



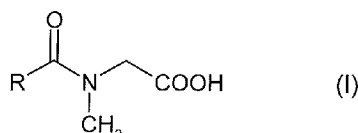
- (51) International Patent Classification:
B03D 1/008 (2006.01) *B03D 1/01* (2006.01)
- (21) International Application Number:
PCT/EP2016/054291
- (22) International Filing Date:
1 March 2016 (01.03.2016)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:
15000927.2 30 March 2015 (30.03.2015) EP
- (71) Applicant: **CLARIANT INTERNATIONAL LTD**
[CH/CH]; Rothausstr. 61, 4132 Muttenz (CH).
- (72) Inventors: **PEDAIN, Klaus-Ulrich**; Hauptstrasse 14,
63128 Dietzenbach-Steinberg (DE). **PITARCH LOPEZ,**
Jesus; Wilhelm-Busch-Str. 35, 60431 Frankfurt am Main
(DE). **LIPOWSKY, Gunter**; Schillerstrasse 8, 68526
Ladenburg (DE). **BEZUIDENHOUT, Jacques Collin**;
Feldbergstrasse 51, 61149 Steinbach (DE).
- (74) Agent: **MIKULECKY, Klaus**; Clariant Produkte
(Deutschland) GmbH, Industriepark Höchst / G 860, 65926
Frankfurt (DE).

- (81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.
- (84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:

— with international search report (Art. 21(3))

(54) Title: COMPOSITION OF FATTY ACIDS AND N-ACYL DERIVATIVES OF SARCOSINE FOR THE IMPROVED FLOTATION OF NONSULFIDE MINERALS



(57) Abstract: This invention relates to a collector composition for the direct froth flotation of nonsulfide ores comprising a) 50 - 99 wt.-% of a mixture of fatty acids and b) 1 - 50 wt.-% of an N-acyl derivative of sarcosine of the formula (I) wherein R is a saturated or unsaturated hydrocarbon chain with 7 to 21 carbon atoms, wherein the mixture of comprises 10.0 - 35.0 wt.-% of fatty acid having a saturated C₁₁ hydrocarbon group, 2.5 - 15.0 wt.-% of fatty acid having a saturated C₁₃ hydrocarbon group, 10.0 - 25.0 wt.-% fatty acid having a monounsaturated C₁₇ hydrocarbon group and 20.0 - 45.0 wt.-% fatty acid having a bisunsaturated C₁₇ hydrocarbon group.



COMPOSITION OF FATTY ACIDS AND N-ACYL DERIVATIVES OF SARCOSINE FOR THE IMPROVED FLOTATION OF NONSULFIDE MINERALS

This invention relates to a novel collector composition comprising a mixture of at least two fatty acids and at least one N-acyl derivative of sarcosine and its use in the direct froth flotation of nonsulfide minerals. The use of the novel collector composition provides improved flotation efficiency.

Froth flotation is a physico-chemical process used to separate mineral particles considered economically valuable from those considered waste. It is based on the ability of air bubbles to selectively attach onto those particles previously rendered hydrophobic. The particle-bubble combinations then rise to the froth phase from where it discharges the flotation cell whilst the hydrophilic particles remain in the flotation cell. Particle hydrophobicity is, in turn, induced by special chemicals called collectors. In direct flotation systems, it is the economically valuable minerals which are rendered hydrophobic by the action of the collector. Similarly, in reverse flotation systems, the collector renders hydrophobicity to those mineral particles considered waste. The efficiency of the separation process is quantified in terms of recovery and grade. Recovery refers to the percentage of valuable product contained in the ore that is removed into the concentrate stream after flotation. Grade refers to the percentage of the economically valuable product in the concentrate after flotation. A higher value of recovery or grade indicates a more efficient flotation system.

The use of mixtures of fatty acids and sarcosine derivatives for the froth flotation of nonsulfide minerals is well-known.

In DD-300730 the use of a collector composition for the froth flotation of fluorite comprising an N-acyl derivative of sarcosine and a saturated or unsaturated fatty acid having a hydrocarbon chain with 14 to 24 carbon atoms is described.

In US-5147528 a process for the direct flotation of phosphate is described where an oxidized intimate mixture of a fatty acid containing 12 to 36 carbon atoms, a tall

oil pitch, an amine derived from a plant, sarcosine and a fuel oil or furnace oil is used as collector.

US-4514290 describes a collector composition comprising a fatty acid or salt thereof, an amidocarboxylic acid or amidosulfonic acid containing an organic hydrophobic group, or a salt thereof, and a partial ester of phosphoric acid and at least one alkoxyated alcohol. Such composition is claimed to show improved efficiency for the froth flotation of minerals containing alkaline earth metals, such as apatite, scheelite, magnesite and barite. The fatty acid in the preferred compositions has 14 to 22 carbon atoms.

WO-2014040686 describes a flotation agent for phosphate ore, comprising at least one fatty acid and at least one N-acyl derivative of sarcosine.

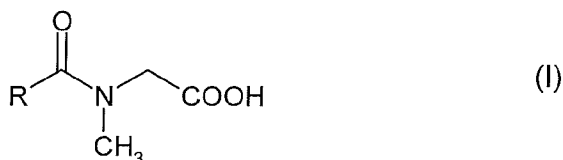
The present invention is related to a novel collector composition comprising a mixture of at least two fatty acids and at least one N-acyl derivative of sarcosine and its use for the beneficiation of nonsulfide minerals. The composition of at least two fatty acids and at least one N-acyl derivative according to the present invention affords in comparison to the collector compositions described by the state-of-the-art an improvement of the flotation efficiency. Under improved flotation efficiency is meant that higher mineral recovery and/or purity are achieved. Furthermore, the collector composition according to the present invention is very easy to prepare by simply mixing and not require any additional treatment, like for example an oxidation step, in order to show excellent improvement of the flotation efficiency.

Unexpectedly it was found that a composition containing 50 - 99 wt.-% of a mixture of at least two fatty acids and 1 - 50 wt.-% of an N-acyl derivative of sarcosine show improved flotation of nonsulfide minerals expressed in terms of higher mineral recovery and/or higher purity.

The instant invention therefore relates to a collector composition comprising

- a) 50 - 99 wt.-% of a mixture of fatty acids and
 b) 1 - 50 wt.-% of an N-acyl derivative of sarcosine of the formula (I)

5



wherein

R is a saturated or unsaturated hydrocarbon chain with 7 to 21 carbon atoms,
 10 wherein the mixture of fatty acids comprises 10.0 - 35.0 wt.-% of fatty acid having
 a saturated C₁₁ hydrocarbon group, 2.5 - 15.0 wt.-% of fatty acid having a
 saturated C₁₃ hydrocarbon group, 10.0 - 25.0 wt.-% fatty acid having a
 monounsaturated C₁₇ hydrocarbon group and 20.0 - 45.0 wt.-% fatty acid having a
 bisunsaturated C₁₇ hydrocarbon group.

15

The inventive collector composition may comprise other fatty acids to balance to
 100 wt.-%. The weight percentages refer to the total fatty acid content of the
 inventive collector composition as being 100 wt.-%.

- 20 In a preferred embodiment the mixture of fatty acids comprises fatty acids having
 1.0 - 6.5 wt.-% of saturated C₇
 1.0 - 4.0 wt.-% of saturated C₉
 10.0 - 35.0 wt.-% of saturated C₁₁
 2.5 - 15.0 wt.-% of saturated C₁₃
 25 1.0 - 7.0 wt.-% of saturated C₁₅
 0.0 - 1.0 wt.-% of monounsaturated C₁₅
 0.0 - 1.0 wt.-% of bisunsaturated C₁₅
 0.5 - 2.0 wt.-% of saturated C₁₇
 10.0 - 25.0 wt.-% of monounsaturated C₁₇
 30 20.0 - 45.0 wt.-% bisunsaturated C₁₇
 0.0 - 2.0 wt.-% trisunsaturated C₁₇
 0.0 - 1.0 wt.-% saturated C₁₉
 0.0 - 4.0 wt.-% monounsaturated C₁₉ hydrocarbon chains, and

0.0 - 7.0 wt.-% other fatty acids

The expression "saturated hydrocarbon chain" means preferably alkyl groups. The expression "monosaturated hydrocarbon chain" means preferably alkenyl groups.

- 5 The expression "bisunsaturated hydrocarbon chains" means alkenyl groups having two double bonds.

Fatty acids are defined in the sense of this invention as mixtures of carboxylic acids bearing a long linear hydrocarbon chain, which can be saturated or
10 unsaturated or multiply unsaturated. Especially effective for the scope of this invention is the use of fatty acids from vegetable oils and tall oil fatty acids. The preferred fatty acids in the sense of this invention are coconut oil fatty acid and tall oil fatty acid. Predominant carboxylic acids in the coconut oil fatty acid are lauric acid (saturated C₁₁ hydrocarbon chain) with a content between 44 and 54 wt.-%
15 and myristic acid (saturated C₁₃ hydrocarbon chain) with a content between 13 and 20 wt.-%. The preferred quality of tall oil fatty acid has an acid value higher than 190 mg KOH/g and a content of rosin acids and unsaponifiables lower than 2.1 wt.-% and 2.0 wt.-%, respectively. Predominant carboxylic acids in the tall oil fatty acid are oleic acid (monounsaturated C₁₇ hydrocarbon chain) with a content
20 between 25 and 50 wt.-% and linoleic acid (bisunsaturated C₁₇ hydrocarbon chain) with a content between 35 and 60 wt.-%.

The characterization of the alkyl chain distribution in fatty acids can be done via gas chromatography after conversion of the carboxylic acids in the volatile methyl
25 ester derivatives according to the AOCS Method Ce 1 - 62, "Fatty Acid Composition by Gas Chromatography" AOCS Official Methods (2005) American Oil Chemists Society.

The mixtures of fatty acids which are especially effective in the collector
30 compositions according to the present invention contain between 0.3 and 1.7 weight parts of fatty acids from vegetable oils to 1 weight part of tall oil fatty acid. The most preferred ratio for the mixture of fatty acids according to this

invention is 1 weight part of coconut oil fatty acid to 1 weight part of tall oil fatty acid.

Especially preferred are N-acyl derivatives of sarcosine where R is a saturated or
5 unsaturated hydrocarbon chain with 11 to 19 carbon atoms. The most preferred
N-acyl derivative of sarcosine is N-oleoylsarcosine.

Especially preferred collector compositions according to the present invention
contain 65 - 99 wt.-% of a mixture of fatty acids (component a) and 1 - 35 wt.-% of
10 an N-acyl derivative of sarcosine (component b). Most preferred collector
compositions contain 80 wt.-% of a 1:1 weight mixture of coconut oil fatty acid and
tall oil fatty acid and 20 wt.-% of N-oleoylsarcosine.

The composition of the invention is for use as collector in direct froth flotation
15 processes of nonsulfide ores. It was found that the composition of the invention is
especially suitable for the direct froth flotation of nonsulfide ores containing
alkaline earth metals, as apatite, calcite, scheelite, fluorspar, magnesite and barite.
Most surprisingly it was found that the composition of the invention is also
especially suitable for the direct froth flotation of ilmenite, a titanium-iron oxide
20 mineral of formula FeTiO_3 which is the most important source for titanium.

Furthermore, the present invention also relates to a process for beneficiation of
nonsulfide minerals, the process comprising the steps of bringing the collector
composition according to the present invention in contact with an aqueous
25 suspension of the nonsulfide mineral and frothing the so formed mineral pulp.
The collector composition according to the present invention is preferably used in
amounts between 100 and 1000 g/t of solid ore for the direct froth flotation of
nonsulfide ore. It is also possible to add other flotation reagents to the mineral
pulp, if these are required. Examples of these reagents are frothers as for example
30 pine oil, polyglycols, polyoxyparaffins or alcohols, depressants as for example
starch, carboxymethylcellulose or sodium silicate and pH-regulators as for
example sodium hydroxide or sodium carbonate.

Examples

1. General procedure for preparation of collector compositions according to
5 this invention:

Distilled coconut fatty acid sample was warmed to 35 °C until it was entirely melted and then added to tall oil fatty acid at room temperature. The fatty acid mixture was then homogenised by a slow stirring action. Finally, N-oleoylsarcosine was slowly added to the fatty acid mixture. The mixture was thereafter homogenised for
10 a further 10 minutes. The procedure is completed when a clear, yellow-coloured liquid solution is obtained.

2. Collector compositions according to this invention prepared following used
15 the procedure describer under 1.

Composition 1:

Component a:

80 wt.-% of a mixture of 1 weight part of distilled coconut fatty acid and
1 weight part of tall oil fatty acid with a hydrocarbon chain distribution
20 as follows:

3.25 wt.-% of saturated C₇ hydrocarbon chain

2.85 wt.-% of saturated C₉

25.65 wt.-% of saturated C₁₁

8.60 wt.-% of saturated C₁₃

25 4.45 wt.-% of saturated C₁₅

0.02 wt.-% of monounsaturated C₁₅

0.20 wt.-% of bisunsaturated C₁₅

1.25 wt.-% of saturated C₁₇

19.10 wt.-% of monounsaturated C₁₇

30 28.5 wt.-% bisunsaturated C₁₇

0.60 wt.-% trisunsaturated C₁₇

0.05 wt.-% saturated C₁₉

1.25 wt.-% monounsaturated C₁₉

4.3 wt.-% others

Component b:

20 wt.-% N-oleoylsarcosine

5 Composition 2:

Component a:

80 wt.-% of a mixture of 0.33 weight part of distilled coconut fatty acid and
1 weight part of tall oil fatty acid with a hydrocarbon chain distribution
as follows:

10 1.62 wt.-% of saturated C₇ hydrocarbon chain

1.42 wt.-% of saturated C₉

12.82 wt.-% of saturated C₁₁

4.30 wt.-% of saturated C₁₃

2.37 wt.-% of saturated C₁₅

15 0.04 wt.-% of monounsaturated C₁₅

0.30 wt.-% of bisunsaturated C₁₅

1.17 wt.-% of saturated C₁₇

24.70 wt.-% of monounsaturated C₁₇

42.15 wt.-% bisunsaturated C₁₇

20 0.90 wt.-% trisunsaturated C₁₇

0.02 wt.-% saturated C₁₉

1.87 wt.-% monounsaturated C₁₉

6.40 wt.-% others

Component b:

25 20 wt.-% N-oleoylsarcosine

Composition 3:

Component a:

80 wt.-% of a mixture of 1.66 weight part of distilled coconut fatty acid and
30 1 weight part of tall oil fatty acid with a hydrocarbon chain distribution
as follows:

4.10 wt.-% of saturated C₇ hydrocarbon chain

3.59 wt.-% of saturated C₉

32.32 wt.-% of saturated C₁₁
10.84 wt.-% of saturated C₁₃
5.53 wt.-% of saturated C₁₅
0.02 wt.-% of monounsaturated C₁₅

5 0.15 wt.-% of bisunsaturated C₁₅
1.29 wt.-% of saturated C₁₇
16.19 wt.-% of monounsaturated C₁₇
21.40 wt.-% bisunsaturated C₁₇
0.44 wt.-% trisunsaturated C₁₇
10 0.06 wt.-% saturated C₁₉
0.92 wt.-% monounsaturated C₁₉
3.21 wt.-% others

Component b:

20 wt.-% N-oleoylsarcosine

15

Composition 4:

Component a:

67 wt.-% of a mixture of 1 weight part of distilled coconut fatty acid and
1 weight part of tall oil fatty acid with a hydrocarbon chain distribution
20 as follows:

3.25 wt.-% of saturated C₇ hydrocarbon chain

2.85 wt.-% of saturated C₉

25.65 wt.-% of saturated C₁₁

8.60 wt.-% of saturated C₁₃

25 4.45 wt.-% of saturated C₁₅

0.02 wt.-% of monounsaturated C₁₅

0.20 wt.-% of bisunsaturated C₁₅

1.25 wt.-% of saturated C₁₇

19.10 wt.-% of monounsaturated C₁₇

30 28.5 wt.-% bisunsaturated C₁₇

0.60 wt.-% trisunsaturated C₁₇

0.05 wt.-% saturated C₁₉

1.25 wt.-% monounsaturated C₁₉

4.3 wt.-% others

Component b:

33 wt.-% N-oleoylsarcosine

5 Composition 5:

Component a:

75 wt.-% of a mixture of 1 weight part of distilled coconut fatty acid and
1 weight part of tall oil fatty acid with a hydrocarbon chain distribution
as follows:

10 3.25 wt.-% of saturated C₇ hydrocarbon chain

2.85 wt.-% of saturated C₉

25.65 wt.-% of saturated C₁₁

8.60 wt.-% of saturated C₁₃

4.45 wt.-% of saturated C₁₅

15 0.02 wt.-% of monounsaturated C₁₅

0.20 wt.-% of bisunsaturated C₁₅

1.25 wt.-% of saturated C₁₇

19.10 wt.-% of monounsaturated C₁₇

28.5 wt.-% bisunsaturated C₁₇

20 0.60 wt.-% trisunsaturated C₁₇

0.05 wt.-% saturated C₁₉

1.25 wt.-% monounsaturated C₁₉

4.3 wt.-% others

Component b:

25 25 wt.-% N-oleoylsarcosine

Composition 6:

Component a:

30 84 wt.-% of a mixture of 1 weight part of distilled coconut fatty acid and
1 weight part of tall oil fatty acid with a hydrocarbon chain distribution
as follows:

3.25 wt.-% of saturated C₇ hydrocarbon chain

2.85 wt.-% of saturated C₉

- 25.65 wt.-% of saturated C₁₁
 8.60 wt.-% of saturated C₁₃
 4.45 wt.-% of saturated C₁₅
 0.02 wt.-% of monounsaturated C₁₅
 5 0.20 wt.-% of bisunsaturated C₁₅
 1.25 wt.-% of saturated C₁₇
 19.10 wt.-% of monounsaturated C₁₇
 28.5 wt.-% bisunsaturated C₁₇
 0.60 wt.-% trisunsaturated C₁₇
 10 0.05 wt.-% saturated C₁₉
 1.25 wt.-% monounsaturated C₁₉
 4.3 wt.-% others
 Component b:
 16 wt.-% N-oleoylsarcosine
 15

3. Comparative collector compositions

Composition 7:

80 wt.-% of tall oil fatty acid with a hydrocarbon chain distribution as follows:

- 20 0 wt.-% of saturated C₇ hydrocarbon chain
 0 wt.-% of saturated C₉
 0 wt.-% of saturated C₁₁
 0 wt.-% of saturated C₁₃
 0.30 wt.-% of saturated C₁₅
 25 0.05 wt.-% of monounsaturated C₁₅
 0.40 wt.-% of bisunsaturated C₁₅
 1.10 wt.-% of saturated C₁₇
 30.30 wt.-% of monounsaturated C₁₇
 55.80 wt.-% bisunsaturated C₁₇
 30 1.20 wt.-% trisunsaturated C₁₇
 0 wt.-% saturated C₁₉
 2.50 wt.-% monounsaturated C₁₉
 8.50 wt.-% others

and

20 wt.-% N-oleoylsarcosine

Composition 8:

- 5 80 wt.-% of distilled coconut fatty acid with a hydrocarbon chain distribution as follows:
- 6.50 wt.-% of saturated C₇ hydrocarbon chain
 - 5.70 wt.-% of saturated C₉
 - 51.30 wt.-% of saturated C₁₁
 - 10 17.20 wt.-% of saturated C₁₃
 - 8.60 wt.-% of saturated C₁₅
 - 0 wt.-% of monounsaturated C₁₅
 - 0 wt.-% of bisunsaturated C₁₅
 - 1.40 wt.-% of saturated C₁₇
 - 15 7.90 wt.-% of monounsaturated C₁₇
 - 1.20 wt.-% bisunsaturated C₁₇
 - 0 wt.-% trisunsaturated C₁₇
 - 0.10 wt.-% saturated C₁₉
 - 0 wt.-% monounsaturated C₁₉
 - 20 0.10 wt.-% others

and

20 wt.-% N-oleoylsarcosine

Composition 9:

- 25 100 wt.-% of tall oil fatty acid with a hydrocarbon chain distribution as follows:
- 0 wt.-% of saturated C₇ hydrocarbon chain
 - 0 wt.-% of saturated C₉
 - 0 wt.-% of saturated C₁₁
 - 0 wt.-% of saturated C₁₃
 - 30 0.30 wt.-% of saturated C₁₅
 - 0.05 wt.-% of monounsaturated C₁₅
 - 0.40 wt.-% of bisunsaturated C₁₅
 - 1.10 wt.-% of saturated C₁₇

- 30.30 wt.-% of monounsaturated C₁₇
 55.80 wt.-% bisunsaturated C₁₇
 1.20 wt.-% trisunsaturated C₁₇
 0 wt.-% saturated C₁₉
 5 2.50 wt.-% monounsaturated C₁₉
 8.50 wt.-% others

Composition 10:

- 100 wt.-% of a mixture of 1 weight part of distilled coconut fatty acid and
 10 1 weight part of tall oil fatty acid with a hydrocarbon chain distribution
 as follows:
- 3.25 wt.-% of saturated C₇ hydrocarbon chain
 2.85 wt.-% of saturated C₉
 25.65 wt.-% of saturated C₁₁
 15 8.60 wt.-% of saturated C₁₃
 4.45 wt.-% of saturated C₁₅
 0.02 wt.-% of monounsaturated C₁₅
 0.20 wt.-% of bisunsaturated C₁₅
 1.25 wt.-% of saturated C₁₇
 20 19.10 wt.-% of monounsaturated C₁₇
 28.5 wt.-% bisunsaturated C₁₇
 0.60 wt.-% trisunsaturated C₁₇
 0.05 wt.-% saturated C₁₉
 1.25 wt.-% monounsaturated C₁₉
 25 4.3 wt.-% others

4. Flotation test results

- 30 Example I: Apatite ore containing 16.1 % P₂O₅, 47.9 % SiO₂, 21.4 % CaO, and
 0.7 % MgO.

A 390 g portion of the ore sample was ground in a laboratory stainless steel mill for 5 minutes at 50 rpm and 66 % solids. This resulted in the following particle size

distribution for the flotation feed: $P_{50} = 17 \mu\text{m}$ and $P_{80} = 47 \mu\text{m}$. On completion of the grinding stage, the milled slurry was transferred to a 2.5 L capacity flotation cell, where the percentage solid was adjusted to approximately 15 % by addition of the appropriate amount of water. The flotation device was a Denver D-12 flotation machine and the impeller speed set to 1100 rpm. The flotation pulp was thereafter conditioned for 4 minutes and 3 minutes with the depressants sodium silicate (Na_2SiO_3 , 550 g/t) and sodium carbonate (Na_2CO_3 , 280 g/t) respectively – in the indicated order. Next the collector mixture, which was freshly prepared as a 1 % solution prior to starting each flotation test, was added and conditioned with the flotation pulp for 3 min. Finally, the air flow rate was set to 2 L/min and the resulting froth collected for 12 minutes.

Collector composition	Coconut oil fatty acid (wt.-%)	Tall oil fatty acid (wt.-%)	N-Oleoyl sarcosine (wt.-%)	Dosage (g/ton)	Grade P_2O_5 (wt.-%)	Recovery P_2O_5 (wt.-%)
1	40	40	20	500	27.86	55.74
2	20	60	20	500	25.47	69.57
3	50	30	20	500	27.77	58.35
7 (C)	-	80	20	500	26.56	54.37
8 (C)	80	-	20	500	30.82	42.91

The results from the flotation tests show that the collector compositions according to this invention (1 to 3) show excellent flotation efficiency and in particular, notably improved mineral recovery in comparison with the reference compositions 7 and 8.

The P_2O_5 grade obtained with the inventive compositions is slightly lower than what was obtained especially with the reference 8 in the laboratory experiments. This difference in grade is considered negligible because industrial flotation plants typically put the rougher concentrate through two, three or even four cleaning steps. In this way, the grade of the final concentrate is typically increased.

Example II: Ilmenite ore containing on approximately 32 % TiO₂

Approximately 1.2 L of sample was collected from the flotation feed stream of an ilmenite flotation plant. The 1.2 L sample, which consisted of approximately 1785 g dry ore and 750 g water, was thereafter transferred to a 3.2 L capacity flotation cell. The collector was thereafter added as-is and conditioned for 10 minutes using a Denver D-12 flotation device with the impeller speed set at 1550 rpm. The percentage solids in the slurry was thereafter reduced from 71 % to 51 % by addition of 1.0 L of industrial water. Hereafter, the air flow rate was set to 8.5 L/min and resulting froth collected for 270 seconds. In the case of the ore in question, a fatty acid and paraffin was used as collector combination. The results are shown below.

Collector composition	Coconut oil fatty acid (wt.-%)	Tall oil fatty acid (wt.-%)	N-Oleoyl sarcosine (wt.-%)	Collector dosage (g/ton)	Paraffin dosage (g/ton)	Grade TiO ₂ (wt.-%)	Recovery TiO ₂ (wt.-%)
1	40	40	20	840	360	37.4	81.1
9 (C)	-	100	-	840	360	36.8	76.0

The flotation results show that a 1:1 replacement of the fatty acid collector resulted in a 5.1 % increase in recovery in combination with a marginal increase in concentrate grade.

Example III: scheelite ore containing 0.20 % WO₃

The ground ore was conditioned with the depressants tannin (25 g/t), sodium silicate (350 g/t) and sodium carbonate (1000 g/t) after which the slurry pH was adjusted to pH 10 by adding the required amount of NaOH solution. The collector was then added as-is and conditioned with the flotation slurry for 2 minutes followed by addition of Clariant frother Flotanol 7026 and conditioning for a further

1 minute. Hereafter sufficient water was added to decrease the percentage solids in the flotation cell from 60% during the conditioning step to 35% in the flotation step. The air flow rate was now set to 5 L/min and the resulting froth collected for 2 minutes.

5

Collector composition	Coconut oil fatty acid (wt.-%)	Tall oil fatty acid (wt.-%)	N-Oleoyl sarcosine (wt.-%)	Dosage (g/ton)	Grade W_2O_3 (wt.-%)	Recovery W_2O_3 (wt.-%)
4	33.5	33.5	33	145	1.16	78.3
5	37.5	37.5	25	194	0.81	84.2
6	42	42	16	151	1.61	78.0
10 (C)	50	50	-	195	1.42	72.7
9 (C)	-	100	-	184	0.87	71.3
9 (C)	-	100	-	369	0.72	78.2

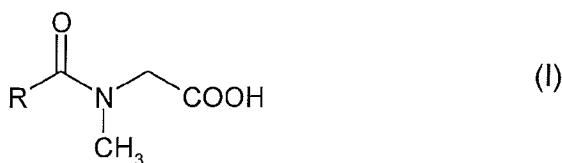
The use of the new collector mixture resulted in a significant increase in WO_3 grade as compared to the comparative product (100 % tall oil fatty acid collector). In addition, a similar WO_3 recovery value was obtained with 151 g/t dosage of the new collector blend as compared with 369 g/t dosage of the comparative tall oil fatty acid collector.

10

Patent claims

1. A collector composition for the direct froth flotation of nonsulfide ores comprising

- 5 a) 50 - 99 wt.-% of a mixture of fatty acids and
 b) 1 - 50 wt.-% of an N-acyl derivative of sarcosine of the formula (I)



wherein

R is a saturated or unsaturated hydrocarbon chain with 7 to 21 carbon atoms, wherein the mixture of comprises 10.0 - 35.0 wt.-% of fatty acid having a saturated C₁₁ hydrocarbon group, 2.5 - 15.0 wt.-% of fatty acid having a saturated C₁₃ hydrocarbon group, 10.0 - 25.0 wt.-% fatty acid having a monounsaturated C₁₇ hydrocarbon group and 20.0 - 45.0 wt.-% fatty acid having a bisunsaturated C₁₇ hydrocarbon group.

20 2. Composition according to claim 1, wherein the fatty acid mixture comprises fatty acids having

- 1.0 - 6.5 wt.-% of saturated C₇ hydrocarbon chain
 1.0 - 4.0 wt.-% of saturated C₉
 10.0 - 35.0 wt.-% of saturated C₁₁
 25 2.5 - 15.0 wt.-% of saturated C₁₃
 1.0 - 7.0 wt.-% of saturated C₁₅
 0.0 - 1.0 wt.-% of monounsaturated C₁₅
 0.0 - 1.0 wt.-% of bisunsaturated C₁₅
 0.5 - 2.0 wt.-% of saturated C₁₇
 30 10.0 - 25.0 wt.-% of monounsaturated C₁₇
 20.0 - 45.0 wt.-% bisunsaturated C₁₇
 0.0 - 2.0 wt.-% trisunsaturated C₁₇
 0.0 - 1.0 wt.-% saturated C₁₉

0.0 - 4.0 wt.-% monounsaturated C₁₉

0.0 - 7.0 wt.-% other fatty acids.

3. The composition of claim 1 and/or 2, wherein the component a) is a mixture
5 of 0.3 to 1.7 weight parts of coconut oil fatty acid to 1 weight part of tall oil fatty
acid.

4. The composition of one or more of claims 1 to 3, wherein the component b)
is N-oleoylsarcosine.

10

5. The composition as claimed in one or more of claims 1 to 4, wherein the
component a) is 80 wt.-% of a mixture of 1 weight part of coconut oil fatty acid and
1 weight part of tall oil fatty acid and component b) is 20 wt.-% of
N-oleoylsarcosine.

15

6. A process for the direct froth flotation of nonsulfide minerals, the process
comprising the steps of bringing the collector composition according to one or
more of claims 1 to 5 in contact with an aqueous suspension of the nonsulfide
mineral and frothing the so formed mineral pulp.

20

7. The process as claimed in claim 6, wherein the nonsulfide minerals is
apatite.

8. The process as claimed in claim 6, wherein the nonsulfide minerals are
25 selected from the group of calcite, scheelite, fluorspar, magnesite and barite.

9. The process as claimed in claim 6, wherein the nonsulfide mineral is
ilmenite.

30 10. A direct froth flotation process according to claims 6 to 9, wherein the
amount of collector composition added is an amount between 100 g and 1000 g
per ton of ore.

11. Use of a composition according to one or more of claims 1 to 5 as collector for the direct flotation of non-sulfide ores in an amount between 100 and 1000 g per ton of ore.

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2016/054291

A. CLASSIFICATION OF SUBJECT MATTER
 INV. B03D1/008 B03D1/01
 ADD.
 According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED
 Minimum documentation searched (classification system followed by classification symbols)
 B03D
 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
 EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 2 708 282 A1 (CLARIANT INT LTD [CH]; CLARIANT S A [BR]) 19 March 2014 (2014-03-19) claims 1-7; examples 1,2 -----	1-11
A	US 5 147 528 A (BULATOVIC SRDJAN [CA]) 15 September 1992 (1992-09-15) cited in the application examples 7,9,11 -----	1-11
A	DE 11 46 824 B (KLOECKNER HUMBOLDT DEUTZ AG) 11 April 1963 (1963-04-11) columns 4-6; figures 7-8 -----	1-11

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the international search 28 April 2016	Date of mailing of the international search report 09/05/2016
--	--

Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Roider, Josef
--	---

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2016/054291

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 2708282	A1	19-03-2014	
		AU 2013314744 A1	26-02-2015
		CA 2885467 A1	20-03-2014
		CN 104755173 A	01-07-2015
		EP 2708282 A1	19-03-2014
		EP 2895272 A1	22-07-2015
		MA 20150297 A1	31-08-2015
		PE 06592015 A1	06-05-2015
		US 2015238976 A1	27-08-2015
		WO 2014040686 A1	20-03-2014

US 5147528	A	15-09-1992	NONE

DE 1146824	B	11-04-1963	
		DE 1146824 B	11-04-1963
		FR 1256702 A	24-03-1961
