

Process for Producing 2-Hydroxy-4-Methylthiobutanoic Acid

BACKGROUND OF THE INVENTION

Field of the Invention

5 [0001]

The present application is filed, claiming the Paris Convention priorities of Japanese Patent Application No. 2008-263736 (filed on October 10, 2008), the entire content of which is incorporated herein by reference.

10 The present invention relates to a process for producing 2-hydroxy-4-methylthiobutanoic acid from 2-hydroxy-4-methylthiobutanenitrile via 2-hydroxy-4-methylthiobutanamide. In particular, the invention relates to a method of using an alloy material for an apparatus for
15 use in hydration of 2-hydroxy-4-methylthiobutanenitrile to obtain 2-hydroxy-4-methylthiobutanamide.

Description of the Related Art

[0002]

20 2-Hydroxy-4-methylthiobutanoic acid useful as an additive to a feed is known to be obtained by a method which comprises the steps of hydrating 2-hydroxy-4-methylthiobutanenitrile in the presence of sulfuric acid to obtain 2-hydroxy-4-methylthiobutanamide, and hydrolyzing 2-
25 hydroxy-4-methylthiobutanamide to obtain 2-hydroxy-4-

methylthiobutanoic acid (cf. JP-A-2007-238555).

[0003]

The use of a conventional SUS 304, SUS 316 or the like
as a material for an apparatus for use in this reaction has
5 difficulties, in view of the conditions such as a
concentration of sulfuric acid and temperature. Thus, a
glass-lined or resin-lined apparatus is used.

However, a glass-lined or resin-lined apparatus with a
complicated structure is hard to manufacture, and also has
10 problems such as its low impact resistance.

SUMMARY OF THE INVENTION

[0004]

An object of the present invention is to provide a
15 highly impact-resistant and corrosion-resistant material
which can be readily fabricated for an apparatus with a
complicated structure, as a material for an apparatus for
use in hydration of 2-hydroxy-4-methylthiobutanenitrile for
obtaining 2-hydroxy-4-methylthiobutanamide, in the process
20 for producing 2-hydroxy-4-methylthiobutanoic acid from 2-
hydroxy-4-methylthiobutanenitrile via 2-hydroxy-4-
methylthiobutanamide.

[0005]

The present invention provides a process for producing
25 2-hydroxy-4-methylthiobutanamide , comprising carrying out

the step of

hydrating 2-hydroxy-4-methylthiobutanenitrile in the presence of sulfuric acid at a temperature of from 40 to 70°C with an apparatus composed of, as a material, an alloy which contains 16.0 to 22.0% by weight of a Cr element, 16.0 to 22.0% by weight of a Mo element, 1.0 to 2.5% by weight of a Ta element and a Ni element as the rest, or an alloy which contains 26.0 to 32.0% by weight of a Mo element and a Ni element as the rest.

10 [0006]

According to the present invention, there is provided a highly impact-resistant and corrosion-resistant material which can be readily fabricated for an apparatus with a complicated structure, as a material for an apparatus for use in hydration of 2-hydroxy-4-methylthiobutanenitrile for obtaining 2-hydroxy-4-methylthiobutanamide, in the process for producing 2-hydroxy-4-methylthiobutanoic acid from 2-hydroxy-4-methylthiobutanenitrile via 2-hydroxy-4-methylthiobutanamide. Thus, 2-hydroxy-4-methylthiobutanamide, and 2-hydroxy-4-methylthiobutanoic acid can be reliably produced without any corrosion or damage of the apparatus.

Detailed Description of the Invention

25 [0007]

2-Hydroxy-4-methylthiobutanoic acid (hereinafter optionally referred to as HMTBA) is produced by hydrating 2-hydroxy-4-methylthiobutanenitrile (hereinafter optionally referred to as HMTBN) in the presence of sulfuric acid
5 usually at a temperature of from about 40 to about 70°C to obtain 2-hydroxy-4-methylthiobutanamide (hereinafter optionally referred to as HMTBAA), and typically adding water to 2-hydroxy-4-methylthiobutanamide to hydrolyze the same usually at a temperature of from about 90 to about
10 130°C.

For example, HMTBN is industrially produced by reacting acrolein with methyl mercaptan to obtain 3-methylthiopropionaldehyde, and reacting 3-methylthiopropionaldehyde with hydrogen cyanide.

15 [0008]

Generally, HMTBAA is produced by feeding HMTBN, water and sulfuric acid to a hydration tank to hydrate HMTBN.

Fed to the hydration tank are usually about 20 to about 70 parts by weight, preferably about 25 to about 50
20 parts by weight of water, per 100 parts by weight of HMTBN, and about 0.5 to about 1 mol in total, preferably about 0.6 to about 0.8 mol in total, of sulfuric acid, per 1 mol of HMTBN.

[0009]

25 It is also possible to feed water as a pre-mixture

thereof with HMTBN and/or sulfuric acid, in other words, as an aqueous HMTBN solution and/or an aqueous sulfuric acid solution, to the hydration tank. Preferably, water is fed as an aqueous sulfuric acid solution to the hydration tank.

5 The hydration reaction is carried out usually at a temperature of about 40 to about 70°C for about 1 to about 3 hours. After the reaction, the reaction solution is aged. In the reaction solution, HMTBA is produced as a result of hydrolysis of a part of HMTBAA.

10 [0010]

The apparatus for use in the hydration refers to a hydration tank, a pipe combined thereto so as to contact a hydration reaction mass, a heat exchanger or the like.

15 In the present invention, as the material for the apparatus for use in the hydration, there is used an alloy which contains 16.0 to 22.0% by weight of a Cr element, 16.0 to 22.0% by weight of a Mo element, 1.0 to 2.5% by weight of a Ta element and a Ni element as the rest, or an alloy which contains 26.0 to 32.0% by weight of a Mo
20 element and a Ni element as the rest.

A commercially available alloy that contains 16.0 to 22.0% by weight of a Cr element, 16.0 to 22.0% by weight of a Mo element, 1.0 to 2.5% by weight of a Ta element and a Ni element as the rest is for example, MAT® 21 (Cr: 19% by
25 weight, Mo: 19% by weight, Ta: 1.8% by weight, and Ni: the

rest). A commercially available alloy that contains 26.0 to 32.0% by weight of a Mo element, and a Ni element as the rest is, for example, Hastelloy® B-2 (Mo: 28% by weight, and Ni: the rest).

5 [0011]

Then, the reaction solution containing HMTBAA as a main component, obtained in the hydration tank, is fed to a hydrolysis tank, and water is added to the reaction solution to thereby hydrolyze HMTBAA to produce HMTBA.

10 About 100 to about 200 parts by weight of water is usually fed per 100 parts by weight of the aqueous sulfuric acid solution in the above-described reaction solution.

In the hydrolysis tank, HMTBAA is hydrolyzed with water and sulfuric acid to produce HMTBA and also to by-
15 produce ammonium bisulfate (NH_4HSO_4) and ammonium sulfate ($(\text{NH}_4)_2\text{SO}_4$). After the addition of water and heating, the hydrolysis reaction is carried out usually at a temperature of from about 90 to about 130°C for about 2 to about 6 hours.

20 In this regard, any of the above-described alloys may be used also as a material for an apparatus for the hydrolysis.

[0012]

The reaction solution containing HMTBA, obtained in
25 the hydrolysis tank, is usually distilled to remove low

boiling point components from the reaction solution.

The distillation is carried out usually at a temperature of from about 80 to about 120°C under a pressure of from about 50 to about 150 kPa to remove by-produced low boiling point components such as dimethyl sulfide, dimethyl disulfide and formic acid. The low boiling point components are removed as a distillate usually at a rate of about 1 to about 4% by weight relative to the reaction solution, as required.

The removal of the low boiling point components may be done after neutralization or phase separation as will be described later.

[0013]

Then, an alkali is added to the reaction solution containing HMTBA, from which the low boiling point components have been removed, to thereby neutralize the reaction solution and to phase-separate the reaction solution into an organic phase containing HMTBA, and an aqueous phase containing water and an inorganic salt (containing ammonium bisulfate and ammonium sulfate). The neutralization and the phase separation are conducted, using a mixer-settler type liquid-liquid extractor in which a stirrer tank and a phase-separation tank are combined as one set.

[0014]

As the alkali, for example, sodium hydroxide, sodium hydrogencarbonate, sodium carbonate or the like is used in the form of an aqueous solution. The alkali is fed usually at a rate of from about 0.5 to about 1.2 mol, preferably
5 from about 0.6 to about 0.8 mol, per 1 mol of ammonium bisulfate in the above-described reaction solution. The addition rate of the alkali may be controlled by a hydrogen-ion concentration (pH) of the reaction solution admixed with the alkali.

10 The neutralization reaction is carried out usually at a temperature of from about 15 to about 120°C, preferably from about 30 to about 110°C, for about 0.1 to 3 hours, preferably about 0.1 to about 2 hours.

[0015]

15 After the neutralization, the reaction solution is left to stand still in a phase-separation tank so as to phase-separate the reaction solution into an organic phase as an upper layer and an aqueous phase as a lower layer (liquid separation). The temperature for this separation
20 is usually from about 30 to about 110°C.

[0016]

The organic phase separated from the aqueous phase contains usually about 40 to about 60% by weight of HMTBA, about 20 to about 30% by weight of water and about 10 to
25 about 30% by weight of an inorganic salt.

This organic phase is concentrated to remove the remaining water. For example, the concentration is done in a concentration tank usually at a temperature of from about 60 to about 150°C under a pressure of from about 1 to about 20 kPa, so that water in the organic phase is reduced to about 5% by weight or less, preferably about 2% by weight or less, more preferably about 1% by weight or less. By this concentration, the sulfate-ion concentration and kinematic viscosity of a product as will be described later can be decreased.

The inorganic salt is precipitated from the organic phase by this concentration, and the resulting organic phase forms a slurry.

[0017]

When the residence time of the organic phase in the concentration tank is set to about 0.5 hour or longer, HMTBA is converted into an oligomer (which is mainly a dimer, containing small amounts of a trimer and a tetramer). By doing so, the solubility of the inorganic salt in the organic phase is lowered, so that the concentration of the inorganic salt in the organic phase can be lowered. Preferably, the temperature, pressure and residence time are selected so that the weight ratio of the monomer of HMTBA to the oligomer of HMTBA can be about 2 to about 4. Such a concentration is effective to increase the particle

size of the inorganic salt precipitated in the organic phase, so that a solid-liquid separation efficiency is improved during a solid-liquid separation as will be described later. Thus, the removal of the inorganic salt is facilitated.

[0018]

Then, the resultant slurry of the organic phase is cooled to usually about 60 to about 120°C, using a heat exchanger or the like. After that, the slurry is separated into a liquid component containing the organic phase and a solid component (or a residue) containing the precipitated inorganic salt.

Generally, the solid-liquid separation is carried out, using a solid-liquid separator such as a decanter type centrifugal separator.

The separated liquid component comprises HMTBA (containing the oligomer produced during the concentration) as a main component. If needed, water is added to the liquid component to provide a product of HMTBA which contains usually about 88 to about 90% by weight of HMTBA, about 10 to about 12.5% by weight of a moisture content and very small amounts of other components.

[0019]

The separated solid component contains usually about 20 to about 60% by weight of HMTBA, and thus is usually

admixed with water to dissolve the inorganic salt, so as to phase-separate the solid component into an aqueous phase containing the inorganic salt and an organic phase. This organic phase is mixed into the former organic phase
5 obtained by the above-described neutralization and phase-separation, for recovery.

Examples

[0020]

10 A hydration reaction mass from 2-hydroxy-4-methylthiobutanenitrile to 2-hydroxy-4-methylthiobutanamide was subjected to a corrosion test for the following metallic material: the hydration reaction mass (HMTBAA: 46% by weight, H_2SO_4 : 18% by weight, HMTBA: 15% by weight,
15 NH_4HSO_4 : 12% by weight, and water: 9% by weight) and a test piece of the following metallic material (a flat plate of 25 mm in length X 20 mm in width X 2 mm in thickness) were put in a glass-made test bottle with a condenser; the gas phase zone was purged with a nitrogen gas; and the test
20 bottle was heated to 60°C and was maintained at the same temperature for 20 hours. After that, the test piece was removed, rinsed and dried. Then, the reduced weight of the test piece was measured to determine a corrosion rate. The results are shown in Table 1.

25 [0021]

(Test Piece)

Test piece 1: MAT® 21 (Cr: 18.4% by weight, Mo: 18.5% by weight, Ta: 1.9% by weight, and Ni: the rest)

5 Test piece 2: Hastelloy® B-2 (Mo: 27.7% by weight, and Ni: the rest)

Test piece 3: Ti-Pd (Pd: 0.17% by weight, and Ti: the rest)

[0022]

[Table 1]

No.	Test piece	Corrosion rate (mm/year)	Corroded state
1	Test piece 1	0.004	Not corroded and not damaged
2	Test piece 2	0.009	Not corroded and not damaged
3	Test piece 3 Ti-Pd	0.11	Active corroded surface, and entirely corroded

10 [0023]

It was confirmed that Test piece 1 or Test piece 2 did not corrode due to the hydration reaction mass. An apparatus with a complicated structure can be readily constructed, using any of these alloys, and 2-hydroxy-4-methylthiobutanamide, and 2-hydroxy-4-methylthiobutanoic acid can be reliably produced without any corrosion or damage of the apparatus.

15

WHAT IS CLAIMED IS:

1. A process for producing 2-hydroxy-4-methylthiobutanamide, comprising

hydrating 2-hydroxy-4-methylthiobutanenitrile in the
5 presence of sulfuric acid at a temperature of from 40 to
70°C with an apparatus composed of, as a material, an alloy
which contains 16.0 to 22.0% by weight of a Cr element,
16.0 to 22.0% by weight of a Mo element, 1.0 to 2.5% by
weight of a Ta element and a Ni element as the rest, or an
10 alloy which contains 26.0 to 32.0% by weight of a Mo
element and a Ni element as the rest.

2. A process according to Claim 1, which further
comprises adding water to the resulting 2-hydroxy-4-
15 methylthiobutanamide and hydrolyzing the same at a
temperature of from 90 to 130°C to produce 2-hydroxy-4-
methylthiobutanoic acid.

3. The process of Claim 1, wherein the alloy is an alloy
20 which contains 16.0 to 22.0% by weight of a Cr element,
16.0 to 22.0% by weight of a Mo element, 1.0 to 2.5% by
weight of a Ta element and a Ni element as the rest.

4. The process of Claim 1, wherein the alloy is an alloy
25 which contains 26.0 to 32.0% by weight of a Mo element, and

a Ni element as the rest.

5. Use of an alloy as a material of an apparatus for use in the process for producing 2-hydroxy-4-methylthiobutanamide by hydrating 2-hydroxy-4-methylthiobutanenitrile in the presence of sulfuric acid at a temperature of from 40 to 70°C.