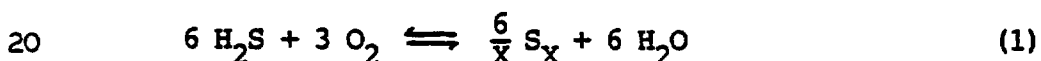


PROCESS FOR THE OXIDATION OF H_2S TO S
AND/OR SO_2 USING A SILICA SUPPORTED CARRIER

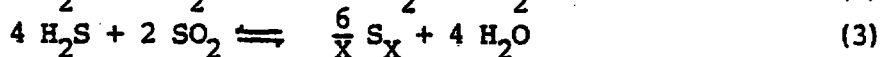
The invention relates to a process for the oxidation of hydrogen sulphide to elemental sulphur and/or sulphur dioxide.

Hydrogen sulphide (H_2S)-containing gases become available, for example, in oil refineries and from processing coke-oven gas. Natural gas may also contain H_2S . H_2S may be removed from H_2S -containing gases by absorption in a regenerable absorbent. Subsequent regeneration of the absorbent yields a gas having a much higher H_2S content than the H_2S -containing starting gas and usually also containing carbon dioxide (CO_2). The H_2S -containing gases obtained by this regeneration are usually fed to a Claus-type process for the production of elemental sulphur. "Elemental sulphur" is hereinafter also referred to as "sulphur".

In a first version of the Claus-type process an H_2S -containing gas is partially combusted with a free oxygen-containing gas in a furnace referred to herein as "thermal zone". In the thermal zone the following reaction takes place with regard to sulphur formation:



which can be considered as occurring in two steps:



In the thermal zone the H_2S is converted to sulphur in accordance with reaction (1), the quantity of free oxygen-containing gas being such that only one third of all H_2S can be oxidized to sulphur dioxide (SO_2) in accordance with reaction (2). The effluent gas from the thermal zone contains H_2S , SO_2 ,

sulphur vapour, nitrogen, water vapour and usually also CO_2 , carbon monoxide (CO), carbonyl sulphide (COS), carbon disulphide (CS_2) and hydrogen.

The effluent gas from the thermal zone has a temperature usually in the range of from 600°C to 1650°C and is cooled to condense most of the sulphur vapour present, to a temperature suitably in the range of from 120°C to 200°C . Thus, sulphur is recovered as a valuable product. The gases are then re-heated, to a temperature usually in the range of from 230°C to 280°C and introduced into a first catalytic zone where H_2S reacts with SO_2 to produce more sulphur in accordance with reaction (3). The effluent from the first catalytic zone is cooled to condense sulphur. The gas stream freed from sulphur is re-heated, to a temperature usually in the range of from 200°C to 280°C and introduced into a second catalytic zone to produce more sulphur. The second catalytic zone may be followed by a third and, if desired, the third by a fourth catalytic zone to produce still more sulphur at a temperature usually in the range of from 200°C to 280°C .

Only gases containing much H_2S - more than, for example, 40% by volume - can be readily combusted in the thermal zone in accordance with reaction (1). Gases containing less H_2S - between, for example, 20 and 40% by volume - can be processed according to a second version of the Claus-type process.

In the second version the H_2S -containing gas is split into a first portion, representing at least one third and not more than two thirds of the total H_2S -containing gas and a second portion containing the balance of the total H_2S -containing gas. The first portion is combusted with a free oxygen-containing gas in an amount stoichiometric with respect to oxidation of one third of the total amount of H_2S according to reaction (2), thus forming a limited amount of sulphur.

The effluent gas from the thermal zone is cooled to condense the major part of the sulphur formed and then introduced, in admixture with the said second portion of the H_2S -containing gas, into a first catalytic zone from where the operation is continued as described for the first version.

Gases containing less than, for example, 20% by volume of H_2S , can be processed according to a third version of the Claus-type process. In the third version the H_2S -containing gas is pre-heated and introduced in admixture with a free oxygen-containing gas into a catalytic zone where H_2S is oxidized to sulphur in accordance with reaction (1), at a temperature usually in the range of from, for example, $200^{\circ}C$ to $280^{\circ}C$. The operation is then continued as described for the first version.

The remaining gases as obtained after condensation of sulphur from the gases leaving the final catalytic zone of any version of the Claus-type process are also referred to herein as "Claus tail gases". These gases contain nitrogen, water vapour, some H_2S and SO_2 and usually also CO_2 , CO , COS , CS_2 and hydrogen. H_2S and SO_2 are present because of the position of the equilibrium of reaction (3) at the temperature maintained in the final catalytic zone referred to hereinbefore. COS and CS_2 are present because at least part thereof has been formed in the thermal zone and they have only partly been hydrolyzed in the catalytic zone or zones mentioned hereinbefore.

Viewed in the light of increasingly stringent requirements with respect to air pollution abatement, various procedures have been developed to remove H_2S from process off-gases, for example from Claus tail gases, and even recover, if possible, H_2S or reaction products of H_2S contained therein.

To remove these sulphur compounds, selective absorption by contacting the Claus tail gases with a suitable absorption solvent, after a hydrogenation treatment of the tail gases, has been

practised. In this procedure, the bulk of the desorbed H_2S after regeneration of the absorption solvent, is returned to the Claus unit, and the final off-gas of this Claus tail gas-treating process, which off-gas contains nitrogen, CO_2 and quite minor amounts of H_2S , is incinerated. By the incineration, H_2S is converted to SO_2 , a material which generally has not been subjected to emission requirements as rigid as those applied to H_2S . However, incineration is costly because of the necessary heat input.

Thus, there exists a need for an economical method for the purification of H_2S -containing streams, particularly off-gas streams of the type mentioned, which method would provide substantial conversion of the H_2S in the off-gas, and concomitantly provide low SO_2 emissions. The method should be economical in that the necessary heat input should be as low as possible as evidenced by relatively low process temperatures. Moreover, the catalysts to be applied should show an activity which remains constant over a long period.

It is an object of the present invention to satisfy that need.

Another object is to catalyze reactions (1) and (2) more effectively than hitherto has been the case.

Accordingly, the invention provides a process for the oxidation of hydrogen sulphide to elemental sulphur and/or sulphur dioxide, in which process a hydrogen sulphide-containing gas is contacted in the presence of a free oxygen-containing gas with a first catalyst composition comprising at least one metal from Group 3b and/or Group 4b of the Periodic Table of the Elements on a silica-containing carrier.

The Periodic Table of the Elements referred to herein is shown on the inside cover of "Handbook of Chemistry and Physics", 63rd edition (1982-1983). The Group 3b metals mentioned hereinbefore are scandium, yttrium, thorium and the lanthanides, i.e.

lanthanum, cerium, praseodymium, neodymium, promethium, samarium, europium, gadolinium, terbium, dysprosium, holmium, erbium, thulium, ytterbium and lutetium. The Group 4b metals mentioned hereinbefore are titanium, zirconium and hafnium. Preferably, the metal from Group 3b and/or Group 4b of the Periodic Table of the Elements is applied as an oxide or sulphate thereof. Most preferably, at least one oxide of titanium, zirconium and/or thorium is applied on the carrier. These oxides are themselves active catalysts or are converted to active catalysts in contact with the starting gas. Very good results have been obtained with titanium dioxide and zirconium dioxide. Modifying the process according to the present invention by replacing the first catalyst composition with pure TiO_2 results in high SO_3 emissions. Surprisingly, SO_3 emissions are low when TiO_2 on a silica-containing carrier is used.

The metals from Group 3b and/or Group 4b are applied in an atomic ratio of metals to silicon which can be varied. As a rule, atomic ratios of metal to silicon in the range of from 0.001 to 1.0 are preferred, atomic ratios in the range of from 0.03 to 0.3 being particularly preferred.

Suitably, at least 50% by weight of silica is applied in the silica-containing carrier. Preferably, at least 75% and, most preferably, at least 90% by weight of silica is applied in the carrier. Among the synthetic silicas commercially available those containing at least 98.0% by weight of silica are generally the most suitable.

According to a very attractive embodiment of the present invention a carrier is used which has been prepared by the following process steps:-

- step a) preparing a silica hydrosol by mixing an aqueous solution of an alkali metal silicate with an aqueous solution of an acid;
- step b) converting the hydrosol into droplet form;
- step c) shaping the droplets in air or in a liquid which is not miscible with water;
- step d) partially pre-drying the hydrogel particles obtained;
- step e) subjecting the partially pre-dried particles to a hydro-

thermal treatment;

step f) decreasing the cation content of the hydrogel particles thus treated in an aqueous medium to less than 10% by weight, calculated on dry material, and

5 step g) drying and optionally calcining the silica particles thus obtained.

The silica support thus prepared has a very high attrition resistance and a very high mean side crushing strength. A description of this method of preparation can be found in European
10 patent application No. 0067459. A suitable way of preparing the first catalyst composition comprises incorporating a compound or compounds of a Group 3b and/or Group 4b metal into the silica when the silica is being prepared, for example during step a) or subsequent to step f) and prior to step g). If desired, a portion of
15 the said compound or compounds may be incorporated into the silica during step a) and the balance subsequent to step f) and prior to step g).

The first catalyst composition may further be prepared by such conventional techniques as, for example, dry-mixing followed
20 by calcination, co-gellation, co-precipitation, impregnation and ion-exchange. For example, a mixture of a titanium salt and a silica sol may be co-gelled, the material then being dried and ground to an appropriate size, or the co-gelled material may be slurried and spray-dried. However, the first catalyst composition
25 may, for example, also be prepared by reacting the hydroxyl groups at the surface of a silica with a titanium salt by the procedure described in U.S. patent specification No. 3,166,542, 3,220,959 or 3,274,120, thus producing a first catalyst composition in which titanium is in chemical combination with silica. Examples of
30 suitable titanium salts are titanium tetrachloride, titanium oxalate and titanyl sulphate, $(TiOSO_4)$; the latter dissolved in a mixture comprising sulphuric acid and water. In yet another technique, a fumed pyrogenic catalyst, in particular a pyrogenic titania-silica composition is prepared by combustion of hydrogen

and oxygen with a mixture of silicon tetrahalide and a titanium halide, "halide" referring to fluoride, chloride, bromide or iodide.

5 Another suitable way of preparing the first catalyst composition comprises impregnating silica with a substantially non-aqueous solution of a titanium compound in a non-basic, essentially inert, oxygen-substituted hydrocarbon as a solvent, removing solvent from the impregnated silica and thereafter calcining the impregnated silica, again producing catalyst com-
10 positions in which titanium is in chemical combination with silica. A description of this method of preparation is found in British patent specification No. 1,332,527.

The first catalyst composition may be subjected to a pre-treatment prior to utilization in the process. As a rule, it is
15 preferable to do so in order to obtain a higher activity. The pre-treatment suitably consists in heating the first catalyst composition in the atmosphere of a non-reducing gas, such as, for example, nitrogen, argon, CO_2 or of a free oxygen-containing gas, such as, for example, air. However, the most suitable method
20 of pre-treatment in general also depends upon the form of chemical combination in which the metal compound is provided. In many instances titanium compounds have to be converted into oxide. This conversion may as a rule suitably be effected by heating in a non-reducing atmosphere, at temperatures in particular in the
25 range of from 250°C to 800°C for periods in the range of from 1 to 18 hours.

The first catalyst composition may be used according to the invention in any convenient physical form, for example, as a powder, flakes, spheres or pellets. Very good results have been
30 obtained with spheres.

The equilibrium position of reaction (1) in a given case depends on the water content, if any, of the H_2S -containing starting gas. It may be desirable to dry the H_2S -containing gases to remove water in order to shift this equilibrium po-
35 sition to the right hand side.

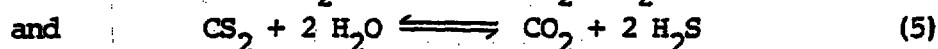
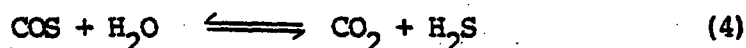
The oxidation of H_2S to sulphur in the presence of the first catalyst composition can be carried out in a wide range of temperatures and space velocities. Suitably, the temperature is in the range of from $150^{\circ}C$ to $350^{\circ}C$ and the space velocity in the range of from 250 to 20,000 Nm^3 gas per m^3 first catalyst composition per hour. It has been found that even when relatively low temperatures and relatively high space velocities are applied thermodynamic equilibrium of reaction (1) is reached, the first catalyst composition being very effective. Preferably, temperatures in the range of from $180^{\circ}C$ to $250^{\circ}C$ and space velocities in the range of from 1000 to 10,000 Nm^3 gas per m^3 first catalyst composition per hour are applied.

As stated hereinbefore gases containing less than, for example, 20% by volume of H_2S can be processed according to the third version of the Claus-type process. The H_2S introduced into the catalytic zone of the third version can suitably be oxidized to sulphur in the presence of the first catalyst composition.

The amount of free oxygen to be used for the oxidation of H_2S to sulphur is important in that a stoichiometric amount of oxygen with respect to reaction (1) is required; the amount of oxygen to be used if no SO_2 is desired should therefore be as close as possible to 50% by volume of the amount of H_2S being used. As stated hereinbefore, the oxidation of H_2S to sulphur can be considered as occurring in an oxidation step (reaction 2) followed by a Claus reaction (reaction 3).

The sulphur-containing gas obtained is cooled to condense sulphur therefrom, leaving an off-gas containing H_2S and SO_2 . If desired, this off-gas may be used as a starting material for further production of elemental sulphur by contacting it with the first catalyst composition at a temperature below the dew point of sulphur, as described in British patent application No. 8318099, filed on even date.

Hydrogen sulphide-containing gases usually contain minor amounts of COS and CS₂. These two compounds are present because at least part thereof has been formed in the thermal zone and they have only partly been converted in the catalytic zone or zones mentioned hereinbefore. During the process according to the present invention they are hydrolyzed according to the reactions



The H₂S thus formed is subsequently oxidized to sulphur according to the present invention.

The oxidation of H₂S to SO₂ in the presence of the first catalyst composition can also be carried out in a wide range of temperatures and space velocities. Suitably, the temperature is in the range of from 150°C to 500°C and the space velocity in the range of from 250 to 20,000 Nm³ gas per m³ first catalyst composition per hour. It has been found that even when relatively low temperatures and relatively high space velocities are applied H₂S is substantially converted to SO₂ according to reaction (2). Preferably, temperatures in the range of from 250°C to 450°C and space velocities in the range of from 1000 to 10,000 Nm³ gas per m³ first catalyst composition per hour are applied.

The amount of free oxygen to be used for the oxidation of H₂S to SO₂ is important in that a stoichiometric excess, preferably a large excess, of free oxygen is desired in order to oxidize substantially all H₂S and any COS and CS₂ present. In general, at least twice and normally up to five times the stoichiometric amount of oxygen required for the oxidation to SO₂ may be supplied. Preferably, an excess of 20 to 280% of the stoichiometric amount of oxygen, based upon all total combustibles, will be supplied. Amounts as high as 100 or even 200 times the stoichiometric amount of free oxygen may be supplied, if desired.

The SO₂ formed may be discharged into the atmosphere or recovered in a manner known to those skilled in the art.

Although the oxidation of H_2S to SO_2 according to the invention is applicable to any H_2S -containing gas of low to moderate H_2S content, the invention is ideally suited to the treatment of H_2S -containing off-gases from various processes from which no further or little recovery of other materials is made, for example to the incineration of a Claus tail gas.

Additionally, as indicated, off-gases may previously have been processed by an off-gas-treating process in order to reduce the contents of combined and elemental sulphur thereof. The invention is eminently suited to the removal of H_2S from off-gases of a Claus tail gas-treating process. Such a process may comprise the step of catalytically reducing SO_2 , COS , CS_2 and sulphur tri-oxide (SO_3) contained in the gas under suitable conditions to H_2S .

So, according to a preferred embodiment of the present invention in which hydrogen sulphide is oxidized to elemental sulphur, the hydrogen sulphide has been obtained by reducing Claus tail gases at a temperature above $175^\circ C$ with a free hydrogen- and/or carbon monoxide-containing gas over a second catalyst composition comprising a sulphided metal from Group 6 and/or Group 8 of the Periodic Table of the Elements and an inorganic oxide carrier supporting the said sulphided metal(s). The hydrogen sulphide present in the reduced Claus tail gases is then oxidized to sulphur in the presence of the first catalyst composition.

The reduction of Claus tail gases in the presence of the second catalyst composition is preferably carried out in the presence of at least the stoichiometric quantity of hydrogen and/or CO for complete conversion of sulphur and SO_2 into H_2S . Generally, 1.3-2.0 times the required stoichiometric quantity is used. Water present in Claus tail gases hydrolyses COS and CS_2 according to reactions (4) and (5), respectively.

Methods of preparation and the most suitable composition of the second catalyst composition and the conditions at which the reduction may be carried out, such as the pressure, the gas hourly

space velocity and the composition of the free hydrogen- and/or free carbon monoxide-containing gas are found in British patent specification No. 1,356,289.

5 The reduced Claus tail gas is preferably dried to remove water prior to the oxidation of H_2S to sulphur, thus moving the equilibrium position of reaction (1) to the right hand side. Drying may be carried out by, for example, condensation of water from the reduced Claus tail gas.

10 According to a preferred embodiment of the present invention elemental sulphur is removed from the gases after having contacted the first catalyst composition and H_2S remaining in the substantially sulphur-free gases is oxidized to sulphur dioxide, preferably in the presence of the first catalyst composition. As in this manner 98.5 to 99.0% of the combined and elemental sulphur can be
15 removed from Claus tail gases, the gases obtained after oxidation to SO_2 over the first catalyst composition can usually be discharged into the atmosphere without objection.

According to another preferred embodiment of the present invention in which hydrogen sulphide is oxidized to sulphur
20 dioxide, the hydrogen sulphide has been obtained by reducing Claus tail gases at a temperature above $175^\circ C$ with a free hydrogen- and/or carbon monoxide-containing gas over a second catalyst composition comprising a sulphided metal from Group 6 and/or Group 8 of the Periodic Table of the elements and an
25 inorganic oxide carrier supporting the said sulphided metal(s), absorbing the H_2S present in the reduced gases followed by desorption of the H_2S and recycle of the desorbed H_2S to the Claus plant, the hydrogen sulphide to be oxidized to sulphur dioxide remaining in the final off-gas from said absorption of
30 H_2S . This hydrogen sulphide is oxidized to sulphur dioxide in the presence of the first catalyst composition.

The pressures employed in the process according to the invention are not critical and may vary within a wide range.

The total pressure may be at or in excess of atmospheric pressure, although sub-atmospheric pressure may be used. Preferably, the pressure is at least atmospheric and may be up to, for example, 10 bar.

5 The free oxygen may be supplied as, for example, pure oxygen, oxygen-enriched air, air, or other gaseous streams containing significant quantities of oxygen and other components which do not interfere significantly with the reaction contemplated.

10 The following Examples further illustrate the invention. The pore volumes of the starting silicas have been measured by means of absorption of liquid water.

Example 1

15 An amount (bulk volume 250 ml, weight 98 g) of silica gel spheres (pore volume 1.10 ml/g, surface area 300 m²/g) was kept for 15 min under a pressure of 0.02 bar abs. The spheres were then impregnated with tetraisopropylorthotitanate under nitrogen of atmospheric pressure. The impregnated spheres were dried at a temperature of 110°C and the temperature of the dried spheres was increased to 250°C at a rate of 100°C/h and to 500°C at a rate of 20 50°C/h. The temperature was kept for one hour at 500°C, increased at a rate of 50°C/h to 550°C and kept for 3 h at 550°C. Then, the spheres were allowed to adopt ambient temperature. The catalyst composition thus prepared is referred to hereinafter as "first catalyst composition A"; some properties thereof are stated in 25 Table I.

Table I

First catalyst composition	Atomic ratio Ti/Si	Size of spheres mm	Pore volume ml/g	Surface area m ² /g
A	0.21	1.6	0.84	221
B	0.08	5.0	0.54	312
C	0.08	5	0.58	303
D	0.05	2.5	1.33	197
	Zr/Si			
E	0.05	5	0.59	313

5 A cylindrical tube having an inside diameter of 4.09 cm was charged over a height of 38.1 cm with a fixed bed of first catalyst composition A. A gaseous feed composed as shown in Table II was conducted in downflow through the fixed bed. Conditions of operation are shown more specifically in Table II. Significant H₂S-removal is obtained at temperatures above 200°C; the results for three different temperatures are presented in Table III.

Table II

Composition, %v

H ₂ S	5.73
O ₂	11.2
N ₂	44.8
CO ₂	11.0
H ₂ O	27.2

Operating conditions

Pressure : atmospheric

Space velocity: 1000 Nl per 1 first catalyst composition A per h

5 Air rate : 30% excess O₂

Table III

Temperature, °C	200	250	300
Composition			
H ₂ S, ppmv	760	700	360
SO ₂ , %v	5.82	5.83	5.84
SO ₃ , ppmv	80	47	56

Example 2

A feed composed as stated in Table IV was conducted with a space velocity of 1000 Nl per 1 first catalyst composition A per hour through the catalyst bed used in Example 1. Three different temperatures were used. The H₂S and SO₂ contents of the off-gas are presented in Table IV.

Table IV

Composition, %v	Feed	Off-gas		
		175°C	200°C	250°C
H ₂ S	6.24	1.64	0.47	0.44
SO ₂	0	1.16	0.48	0.45
O ₂	3.06	x)	x)	x)
N ₂	49.1	48.4	47.8	47.8
CO ₂	12.0	11.8	11.7	11.7
H ₂ O	29.6	33.6	34.4	34.4
S	0	3.35	5.13	5.19

Total sulphur

conversion, %

54

84

85

x) not determined

Table IV shows that significant conversion of H₂S to sulphur takes place at temperatures above 175°C.

Example 3

Example 1 was modified in that a gas composed as stated in Table V was conducted with a space velocity of 2000 Nl per 1 first catalyst composition per hour through the bed of first catalyst composition A.

Table V

Composition, %v on water-free gas

H ₂ S	1.01
N ₂	81.5
CO ₂	17.0
O ₂	0.50

Table VI states which combinations of temperature and water content were used and also presents the composition of the off-gas and the total sulphur conversion.

Table VI

Temperature, °C	250	200	200	200	250
Water content of feed, %v	30	30	7.4	0	0
Off-gas composition, %v on water-free basis					
H ₂ S	0.62	0.61	0.41	0.13	0.47
SO ₂	0.40	0.38	0.23	0.12	0.27
S	0	0.02	0.37	0.76	0.27
Total sulphur conversion, %	0	2	37	76	27

The thermodynamic equilibrium was reached in each of these five experiments. Table VI clearly shows that removal of water from the feed has a very favourable effect on the total sulphur conversion.

Example 4

- 5 An amount (46.19 g) of tetraisopropylorthotitanate was added dropwise under nitrogen to water (150 ml) and the precipitate thus formed was filtered off and washed three times with water (10 ml each time). The TiO₂ thus obtained was added to a solution of oxalic acid (12.6 g) in water (80 ml). Then, oxalic acid (10.23 g)
- 10 was added. The mixture thus obtained was filtered off, yielding 110 ml of an opale liquid. An amount (84.5 g, bulk volume 200 ml) of silica gel spheres (pore volume 1.02 ml, surface area 349 m²/g) was impregnated with this solution (110 ml). The impregnated spheres were dried at a temperature of 120°C and the temperature
- 15 of the dried spheres was increased to 320°C at a rate of 100°C/h, kept for 0.5 h at 320°C, increased at a rate of 100°C/h to 600°C and kept for one hour at 600°C. Then, the spheres were allowed to adopt ambient temperature.

The first catalyst composition thus prepared is referred to as "first catalyst composition B"; some properties thereof are stated in Table I.

5 A cylindrical tube having an inside diameter of 4.09 cm was charged over a height of 38.4 cm with a fixed bed of first catalyst composition B. A gaseous feed composed as shown in Table VII was conducted in downflow through the fixed bed at a space velocity of 1000 Nl per l first catalyst composition B per hour. Table VII presents the composition of the off-gas.

Table VII

Compound	Composition, % by volume	
	feed	off-gas
H ₂ S	0.98	0.12
SO ₂	0	0.05
SO ₃	0	< 0.0005
H ₂ O	4.5	5.4
CO ₂	11.7	11.7
O ₂	0.5	< 0.0001
N ₂	82.3	82.0
S	0	0.8

10 The temperature and a further result are stated in Table VIII.

Table VIII

Example	First catalyst composition	Temperature, °C	Approach of thermodynamic equilibrium reaction (1), %	Content in off-gas, ppmv, of	
				H ₂ S	SO ₃
4	B	250	100	-	-
5	B	300	100	-	-
6	C	250	100	-	-
7	D	250	100	-	-
8	E	250	100	-	-
9	E	300	100	-	-
10	B	250	-	30	not determined
11	B	300	-	<3	7
12	B	350	-	<3	<5
13	C	300	-	6	5
14	C	350	-	<3	<5
15	D	300	-	8	<5
16	D	350	-	<3	<5
17	E	300	-	<3	<5

Example 5

The experiment described in Example 4 was modified in that a temperature of 300°C was applied. Table VIII presents the results.

Example 6

- 5 TiCl₄ (38.85 g) was added dropwise at a temperature of 4 to 13°C to water (150 ml). An amount (146 ml) of the solution thus obtained was used for impregnation of silica gel spheres (128.4 g, bulk volume 300 ml, pore volume 1.02 ml/g, surface area 349 m²/g). The impregnated spheres were then treated as described for first
- 10 catalyst composition B except that a temperature of 500°C instead

of 600°C was used. The first catalyst composition thus prepared is referred to as "first catalyst composition C"; some properties thereof are stated in Table I.

The experiment described in Example 4 was modified in that first catalyst composition C was used. The result is presented in Table VIII.

Example 7

An aqueous solution of sodium silicate was mixed with a solution obtained by dissolving TiO_2 in dilute aqueous sulphuric acid, using a volume ratio of the latter to the former solution of 0.75. The hydrosol thus obtained was shaped into droplet form, the droplets were shaped in air and pre-dried at 120 °C for 7.5 min. The pre-dried particles were quenched with water to 20 °C and the temperature of the particles was increased over a period of 2.5 h from 20 °C to 150 °C in the presence of liquid water and kept for 1 h at 150 °C. The particles were allowed to adopt a temperature of 20 °C, washed with dilute sulphuric acid, dried for 20 h at 120 °C, heated over a period of 5 min from 20 °C to 875 °C and kept for 10 min at 875 °C. Then, the particles were allowed to adopt ambient temperature. The catalyst composition thus prepared is referred to as "first catalyst composition D"; some properties thereof are stated in Table I.

The experiment described in Example 4 was modified in that first catalyst composition D was used.

The result is presented in Table VIII.

Example 8

An amount (90.7 g, bulk volume 200 ml) of silica spheres having the same properties as those used for the preparation of catalyst composition C was impregnated with 92.5 ml of an aqueous solution of zirconium acetate containing 7.2 g of this salt. The impregnated spheres were further treated as described for first catalyst composition C. The catalyst composition thus prepared is referred to as "first catalyst composition E"; some properties thereof are stated in Table I.

The experiment described in Example 4 was modified in that first catalyst composition E was used. The result is presented in Table VIII.

Example 9

5 The experiment described in Example 8 was modified in that a temperature of 300°C was used. The result is presented in Table VIII.

Comparative Experiment

10 The experiment described in Example 4 was modified in that the bed of first catalyst composition B was replaced with silica spheres.

Reaction (1) had reached only 54% of the thermodynamic equilibrium.

Examples 10, 11 and 12

15 The experiment described in Example 4 was modified in that a gaseous feed composed as shown in Table IX was used at the temperatures shown in Table VIII. Table VIII also presents the results.

Table IX

Composition,	%vol
H ₂ S	1.0
N ₂	balance
CO ₂	11.4
H ₂ O	12.0
O ₂	3.6

Examples 13 and 14

20 The experiments described in Examples 11 and 12 were modified in that first catalyst composition C was used. Table VIII presents the results.

Examples 15 and 16

25 The experiments described in Examples 11 and 12 were modified in that first catalyst composition D was used. Table VIII presents the results.

Example 17

The experiment described in Example 11 was modified in that first catalyst composition E was used. Table VIII presents the results.

5 Comparative Experiment

The experiment described in Example 12 was modified in that first catalyst composition B was replaced with pure titanium dioxide. The off-gas contained less than 4 ppmv of H_2S and 50 ppmv of SO_3 .

C L A I M S

1. A process for the oxidation of hydrogen sulphide to elemental sulphur and/or sulphur dioxide, in which process a hydrogen sulphide-containing gas is contacted in the presence of a free oxygen-containing gas with a first catalyst composition comprising
5 at least one metal from Group 3b and/or Group 4b of the Periodic Table of the Elements on a silica-containing carrier.
2. A process as claimed in claim 1, in which the said metal or metals is (are) applied as oxide or sulphate thereof.
3. A process as claimed in claim 2, in which at least one oxide
10 of titanium, zirconium and/or thorium is applied on the carrier.
4. A process as claimed in any one of the preceding claims, in which the said metal or metals is (are) applied in the catalyst composition in an atomic ratio of metal or metals to silicon in the range of from 0.001 to 1.0.
- 15 5. A process as claimed in claim 4, in which the atomic ratio being applied is in the range of from 0.03 to 0.3.
6. A process as claimed in any one of the preceding claims, in which at least 75% by weight of silica is applied in the carrier.
7. A process as claimed in claim 6, in which at least 90% by
20 weight of silica is applied in the carrier.
8. A process as claimed in any one of the preceding claims, in which the carrier has been prepared by the following process steps:-
 - step a) preparing a silica hydrosol by mixing an aqueous
25 solution of an alkali metal silicate with an aqueous solution of an acid;
 - step b) converting the hydrosol into droplet form;
 - step c) shaping the droplets in air or in a liquid which is not miscible with water;
 - 30 step d) partially pre-drying the hydrogel particles obtained;
 - step e) subjecting the partially pre-dried particles to a hydrothermal treatment;

22



- 23 -

step f) decreasing the cation content of the hydrogel particles thus treated in an aqueous medium to less than 10% by weight, calculated on dry material, and

step g) drying and optionally calcining the silica particles thus obtained.

9. A process as claimed in any one of the preceding claims, in which hydrogen sulphide is oxidized to sulphur dioxide at a temperature in the range of from 250°C to 450°C.

10. A process as claimed in any one of the preceding claims, in which hydrogen sulphide is oxidized to sulphur dioxide supplying the free oxygen-containing gas in an amount of up to five times the stoichiometric amount of oxygen required for oxidation to sulphur dioxide.

11. A process as claimed in claim 10, in which the free oxygen-containing gas is supplied in an excess of 20% to 280% of the stoichiometric amount of oxygen.

12. A process as claimed in any one of claims 1-8, in which hydrogen sulphide is oxidized to elemental sulphur at a temperature in the range of from 150°C to 350°C.

13. A process as claimed in claim 12, in which hydrogen sulphide is oxidized to elemental sulphur at a temperature in the range of from 180°C to 250°C.

14. A process as claimed in any one of claims 1 to 8, 12 or 13, in which hydrogen sulphide is oxidized to elemental sulphur, which hydrogen sulphide has been obtained by reducing Claus tail gases as hereinbefore defined at a temperature above 175°C with a free hydrogen- and/or carbon monoxide-containing gas over a second catalyst composition comprising a sulphided metal from Group 6 and/or Group 8 of the Periodic Table of the Elements and an inorganic oxide carrier supporting the said sulphided metal(s).

15. A process as claimed in claim 14, in which the reduced gas prior to the oxidation of hydrogen sulphide to sulphur is dried to remove water.



16. A process as claimed in claim 14 or 15, in which elemental sulphur is removed from the gases after having contacted the first catalyst composition and H_2S remaining in the substantially sulphur-free gases is oxidized to sulphur dioxide by a process as
5 claimed in any one of claims 1 to 10.

17. A process as claimed in any one of claims 1 to 11, in which hydrogen sulphide is oxidized to sulphur dioxide, the hydrogen sulphide has been obtained by reducing Claus tail gases at a temperature above $175^{\circ}C$ with a free hydrogen- and/or carbon
10 monoxide-containing gas over a second catalyst composition comprising a sulphided metal from Group 6 and/or Group 8 of the Periodic Table of the Elements and an inorganic oxide carrier supporting the said sulphided metal(s), absorbing the H_2S present in the reduced gases followed by desorption of the H_2S
15 and recycle of the desorbed H_2S to the Claus plant, the hydrogen sulphide to be oxidized to sulphur dioxide remaining in the final off-gas from said absorption of H_2S .

18. A process as claimed in claim 1, substantially as herein-before described with reference to the Examples.

20 19. Elemental sulphur whenever produced by a process as claimed in any one of claims 1-8 and 12-18.
