AUSTRALIA Patents Act 1990 68170 P/00/001

## **PATENT REQUEST: STANDARD PATENT / PATENT OF ADDITION**

We, being the person identified below as the Applicant, request the grant of a patent to the person identified below as the Nominated Person, for an invention described in the accompanying standard complete specification, Full application details follow.

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(54) Invention Title	Gassed emulsion explosives					
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[31] Application Number	[33] Country	Country Code	[32] Date of Application			
93/6052	Republic of South Africa	ZA	18 November 1993			
93/9275	Republic of South Africa	ZA	10 December 1993			
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Frank Di Giantomasso **Registered Patent Attorney** for the Applicant

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P/00/008 Section 29(1) Regulation 3.1(2)

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### NOTICE OF ENTITLEMENT

(To be filed before acceptance)

We, Sasol Chemical Industries (Proprietary) Limited (Reg. No. 68/13914/07), of I Sturdee Avenue, Rosebank, Johannesburg 2196, Republic of South Africa, being the person nominated for the grant of the patent and being the applicant in respect of Application No. 77589/94, state the following:

The person nominated for the grant of the patent:

has entitlement from the actual inventors Mark Owen Delagey and Hermanus Joachim De Klerk by assignments.

The person nominated for the grant of the patent:

is the applicant of the basic applications listed on the patent request form.

'Ehe basic applications listed on the patent request form:

are the first applications made in a Convention country in respect of the invention.

:::: CHARLES JOSEPH GÖEDHALS SASOL CHEMICAL INDUSTRIES (PTY) LTD (REG. NO. 68/13914/07)

(Date)

(If the applicant is a Company or other legal entity, also indicate the name and standing of the authorized signatory.)

#### (12) PATENT ABRIDGMENT (11) Document No. AU-B-77589/94 (19) AUSTRALIAN PATENT OFFICE (10) Acceptance No. 681702 (54) Titia GASSED EMULSION EXPLOSIVES International Patent Classification(s) (51)<sup>6</sup> C06B 043/00 (21) Application No.: 77589/94 (22) Application Date: 02.11.94 (30) Priority Data (31) (33) Country Number (32) Date ZA SOUTH AFRICA 93/6052 18,11,93 93/9275 10.12.93 ZA SOUTH AFRICA Publication Date : 25.05.95 (43) (44) Publication Date of Accepted Application: 04.09.97 (71) Applicant(s) SASOL CHEMICAL INDUSTRIES (PROPRIETARY) LIMITED (72) inventor(s) MARK OWEN DELAGEY; HERMANUS JOACHIM DE KLERK (74) Attorney or Agent FREEHILLS PATENT ATTORNEYS, Level 47, 101 Collins Street, MELBOURNE VIC 3000 (57) Claim A PIBSA-based emulsion explosive including a phosphate ester (as herein defined) 1.

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- 16. A gassed emulsion explosive having the following constitution expressed in mass percentages: calcium nitrate 0.35%; PIBSA 0.1-2.0%; SMO 0.0-2.0%; phosphate ester (as herein defined) 0.1-2.0%; oil (or oil and wax) 2.5-20%; water 1-24%; gassing reagent 0.4-5% and the balance ammonium nitrate, and of which the pH is between 2.0 and 6.5.
- 17. A gassed emulsion explosive having the following constitution expressed in mass percentages: calcium nitrate 0-15%; PIBSA 0.1-1.2%; SMO 0.1-0.6%; phosphate ester (as herein defined) 0.1-0.7%; oil (or oil and wax) 6-9%; water 7-20%; gassing reagent 0.8-2% and the balance ammonium nitrate, and of which the pH is between 4.5 and 5.8.

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# ORIGINAL COMPLETE SPECIFICATION STANDARD PATENT

Invention Title:

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ENEMETE CONTROLS &

GASSED EMULSION EXPLOSIVES

The following statement is a full description of this invention, including the best method of performing it known to us:

#### **GASSED EMULSION EXPLOSIVES**

This invention relates to gassed emulsion explosives.

Emulsion explosives usually consist of a water-in-oil emulsion, containing oxidisers. Historically they were manufactured with the aid of emulsifiers based on sugar esters.
More recently the generic family of PIBSA (polyisobutylenesuccinicanhydride) based emulsifiers found application in this respect. PIBSA-based emulsifiers result in more stable emulsions, which can be made under a wide range of process parameters, and which exhibit the required shelf life.

It is common practice in the field of emulsion explosives to manipulate the density of the explosive by controlled gassing thereof. Gassing can be accomplished by chemical or mechanical means.

Chemical gassing can be based on a number of reactions, such as the reaction of a metal with a strong acid to form hydrogen, but the most commonly used reaction is the catalytic decomposition of sodium nitrite in an acid medium to form nitrogen.

5 PIBSA-based emulsifiers are generally able to react with the acid required for chemical gassing, usually with a deleterious effect on the emulsification properties of the emulsifier. The commonly used chemical gassing reaction can thus not be used to gas the known PIBSA-based explosive emulsions.

It is also known that porous ammonium nitrate prills (PPAN), which are used in an admixture with fuel oil to form ammonium-nitrate-fuel-oil explosives (ANFO), can be stabilised with naphthalene sulphonate against the thermal degradation of the prill at 32°C. Such a sulphonate is a surfactant which has an opposite emulsifying action to the emulsifiers commonly used in emulsion explosive technology. The customary admixture of ANFO, made from prills stabilised in this way, with emulsion explosives to form a product commonly known in the explosives art as "Heavy ANFO", results in the degradation of the emulsion part of the admixture and the failure of the explosive due to the incompatibility of the sulphonate and the emulsifier.

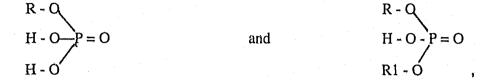
Accordingly, investigations have been carried out in respect of a PIBSA-based emulsion explosive with which on or more of the aforesaid problems can be overcome or at least minimised.

According to the invention, a PIBSA-based emulsion explosive is provided which includes a mixture of mono-esters and di-esters of orthophosphoric acid. Whenever used in this specification and the appended claims, the term "phosphate ester" is defined to mean a mixture of mono-esters and di-esters of orthophosphoric acid.

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Applicant has found that such an explosive can be gassed within a reasonable time (less than 30 minutes at 10°C), and that it remains stable for at least 28 days after being charged into a drill hole and kept there at ambient temperatures (-10°C to 40°C).

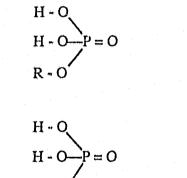
Preferably the mixture of esters includes compounds with the following configuration:



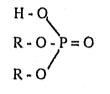
wherein R and R1 are carbon chains with from 6 to 18 carbon atoms, with R and R1 being the same or different.

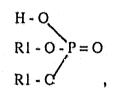
10 Applicant has found the most suitable mono-and di-esters to be those formed from alcohols wherein R is a carbon chain of from 7 to 9, and R1 from 10 to 18 carbon atoms. Most preferably R is a carbon chain of 8, and R1 of from C12 to C16 carbon atoms.

The following compounds may accordingly be present in a typical ester mixture according to the invention:



R1 - C







wherein R and R1 have the same values as given above.

25 Applicant has found that such a mixture may also contain C7, C9, C11 and C17 carbon chains as well as some unreacted orthophosphoric acid and phosphoric acid dimers.

The concentration of the phosphate ester in the total emulsion explosive is preferably between 0.02 and 2.0%, more preferably between 0.05 and 0.7% mass per mass.



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The phosphate ester can be added at any step or to any of the two phases of the water-in-oil emulsion explosive, but preferably the phosphate ester is dissolved in the oil phase of the explosive before emulsification.

Further according to the invention the explosive according to the invention may include an additional emulsifier to the main PIBSA-based emulsifier in a concentration less than that of the PIBSA-based emulsifier. This emulsifier, which acts as a co-emulsifier, preferably consists of a fatty acid ester of sorbitan, more specifically sorbitan mono-oleate (SMO).

Applicant has found the following with such an emulsion explosive containing a phosphate ester according to the invention:

1. If the phosphate ester is present in excess compared to the PIBSA-based emulsifier, it can act as an emulsifier in its own right, forming an oil-in-water emulsion but, when it is present in the preferred minor percentage, it acts as a co-emulsifier for the water-in-oil emulsion which comprises the emulsion explosive.

If a gassing solution is added to the emulsion explosive without the addition of supplementary acid, the resultant gas evolution is due to the presence of free orthophosphoric acid and dimers thereof, as well as to the mono-esters which contain free OH groups and, to a lesser extent, to the single OH group of the di-esters.

The rate of gassing of the emulsion explosive is acceptable down to a temperature of 10°C; the gassing to reduce the density of the explosive taking place within 30 minutes at a pH of 5. This pH is preferably obtained by the addition of either nitric acid or ammonium hydroxide to the aqueous oxidiser phase, but it can also be attained by using any other suitable acid or alkali.

The gassed emulsion explosive retains its explosive and rheological properties for at least 28 days after admixing with sulphonate stabilised PPAN prills and pumped into a borehole. This is referred to as a 28 day sleep time.

5. The gas bubbles are smaller and of a more uniform size distribution than in conventionally gassed, SMO based, emulsion explosives. The gas bubbles retain their size even after applying a standardised stirring test which leads to the collapse of bubbles in unstabilised emulsion explosives.

6. Stabilisation of the gas bubbles is independent of the method by which the bubbles have been produced. The method could be by chemical gassing as described, or by mechanical means, e.g. by beating in bubbles with an aerating stirring mechanism.



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7. The stabilisation effect is the most pronounced when the PIBSA has a molecular weight of between 500 and 1,300 and when the ester derivative is synthesised with tri-ethanolamine.

The gassing reagent is preferably based on the catalytic acid decomposition of sodium nitrite. It can consist of a mixture of sodium nitrite, urea or thiourea and water in suitable concentrations. In a non-limiting example the gassing reagent may include a mixture of a thiourea sodium nitrite and water in the mass ratio of 5:10:85.

The gassing reagent can be added to the explosive as an aqueous solution or as a water-in-oil emulsion. If the latter method is used, even more uniform gas bubbles are obtained. If this emulsion also contains the phosphate ester, the greatest uniformity and stabilisation of bubbles is obtained.

Applicant has also found that in so far as such gassing is concerned, the phosphate ester serves as acid catalyst for the gassing reaction from sodium nitrite and that it also serves as a buffer and source of acid to facilitate gassing at pH5, which is neutral for an ammonium nitrate solution.

Preferably the composition of the gassed emulsion explosive according to the invention, expressed in mass percentages, is as follows:

	GENERAL	PREFERRED	MOST PREFERRED	
Calcium nitrate	0 - 35%	0 - 15%	0 %	0%
PIBSA	0.1 - 2.0%	0.1 - 1.2%	0.69%	0.81%
SMO	0.0 - 2.0%	0.1 - 0.6%	0.30%	0.0%
Phosphate ester	0.1 - 2.0%	0.1 - 0.7%	0.31%	0.31%
Oil (or oil and wax)	2.5 - 20%	6 - 9%	6.8%	6.98%
Water	1 - 24%	7 - 20%	18.0%	18.0%
Gassing reagent	0.4 - 5%	0.8 - 2%	1.7%	1.7%
Ammonium nitrate	Balance	Balance	72.2%	72.2%

The pH range is generally from 2.0 - 6.5, preferably 3.6 - 5.8, and most preferably 5.

It has been found that the most preferred formulation given above gasses particularly well.

Applicant has also found that if the explosive composition includes PPAN, stabilised in Applicant known in the explosives and including naphthalene sulphonate stabilised

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PPAN to form heavy ANFO, such PPAN is compatible with the emulsifier being utilised, either with or without the addition of calcium nitrate to the oxidiser solution.

The invention accordingly also includes within its scope such an explosive composition including such PPAN, either on its own or with fuel oil to form ANFO.

5 The invention furthermore also includes within its scope a mixture of such a gassed emulsion explosive and 0-90% (mass percent) of either ammonium nitrate prills or ANFO made from ammonium nitrate prills and fuel oil in the typical ratio of 94:6 stabilised in any way known in the explosive art, to form gassed heavy ANFO.

When so admixed with 40% ANFO, the resultant heavy ANFO has a sleep time of at least 28 days.

The phosphate ester in this instance can be added to the fuel oil component of the ANFO.

It will be appreciated that the invention also includes within its scope a method of manufacturing gassed emulsion explosives and gassed heavy ANFO as described herein.

It will be appreciated further that there are no doubt many variations in detail possible with a gassed emulsion explosives and/or gassed heavy ANFO according to the invention without departing from the spirit and/or scope of the appended claims and all such variations are included in the scope of invention.



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

- 1. A PIBSA-based emulsion explosive including a phosphate ester (as herein defined)
- 2. The emulsion explosive of claim 1 wherein the phosphate ester includes compounds with the following configuration:

R - O H - O = Oand H - O

R - 0 H - 0 - P = 0

wherein R and R1 are carbon chains with from 6 to 18 carbon atoms, with R and R1 being the same or different.

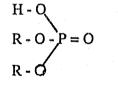
The emulsion explosive of claim 2 wherein the phosphate ester is formed from alcohols in which R is a carbon chain of from 7 to 9, preferably 8, carbon atoms and R1 is a carbon chain from 10 to 18, preferably 12 to 16, carbon atoms.

The emulsion explosive of claims 2 or 3 wherein the phosphate ester contains the following compounds:

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H - Q H - O P = OR - 0

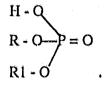
H-Q H - O ) = O R1 - C



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H-Q R1 - O -2 = 0R1 - 0



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The emulsion explosive of any one of the preceding claims wherein the concentration of the phosphate ester in the total emulsion explosive is from 0.02 to 2,0% mass/mass.

The emulsion explosive of any one of the preceding claims wherein the concentration of the phosphate ester in the total emulsion explosive is from 0.05 to 0.7% mass per mass.

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- 7. The emulsion explosive of any one of the preceding claims including an additional emulsifier to the main PIBSA-based emulsifier in a concentration less than that of the PIBSA-based emulsifier.
- 8. The emulsion explosive of claim 7 wherein the additional emulsifier consists of a fatty acid ester of sorbitan.
- 9. The emulsion explosive of claim 8 wherein the sorbitan ester includes sorbitan mono-oleate.

10. The emulsion explosive of any one of the preceding claims which is gassed.

11. The emulsion explosive of claim 10 wherein the gassing is effected chemically.

12. The emulsion explosive of claim 10 wherein the gassing is effected mechanically.

- 13. The emulsion explosive of any one of claims 1 to 9 including a gassing reagent which operates on the catalytic acid decomposition of sodium nitrite.
- 14. The emulsion explosive of claim 13 wherein the gassing reagent includes a mixture of sodium nitrite, urea or thiourea and water in a suitable concentration.

15. The emulsion explosive of claim 14 wherein the gassing reagent includes a mixture of a thiourea, sodium nitrite and water in the mass ratio of 5:10:85.

16. A gassed emulsion explosive having the following constitution expressed in mass percentages: calcium nitrate - 0.35%; PiBSA - 0.1-2.0%; SMO - 0.0-2.0%; phosphate ester (as herein defined) - 0.1-2.0%; oil (or oil and wax) - 2.5-20%; water - 1-24%; gassing reagent - 0.4-5% and the balance ammonium nitrate, and of which the pH is between 2.0 and 6.5.

- 17. A gassed emulsion explosive having the following constitution expressed in mass percentages: calcium nitrate 0-15%; PIBSA 0.1-1.2%; SMO 0.1-0.6%; phosphate ester (as herein defined) 0.1-0.7%; oil (or oil and wax) 6-9%; water 7-20%; gassing reagent 0.8-2% and the balance ammonium nitrate, and of which the pH is between 4.5 and 5.8.
- 18. A gassed emulsion explosive having the following constitution expressed in mass percentages: calcium nitrate 0%; PIBSA 0.69%; SMO 0.30%; phosphate ester (as herein defined) 0.31%; oil (or oil and wax) 6.8%; water 18.0%; gassing reagent 1.7% and ammonium nitrate 72.2%; and of which the pH is 5.
  - A gassed emulsion explosive having the following constitution expressed in mass percentages: calcium nitrate 0%; PIBSA 0.81%; SMO 0.0%; phosphate ester (as herein defined) 0.31%; oil (or oil and wax) 6.98%; water 18.0%; gassing reagent 1.7% and ammonium nitrate 72.2%; and of which the pH is 5.

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- 20. The gassed emulsion explosive of any one of claims 10 to 19 together with 0-90% (mass percent) of either ammonium nitrate prills or ANFO made from ammonium nitrate prills and fuel oil in the ratio of 94:6 to form heavy ANFO.
- 21. The gassed heavy ANFO of claim 20 of which the ammonium nitrate prills have been stabilised in any suitable manner known in the explosive art.
- 22. The gassed heavy ANFO of claim 20 including naphthalene sulphonate stabilised PPAN.
- 23. An emulsion explosive substantially as herein described with reference to the examples.

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24. A gassed emulsion explosive substantially as herein described with reference to the examples.

Sasol Chemical Industries (Proprietary) Limited

18 June 1997



#### ABSTRACT

A PIBSA-based emulsion explosive including mixture of mono-esters and di-esters of orthophosphoric acid, preferably made from long chain alcohols having from 6 to 18 carbon atoms. The explosive may be gassed chemically or mechanically and can be used in the manufacture of heavy ANFO.

