quartz.

The conditioning with *Bacillus cereus* prior to xanthate may enhance the xanthate-gold adsorption while it decreases xanthate-quartz adsorption. The exopolymers, or metabolites, interact with the organism and the minerals in a variety of ways. The metabolites of *Bacillus cereus*, such as the polysaccharides, proteins, and organic acids, are responsible for the surface modification.

The metabolites may form a type of polymeric bridge in potassium butyl xanthate, leading to higher surface modification. Many of the interacting mechanisms are still unknown and need to explain how the bacteria adhere to the solid surface. This is confirmed by the different results for the mode of addition, the bacteria first or the xanthate first [28].
3.3. Binary mixture bio-flotation

The flotation study of binary mixture aims to determine the effect of interaction between gold and quartz, which may be occurred during their conditioning together rather than individual conditioning. A synthetic binary mixture composed of 0.05% of pure gold (500 g/t) with quartz was employed for this investigation.

3.3.1. Effect of Bacillus cereus

The effect of Bacillus cereus concentration (cell count) on the gold grade (g/t) and recovery, in the presence of $5 \times 10^{-3}$ M potassium butyl xanthate and $5 \times 10^{-3}$ M pine oil at 25°C and pH 7 showed that the gold grade is 9863 g/t in the presence of $1 \times 10^8$ bacteria cells, Fig. 9. Then it is dramatically decreased to 758 g/t in the presence of $3.75 \times 10^8$ bacteria cells. On the other hand, the recovery is increased from 39.4 to 80% in the presence of $1 \times 10^8$ bacteria cells. Then it is slightly increased up to 88% in the presence of $3.75 \times 10^8$ bacteria cells. Although the maximum grade (9863 g/t) with 39.4% recovery is achieved at $5 \times 10^7$ bacteria cells, $1 \times 10^8$ bacterial cells are considered the best result at which the grade was 8324 g/t with 80% recovery. As mentioned above, the adsorption of bacteria on the gold enhanced its flotation, which provides a more hydrophobic surface and stable agglomeration [28].

3.3.2. Effect of pH

Figure 8. Floatability of pure quartz and gold particles in the presence of Bacillus cereus and potassium butyl xanthate.

Figure 9. Effect of Bacillus cereus concentration on the gold content and recovery of the floated fraction from the binary mixture.
Neutral pH is more commonly used for native gold recovery. Native gold shows its maximum floatability in the water at pH = 6-9. The floatability is totally inhibited at pH > 10. The effect of pH on the grade and recovery of gold in the presence of bacteria, potassium butyl xanthate, and pine oil showed that the gold grade is increased from 7435 to 10442 g/t with increasing the pH from 5 to 7, Fig. 10. Then it is dramatically decreased to 2147 g/t at pH 9.

On the other hand, the recovery is increased from 46.1 to 90.2% at pH 7 - 8, and then it is decreased to 39.5% at pH 9. It was reported that the uncharged and water-insoluble collectors are the most effective collectors for the gold flotation at the natural pH. The xanthates are normally used in slightly alkaline pulps since they decompose in an acid medium and in a very alkaline environment. The hydroxyl ions (OH⁻) can displace the xanthate ions from the surface of the mineral [29].

\[ \text{OH}^- \] can displace the xanthate ions from the surface of the mineral [29].

**Figure 10.** Effect of pH on the gold content and recovery of the floated fraction from the binary mixture.

### 3.3.3. Effect of conditioning time

The effect of conditioning time on the gold grade and recovery in the presence of bacteria cells, potassium butyl xanthate, and pine oil at 25°C and pH 7 is investigated. Fig. 11 shows that the gold grade is increased from 7342 to 10442 g/t with increasing conditioning time from 5 to 10 min, and then it is decreased again to 3292 g/t after 15 min. On the other hand, the recovery is increased from about 28 to 90% by increasing the conditioning time from 5 to 10 min. Then it is slightly increased to 94% after 15 min. This is because the attachment of the particle with the air bubble needs time (induction time). Induction time is associated with the thin water film's properties that separate the particle and bubble just before attaching. For a hydrophobic surface, induction time is short, a few milliseconds. Suppose it is less than the time the particle and bubble are in contact. In that case, attachment is successful, and the particle is floated. The induction time is very large for hydrophobic particles. The hydrophobicity of gold depends on the time due to its higher Hamaker constant \((45\times10^{-20} \text{ J})\). It results from strong water dispersion forces, whereas its hydrophobicity develops upon physical adsorption of water vapor [17].

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3.3.4. Effect of temperature

The metabolism of bacteria is directly associated with enzymes' presence, which means that the biofilm formation depends on the enzymes' reaction rates. A pivotal factor that affects the enzymatic reaction rate is the temperature, as it is correlated to the formation of cells forming a biofilm [30]. The effect of temperature on both grade and recovery of gold in the presence of bacteria cells, potassium butyl xanthate, and pine oil at pH 7 is investigated. Fig. 12 shows that the gold grade increases from 9194 to 10612 g/t, increasing temperature from 20 to 35°C. Then it is decreased again to 7706 g/t at 40°C. On the other hand, the recovery is increased from 63 to 95% with increasing temperature from 20 to 35°C, and then it is decreased to 71% at 40°C. This is due to the microbial activity usually increases with increasing temperature. The higher the effect of bacteria may be due to increasing the biofilm thickness at 35°C compared with that at 20°C. It is ascribed to the number of appendages. It may also be due increase in the surface area and the likelihood of bacterial adhesion at 35°C [31-32]. The high temperature has little impact on removing the biofilm, so the biofilm adherent to the surface increases [34].

Figure 11. Effect of conditioning time on the gold content and recovery of the floated fraction from the binary mixture.

Figure 12. Effect of temperature on the gold content and recovery of the floated fraction from the binary mixture.
4. Conclusions

*Bacillus cereus* is used as a surface modifier to enhance the flotation selectivity of gold. The gold surface is strongly affected as a result of bacterial interaction, while the quartz doesn’t affect it. The gold surface is strongly affected by bacterial interaction, while there is no significant change of the zeta potential of quartz minerals after treatment. FTIR and SEM proved the selective adhesion of *Bacillus cereus* on the gold surface.

The maximum floatability of pure gold metal and quartz individually after treatment with *Bacillus cereus* is achieved at pH 7-8 for the gold and pH 7 for quartz. The maximum floatability is achieved after 10 min conditioning time at 35°C and pH 7.

The gold flotation from a binary mixture (500 g/t gold) produced a concentrate of 10612 g/t gold with 95% recovery in the presence of 1×10^8 bacteria cells, 5×10^{-3} M potassium butyl xanthate and 5×10^{-3} M pine oil at 35°C and pH 7 for 10 min.

**Funding**

This research received no external funding.

**Acknowledgments**

This research has no acknowledgment.

**Conflicts of Interest**

The authors declare no conflict of interest.

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