



**Notification of an active substance
according to Article 4 and Article 5 (2) (a)
of Commission Regulation (EC) No. 1112/2002**

Basic Notification, Part 1

Reference No SIC-IT-001

1 Identification data on the notifier

1.1 Manufacturer of the active substance *as defined in Article 2 (b)*

1.1.1 Name SICIT 2000 S.p.A.
1.1.2 Address Via Arzignano 80
I-36072 CHIAMPO (VI)

1.1.3 Location of plant Via Arzignano 80
I-36072 CHIAMPO(VI)

1.2 Producer of the active substance *as defined in Article 2 (a)*

1.2.1 Name SICIT 2000 S.p.A.
1.2.2 Address Via Arzignano 80
I-36072 CHIAMPO (VI)

1.2.3 Telephone No ++39 444 623132
1.2.4 Telefax No ++39 444 625903
1.2.5 E-mail Address sicitspa@tin.it

1.2.6	Contact Address	Massimo Neresini SICIT 2000 S.p.A. Via Arzignano 80 I-36072 CHIAMPO (VI)
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1.2.7	Telefon No	++39 444 623132
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1.2.8	E-mail Address	sicitspa@tin.it
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1.2.9	Alternative	Oreste Odellii
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1.2.10	Telefon No	++39 444 623132
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1.2.11	E-mail Address	sicitspa@tin.it
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2 Information to facilitate identification of the active substance

2.1.1	Common name (proposed or ISO-accepted)	HYDROLYSED PROTEINS
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2.1.2	Variant(s) (where relevant)	None
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2.1.3	Synonyms of common name	None
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2.1.4	Species (for microorganism)	Not applicable
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2.1.5	Subspecies (where relevant)	Not applicable
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2.2.1	Chemical name (IUPAC nomenclature)	Not applicable
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2.2.2	Chemical name (CAS nomenclature)	Not applicable
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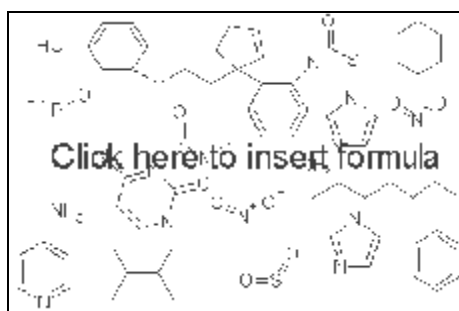
2.3.1	CAS number	Not applicable
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2.3.2	CIPAC number	Not applicable
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2.3.3	EEC number	Not applicable
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2.4.1	Empirical formula	Not applicable
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2.4.2	Structural formula	
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2.4.3	Molecular mass	Not applicable
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2.5	Any other information, e.g. example method of manufacture/extraction or origin of materials from which the substance is manufactured	
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The hydrolysed proteins are obtained from cutaneous and subcutaneous hides tissues. The raw material is hydrolysed in acid and/or alkaline conditions. During the manufacturing process there is the sterilization treatment in the conditions temperature > 133°C, time > 20 minutes and pressure > 3 bar. In addition the manufacturing provides for the refining, the filtering and finally the concentration.

2.6	Specification of purity of the a. s.	The active substance is characterized by the content in organic nitrogen that derives only by aminoacids and peptides and so the organic nitrogen is indicating the purity. The content of the active substance expressed as Norgx6,25 (conventional factor) as minimum is 240 g/kg
2.7	Classification and labelling of the a. s.	
2.7.1	Hazard symbols	Non toxic-Non oral noxious-Non topic noxious, Non irritant for eyes and skin
2.7.2	Risk phrases	None
2.7.3	Safety phrases	Obey reasonable safety precautions and practise according to good manufacturing procedures. In case of accidental release, collect the product for the re-use how much as possible and limitate the pouring area. The product is completely degradable but if present in copious quantities can pollute ground and surface water: It is necessary to prevent concentrated product from penetrating into ground and surface waters. The product is used as fertilizer and consequently does not cause negative transformations in the environment if it is used following the suggested dosages and the suggested conditions.

3. Further information

- 3.1 For each Member State a list of crops/uses for which plant protection products containing the active substance are currently authorised or used.

Please refer to BN-2

file name:

SICIT 001 BN2

HYDROLYSED PROTEINS

- 3.2 Further information on the active substance as set out in Annex II to the Directive points 3.1 to 3.5.

Please refer to BN-3

file name:

SICIT 001 BN3

HYDROLYSED PROTEINS

- 3.3 Date and reference number of the most recent review of the active substance in a Member State of the European Union:

Date:

Reference No:

Country:

- 3.4 Date and reference number of the most recent review of the active substance in an OECD-Country:

Date:

Reference No:

Country:

4 Undertaking

- 4.1 The notifier undertakes to submit to the designated coordinating authority of the designated rapporteur Member State the dossiers within the deadline provided for in the Regulation to be adopted according to Article 6 (4) of this Regulation. Whenever the newly adopted Regulation mentions several notifiers for this active substance, the notifier agrees to make all reasonable efforts to present a single dossier collectively with the other notifiers.
- 4.2 The notifier undertakes to pay a fee as provided for in Article 7 at the time of the submission of the notification to the body designated in Annex V.
- 4.3 The notifier declares that he is aware that he will be charged a fee by Member States at the time of the submission of the full dossier for active substances covered by the Regulation meant in Article 6 (4).
- 4.4 The notifier confirms that the above information is honest and correct.
- 4.5 The notifier declares that an authorisation by the manufacturer to act as his sole representative for the purpose of complying with this Regulation is enclosed if necessary.

Signature *(of the person competent to act for the company mentioned under 1.1)*

Name: Massimo Neresini

Date: 29/10/02

Payment

Effected by transfer: Yes

Paid by: SICIT 2000 S.p.A.

Reference: BBA-RENDER-4;SIC-IT-001

Amount € 750

Date of transfer: 29-10-02