

Method for Reclaiming Metal Sulfate-Containing Waste
Sulfuric Acid

✓
19053
copy.

BACKGROUND OF THE INVENTION

The present invention relates to a method for reclaiming waste sulfuric acid discharged from, for instance, a titanium oxide-production process.

The so-called sulfuric acid method has been well-known as a method for preparing titanium oxide (titania) widely used as a coloring material. A large amount of waste sulfuric acid is discharged mainly from the solid/liquid separation step for titanium hydroxide during the titanium oxide-production process and the waste sulfuric acid contains, for instance, Fe, Ti, Cr and V originated from ilmenite and titanium slug as raw materials for titanium.

In the past, the waste sulfuric acid was neutralized and then buried under the ground or dumped at sea in the form of gypsum. However, there has been required for the development of other techniques for treating the waste sulfuric acid from the viewpoint of various problems, for instance, those concerning environmental pollution, insurance of the filled-up land for burying such gypsum and enormous expenses necessary for the post-treatment of the waste sulfuric acid.

There have been a variety of proposals in response to such request. For instance, Japanese Un-examined Patent Publication (hereunder referred to as "J.P. KOKAI") Nos. Hei 3-80103 and Hei 3-88718 each discloses a method for removing impurities such as

This Sheet Has Been Replaced



REPLACED SHEETS		SG
☒	Description : paged to paged	14
☐	Claim(s) : paged to paged	
☒	Drawing(s) : paged 1/1 to paged	
☐	Abstract : paged to paged	
DATE OF RECEIPT		

Fe ions, which makes use of a solvent-extraction process. In these methods, however, the strongly acidic sulfuric acid solution per se is directly subjected to a solvent-extraction treatment. Therefore, Fe ions are, in fact, extracted in poor efficiency and the iron-containing component thus recovered also has low purity, in spite of the assertions of the inventors of the foregoing patents. Moreover, in these methods, trivalent iron ions are directly extracted and accordingly, the use of an acidic extraction agent would require a multistage extraction treatment because of the low extraction rate thereof.

Under such circumstances, the inventors of this invention have developed a method for effectively recovering a highly concentrated sulfuric acid solution and filed a patent application, EP-A1-0541002, in which the aforementioned solvent-extraction method is improved so as to enhance the bivalent iron ion-removing efficiency. However, the sulfuric acid solution thus recovered is colored dark green because of the presence of residual Cr and V which are converted into colored oxides and it is difficult to reuse the colored sulfuric acid solution in the process for preparing titanium oxide used, in particular, as a white pigment.

Moreover, there has also been proposed a diffusive dialysis method, as a technique for treating the waste sulfuric acid, which makes use of an anion exchange membrane. However, if the waste sulfuric acid is treated through the diffusive dialysis, particles are deposited on the surface of the ion exchange membrane immediately after putting the dialyzer in operation and this leads to the quality-deterioration of the

This Sheet Has Been Replaced



membrane and the breakage of the membrane due to deposition of the particles within the membrane. For this reason, if the foregoing phenomenon occurs during the operation of the device over a long time, the deposits should be removed by washing the anionic exchange membrane with an agent capable of washing them away.

To eliminate the problem of poor efficiency, J.P. KOKAI No. Hei 2-153806 proposes a method for recovering sulfuric acid present in a titanium-containing waste sulfuric acid solution while setting the solution against water in a multistage diffusive dialysis tank equipped with an anion exchange membrane wherein the solution and water are countercurrently supplied to the dialysis tank. In other words, the former is passed, through the tank, from the front stage to the rear stage, while the latter is passed therethrough from the rear stage to the front stage.

This method permits the elimination of the step for washing the anion exchange membrane and a decrease in the degree of quality-deterioration of the membrane, but the membrane should be washed every 10 days as is described in Examples of the patent. Therefore, the method should further be improved in its efficiency.

SUMMARY OF THE INVENTION

It is an object of the present invention to solve the foregoing problems and to provide a method for efficiently and easily regenerating a sulfuric acid solution having a high concentration and a low impurity content from metal sulfate-

This Sheet Has Been Replaced



containing waste sulfuric acid and waste liquor formed during acid cleaning which are discharged from the process for preparing titanium (IV) oxide according to the sulfuric acid method.

According to the present invention, the foregoing object can be accomplished by removing titanium present in a metal sulfate-containing waste sulfuric acid solution through a solvent-extraction treatment and then subjecting the solution, from which titanium is extracted, to a diffusive dialysis treatment.

It is preferred to backwardly extract the titanium thus extracted by bringing the organic solvent obtained through the solvent-extraction and containing titanium into contact with an alkaline solution to thus convert the titanium into an insoluble compound through the backward extraction; to bring the organic solvent obtained after the removal of titanium into contact with the remaining solution obtained after the diffusive dialysis treatment for regenerating the organic solvent through the backward extraction of the alkali ions into the aqueous phase; and to reuse the regenerated organic solvent in the solvent-extraction process for reclaiming the waste sulfuric acid.

Moreover, it is desirable that the waste sulfuric acid solution be optionally filtered with cooling to control the concentration of free sulfuric acid present therein prior to the solvent-extraction of the solution.

The method for reclaiming metal sulfate-containing waste sulfuric acid according to the present invention basically

This Sheet Has Been Replaced



comprises the step of effectively removing principal impurities such as iron, titanium, chromium and vanadium to thus reclaim the solution into highly concentrated sulfuric acid which can widely be reused.

Among these impurities, titanium ions are included in, for instance, the waste sulfuric acid solution discharged from the titanium oxide-production process in an amount ranging from 2 to 10 g/l (these ions are dissolved therein mainly in the form of $TiO-SO_4$ and partially in the form of $Ti_2(SO_4)_3$). Easily hydrolyzable tetravalent titanium ions, among others, can form titanium oxide particles which may become a cause of quality-deterioration and breakage of an ion exchange membrane through deposition thereof on the surface of the ion exchange membrane and within the membrane during the diffusive dialysis treatment. For this reason, the removal of these ions is indispensable for industrially applying the diffusive dialysis method to the reclamation of the metal sulfate-containing waste sulfuric acid. The inventors of this invention have found out that the formation of titanium oxide particles on an ion exchange membrane can substantially be inhibited if the titanium content in the solution is, in advance, reduced to a level of not more than 0.5 g/l through solvent-extraction. The titanium components remaining even after the solvent-extraction are mainly composed of trivalent titanium ions which are less hydrolyzable as compared with tetravalent titanium ions. Therefore, the waste solution obtained after the solvent-extraction treatment may directly be subjected to a diffusive dialysis treatment without impairing the intended effect of the

This Sheet Has Been Replaced



method to thus give a sulfuric acid solution having a low content of impurities such as chromium and iron. In particular, it is desirable to remove solid contents present in the waste solution obtained after the solvent-extraction through micro filtration prior to the diffusive dialysis thereof. Moreover, if the organic solvent still remains in the waste solution, it is desirable to remove the remaining organic solvent by bringing the waste solution into contact with, for instance, activated carbon.

If the method of the present invention is used for reclaiming waste sulfuric acid, the method can be carried out according to the batchwise process in which a complete waste solution as a waste acid solution is treated or the so-called continuous recovery process which comprises the steps of continuously withdrawing a waste sulfuric acid solution from, for instance, a titanium oxide-production process, recovering the acid and simultaneously separating and removing metal salts separated from the solution, but the continuous recovery process is generally preferred since the organic solvent used in the solvent-extraction treatment can be regenerated by removing titanium components through backward extraction with an alkali solution.

The extraction of titanium ions is carried out using, for instance, an acidic organophosphoric acid compound, a carboxylic acid, a sulfonic acid or hydroxy oxime, as has been well-known in the art. Among the acidic organophosphoric acid compounds, acidic phosphoric acid diesters are particularly preferred.

Titanium ions extracted with, for instance, an acidic



organophosphoric acid compound are backwardly extracted with, for instance, an alkali solution such as an ammonium carbonate solution and recovered in the form of, for instance, hydroxide. The titanium compound can be fired to give titanium oxide after the recovery thereof. The product can be used as a raw material for titanium.

Ferrous sulfate can be crystallized and the sulfuric acid concentration of the waste solution can be controlled to a level favorable for the diffusive dialysis treatment by filtering the waste sulfuric acid solution with cooling prior to the solvent-extraction treatment thereof.

In the method of the present invention, titanium components present in the metal sulfate-containing waste sulfuric acid solution is removed through the solvent-extraction treatment prior to the diffusive dialysis treatment of the waste solution and thus the waste sulfuric acid solution can quite efficiently and easily be reclaimed to give a sulfuric acid solution having a low content of impurities such as iron, while using the diffusive dialysis method which conventionally has a difficulty in efficiency. Moreover, the diffusive dialysis method permits the removal of components such as chromium and vanadium which cannot be removed by the solvent-extraction method. The removal of such components is quite beneficial from the viewpoint of environmental sanitation.

In particular, the method of the present invention has high industrial value. This is because the method permits effective recovery of sulfuric acid from a metal sulfate-containing waste sulfuric acid solution in a closed system and

This Sheet Has Been Replaced



the reuse thereof and also permits the prevention of any environmental pollution by bringing an organic solvent containing extracted titanium and obtained through a solvent-extraction treatment into contact with an equal amount of an alkali solution to thus backwardly extract titanium and to convert it into an insoluble compound; bringing the organic solvent obtained after the titanium extraction into contact with the liquid remaining after the dialysis formed during a diffusive dialysis treatment to thus regenerate the organic solvent through backward extraction of alkali ions into the aqueous phase; and using the regenerated organic solvent in the solvent-extraction process for reclaiming the waste sulfuric acid solution. In addition, titanium, scandium or the like can be recovered as by-products and therefore, the method of the present invention is likewise quite beneficial from the viewpoint of effective use of resources.

If the waste sulfuric acid solution is filtered with cooling prior to the solvent-extraction, iron sulfate can easily be obtained as well.

BRIEF DESCRIPTION OF THE DRAWINGS

The only figure attached hereto is a graph showing the relation between the titanium concentration of the solution obtained by the solvent-extraction treatment and the recovery of free sulfuric acid or the rate of chromium removal observed during the diffusive dialysis treatment.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

This Sheet Has Been Replaced



Examples of the present invention will hereunder be described, but the present invention is not restricted to these specific Examples.

Example 1

An amount of 10 ℓ of a waste sulfuric acid solution discharged from a titanium oxide-production process carried out according to the sulfuric acid method which had an Fe content of 30g/ ℓ (as expressed in terms of the amount of bivalent and trivalent iron ions), a Ti content of 2 g/ ℓ (titanium ions), a Cr content of 0.1 g/ ℓ (chromium ions), a V content of 0.15 g/ ℓ (vanadium ions), an Sc content of 14mg/ ℓ (scandium ions) and a sulfate content of 310 g/ ℓ (sulfate ions) was brought into contact with 10 ℓ of 1 mol/ ℓ solution of di-2-ethylhexylphosphoric acid in kerosine over 20 minutes to thus extract titanium ions and scandium ions present in the aqueous phase into the organic phase. The solvent-extraction treatment gave 10 ℓ of a solvent-extracted solution which had an Fe content of 30g/ ℓ (iron ions), a Ti content of 0.01g/ ℓ (titanium ions), a Cr content of 0.1 g/ ℓ (chromium ions), a V content of 0.15 g/ ℓ (vanadium ions), an Sc content of not more than 1 mg/ ℓ (scandium ions) and a sulfate content of 310 g/ ℓ (sulfate ions). The rate of recovery and removal of titanium and scandium was found to be 99%.

Then the solution thus treated was supplied to a dialysis chamber of a diffusive dialysis device at a flow rate of 240 ml/hr at ordinary temperature, while water was supplied to a recovery chamber of the device separated from the dialysis



chamber by an anion exchange membrane at a flow rate of 150 ml/hr, the solution and water were circulated through each corresponding chamber over 20 hours. As a result, there were recovered 2.98 ℓ of a recovered solution which had an Fe content of 2.1 g/ ℓ (as expressed in terms of the amount of Fe ions), a Ti content of not more than 0.001g/ ℓ (Ti ions), a Cr content of 0.006 g/ ℓ (Cr ions), a V content of 0.014g/ ℓ (V ions) and a sulfate content of 305g/ ℓ (sulfate ions) and 4.6 ℓ of a solution remaining after the dialysis which had an Fe content of 29.8 g/ ℓ (Fe ions), a Ti content of 0.01g/ ℓ (Ti ions), a Cr content of 0.1 g/ ℓ (Cr ions), a V content of 0.147g/ ℓ (V ions) and a sulfate content of 125.8g/ ℓ (sulfate ions). The recovery of free sulfuric acid, the rate of chromium removal and the rate of vanadium removal were found to be 70%, 96% and 94%, respectively.

Moreover, 10 ℓ of the organic phase recovered through the foregoing solvent-extraction treatment was brought into contact with an aqueous phase comprising 0.2mol/ ℓ of ammonium carbonate aqueous solution and 2 mol/ ℓ of aqueous ammonia over 20 minutes, then precipitates formed in the organic phase were filtered off and dried to thus give 58.6 g of a mixture comprising titanium and scandium (having a Ti content of 32.4% and an Sc content of 0.18%). The recoveries of Ti and Sc were found to be 95% and 75%, respectively.

The foregoing organic solvent (12 ℓ) whose volume was changed through the backward extraction was brought into contact with the same volume of the solution remaining after the diffusive dialysis treatment over 20 minutes to backwardly



extract the alkali ions present in the organic phase into the aqueous phase and to thus give 10 ℓ of the organic solvent. The organic solvent thus recovered was again brought into contact with a waste sulfuric acid solution discharged during a titanium oxide-production process and as a result, it was found that the recoveries of titanium and scandium reached 99%.

Example 2

An amount of 10 ℓ of a waste sulfuric acid solution discharged from a titanium oxide-production process carried out according to the sulfuric acid method which had an Fe content of 36.5g/ ℓ (as expressed in terms of the amount of bivalent and trivalent iron ions), a Ti content of 3 g/ ℓ (titanium ions), a Cr content of 0.2 g/ ℓ (chromium ions), a V content of 0.6g/ ℓ (vanadium ions), an Sc content of 20mg/ ℓ (scandium ions) and a sulfate content of 370 g/ ℓ (sulfate ions) was cooled to 0 $^{\circ}\text{C}$ to give 960 g of $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ (having an Fe content of 18.8%, a Ti content of 0.04%, a Cr content of 0.002%, a V content of 0.002% and a sulfate ion content of 33.9%).

After crystallizing $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ through the cooling treatment, the waste solution was filtered and 9.5 ℓ of the resulting filtrate which had an Fe content of 19.4g/ ℓ (iron ions), a Ti content of 3.1 g/ ℓ (titanium ions), a Cr content of 0.2 g/ ℓ (chromium ions), a V content of 0.6g/ ℓ (vanadium ions), an Sc content of 20mg/ ℓ (scandium ions) and a sulfate content of 355 g/ ℓ (sulfate ions) was brought into contact with 9.5 ℓ of 1 mol/ ℓ solution of di-2-ethylhexylphosphoric acid in kerosine over 20 minutes to thus extract titanium ions



and scandium ions present in the aqueous phase into the organic phase. The solvent-extraction treatment gave 9.5ℓ of a solvent-extracted solution which had an Fe content of 19.4g/ℓ (iron ions), a Ti content of 0.01g/ℓ (titanium ions), a Cr content of 0.2 g/ℓ (chromium ions), a V content of 0.6g/ℓ (vanadium ions), an Sc content of not more than 1 mg/ℓ (scandium ions) and a sulfate content of 355 g/ℓ (sulfate ions). The rate of recovery or removal of titanium, scandium was found to be 99%.

Then the solution thus treated was supplied to a dialysis chamber of a diffusive dialysis device at a flow rate of 230 ml/hr at ordinary temperature, while water was supplied to a recovery chamber of the device separated from the dialysis chamber by an anion exchange membrane at a flow rate of 250 ml/hr to thus circulate the solution and water through each corresponding chamber over 20 hours. As a result, there were recovered 2.45ℓ of a recovered liquid which had an Fe content of 1.8 g/ℓ (as expressed in terms of the amount of Fe ions), a Ti content of 0.001g/ℓ (Ti ions), a Cr content of 0.01g/ℓ (Cr ions), a V content of 0.05 g/ℓ (V ions) and a sulfate content of 246g/ℓ (sulfate ions) and 2.37ℓ of a solution remaining after the dialysis which had an Fe content of 16.9 g/ℓ (Fe ions), a Ti content of 0.008 g/ℓ (Ti ions), a Cr content of 0.18g/ℓ (Cr ions), a V content of 0.53 g/ℓ (V ions) and a sulfate content of 90.2 g/ℓ (sulfate ions). The recovery of free sulfuric acid, the rate of chromium removal and the rate of vanadium removal were found to be 80%, 92% and 91%, respectively.

Moreover, 9.5 ℓ of the organic phase recovered through

This Sheet Has Been Replaced



the foregoing solvent-extraction treatment was brought into contact with an aqueous phase comprising 0.2mol/ℓ of ammonium carbonate aqueous solution and 2 mol/ℓ of aqueous ammonia over 20 minutes to backwardly extract titanium ions and scandium ions present in the organic phase into the aqueous phase in the form of hydroxides. After allowing the aqueous phase to stand for 20 minutes, the precipitates formed therein were filtered off and dried to thus give 85.8 g of a mixture comprising titanium and scandium (having a Ti content of 33.2% and an Sc content of 0.2%). The recoveries of Ti and Sc were found to be 95% and 85%, respectively.

The foregoing organic solvent (11.4 ℓ) whose volume was changed through the backward extraction was brought into contact with the same volume of the solution remaining after the diffusive dialysis treatment over 20 minutes to backwardly extract the alkali ions present in the organic phase into the aqueous phase and to thus give 9.5ℓ of the organic solvent. The organic solvent thus recovered was again brought into contact with a waste sulfuric acid solution discharged during a titanium oxide-production process and as a result, it was found that the recoveries of titanium and scandium reached 99%.

Example 3

A waste sulfuric acid solution discharged from a titanium oxide-production process performed according to the sulfuric acid method which had an Fe content of 30g/ℓ (as expressed in terms of the amount of bivalent and trivalent Fe ions), a Ti content of 2 g/ℓ (Ti ions), a Cr content of 0.2 g/ℓ (Cr ions)



and a sulfate content of 300 g/ℓ (sulfate ions) was continuously brought into contact with an organic solvent comprising a solution of di-2-ethylhexylphosphoric acid (D2EHPA) in kerosine, whose concentration was variously changed, under the same conditions used in Example 1, over 30 days. Table 1 given below shows titanium concentrations of the solutions obtained through the foregoing solvent-extraction treatment.

Table 1

<u>D2EHPA Conc. of the Organic Solv. Brought into Contact With the Waste Solution (mol/ℓ)</u>	<u>Ti Conc. of the Solution Treated (g/ℓ)</u>
2	0.01
1.5	0.06
1	0.1
0.5	0.3
0.1	1.4
untreated	2

Fig. 1 shows the results observed when the foregoing solution obtained after the solvent-extraction was subjected to the diffusive dialysis treatment. These results indicate that sulfuric acid can efficiently be recovered from waste sulfuric acid solutions by the diffusive dialysis treatment if titanium concentration of the waste solution is reduced to a level of not more than 0.5 g/ℓ. Moreover, it can also be confirmed that the diffusive dialysis treatment permits the effective removal of chromium.

This Sheet Has Been Replaced





What is claimed is:

1. A method for reclaiming a metal sulfate-containing waste sulfuric acid solution comprising the steps of subjecting the waste sulfuric acid solution to a solvent-extraction treatment to remove titanium present therein and then subjecting the solution obtained after the extraction to a diffusive dialysis treatment.

2. The method for reclaiming a waste sulfuric acid solution as set forth in claim 1 wherein an organic solvent used in the titanium-extraction through the solvent-extraction treatment is brought into contact with an alkali solution to backwardly extract the titanium and to convert it into an insoluble compound; the organic solvent obtained after the titanium-extraction is brought into contact with an aqueous solution remaining after the diffusive dialysis treatment to backwardly extract the alkali ions into the aqueous phase to regenerate the organic solvent; and the regenerated organic solvent is used in the solvent-extraction process for reclaiming the waste sulfuric acid solution.

3. The method for reclaiming a waste sulfuric acid solution as set forth in claim 1 or 2 wherein the waste sulfuric acid solution is cooled to precipitate iron sulfate, then the precipitates are filtered off to control the free sulfuric acid content in the waste solution to 15 to 35% by weight and thereafter the filtrate is subjected to a solvent-extraction treatment.

THIS SHEET HAS BEEN
REPLACED



ABSTRACT

For the purpose of efficiently and easily regenerating a sulfuric acid solution having a high concentration and a low impurity content from a metal sulfate-containing waste sulfuric acid solution and waste liquor formed during acid cleaning which are discharged from titanium (IV) oxide-production process carried out according to the sulfuric acid method, titanium present in the metal sulfate-containing waste sulfuric acid solution is removed through a solvent-extraction treatment and then the solution obtained after the solvent-extraction is subjected to a diffusive dialysis treatment.

THIS SHEET HAS BEEN
REPLACED

This Sheet Has Been Replaced

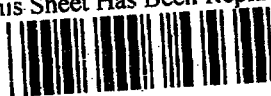
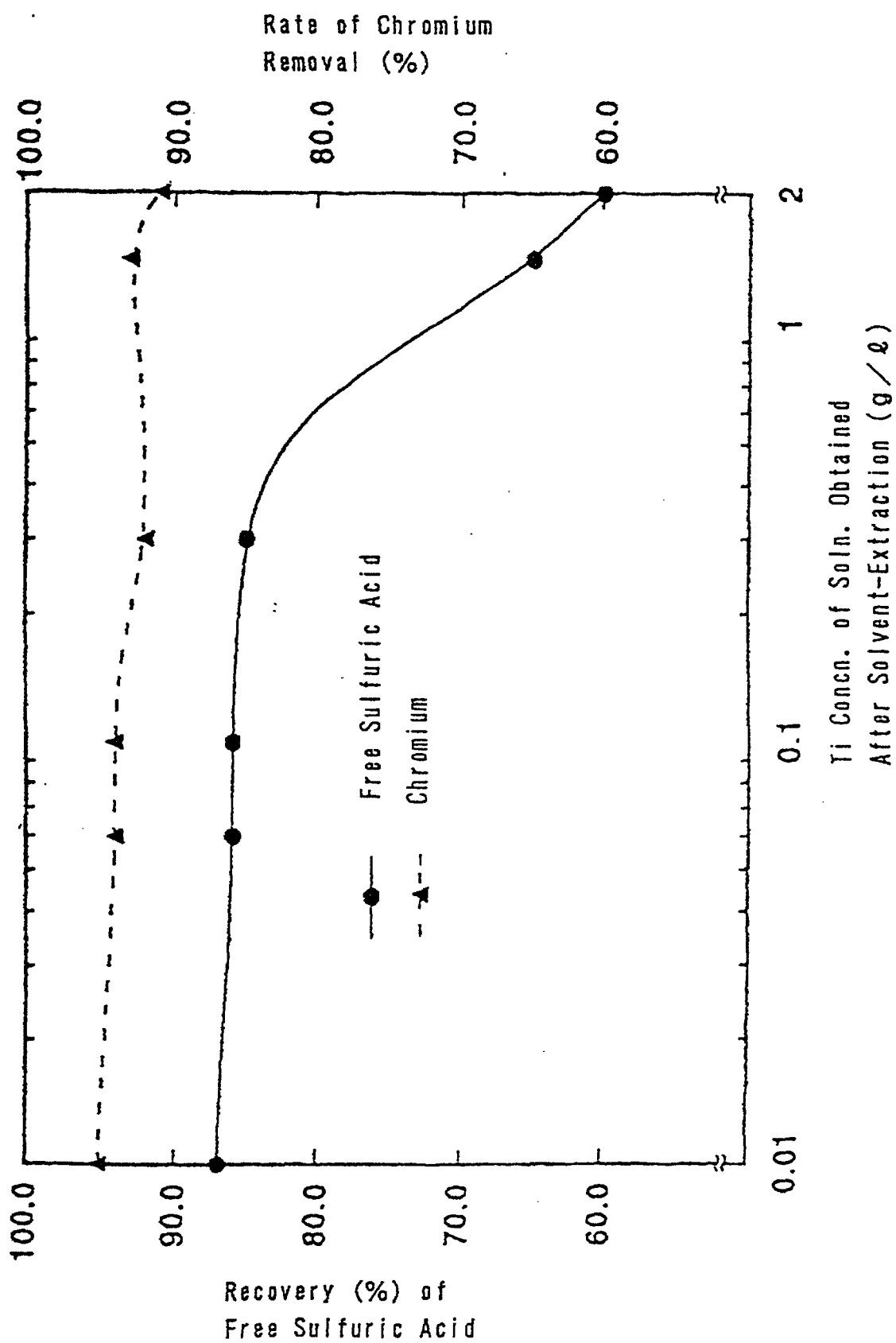
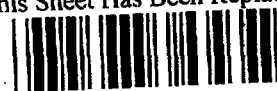


Fig. 1



This Sheet Has Been Replaced



What is claimed is:

1. A method for reclaiming a metal sulfate-containing waste sulfuric acid solution comprising the steps of subjecting the waste sulfuric acid solution to a solvent-extraction treatment to remove titanium present therein and then subjecting the solution obtained after the extraction to a diffusive dialysis treatment.
2. The method for reclaiming a waste sulfuric acid solution as set forth in claim 1 wherein an organic solvent used in the titanium-extraction through the solvent-extraction treatment is brought into contact with an alkali solution to backwardly extract the titanium and to convert it into an insoluble compound; the organic solvent obtained after the titanium-extraction is brought into contact with an aqueous solution remaining after the diffusive dialysis treatment to backwardly extract the alkali ions into the aqueous phase to regenerate the organic solvent; and the regenerated organic solvent is used in the solvent-extraction process for reclaiming the waste sulfuric acid solution.
3. The method for reclaiming a waste sulfuric acid solution as set forth in claim 1 or 2 wherein the waste sulfuric acid solution is cooled to precipitate iron sulfate, then the precipitates are filtered off to control the free sulfuric acid content in the waste solution to 15 to 35% by weight and thereafter the filtrate is subjected to a solvent-extraction treatment.



THIS SHEET HAS BEEN
REPLACED

REPLACEMENT SHEET

25 FEB 1997

This Sheet Has Been Replaced



ABSTRACTMETHOD FOR RECLAIMING METAL SULFATE-CONTAINING WASTE SULFURIC ACID

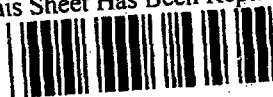
For the purpose of efficiently and easily regenerating a sulfuric acid solution having a high concentration and a low impurity content from a metal sulfate-containing waste sulfuric acid solution and waste liquor formed during acid cleaning which are discharged from titanium (IV) oxide-production process carried out according to the sulfuric acid method, titanium present in the metal sulfate-containing waste sulfuric acid solution is removed through a solvent-extraction treatment and then the solution obtained after the solvent-extraction is subjected to a diffusive dialysis treatment. (Fig. 1)

THIS SHEET HAS BEEN
REPLACED

REPLACEMENT SHEET

25 FEB 1997

This Sheet Has Been Replaced



Method for Reclaiming Metal Sulfate-Containing Waste
Sulfuric Acid

BACKGROUND OF THE INVENTION

The present invention relates to a method for reclaiming waste sulfuric acid discharged from, for instance, a titanium oxide-production process.

The so-called sulfuric acid method has been well-known as a method for preparing titanium oxide (titania) widely used as a coloring material. A large amount of waste sulfuric acid is discharged mainly from the solid/liquid separation step for titanium hydroxide during the titanium oxide-production process and the waste sulfuric acid contains, for instance, Fe, Ti, Cr and V originated from ilmenite and titanium slug as raw materials for titanium.

In the past, the waste sulfuric acid was neutralized and then burried under the ground or dumped at sea in the form of gypsum. However, there has been required for the development of other techniques for treating the waste sulfuric acid from the viewpoint of various problems, for instance, those concerning environmental pollution, insurance of the filled-up land for burrying such gypsum and enomous expenses necessary for the post-treatment of the waste sulfuric acid.

There have been a variety of proposals in response to such request. For instance, Japanese Un-examined Patent Publication (hereunder referred to as "J.P. KOKAI") Nos. Hei 3-80103 and Hei 3-88718 each discloses a method for removing impurities such as

This is the replacement sheet

Date: - 6 MAY 1997

REPLACEMENT SHEETS		SG
☒	Description : paged 1 to paged 14	
☒	Claim(s) : paged 15 to paged 16	
☒	Drawing(s) : paged 17 to paged 18	
☒	Abstract : paged 19 to paged 20	
		06 MAY 1997
DATE OF RECEIPT		

Fe ions, which makes use of a solvent-extraction process. In these methods, however, the strongly acidic sulfuric acid solution per se is directly subjected to a solvent-extraction treatment. Therefore, Fe ions are, in fact, extracted in poor efficiency and the iron-containing component thus recovered also has low purity, in spite of the assertions of the inventors of the foregoing patents. Moreover, in these methods, trivalent iron ions are directly extracted and accordingly, the use of an acidic extraction agent would require a multistage extraction treatment because of the low extraction rate thereof.

Under such circumstances, the inventors of this invention have developed a method for effectively recovering a highly concentrated sulfuric acid solution and filed a patent application, EP-A1-0541002, in which the aforementioned solvent-extraction method is improved so as to enhance the bivalent iron ion-removing efficiency. However, the sulfuric acid solution thus recovered is colored dark green because of the presence of residual Cr and V which are converted into colored oxides and it is difficult to reuse the colored sulfuric acid solution in the process for preparing titanium oxide used, in particular, as a white pigment.

Moreover, there has also been proposed a diffusive dialysis method, as a technique for treating the waste sulfuric acid, which makes use of an anion exchange membrane. However, if the waste sulfuric acid is treated through the diffusive dialysis, particles are deposited on the surface of the ion exchange membrane immediately after putting the dialyzer in operation and this leads to the quality-deterioration of the

This is the replacement sheet

- 6 MAY 1997

Date:

membrane and the breakage of the membrane due to deposition of the particles within the membrane. For this reason, if the foregoing phenomenon occurs during the operation of the device over a long time, the deposits should be removed by washing the anionic exchange membrane with an agent capable of washing them away.

To eliminate the problem of poor efficiency, J.P. KOKAI No. Hei 2-153806 proposes a method for recovering sulfuric acid present in a titanium-containing waste sulfuric acid solution while setting the solution against water in a multistage diffusive dialysis tank equipped with an anion exchange membrane wherein the solution and water are countercurrently supplied to the dialysis tank. In other words, the former is passed, through the tank, from the front stage to the rear stage, while the latter is passed therethrough from the rear stage to the front stage.

This method permits the elimination of the step for washing the anion exchange membrane and a decrease in the degree of quality-deterioration of the membrane, but the membrane should be washed every 10 days as is described in Examples of the patent. Therefore, the method should further be improved in its efficiency.

SUMMARY OF THE INVENTION

It is an object of the present invention to solve the foregoing problems and to provide a method for efficiently and easily regenerating a sulfuric acid solution having a high concentration and a low impurity content from metal sulfate-

This is the replacement sheet

- 6 MAY 1997

Date:

containing waste sulfuric acid and waste liquor formed during acid cleaning which are discharged from the process for preparing titanium (IV) oxide according to the sulfuric acid method.

According to the present invention, the foregoing object can be accomplished by removing titanium present in a metal sulfate-containing waste sulfuric acid solution through a solvent-extraction treatment and then subjecting the solution, from which titanium is extracted, to a diffusive dialysis treatment.

It is preferred to backwardly extract the titanium thus extracted by bringing the organic solvent obtained through the solvent-extraction and containing titanium into contact with an alkaline solution to thus convert the titanium into an insoluble compound through the backward extraction; to bring the organic solvent obtained after the removal of titanium into contact with the remaining solution obtained after the diffusive dialysis treatment for regenerating the organic solvent through the backward extraction of the alkali ions into the aqueous phase; and to reuse the regenerated organic solvent in the solvent-extraction process for reclaiming the waste sulfuric acid.

Moreover, it is desirable that the waste sulfuric acid solution be optionally filtered with cooling to control the concentration of free sulfuric acid present therein prior to the solvent-extraction of the solution.

The method for reclaiming metal sulfate-containing waste sulfuric acid according to the present invention basically

This is the replacement sheet

Date: - 6 MAY 1997

comprises the step of effectively removing principal impurities such as iron, titanium, chromium and vanadium to thus reclaim the solution into highly concentrated sulfuric acid which can widely be reused.

Among these impurities, titanium ions are included in, for instance, the waste sulfuric acid solution discharged from the titanium oxide-production process in an amount ranging from 2 to 10 g/ℓ (these ions are dissolved therein mainly in the form of TiO-SO_4 and partially in the form of $\text{Ti}_2(\text{SO}_4)_3$). Easily hydrolyzable tetravalent titanium ions, among others, can form titanium oxide particles which may become a cause of quality-deterioration and breakage of an ion exchange membrane through deposition thereof on the surface of the ion exchange membrane and within the membrane during the diffusive dialysis treatment. For this reason, the removal of these ions is indispensable for industrially applying the diffusive dialysis method to the reclamation of the metal sulfate-containing waste sulfuric acid. The inventors of this invention have found out that the formation of titanium oxide particles on an ion exchange membrane can substantially be inhibited if the titanium content in the solution is, in advance, reduced to a level of not more than 0.5 g/ℓ through solvent-extraction. The titanium components remaining even after the solvent-extraction are mainly composed of trivalent titanium ions which are less hydrolyzable as compared with tetravalent titanium ions. Therefore, the waste solution obtained after the solvent-extraction treatment may directly be subjected to a diffusive dialysis treatment without impairing the intended effect of the

This is the replacement sheet

Date: - 6 MAY 1997

method to thus give a sulfuric acid solution having a low content of impurities such as chromium and iron. In particular, it is desirable to remove solid contents present in the waste solution obtained after the solvent-extraction through micro filtration prior to the diffusive dialysis thereof. Moreover, if the organic solvent still remains in the waste solution, it is desirable to remove the remaining organic solvent by bringing the waste solution into contact with, for instance, activated carbon.

If the method of the present invention is used for reclaiming waste sulfuric acid, the method can be carried out according to the batchwise process in which a complete waste solution as a waste acid solution is treated or the so-called continuous recovery process which comprises the steps of continuously withdrawing a waste sulfuric acid solution from, for instance, a titanium oxide-production process, recovering the acid and simultaneously separating and removing metal salts separated from the solution, but the continuous recovery process is generally preferred since the organic solvent used in the solvent-extraction treatment can be regenerated by removing titanium components through backward extraction with an alkali solution.

The extraction of titanium ions is carried out using, for instance, an acidic organophosphoric acid compound, a carboxylic acid, a sulfonic acid or hydroxy oxime, as has been well-known in the art. Among the acidic organophosphoric acid compounds, acidic phosphoric acid diesters are particularly preferred.

Titanium ions extracted with, for instance, an acidic

This is the replacement sheet

Date: - 6 MAY 1997

organophosphoric acid compound are backwardly extracted with, for instance, an alkali solution such as an ammonium carbonate solution and recovered in the form of, for instance, hydroxide. The titanium compound can be fired to give titanium oxide after the recovery thereof. The product can be used as a raw material for titanium.

Ferrous sulfate can be crystallized and the sulfuric acid concentration of the waste solution can be controlled to a level favorable for the diffusive dialysis treatment by filtering the waste sulfuric acid solution with cooling prior to the solvent-extraction treatment thereof.

In the method of the present invention, titanium components present in the metal sulfate-containing waste sulfuric acid solution is removed through the solvent-extraction treatment prior to the diffusive dialysis treatment of the waste solution and thus the waste sulfuric acid solution can quite efficiently and easily be reclaimed to give a sulfuric acid solution having a low content of impurities such as iron, while using the diffusive dialysis method which conventionally has a difficulty in efficiency. Moreover, the diffusive dialysis method permits the removal of components such as chromium and vanadium which cannot be removed by the solvent-extraction method. The removal of such components is quite beneficial from the viewpoint of environmental sanitation.

In particular, the method of the present invention has high industrial value. This is because the method permits effective recovery of sulfuric acid from a metal sulfate-containing waste sulfuric acid solution in a closed system and

This is the replacement sheet

Date: 6 MAY 1997

the reuse thereof and also permits the prevention of any environmental pollution by bringing an organic solvent containing extracted titanium and obtained through a solvent-extraction treatment into contact with an equal amount of an alkali solution to thus backwardly extract titanium and to convert it into an insoluble compound; bringing the organic solvent obtained after the titanium extraction into contact with the liquid remaining after the dialysis formed during a diffusive dialysis treatment to thus regenerate the organic solvent through backward extraction of alkali ions into the aqueous phase; and using the regenerated organic solvent in the solvent-extraction process for reclaiming the waste sulfuric acid solution. In addition, titanium, scandium or the like can be recovered as by-products and therefore, the method of the present invention is likewise quite beneficial from the viewpoint of effective use of resources.

If the waste sulfuric acid solution is filtered with cooling prior to the solvent-extraction, iron sulfate can easily be obtained as well.

BRIEF DESCRIPTION OF THE DRAWINGS

The only figure attached hereto is a graph showing the relation between the titanium concentration of the solution obtained by the solvent-extraction treatment and the recovery of free sulfuric acid or the rate of chromium removal observed during the diffusive dialysis treatment.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

This is the replacement sheet

Date: - 6 MAY 1997

Examples of the present invention will hereunder be described, but the present invention is not restricted to these specific Examples.

Example 1

An amount of 10 ℓ of a waste sulfuric acid solution discharged from a titanium oxide-production process carried out according to the sulfuric acid method which had an Fe content of 30g/ℓ (as expressed in terms of the amount of bivalent and trivalent iron ions), a Ti content of 2 g/ℓ (titanium ions), a Cr content of 0.1 g/ℓ (chromium ions), a V content of 0.15 g/ℓ (vanadium ions), an Sc content of 14mg/ℓ (scandium ions) and a sulfate content of 310 g/ℓ (sulfate ions) was brought into contact with 10 ℓ of 1 mol/ℓ solution of di-2-ethylhexylphosphoric acid in kerosine over 20 minutes to thus extract titanium ions and scandium ions present in the aqueous phase into the organic phase. The solvent-extraction treatment gave 10 ℓ of a solvent-extracted solution which had an Fe content of 30g/ℓ (iron ions), a Ti content of 0.01g/ℓ (titanium ions), a Cr content of 0.1 g/ℓ (chromium ions), a V content of 0.15 g/ℓ (vanadium ions), an Sc content of not more than 1 mg/ℓ (scandium ions) and a sulfate content of 310 g/ℓ (sulfate ions). The rate of recovery and removal of titanium and scandium was found to be 99%.

Then the solution thus treated was supplied to a dialysis chamber of a diffusive dialysis device at a flow rate of 240 ml/hr at ordinary temperature, while water was supplied to a recovery chamber of the device separated from the dialysis

This is the replacement sheet

Date: - 6 MAY 1997

chamber by an anion exchange membrane at a flow rate of 150 ml/hr, the solution and water were circulated through each corresponding chamber over 20 hours. As a result, there were recovered 2.98 ℓ of a recovered solution which had an Fe content of 2.1 g/ ℓ (as expressed in terms of the amount of Fe ions), a Ti content of not more than 0.001g/ ℓ (Ti ions), a Cr content of 0.006 g/ ℓ (Cr ions), a V content of 0.014g/ ℓ (V ions) and a sulfate content of 305g/ ℓ (sulfate ions) and 4.6 ℓ of a solution remaining after the dialysis which had an Fe content of 29.8 g/ ℓ (Fe ions), a Ti content of 0.01g/ ℓ (Ti ions), a Cr content of 0.1 g/ ℓ (Cr ions), a V content of 0.147g/ ℓ (V ions) and a sulfate content of 125.8g/ ℓ (sulfate ions). The recovery of free sulfuric acid, the rate of chromium removal and the rate of vanadium removal were found to be 70%, 96% and 94%, respectively.

Moreover, 10 ℓ of the organic phase recovered through the foregoing solvent-extraction treatment was brought into contact with an aqueous phase comprising 0.2mol/ ℓ of ammonium carbonate aqueous solution and 2 mol/ ℓ of aqueous ammonia over 20 minutes, then precipitates formed in the organic phase were filtered off and dried to thus give 58.6 g of a mixture comprising titanium and scandium (having a Ti content of 32.4% and an Sc content of 0.18%). The recoveries of Ti and Sc were found to be 95% and 75%, respectively.

The foregoing organic solvent (12 ℓ) whose volume was changed through the backward extraction was brought into contact with the same volume of the solution remaining after the diffusive dialysis treatment over 20 minutes to backwardly

This is the replacement sheet

Date: 6 MAY 1997

extract the alkali ions present in the organic phase into the aqueous phase and to thus give 10 ℓ of the organic solvent. The organic solvent thus recovered was again brought into contact with a waste sulfuric acid solution discharged during a titanium oxide-production process and as a result, it was found that the recoveries of titanium and scandium reached 99%.

Example 2

An amount of 10 ℓ of a waste sulfuric acid solution discharged from a titanium oxide-production process carried out according to the sulfuric acid method which had an Fe content of 36.5g/ℓ (as expressed in terms of the amount of bivalent and trivalent iron ions), a Ti content of 3 g/ℓ (titanium ions), a Cr content of 0.2 g/ℓ (chromium ions), a V content of 0.6g/ℓ (vanadium ions), an Sc content of 20mg/ℓ (scandium ions) and a sulfate content of 370 g/ℓ (sulfate ions) was cooled to 0 °C to give 960 g of $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ (having an Fe content of 18.8%, a Ti content of 0.04%, a Cr content of 0.002%, a V content of 0.002% and a sulfate ion content of 33.9%).

After crystallizing $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ through the cooling treatment, the waste solution was filtered and 9.5ℓ of the resulting filtrate which had an Fe content of 19.4g/ℓ (iron ions), a Ti content of 3.1 g/ℓ (titanium ions), a Cr content of 0.2 g/ℓ (chromium ions), a V content of 0.6g/ℓ (vanadium ions), an Sc content of 20mg/ℓ (scandium ions) and a sulfate content of 355 g/ℓ (sulfate ions) was brought into contact with 9.5 ℓ of 1 mol/ℓ solution of di-2-ethylhexylphosphoric acid in kerosine over 20 minutes to thus extract titanium ions

This is the replacement sheet

Date: - 6 MAY 1997

and scandium ions present in the aqueous phase into the organic phase. The solvent-extraction treatment gave 9.5ℓ of a solvent-extracted solution which had an Fe content of 19.4g/ℓ (iron ions), a Ti content of 0.01g/ℓ (titanium ions), a Cr content of 0.2 g/ℓ (chromium ions), a V content of 0.6g/ℓ (vanadium ions), an Sc content of not more than 1 mg/ℓ (scandium ions) and a sulfate content of 355 g/ℓ (sulfate ions). The rate of recovery or removal of titanium, scandium was found to be 99%.

Then the solution thus treated was supplied to a dialysis chamber of a diffusive dialysis device at a flow rate of 230 ml/hr at ordinary temperature, while water was supplied to a recovery chamber of the device separated from the dialysis chamber by an anion exchange membrane at a flow rate of 250 ml/hr to thus circulate the solution and water through each corresponding chamber over 20 hours. As a result, there were recovered 2.45ℓ of a recovered liquid which had an Fe content of 1.8 g/ℓ (as expressed in terms of the amount of Fe ions), a Ti content of 0.001g/ℓ (Ti ions), a Cr content of 0.01g/ℓ (Cr ions), a V content of 0.05 g/ℓ (V ions) and a sulfate content of 246g/ℓ (sulfate ions) and 2.37ℓ of a solution remaining after the dialysis which had an Fe content of 16.9 g/ℓ (Fe ions), a Ti content of 0.008 g/ℓ (Ti ions), a Cr content of 0.18g/ℓ (Cr ions), a V content of 0.53 g/ℓ (V ions) and a sulfate content of 90.2 g/ℓ (sulfate ions). The recovery of free sulfuric acid, the rate of chromium removal and the rate of vanadium removal were found to be 80%, 92% and 91%, respectively.

Moreover, 9.5 ℓ of the organic phase recovered through

This is the replacement sheet

Date: - 6 MAY. 1997

the foregoing solvent-extraction treatment was brought into contact with an aqueous phase comprising 0.2mol/ℓ of ammonium carbonate aqueous solution and 2 mol/ℓ of aqueous ammonia over 20 minutes to backwardly extract titanium ions and scandium ions present in the organic phase into the aqueous phase in the form of hydroxides. After allowing the aqueous phase to stand for 20 minutes, the precipitates formed therein were filtered off and dried to thus give 85.8 g of a mixture comprising titanium and scandium (having a Ti content of 33.2% and an Sc content of 0.2%). The recoveries of Ti and Sc were found to be 95% and 85%, respectively.

The foregoing organic solvent (11.4 ℓ) whose volume was changed through the backward extraction was brought into contact with the same volume of the solution remaining after the diffusive dialysis treatment over 20 minutes to backwardly extract the alkali ions present in the organic phase into the aqueous phase and to thus give 9.5ℓ of the organic solvent. The organic solvent thus recovered was again brought into contact with a waste sulfuric acid solution discharged during a titanium oxide-production process and as a result, it was found that the recoveries of titanium and scandium reached 99%.

Example 3

A waste sulfuric acid solution discharged from a titanium oxide-production process performed according to the sulfuric acid method which had an Fe content of 30g/ℓ (as expressed in terms of the amount of bivalent and trivalent Fe ions), a Ti content of 2 g/ℓ (Ti ions), a Cr content of 0.2 g/ℓ (Cr ions)

This is the replacement sheet

Date: - 6 MAY 1997

and a sulfate content of 300 g/ℓ (sulfate ions) was continuously brought into contact with an organic solvent comprising a solution of di-2-ethylhexylphosphoric acid (D2EHPA) in kerosine, whose concentration was variously changed, under the same conditions used in Example 1, over 30 days. Table 1 given below shows titanium concentrations of the solutions obtained through the foregoing solvent-extraction treatment.

Table 1

<u>D2EHPA Conc. of the Organic Solv. Brought into Contact With the Waste Solution (mol/ℓ)</u>	<u>Ti Conc. of the Solution Treated (g/ℓ)</u>
2	0.01
1.5	0.06
1	0.1
0.5	0.3
0.1	1.4
untreated	2

Fig. 1 shows the results observed when the foregoing solution obtained after the solvent-extraction was subjected to the diffusive dialysis treatment. These results indicate that sulfuric acid can efficiently be recovered from waste sulfuric acid solutions by the diffusive dialysis treatment if titanium concentration of the waste solution is reduced to a level of not more than 0.5 g/ℓ. Moreover, it can also be confirmed that the diffusive dialysis treatment permits the effective removal of chromium.

This is the replacement sheet

Date: - 6 MAY 1997

What is claimed is:

1. A method for reclaiming a metal sulfate-containing waste sulfuric acid solution comprising the steps of subjecting the waste sulfuric acid solution to a solvent-extraction treatment to remove titanium present therein and then subjecting the solution obtained after the extraction to a diffusive dialysis treatment.
2. The method for reclaiming a waste sulfuric acid solution as set forth in claim 1 wherein an organic solvent used in the titanium-extraction through the solvent-extraction treatment is brought into contact with an alkali solution to backwardly extract the titanium and to convert it into an insoluble compound; the organic solvent obtained after the titanium-extraction is brought into contact with an aqueous solution remaining after the diffusive dialysis treatment to backwardly extract the alkali ions into the aqueous phase to regenerate the organic solvent; and the regenerated organic solvent is used in the solvent-extraction process for reclaiming the waste sulfuric acid solution.
3. The method for reclaiming a waste sulfuric acid solution as set forth in claim 1 or 2 wherein the waste sulfuric acid solution is cooled to precipitate iron sulfate, then the precipitates are filtered off to control the free sulfuric acid content in the waste solution to 15 to 35% by weight and thereafter the filtrate is subjected to a solvent-extraction treatment.



* C L N S *

This is the replacement sheet

Date: 6 MAY 1997

ABSTRACT

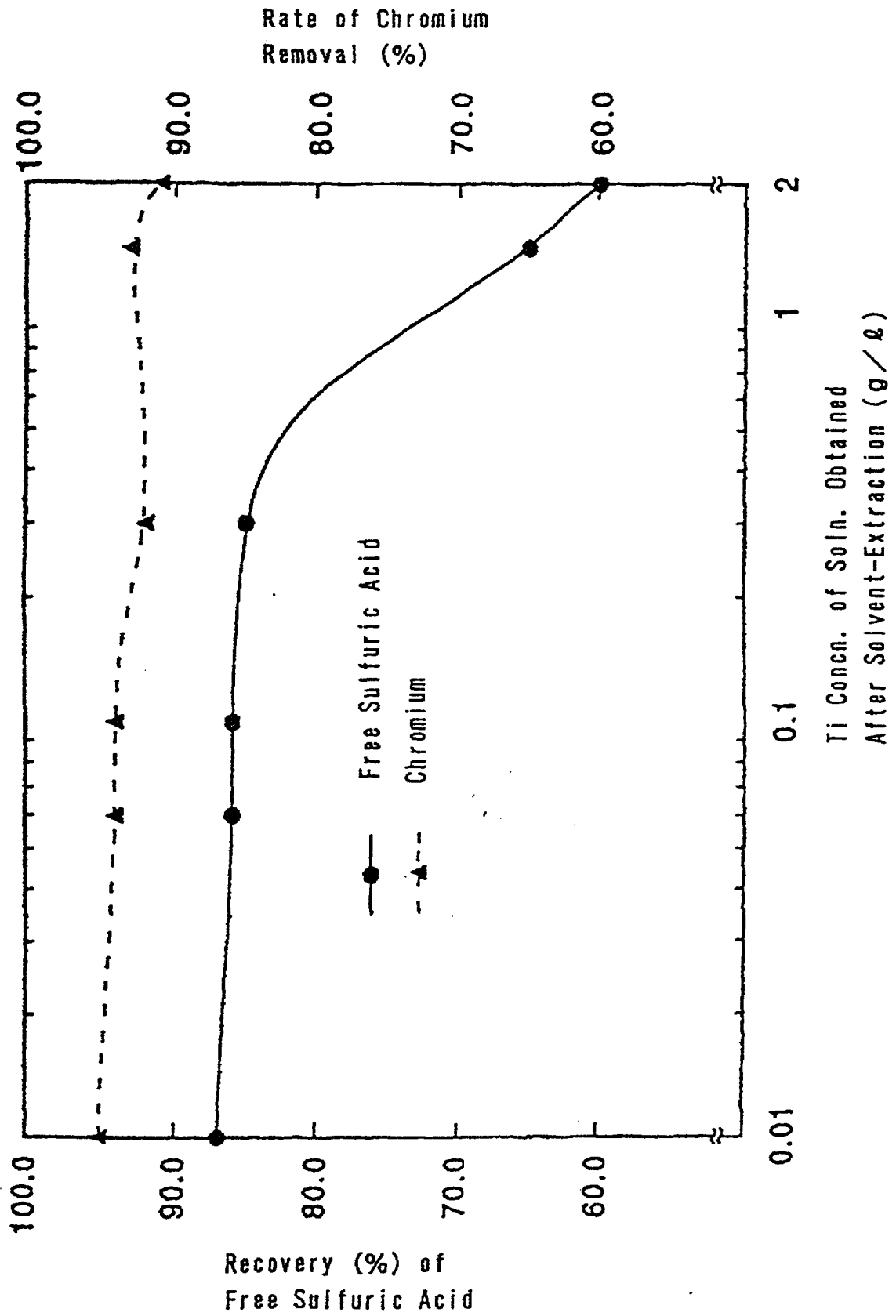
METHOD FOR RECLAIMING METAL SULFATE-CONTAINING WASTE SULFURIC ACID

For the purpose of efficiently and easily regenerating a sulfuric acid solution having a high concentration and a low impurity content from a metal sulfate-containing waste sulfuric acid solution and waste liquor formed during acid cleaning which are discharged from titanium (IV) oxide-production process carried out according to the sulfuric acid method, titanium present in the metal sulfate-containing waste sulfuric acid solution is removed through a solvent-extraction treatment and then the solution obtained after the solvent-extraction is subjected to a diffusive dialysis treatment. (Fig. 1)

This is the replacement sheet

Date: 5-6 MAY 1997

Fig. 1



This is the replacement sheet

Date: 8 MAY 1997