#### COMMONWEALTH OF AUSTRALIA

Patents Act 1952-1969

### CONVENTION APPLICATION FOR A PATENT

(1) Here insert (in	XX(1) DRAGERWERK	AKTIENGESELLSCHA	T		
full) Name or Names of	we of Moislinger	Allee 53-55, D-24	400 Lubeck, Federal	Republic of	
Applicant or Applicants, followed by	Germany	•			
Address(es).					
(2) Here insert Title	hereby apply for the grant of	a Patent for an invent	ion entitled: <sup>(2)</sup>		
of Invention.	CO <sub>2</sub> ABSORBER MATERIAL	•••••	••••		
		•••••	•••••		
		••••••			
	which is described in the acco	mpanying complete sp	pecification. This applica	tions is a Covention	
(3) Here insert number(s)	application and is based on th				
of basic application(s).					
(4) Here insert Name of basic Country or Countries, and	for a patent or similar protection made in <sup>(4)</sup> Federal Republic of Germany on 14th December 1988				
basic date or dates.					
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		••••••	•••••••••••••••••••••••••••••••••••••••	•••••••	
	Our address for service is WATERMARK PATENT & TRADEMARK ATTORNEYS				
	290 Burwood Road, Hawthorn, Victoria, Australia.				
2					
	DATED this	2th day of	December	19.89	
(5) Signature(s) of Application)	13/12/89	(5) DRAGE	DRAGERWERK AKTIENGESELLSCHAFT		
		<b></b>	Mist. 1		
Seal of Company and Signatures of		by	your last	***************************************	
its Officers as prescribed by		Ia	n A. Scott		
its Articles of Association.		Regis	stered Patent Attorn	ıey	

To: THE COMMISSIONER OF PATENTS.

#### COMMONWEALTH OF AUSTRALIA

Patents Act 1952-1969

# DECLARATION IN SUPPORT OF A CONVENTION APPLICATION FOR A PATENT OR PATENT OF ADDITION

(1) Here insert (in full) Name of	In support of the Convention Application made by(1)			
Company.	DRÄGERWERK AKTIENGESELLSCHAFT			
	(hereinafter referred to as the applicant) for a Patent			
(2) Here insert title	for an invention entitled:(2)			
of Invention.	CO <sub>2</sub> ABSORBER MATERIAL.			
(3) Here	We k <sup>(3)</sup> Dr. Harald Helmreich, Dr. Peter Naumann			
insert full Name and Address,	Drägerwerk Aktiengesellschaft			
of Company official authorized	of Moistinger Atlee 53-55 D-2400 Lübeck			
to make declaration.	do solemnly and sincerely declare as follows:			
	1. We are authorised by the applicant for the patent			
	to make this declaration on its behalf.			
(4) Here	2. The basic application as defined by Section 141 of the Act was			
Country or	made in <sup>(4)</sup> Federal Republic of Germany			
followed by date or dates and basic	on the			
Applicant or Applicants.	DRÄGERWERK AKTIENGESELLSCHAFT			
	on the 18th day of August 19.89, by			
(5) Here insert (in full) Name	3. <sup>(5)</sup> please see enclosure			
and Address of Actual Inventor or				
Inventors.				
	is/are the actual inventors of the invention and the facts upon which the applicant			
	is entitled to make the application are as follow:			
	The applicant is the assignee ofthes.a.i.dac.t.ua.linventor.s			
	4. The basic application referred to in paragraph 2 of this Declaration			
	wasthe first application made in a Convention country in			
	respect of the invention the subject of the application.			
	DECLARED at Lübeck			
	this 18th day of August 19.89			
(6) Signature.	Dr. Harald Helmreich Dr. Peter Naumann			
	Dr. Harald Helmreich Dr. Peter Naumann To: The Commissioner of Patents.			

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#### Enclosure to Declaration (Form 8)

#### List of Inventors (P 38 42 048.1)

- 1. Dr. Carl Ernst van der Smissen
  Am Traveeck 30
  D-2400 Lübeck
  -German-
- 2. Kai vom Hofe Fleischhauerstraße 38 D-2400 Lübeck -German-
- 3. Dr. Herbert Röhl
  Emanuel-Geibel-Weg 5
  D-2067 Reinfeld
  -German-
- 4 Dr. Horst Wezurek Mühlenweg 12a D-2419 Ziethen -German-

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## (12) PATENT ABRIDGMENT (11) Document No. AU-B-46112/89 (19) AUSTRALIAN PATENT OFFICE (10) Acceptance No. 621171

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(71) Applicant(s)

DRAGERWERK AKTIENGESELLSCHAFT

(72) Inventor(s)
DR. CARL ERNST VAN DER SMISSEN; KAI VOM HOFE; DR. HERBERT ROHL; DR. HORST WEZUREK

(74) Attorney or Agent WATERMARK PATENT & TRADEMARK ATTORNEYS, Locked Bag 5, HAWTHORN VIC 3122

(57) The "respiratory lime" obtained has particularly good properties with respect to cohesiveness and CO<sub>2</sub> absorbing capacity if sodium hexametaphosphate or potassium orthophosphate are used as the salt. These materials are also favourable in terms of their price and in view of the fact that they are physiologically harmless.

Alkaline salts of other oxyacids such as sulphuric acid, permanganic acid, chromic acid and vanadic acid, as well as acidic alkaline salts of orthophosphoric acid, can also be used with similar success.

#### CLAIM

1. A CO<sub>2</sub> adsorbent mass comprising a kneadable mixture of calcium hydroxide and water and between 0.5 wt.% and 15 wt.% of an alkali salt of an oxyacid from the group consisting of alkaline metal hexametaphosphate and alkaline metal orthophosphate as an additive to increase the strength of the mixture.

621171

Form 10

## COMMONWEALTH OF AUSTRALIA PATENTS ACT 1952-69

### COMPLETE SPECIFICATION

(ORIGINAL)

	Class Int. Class	
Application Number:		
Lodged:		
Complete Specificatio	n Lodged	
	Accepted:	
	Published:	
Priority:		
Related Art :		
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4 K 9 D		
• ៧ឆ្នំme of Applicant :	URAGERWERK AKTIENGESELLSCHAFT	
Address of Applicant	Moislinger Allee 53–55, D–2400 Lubeck, Federal Republic of Germany	У
Actual Inventor :	CARL ERNST VAN DER SMISSEN, KAI VOM HOFE, HERBERT ROHL and HORST W	√EZUREK
Address for Service :	WATERMARK PATENT & TRADEMARK ATTORNEYS. X50 X12EN STREET, WEED VINE, X13EN X2EN X2EN X2EN X2EN X2EN X2EN X2EN X2	
Complete Specificatio	n for the invention entitled:	
	CO <sub>2</sub> ABSORBER MATERIAL	

us

#### CO<sub>2</sub> Absorber Material

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Absorber materials for CO2 are known. used for bonding exhaled CO2 in protective respiratory circulatory apparatus, and are called "respiratory "Respiratory lime" consists of calcium hy/roxide containing a certain proportion of water, usually 14 to 18%. For increasing reactivity, the known "respiratory lime" contains, at most, a few percent of sodium hydroxide or potassium hydroxide. order for it to be used in protective respiratory apparatus, the "respiratory lime" must be granulated. The particle size of the granules varies, depending on the intended use, from 2 mm to 8 mm. Very often fractured granules or shaped particles with diameters of about 4 mm are used. The known types of "respiratory lime" have generally good CO2-bonding properties. However, the granules tend to break when carried in mechanically stressed protective respiratory apparatus, e.g. in escape apparatus, even after a short time, because of their lack of cohesion, whereby the apparatus becomes extremely dusty and its functioning capacity is reduced. The use of the known types of "respiratory lime" in escape apparatus has therefore great risks.

Methods are known for increasing the cohesiveness of "respiratory lime". However, these methods always result in a marked reduction in reactivity. The use of alkali silicates as an additive to calcium hydroxide paste is one such method. Thus, liquid alkali silicates (water glass) effect a strengthening of the structure of the calcium hydroxide particles; however, its porosity is reduced and an outer layer of greatly reduced reactivity is formed on the particles. The result is that, for maintaining a sufficient CO<sub>2</sub> absorption performance, more "respiratory lime" must be used, whereby the protective respiratory apparatus

become larger, heavier and more bulky. In the particular case of an escape apparatus, this is undesirable.

The object of the present invention is to provide additive materials which increase the cohesiveness of "respiratory lime" particles, without impairing the CO<sub>2</sub> absorbing capacity of the "respiratory lime".

This is achieved in that 0.5% by weight to 15% by weight of an alkaline salt (i.e. alkali metal salt or alkaline earth metal salt) of an oxygen-containing acid (i.e. an "oxyacid") is added to the "respiratory lime" paste before granulating to give a kneadable mixture.

Thus, according to the present invention, there is provided a material for absorbing CO<sub>2</sub>, comprising a mixture of calcium hydroxide and water, the mixture containing from 0.5% by weight to 15% by weight of an alkaline salt of an oxyacid.

The "respiratory lime" obtained has particularly good properties with respect to cohesiveness and  $\rm CO_2$  absorbing capacity if sodium hexametaphosphate or potassium orthophosphate are used as the salt. These materials are also favourable in terms of their price and in view of the fact that they are physiologically harmless.

Alkaline salts of other oxyacids such as sulphuric acid, permanganic acid, chromic acid and vanadic acid, as well as acidic alkaline salts of orthophosphoric acid, can also be used with similar success.

The invention will now be illustrated by the following Examples.

#### Example 1

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1 kg of calcium oxide was stirred homogeneously with 2 litres of water containing 50 g of sodium hexametaphosphate. After cooling, the highly viscous pulp, in accordance with DE-C-2715635, was placed on a flexible band having hemispherical depressions therein,

and dried in an air drier to a water content of 16%. The "respiratory lime" was then discharged in the form of hemispheres. 600 ml of this "respiratory lime" fulfilled the requirements in respect of CO<sub>2</sub> absorbing capacity, of an absorber of a 15 minute escape apparatus even when the apparatus was previously shaken, corresponding to many years of use as escape apparatus.

#### Example 2

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1 kg of calcium oxide was stirred homogeneously with 2 litres of water containing 60 g of potassium orthophosphate. After cooling, the lime paste was extruded through 6 mm nozzles and then dried. The resulting fibrous "respiratory lime" was broken and sifted to obtain a fraction having a size of 2.5 mm to 4.0 mm. It was then humidified to a water content of 16%. The resulting fractured and granulated "respiratory lime" could be used in the same way as the hemispherical shaped lime described in Example 1.

#### THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

- 1. A CO<sub>2</sub> adsorbent mass comprising a kneadable mixture of calcium hydroxide and water and between 0.5 wt.% and 15 wt.% of an alkali salt of an oxyacid from the group consisting of alkaline metal hexametaphosphate and alkaline metal orthophosphate as an additive to increase the strength of the mixture.
- 2. A CO<sub>2</sub> adsorbent mass according to claim 1, wherein sodium hexametaphosphate is used as the alkali salt of oxyacid.
- 3. A CO<sub>2</sub> adsorbent mass according to claim 1, wherein potassium hexametaphosphate is used as the alkali salt of an oxyacid.
- 4. A CO<sub>2</sub> adsorbent mass according to claim 1, wherein sodium orthophosphate is used as the alkali salt of an oxyacid.
- 5. A CO<sub>2</sub> adsorbent mass according to claim 1, wherein a potassium orthophosphate is used as the alkali salt of an oxyacid.
- 6. ACO<sub>2</sub> adsorbent mass substantially as hereinbefore described with reference to the Examples.

Dated this 4th day of November, 1991.

DRAGERWERK AKTIENGESELLSCHAFT

WATERMARK PATENT & TRADEMARK ATTORNEYS FLOOR 2, "THE ATRIUM", 290 BURWOOD ROAD, HAWTHORN VICTORIA 3122, AUSTRALIA.

