



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵ : C12N 9/54, C11D 3/386	A1	(11) International Publication Number: WO 93/18140 (43) International Publication Date: 16 September 1993 (16.09.93)
(21) International Application Number: PCT/DK93/00074 (22) International Filing Date: 3 March 1993 (03.03.93) (30) Priority data: 287/92 4 March 1992 (04.03.92) DK (71) Applicant (for all designated States except US): NOVO NORDISK A/S [DK/DK]; Novo Allé, DK-2880 Bagsvaerd (DK). (72) Inventors; and (75) Inventors/Applicants (for US only) : OUTTRUP, Helle [DK/DK]; Syvendehusvej 46, DK-2750 Ballerup (DK). DAMBMANN, Claus [DK/DK]; Hoeje Gladsaxe 61, 7. t.h., DK-2860 Soeborg (DK). AASLYNG, Dorrit, Anita [DK/DK]; Fyrren 8, Svogersley, DK-4000 Roskilde (DK). LINDEGAARD, Poul [DK/DK]; Oester Soegade 52, DK-2100 Copenhagen ø. (DK).		(74) Common Representative: NOVO NORDISK A/S; Patent Department, PeV, Novo Allé, DK-2880 Bagsvaerd (DK). (81) Designated States: FI, JP, KR, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i>
(54) Title: NOVEL PROTEASES (57) Abstract This invention is in the field of detergent proteases derived from strains of a new <i>Bacillus sp.</i>. More specifically, the invention is directed towards a protease derived from a strain of <i>Bacillus sp.</i> PD138. Moreover, the invention is directed towards a process for the preparation of the protease, the use of the protease as detergent enzyme, and detergent compositions comprising the protease of the invention.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	MR	Mauritania
AU	Australia	GA	Gabon	MW	Malawi
BB	Barbados	GB	United Kingdom	NL	Netherlands
BE	Belgium	GN	Guinea	NO	Norway
BF	Burkina Faso	GR	Greece	NZ	New Zealand
BG	Bulgaria	HU	Hungary	PL	Poland
BJ	Benin	IE	Ireland	PT	Portugal
BR	Brazil	IT	Italy	RO	Romania
CA	Canada	JP	Japan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SK	Slovak Republic
CI	Côte d'Ivoire	LJ	Liechtenstein	SN	Senegal
CM	Cameroon	LK	Sri Lanka	SU	Soviet Union
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	MC	Monaco	TG	Togo
DE	Germany	MG	Madagascar	UA	Ukraine
DK	Denmark	ML	Mali	US	United States of America
ES	Spain	MN	Mongolia	VN	Viet Nam
FI	Finland				

NOVEL PROTEASES

TECHNICAL FIELD

This invention is in the field of detergent proteases derived from strains of *Bacillus* sp. More specifically, the invention is directed towards a novel alkaline
5 protease derived from a strain of *Bacillus* sp. PD138. Moreover, the invention is directed towards a process for the preparation of the protease, the use of the protease as detergent enzyme, and detergent compositions comprising the protease of the invention.

BACKGROUND ART

10 Detergent enzymes have been marketed for more than 20 years and are now well established as normal detergent ingredients in both powder and liquid detergents all over the world. With the trend towards lower temperature washing, detergent enzyme consumption has increased during late years. Enzymes used in washing formulations comprise proteases, lipases, amylases, cellulases, as well as
15 other enzymes, or mixtures hereof. Commercially most important are proteases.

Detergent proteases have been developed by isolation of proteases found in nature followed by testing in detergent formulations. Most detergent proteases are obtained from members of the genus *Bacillus*. Currently new types of proteases enter the market, offering the possibility of giving a better cost/per-
20 formance ratio at various specified conditions.

Examples of commercial protease products are ALCALASE™, ESPERASE™ and SAVINASE™, all supplied by Novo Nordisk A/S, Denmark. These and similar enzyme products from other commercial sources are active in detergent solutions, i.e. at pH values in the range of from 8 to 11 and in the presence of
25 sequestering agents, surfactants and bleaching agents such as sodium borate. The ALCALASE™ protease is produced by strains of the species *Bacillus licheniformis*. The ESPERASE™ and SAVINASE™ proteases are obtained by cultivation of strains of alkalophilic *Bacilli*.

SUMMARY OF THE INVENTION

It is an object of the present invention to provide novel detergent proteases with improved washing performance at high pH and high ion strength.

Accordingly, in its first aspect, the invention provides a protease having
5 an apparent molecular weight of 28 kD, pI above 9.5, pH optimum at pH 11 or above (at 25°C), temperature optimum in the range of from 45 to 55°C (at pH 9.5), and immunochemical properties identical or partially identical to those of a protease derived from the strain *Bacillus sp.* PD138, NCIMB 40338.

In a second aspect, the invention relates to an isolated biologically pure
10 culture of a strain of *Bacillus sp.* represented by the strain *Bacillus sp.* PD138. In a more specific aspect, the invention relates to a strain of *Bacillus sp.* PD138, NCIMB 40338, or a mutant or a variant thereof.

In a third aspect, the invention provides a process for the preparation of the protease, which process comprises cultivation of a protease producing strain
15 of *Bacillus sp.* PD138 in a suitable nutrient medium, containing carbon and nitrogen sources and inorganic salts, followed by recovery of the desired enzyme. In a more specific aspect, *Bacillus sp.* PD138, NCIMB 40338, or a mutant or a variant thereof, or another host organism carrying the gene encoding a protease having immunochemical properties identical or partially identical to those of the protease
20 derived from *Bacillus sp.* PD138, is cultivated.

In a fourth aspect, the use of the enzyme as detergent enzyme is claimed. In more specific aspects, the invention provides detergent compositions and detergent additives comprising the protease.

In a fifth aspect, the invention provides a washing process comprising
25 addition of the protease.

BRIEF DESCRIPTION OF DRAWINGS

The present invention is further illustrated by reference to the accompanying drawings, in which

Fig. 1 shows the relation between temperature and the proteolytic activity of an enzyme according to the invention (the enzyme preparation obtained according to Ex. 1, with 2% of casein as substrate and at pH 9.5; ■ Buffer, □ Buffer + 0.1% of STPP).

5 Fig. 2 shows the relation between pH and the proteolytic activity of an enzyme according to the invention (the enzyme preparation obtained according to Ex. 1, with 1% of casein as substrate and at 25°C).

DETAILED DISCLOSURE OF THE INVENTION

The Microorganism

10 The novel microorganism of the invention, able to produce an enzyme of the invention, is represented by the strain that was isolated from a sample of Jamaican soil. *Bacillus* sp. PD138, has been deposited according to the Budapest Treaty on the International Recognition of the Deposits of Microorganisms for the Purpose of Patent Procedures, on 3 December 1990 at National Collections of
15 Industrial and Marine Bacteria Ltd. (NCIMB), 23 St. Machar Drive, Aberdeen AB2 1RY, Scotland, UK, under Accession No. NCIMB 40338.

The microorganism of this invention is an aerobic, spore forming bacterium belonging to the genus *Bacillus*. Morphologically it can be described as motile rods with a diameter of 0.7 - 0.9 micron, and a length of 2 - 3 micron. The
20 spores are round to ellipsoid, not swelling the sporangium, central to subterminal. Optimum temperature for growth is within 25 - 37°C, and optimal pH for growth is within 7 - 9, no growth at pH 9.7, and no growth at 50°C. The microorganism forms yellowish to yellow colonies, round and smooth, on nutrient agar slants, and no diffusion of pigment into the agar is observed.

25 Cultivation of the Microorganism

The microorganism of the invention can be cultivated under aerobic conditions in a nutrient medium containing assimilable carbon and nitrogen together with other essential nutrients, the medium being composed in accordance with the principles of the known art.

Suitable carbon sources are carbohydrates such as sucrose, glucose and starch, or carbohydrate containing materials such as cereal grain, malt, rice and sorghum. The carbohydrate concentration incorporated in the medium may vary widely, e.g. up to 25% and down to 1 - 5%, but usually 8 - 10% will be suitable, the 5 percentages being calculated as equivalents of glucose.

The nitrogen source in the nutrient medium may be of inorganic and/or organic nature. Suitable inorganic nitrogen sources are nitrates and ammonium salts. Among the organic nitrogen sources quite a number are used regularly in fermentation processes involving the cultivation of bacteria. Illustrative examples are soybean 10 meal, cotton seed meal, peanut meal, casein, corn, corn steep liquor, yeast extract, urea, and albumin. In addition, the nutrient medium should also contain usual trace substances.

The novel *Bacillus* species of this invention are slightly alkalophilic. Therefore, the cultivation is preferably conducted at slightly alkaline pH values, which 15 can be obtained by addition of suitable buffers such as sodium bicarbonate, pH 9.0, after sterilization of the growth medium. For cultivation in tank fermentors it is necessary to use artificial aeration. The rate of aeration is similar to that used in conventional tank fermentation.

After fermentation, liquid enzyme concentrates may be produced by 20 removal of coarse material from the broth or, if desired, concentration of the broth by evaporation at low temperature or by reverse osmosis. Finally, preservatives may be added to the concentrate.

Solid enzyme preparations may be prepared from the purified and/or concentrated broth by precipitation with salts, such as Na_2SO_4 or water-miscible 25 solvents, such as ethanol or acetone. Removal of the water in the broth by suitable drying methods, such as spray-drying, may also be employed.

Assay for Proteolytic Activity

The proteolytic activity is determined with casein as substrate. One Casein Protease Unit (CPU) is defined as the amount of enzyme liberating 1 mM of 30 primary amino groups (determined by comparison with a serine standard) per minute under standard conditions, i.e. incubation for 30 minutes at 25°C and pH 9.5.

A folder AF 228, describing the analytical method, is available upon request to Novo Nordisk A/S, Denmark, which folder is hereby included by reference.

The Enzyme

The enzyme of the invention is a novel detergent protease. It is an
5 alkaline protease, obtainable by cultivation of a microorganism of the invention, preferably *Bacillus* sp. PD138, NCIMB 40338, or a mutant or a variant thereof, in a suitable nutrient medium, containing carbon and nitrogen sources and inorganic salts. The enzyme can also be obtained by recombinant DNA-technology.

The protease of the invention can be described by the following
10 characteristics.

Physical-Chemical Properties

A molecular weight of 28 kD, determined by SDS-PAGE. A pI above
9.5 could not be accurately determined by isoelectric focusing on LKB Ampholine®
PAG plates. The protease activity is inhibited by PMSF, α -1-antitrypsin, and Turkey-
15 egg-white proteinase inhibitor. EDTA and soybean-protein inhibitor do not influence the protease activity.

The temperature activity relationship was determined with 2% of casein
as substrate and at pH 9.5 in the presence (white squares) and absence (black
squares) of 0.1% of sodium tripolyphosphate (STPP, a common ingredient in many
20 commercial detergents). The assay for proteolytic activity described previously was used with the modification that the incubation temperature was varied in the interval of from 15 to 70°C.

The result is shown in Fig. 1. It appears from the figure that the enzyme
possesses proteolytic activity from temperatures below 15°C to above 70°C, and has
25 a temperature optimum within the range of 45 to 55°C, around 50°C.

The dependence of activity on pH was determined by the same
procedure using buffers adjusted to predetermined pH values in the pH range of
from 6 to 11.

The result is shown in Fig. 2. It appears from this figure that the enzyme possesses proteolytic activity at pH values below 6 to above 11, with a pH optimum above pH 10, more specifically with a pH optimum of at least pH 11.

Furthermore, it was found that the protease of the invention is stable
5 for 60 minutes at 25°C under washing conditions, in European type and American type detergents.

The protease of the invention possesses especial potentials in detergents with high ionic strength and high pH. From the wash performance test, cf. Ex. 2, during wash at pH 11 in 9° dH (German Hardness) water, a washing
10 performance far superior to that of the alkaline Bacillus protease SAVINASE™ (Novo Nordisk A/S, Denmark) appeared.

Immunochemical Properties

The protease of the invention has immunochemical properties identical or partially identical (i.e. at least partially identical) to those of a protease derived
15 from the strain Bacillus sp. PD138, NCIMB 40338.

The immunochemical properties can be determined immunologically by cross-reaction identity tests. The identity tests can be performed by the well-known Ouchterlony double immunodiffusion procedure or by tandem crossed immunoelectrophoresis according to *N. H. Axelsen*; Handbook of Immuno-
20 precipitation-in-Gel Techniques; Blackwell Scientific Publications (1983), chapters 5 and 14. The terms "antigenic identity" and "partial antigenic identity" are described in the same book, chapters 5, 19 and 20.

Monospecific antiserum was generated according to the above mentioned method by immunizing rabbits with the purified protease of the invention.
25 The immunogen was mixed with Freund's adjuvant and injected subcutaneously into rabbits every second week. Antiserum was obtained after a total immunization period of 8 weeks, and immunoglobulin was prepared therefrom as described by *N. H. Axelsen, supra*.

Ouchterlony double immunodiffusion tests showed no cross reaction
30 between the protease of the invention and the known alkaline serine proteases

ALCALASE™, SAVINASE™, ESPERASE™, subtilisin Novo (available from Novo Nordisk A/S), and KAZUSASE™ (available from SHOWA DENKO).

Detergent Compositions

According to the the invention, the protease may typically be added as
5 a component of a detergent composition. As such, it may be included in the detergent composition in the form of a detergent additive. The detergent composition as well as the detergent additive may additionally comprise one or more other enzymes conventionally used in detergents, such as lipases, amylases, cellulases, oxidases or peroxidases.

10 In a specific aspect, the invention provides a detergent additive. The enzymes may be included in a detergent composition by adding separate additives containing one or more enzymes, or by adding a combined additive comprising all of these enzymes.

Preferably, the detergent additive, i.e. a separated additive or a
15 combined additive, is provided in the form of a granulate, preferably a non-dusting granulate, a liquid, in particular a stabilized liquid, a slurry, or in a protected form.

Dust free granulates may be produced, e.g. as disclosed in US 4,-
106,991 and US 4,661,452 (both to Novo Industri A/S) and may optionally be coated
by methods known in the art. The detergent enzymes may be mixed before or after
20 granulation.

Liquid enzyme preparations may, for instance, be stabilized by adding
a polyol such as propylene glycol, a sugar or sugar alcohol, lactic acid or boric acid,
according to established methods. Other enzyme stabilizers are well known in the
art.

25 Protected enzymes may be prepared according to the method disclosed in EP 238,216 A.

The detergent composition of the invention may be in any convenient form, e.g. as powder, granules or liquid. A liquid detergent may be aqueous, typically containing up to 90% of water and 0-20% of organic solvent.

The detergent composition comprises a surfactant which may be anionic, non-ionic, cationic, amphoteric or a mixture of these types. The detergent will usually contain 0 -50% of anionic surfactant such as linear alkyl benzene sulphonate (LAS), alpha-olefin sulphonate (AOS), alkyl sulphate (AS), alcohol ethoxy sulphate (AES) or soap. It may also contain 0 - 40% of non-ionic surfactant such as nonyl phenol ethoxylate or alcohol ethoxylate. Furthermore, it may contain a polyhydroxy fatty acid amide surfactant (e.g. as described in WO 92/06154).

The pH (measured in aqueous detergent solution) will usually be neutral or alkaline, e.g. 7 - 11. The detergent may contain 1 - 40% of a detergent builder such as zeolite, phosphate, phosphonate, citrate, NTA, EDTA or DTPA, alkenyl succinic anhydride, or silicate, or it may be unbuilt (i.e. essentially free of a detergent builder). It may also contain other conventional detergent ingredients, e.g. fabric conditioners, foam boosters, bleaching agents, e.g. perborate, percarbonate, tetraacetyl ethylene diamine (TAED), or nonanoyloxybenzene sulfonate (NOBS), anti-corrosion agents, soil-suspending agents, sequestering agents, anti-soil redeposition agents, stabilizing agents for the enzyme(s), foam depressors, dyes, bactericides, optical brighteners or perfumes.

Particular forms of detergent composition within the scope of the invention include:

a) A detergent composition formulated as a detergent powder containing phosphate builder, anionic surfactant, nonionic surfactant, silicate, alkali to adjust to desired pH in use, and neutral inorganic salt.

b) A detergent composition formulated as a detergent powder containing zeolite builder, anionic surfactant, nonionic surfactant, acrylic or equivalent polymer, silicate, alkali to adjust to desired pH in use, and neutral inorganic salt.

c) A detergent composition formulated as an aqueous detergent liquid comprising anionic surfactant, nonionic surfactant, humectant, organic acid, caustic alkali, with a pH in use adjusted to a value between 7 and 10.5.

d) A detergent composition formulated as a nonaqueous detergent liquid comprising a liquid nonionic surfactant consisting essentially of linear alkoxylated primary alcohol, phosphate builder, caustic alkali, with a pH in use adjusted to a value between about 7 and 10.5.

e) A detergent composition formulated as a detergent powder in the form of a granulate having a bulk density of at least 600 g/l, containing anionic surfactant and nonionic surfactant, low or substantially zero neutral inorganic salt, phosphate builder, and sodium silicate.

5 f) A detergent composition formulated as a detergent powder in the form of a granulate having a bulk density of at least 600 g/l, containing anionic surfactant and nonionic surfactant, low or substantially zero neutral inorganic salt, zeolite builder, and sodium silicate.

g) A detergent composition formulated as a detergent powder containing
10 ing anionic surfactant, nonionic surfactant, acrylic polymer, fatty acid soap, sodium carbonate, sodium sulphate, clay particles, and sodium silicate.

h) A liquid compact detergent comprising 5 - 65% by weight of surfactant, 0 - 50% by weight of builder and 0 - 30% by weight of electrolyte.

It is at present contemplated that, in the detergent composition of the
15 invention, the [protease] may be added in an amount corresponding to [0.001 - 100] mg of enzyme per liter of wash liquor.

The invention is further illustrated in the following examples, which are not intended to be in any way limiting to the scope of the invention as claimed.

EXAMPLE 1

20 *Bacillus* sp. PD138, NCIMB 40338, was cultivated at 25°C on a rotary shaking table (300 r.p.m.) in 500 ml baffled Erlenmeyer flasks containing 100 ml of medium of the following composition (per litre):

25	Potato starch	100 g
	Ground barley	50 g
	Soybean flour	20 g
	Na ₂ HPO ₄ x 12 H ₂ O	9 g
	Pluronic®	0.1 g
	Sodium caseinate	10 g

The starch in the medium is liquified with α -amylase, and the medium is sterilized by heating at 120°C for 45 minutes.

After sterilization the pH of the medium is adjusted to 9.0 by addition of 10 ml of a 1 M solution of sodium bicarbonate.

5 After 4 days of incubation the proteolytic activity of the culture was determined using the method described above.

After cultivation, the enzyme activity of the broth was 20 CPU/l.

After separation of the solid material the protease was purified by a conventional chromatographic method.

10 Yield from 1 l of culture broth was 50 ml with 100 CPU/l. Purity was more than 90% as judged by SDS-PAGE.

The characteristics of the preparation prepared in accordance with this Example have been referred to earlier in this specification, and reference is made hereto.

15

EXAMPLE 2

Wash Performance

The wash performance tests were accomplished on grass soiled cotton, at 40°C, isothermally for 10 minutes.

The tests were performed at enzyme concentrations of 0.01, 0.025, 20 0.05, 0.1, 0.5, and 1.0 mg of enzyme protein per litre.

5.0 g/l of a commercial European compact powder detergent were used. The detergent was dissolved in approx. 9°dH (German Hardness) water, and pH was adjusted to 11. The textile/wash liquor ratio was 6 g of textile per litre of wash liquor.

25 Subsequent to washing, the cloths were flushed in running tap water and air-dried. The remission (%R) at 460 nm was determined.

As a measure of the wash performance differential remission, ΔR , was used being equal to the remission after wash with enzyme added, minus the remission after wash with no enzyme added.

30 The results of these tests are shown in Table 1 below (mean of 2 tests).

Table 1

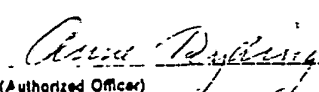
	Enzyme Concentration (mg enzyme protein/litre)	delta R	
		PD138	SAVINASE™
5	0.01	5.7	3.2
	0.025	13.5	-
	0.05	19.4	8.1
	0.10	22.0	14.8
	0.50	22.6	22.0
10	1.0	-	22.2

(- not done)

The differential remission values show that the protease of the invention has especial potentials in detergents with high ionic strength and high pH, where it
 15 possesses good washability.

International Application No: PCT/

/

MICROORGANISMS	
Optional Sheet in connection with the microorganism referred to on page <u>1</u> , line <u>3</u> of the description ¹	
A. IDENTIFICATION OF DEPOSIT ¹	
Further deposits are identified on an additional sheet ☒ ²	
Name of depositary institution ⁴ NATIONAL COLLECTIONS OF INDUSTRIAL & MARINE BACTERIA LTD.	
Address of depositary institution (including postal code and country) ⁴ 23 St. Machar Drive, Aberdeen AB2 1RY, Scotland	
Date of deposit ⁵ 3 December 1990	Accession Number ⁶ NCIMB 40338
B. ADDITIONAL INDICATIONS ⁷ (leave blank if not applicable). This information is continued on a separate attached sheet ☐	
In respect of those designations in which a European patent is sought, a sample of the deposited micro-organism will be made available only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC) until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or is deemed to be withdrawn.	
C. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE ⁸ (if the indications are not for all designated States)	
D. SEPARATE FURNISHING OF INDICATIONS ⁹ (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later ⁹ (Specify the general nature of the indications e.g., "Accession Number of Deposit")	
E. ☒ This sheet was received with the international application when filed (to be checked by the receiving Office)	
 (Authorized Officer)	
☐ The date of receipt (from the applicant) by the International Bureau ¹⁰	
 (Authorized Officer)	

CLAIMS

1. A protease characterized by having the following properties:
 - (a) an apparent molecular weight of 28 kD;
 - (b) pI above 9.5;
 - 5 (c) pH optimum above pH 10 (at 25°C and with casein as substrate);
 - (d) temperature optimum in the range of from 45 to 55°C (at pH 9.5 and with casein as substrate);
 - (e) immunochemical properties identical or partially identical to those of a protease derived from the strain *Bacillus sp.* PD138, NCIMB 40338.
- 10 2. A protease according to claim 1, the protease being obtainable from a strain of *Bacillus sp.*, represented by the strain *Bacillus sp.* PD138, or from another host organism carrying the gene encoding a protease having immunochemical properties identical or partially identical to those of the protease derived from the strain *Bacillus sp.* PD138, NCIMB 40338.
- 15 3. The protease of either of claims 1 - 2, being obtainable from the strain *Bacillus sp.* PD138, NCIMB 40338, or a mutant or a variant thereof.
4. An isolated biologically pure culture of a strain of *Bacillus sp.*, represented by the strain *Bacillus sp.* PD138, NCIMB 40338.
5. A culture according to claim 4, the strain being *Bacillus sp.* PD138,
20 NCIMB 40338, or a mutant or a variant thereof.
6. A process for the preparation of a protease according to any of claims 1 - 3, which process comprises cultivation of a protease producing strain of *Bacillus sp.* according to either of claims 4 - 5, in a suitable nutrient medium, containing carbon and nitrogen sources and inorganic salts, followed by recovery
25 of the desired enzyme.

7. A process according to claim 6, in which *Bacillus sp.* PD138, NCIMB 40338, or a mutant or a variant thereof, or another host organism carrying the gene encoding a protease having immunochemical properties identical or partially identical to those of the protease derived from *Bacillus sp.* PD138, is cultivated.

5 8. The use of a protease according to any of claims 1 - 3 as a detergent enzyme.

9. A detergent composition comprising a protease according to any of claims 1 - 3.

10 10. A detergent composition according to claim 9, which further comprises one or more other enzymes, in particular an amylase, a lipase, a cellulase, a peroxidase, and an oxidase.

11. A detergent additive comprising a protease according to any of claims 1 - 3, provided in the form of a non-dusting granulate, a liquid, in particular a stabilized liquid, a slurry, or a protected enzyme.

15 12. A washing process comprising addition of a protease according to any of claims 1 - 3.

13. A washing process according to claim 12, comprising addition of a detergent composition according to either of claims 9 - 10.

20 14. A washing process according to claim 12, comprising addition of a detergent additive according to claim 11.

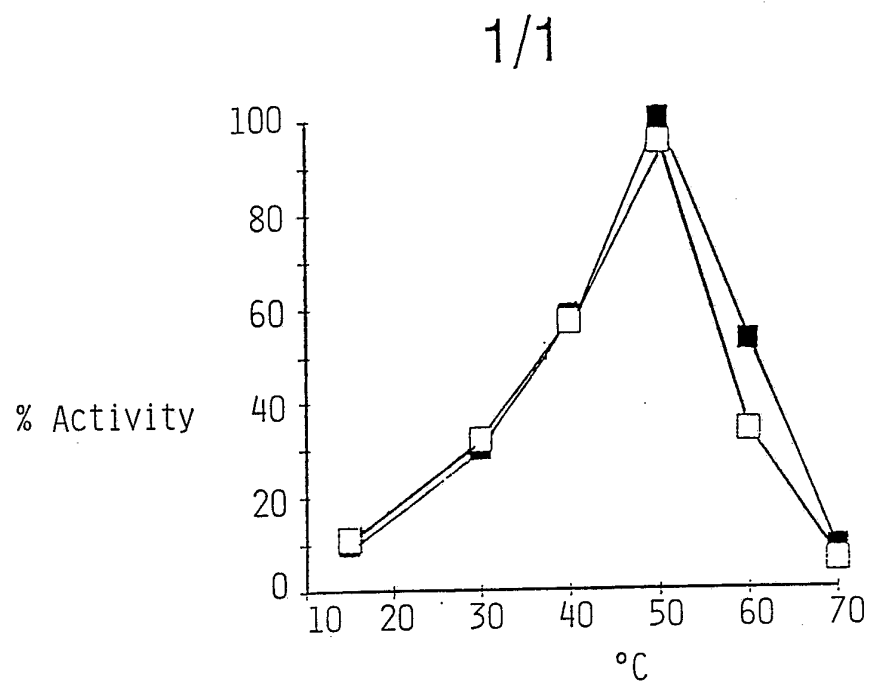


Fig. 1

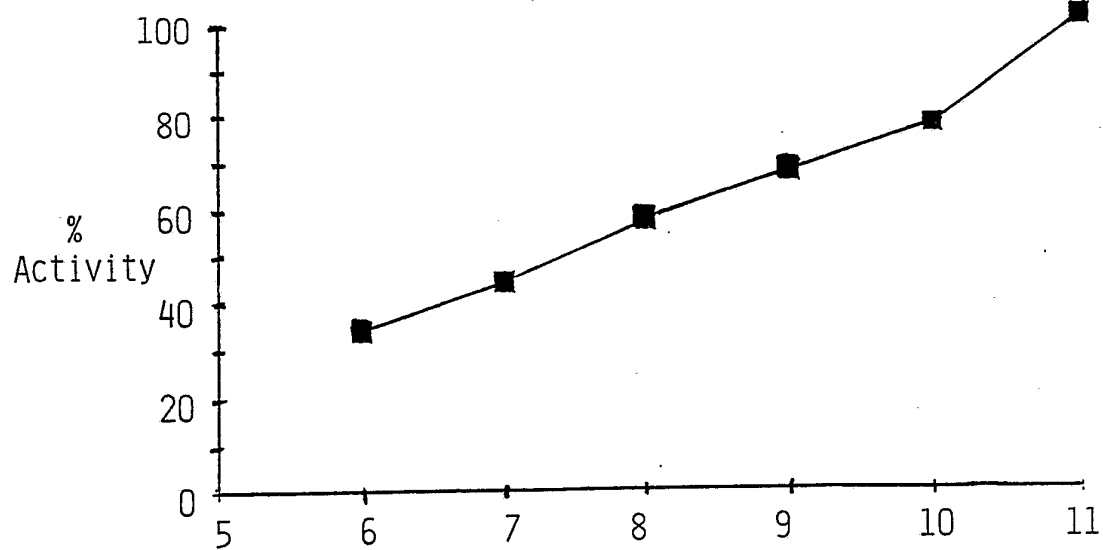


Fig. 2

INTERNATIONAL SEARCH REPORT

International application No.

PCT/DK 93/00074

A. CLASSIFICATION OF SUBJECT MATTER

IPC5: C12N 9/54, C11D 3/386

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC5: C12N, C11D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	EP, A1, 0495401 (KAO CORPORATION), 22 July 1992 (22.07.92), claim 1 --	1-14
X	US, A, 4764470 (GENEX CORPORATION), 16 August 1988 (16.08.88), claim 6 --	1-14
X	US, A, 4480037 (SHOWA DENKO KABUSHIKI KAISHA), 30 October 1984 (30.10.84), claim 1 --	1-14
X	DE, C2, 3328619 (SHOWA DENKO K.K.), 28 February 1985 (28.02.85), claim 1 --	1-14

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

11 June 1993

Date of mailing of the international search report

16 -06- 1993

Name and mailing address of the ISA/
Swedish Patent Office
Box 5055, S-102 42 STOCKHOLM
Facsimile No. +46 8 666 02 86

Authorized officer

Yvonne Siösteen
Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/DK 93/00074

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US, A, 4771003 (GENEX CORPORATION), 13 Sept 1988 (13.09.88), claim 7 -- -----	1-14

INTERNATIONAL SEARCH REPORT

Information on patent family members

30/04/93

International application No.

PCT/DK 93/00074

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP-A1- 0495401	22/07/92	JP-A- 4349882 JP-A- 4281782	04/12/92 07/10/92
US-A- 4764470	16/08/88	EP-A- 0232169	12/08/87
US-A- 4480037	30/10/84	AU-B- 556045 AU-A- 1097083 CA-A- 1193213 FR-A,B- 2521163 GB-A,B- 2116561 JP-C- 1614949 JP-A- 58134990 JP-B- 60055118	23/10/86 18/08/83 10/09/85 12/08/83 28/09/83 15/08/91 11/08/83 03/12/85
DE-C2- 3328619	28/02/85	BE-A- 897479	01/12/83
US-A- 4771003	13/09/88	EP-A- 0220921	06/05/87