



Product Service

# EC Certificate

## Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in Class IIa, IIb or III)

No. G2 17 07 53618 024

**Manufacturer:** ZHERMACK S.p.A

Via Bovazecchino, 100  
45021 Badia Polesine (RO)  
ITALY



**Facility(ies):**

ZHERMACK S.p.A  
Via Bovazecchino, 100, 45021 Badia Polesine (RO), ITALY

**Product  
Category(ies):**

Disinfectants for medical devices

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

**Report No.:** ITA949069

**Valid from:** 2017-10-18

**Valid until:** 2022-10-17



**Date,** 2017-08-03

Stefan Preiß

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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## Zertifizierungsvertrag

Grundlage für die Zertifikatserteilung ist die Prüf- und Zertifizierungsordnung von TÜV SÜD Product Service.

Mit Erhalt des Zertifikates erkennt der Zertifikatsinhaber die jeweils gültige Fassung der Prüf- und Zertifizierungsordnung an ([www.tuev-sued.de/ps\\_regulations](http://www.tuev-sued.de/ps_regulations)) und wird somit Partner im Zertifiziersystem von TÜV SÜD Product Service.

### Prinzipielle Voraussetzung für die Gültigkeit des Zertifikates:

– Gültigkeit der zitierten normativen Prüfgrundlage(n) ist gegeben

und zusätzlich bei Zertifikaten mit Berechtigung zur Verwendung eines Prüfzeichens bzw. bei Zertifikaten für QM-Systeme:

– Voraussetzungen für vorschriftsmäßige Fertigung werden eingehalten.

– Die Fertigungs- bzw. Betriebsstätten werden regelmäßig überwacht.

## Certification contract

Certification is based on the TÜV SÜD Product Service Testing and Certification Regulations.

On receipt of the certificate the certificate holder agrees to the current version of the Testing and Certification Regulations ([www.tuev-sued.de/ps\\_regulations](http://www.tuev-sued.de/ps_regulations)) and thus becomes partner in the TÜV SÜD Product Service Certification System.

### Requirements for the validity of the certificate in principle:

– Validity of the quoted test standard(s)

In addition for certificates with the right to use a certification mark and for QM certificates:

– Conditions for an adequate manufacturing are maintained

– Regular surveillance of the facility is performed

Akkreditierungen / Benennungen (Status 14.10.2013) /  
Accreditations / notifications (as of 2013-10-14)

## Deutschland / Germany

Produktsicherheitsgesetz (ProdSG) /  
Product Safety Act (ProdSG)

## Europa / Europe

- Niederspannungsrichtlinie 2006/95/EG
- Spielzeugrichtlinie 2009/48/EG
- Richtlinie für aktive medizinische Implantate 90/385/EWG
- Richtlinie für Medizinprodukte 93/42/EWG
- Richtlinie für In-vitro-Diagnostika 98/79/EG
- Richtlinie für Gasverbrauchseinrichtungen 2009/142/EG
- Richtlinie für persönliche Schutzausrüstungen 89/686/EWG
- EMV-Richtlinie 2004/108/EG
- Richtlinie für Sportboote 94/25/EG + 2003/44/EG
- Richtlinie für Maschinen 2006/42/EG
- Richtlinie für Ex-Schutz Geräte 94/9/EG
  
- Low Voltage Directive 2006/95/EC
- Toys Directive 2009/48/EC
- Directive for Active Implantable Medical Devices 90/385/EEC
- Directive for Medical Devices 93/42/EEC
- Directive on In Vitro Diagnostic Medical Devices 98/79/EC
- Directive for Gas Appliances 2009/142/EC
- Directive for Personal Protective Equipment 89/686/EEC
- EMC Directive 2004/108/EC
- Directive for Recreational Craft 94/25/EC + 2003/44/EC
- Directive for Machinery 2006/42/EC
- Directive for Ex Safe Equipment 94/9/EC
  
- ENEC Agreement for luminaires, household and IT equipment

## USA

- Nationally Recognized Testing Laboratory (NRTL) to 29 CFR 1910.7 by OSHA
- Accredited for FDA 510(k) Third Party Review
- Conformity Assessment Body to the MRA for Medical Devices; FDA QSR Reg Inspections, FDA 510(k) Third Party Review

## Asien-Pazifik Region / Asia Pacific

- Recognized Certification Body to Electrical Products (Safety Regulation); Hong Kong
- Konformitätsbewertungsstelle / Conformity Assessment Body to the MRA for Medical Devices; Australien / Australia
- Konformitätsbewertungsstelle / Conformity Assessment Body to the MRA for Medical Devices; Neuseeland / New Zealand

## Weltweit / Worldwide

- NCB im CB-Scheme des IECEE / NCB in the CB Scheme of IECEE
- ExCB im IECEX-Scheme des IECEE / ExCB in the IECEX Scheme of IECEE
- Zertifizierstellen durch DAkkS akkreditiert  
DE-ZE-11321-01, DE-ZM-11321-09 und DE-ZM-11321-01.  
Certification Bodies accredited by DAkkS  
DE-ZE-11321-01, DE-ZM-11321-09 and DE-ZM-11321-01.

### DECLARATION OF CONFORMITY

|   |  |                            |                   |
|---|--|----------------------------|-------------------|
| Manufacturer:   | ZHERMACK SPA<br>VIA BOVAZECCHINO, 100<br>45021 BADIA POLESINE (RO) ITALY   |                            |                   |
| Product:  | <b>ITEM CODE</b>   | <b>NAME</b>                | <b>PACKAGING</b>  |
|   | C810001  | Zeta 1 Ultra               | 1 bottle 5000 ml  |
|   | C810000  | Zeta 1 Ultra               | 1 bottle 1000 ml  |
|   | C810002  | Zeta 1 Ultra               | 1 bottle 125 ml   |
|   | C810011  | Zeta 2 Sporex              | 1 jar 900 g       |
|   | C810012  | Zeta 2 Enzyme              | 1 jar 1200 g      |
|   | C810021  | Zeta 3 Ultra               | 1 bottle 750 ml   |
|   | C810023  | Zeta 3 Soft                | 1 bottle 750 ml   |
|   | C810024  | Zeta 3 Soft                | 2 bottles 2500 ml |
|   | C810029  | Zeta 3 Soft                | 1 bottle 125 ml   |
|   | C810027  | Zeta 3 Soft                | 1 bottle 750 ml   |
|   | C810028  | Zeta 3 Soft                | 2 bottles 2500 ml |
|   | C810032  | Zeta 3 Soft                | 1 bottle 125 ml   |
|   | C810025  | Zeta 3 Foam                | 1 bottle 750 ml   |
|   | C810026  | Zeta 3 Foam                | 1 bottle 3000 ml  |
|   | C810062  | Zeta 3 Wipes Total         | 1 bag 120 wipes   |
|   | C810063  | Zeta 3 Wipes Total         | 1 tub 120 wipes   |
|   | C810064  | Zeta 3 Wipes Pop-up        | 1 bag 100 wipes   |
|   | C800061  | Zeta 5 Unit                | 1 bottle 5000 ml  |
|   | C810048  | Zeta 7 Solution            | 1 bottle 1000 ml  |
| C810052   | Zeta 7 Solution  | 1 bottle 125 ml            |                   |
| C810050   | Zeta 7 Spray   | 1 bottle 750 ml            |                   |
| C810053   | Zeta 7 Spray   | 1 bottle 125 ml            |                   |
| C810040   | Zeta 5 Power Act   | 1 bottle 1000 ml           |                   |
| C810038   | Zeta 5 Power Act   | 1 box 50 unit dose (10 ml) |                   |
| Classification:   | CLASS IIa: Zeta 3 Wipes Pop-up, Zeta 5 Unit, Zeta 7 Solution, Zeta 7 Spray, Zeta 5 Power Act.<br>CLASS IIb: Zeta 1 Ultra, Zeta 2 Sporex, Zeta 2 Enzyme, Zeta 3 Ultra, Zeta 3 Soft, Zeta 3 Foam, Zeta 3 Wipes Total.<br>RULE 15 ANNEX IX OF THE MDD 93/42/CEE amended by 2007/47/CE |                            |                   |
| Intended Use :  | Disinfectants for Medical Device   |                            |                   |
| WE HEREWITH DECLARE THAT THE ABOVE MENTIONED PRODUCTS MEET THE PROVISIONS OF THE COUNCIL DIRECTIVE 93/42/CEE AMENDED BY 2007/47/CE FOR MEDICAL DEVICES, IN ACCORDANCE WITH THE ANNEXES I, (II (excluding 4) and/or V and/or VII based on the classification), X, XII. ALL SUPPORTING DOCUMENTATION IS RETAINED UNDER THE PREMISES OF THE MANUFACTURER. THE MANUFACTURER IS EXCLUSIVELY RESPONSIBLE FOR THE DECLARATION OF CONFORMITY. |  |                            |                   |
| Harmonized Standards applied:   | UNI EN ISO 13485 :2012<br>UNI CEI EN ISO 14971 :2012<br>UNI EN 1041:2013<br>UNI EN 980:2009  |                            |                   |
| Technical standards ref.:   | Zeta 1 Ultra: EN 13727:2012, EN 14561:2006 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013, EN 14562:2006 (C. albicans, A.niger), EN 14348:2005, EN 14563:2008 (M. terrae, M. avium), EN 14476:2013 (Poliovirus, Adenovirus).  |                            |                   |

| REVISION | DATE       | DESCRIPTION  |
|----------|------------|--|
| 22       | 01/04/16   | Reformulation of Intended Use in alignment with products labeling  |
| 23       | 01/02/17   | Addition of codes C810038/ C810040 Zeta 5 Power Act and related data. Removal of the withdrawn code C810030. General review. |
| 24       | 15/03/17   | Correction of typing error in packaging description of code C810011 (90 g instead of 900 g).                                 |
| 25       | 18/10/2017 | Obtaining renewed certificate that replaces the old ones (G2 15 05 53618 021)  |

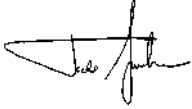
|  |  |
|--|--|
|  | <p>Zeta 3 Ultra: EN 13727:2012, EN 14561:2006 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013, EN 14562:2006 (C. albicans, A. niger), EN 14348:2005, EN 14563:2008 (M. terrae, M. avium), EN 14476:2013 (Poliovirus, Adenovirus).</p> <p>Zeta 3 Soft: EN 13727:2012, EN 14561:2006 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013, EN 14562:2006 (C. albicans), EN 14348:2005, EN 14563:2008 (M. terrae), EN 14476:2013 (Poliovirus, Adenovirus, Parvovirus).</p> <p>Zeta 3 Foam: EN 13727:2012, EN 14561 :2006 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013, EN 14562:2006 (C. albicans), EN 14348:2005, EN 14563:2008 (M. terrae), EN 14476:2013 (Poliovirus, Adenovirus, Parvovirus).</p> <p>Zeta 3 Wipes Total: EN 13727:2012, EN 14561:2006 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013, EN 14562:2006 (C. albicans), EN 14348:2005, EN 14563:2008 (M. terrae), EN 14476:2013 (Poliovirus, Adenovirus, Parvovirus). Reduction Factor efficacy of wipes (prEN 16615:2013, 10 min contact time): R&gt;4Log vs. S. aureus, E. hirae, R&gt;3Log vs. C. albicans, P. aeruginosa.</p> <p>Zeta 3 Wipes Pop-up: EN 14476:2013 (HBV, HCV, Adenovirus, Coronavirus, Norovirus, VRS, Polyomavirus, H1N1), EN 14476 (HSV), EN 14476 (Rotavirus); EN 13697, EN 1276, EN 14561, EN 14561 (MRSA); EN 13697 (A. niger), EN 14562 (A. fumigatus); EN 1650, EN 13624, EN 13697, EN 14562 (C. albicans); EN 14348, EN 14563 (M. terrae).</p> <p>Zeta 2 Sporex: EN 13727:2012, EN 14561:2006 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013, EN 14562:2006 (C. albicans, A. niger), EN 14348:2005, EN 14563:2008 (M. terrae, M. avium), EN 14476:2013 (Poliovirus, Adenovirus, Parvovirus), EN 13704:2002 (B. cereus).</p> <p>Zeta 2 Enzyme: EN 13727:2012, EN 14561:2006 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013, EN 14562:2006 (C. albicans), EN 14348:2005, EN 14563:2008 (M. terrae, M. avium), EN 14476:2013 (Poliovirus, Adenovirus, Parvovirus).</p> <p>Zeta 7 Solution: EN 13727:2012 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013 (C. albicans), EN 14348:2005, EN 14563:2008 (M. terrae), EN 14476:2013 (Poliovirus, Adenovirus, Parvovirus).</p> <p>Zeta 7 Spray: EN 13727:2012 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013 (C. albicans), EN 14348:2005, EN 14563:2008 (M. terrae), EN 14476:2013 (Poliovirus, Adenovirus, Parvovirus).</p> <p>Zeta 5 Power act (bottle/unit dose): EN 13727:2012, EN 14561:2006 (S. aureus, P. aeruginosa, E. hirae), EN 13624:2013, EN 14562:2006 (C. albicans), EN 14348:2005 (M. terrae), DVV/RKI:2014, DVV:2012, prEN 16777:2014 (all enveloped viruses, including the blood-borne viruses such as HIV, HBV and HCV, other enveloped viruses such as Herpes simplex virus and virus families such as orthomyxoviridae (including all human and animal influenza viruses like H5N1 and H1N1), filoviridae (ebola virus) and paramyxoviridae (measles virus)).</p> |
| EC certificate :                       | <p>According to annex II excluding (4) of the Directive 93/42/EEC<br/>N° G1 15 04 53618 020 valid until 2020/05/05<br/>TUV SUD PS (0123)<br/>Ridlerstrasse 65, 80339 Munchen – Germany</p> <p>Only for Zeta 3 Wipes Pop-up:<br/>According to annex V of the Directive 93/42/EEC<br/>N°G2 17 07 53618 024 valid until 2022/10/17<br/>TUV SUD PS (0123)<br/>Ridlerstrasse 65, 80339 Munchen – Germany</p>  |
| Validity of Declaration of conformity: | Valid until 2020/05/05   |
| Start of CE marking:                   | <p>Lot number/Date of first CE marking:</p> <p>C810000 Zeta 1 Ultra 1000 ml: <b>74399</b></p> <p>C810040 Zeta 5 Power act 1000 ml: <b>255429</b></p> <p>C810038 Zeta 5 Power act 50 unit dose (10 ml): <b>255429</b></p> <p>C810001 Zeta 1 Ultra 5000 ml: <b>74399</b></p> <p>C810021 Zeta 3 Ultra 750 ml: <b>75574</b></p> <p>C810023 Zeta 3 Soft 750 ml: <b>74398</b></p> <p>C810024 Zeta 3 Soft 5000 ml: <b>74398</b></p> <p>C810027 Zeta 3 Soft 750 ml: <b>91773</b></p> <p>C810028 Zeta 3 Soft 5000 ml: <b>91773</b></p> <p>C810025 Zeta 3 Foam 750 ml: <b>74768</b></p>  |

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C810026 Zeta 3 Foam 3000 ml: **74768**  
C810062 Zeta 3 Wipes Total Bag 120 pcs: **164717**  
C810063 Zeta 3 Wipes Total Tub 120 pcs: **164717**  
C810064 Zeta 3 Wipes Pop-up Bag 100 pcs: **3107461**  
C810011 Zeta 2 Sporex 900 g: **74408**  
C810012 Zeta 2 Enzyme 1200 g: **75707**  
C810048 Zeta 7 Solution 1000 ml: **75762**  
C810050 Zeta 7 Spray 750 ml: **75438**  
C800061 Zeta 5 Unit: **14/06/1998**

Badia Polesine, 18/10/2017

Paolo Ambrosini



General Manager  
**Zhermack S.p.A.**

| REVISION | DATE       | DESCRIPTION  |
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## Instructiuni pentru siguranta

### 1. Identificarea / prepararea substantei si prezentarea companiei

1.1 Identificarea substantei sau codul de preparare:

Cod: C810061-C810060  
Denumirea produsului ZETA 3 SERVETELE

1.2 Utilizarea substantei /

Dezinfectant pentru suprafete si echipament dentar.

1.3 Prezentarea companiei

Nume Zhermack S.p.a  
Adresa completa Via Bovazecchino  
Localitate, tara 45021 Badia Polesine (RO)  
Italy Tel. +39 0425-597611 Fax +39 0425-53596

Adresa de e-mail a persoanei responsabile

[tania.demetri@zhermack.com](mailto:tania.demetri@zhermack.com)

cu instructiunile de siguranta

1.4 Telefon de urgenta

Pentru intrebari urgente sunati la +39 0425-597611

### 2. Identificarea pericolului

2.1 Substanta/Clasificarea preparatului

F – Puternic inflamabil

Incadrari de risc

11 – Puternic inflamabil

Informatii legate de pericolele speciale pentru oameni si mediul inconjurator

Inhalarea cauzeaza efecte narcotice.

### 3. Compozitie / Informatii despre ingrediente

#### Caracterizare chimica

Servetele imbibate cu dezinfectant cu alcool

#### Ingrediente periculoase

CAS-No EC-No. Denumirea clasificarii [%]

64-17-5 200-578-6 Etanol < 45 F, R 11

### 4. Masuri de prin ajutor

(pentru informatii suplimentare cititi capitolul 16: Alte informatii)

#### Sfaturi generale

Nu sunt necesare masuri speciale.

#### In cazul inhalatiei

Scoateti persoana afectata la aer. Daca simptomele persista, chemati un doctor.

#### In cazul contactului cu pielea

Nu sunt necesare masuri speciale.

#### In cazul contactului cu ochii

Spalati imediat cu multa apa, deasemeni sub pleoape. Daca simptomele persista, chemati un doctor.

#### In caz de ingerare

Clatiti gura cu apa. Nu induceti voma. Cereti sfatul medicului si aratati-i ambalajul sau eticheta.

### 5. Masuri contra incendiilor

(pentru informatii suplimentare cititi capitolul 16: Alte informatii)

#### Medii corespunzatoare de stingere a focului

Spuma rezistenta la alcool, dioxid de carbon (CO<sub>2</sub>), medii de stingere uscata, jet de apa

#### Medii de stingere a focului ce nu trebuiesc folosite din masuri de siguranta

Volum mare de apa

#### Masuri speciale pentru expunerea la pericol, luate din cauza substantei sau preparatului, produselor de combustie, gazelor rezultate

In caz de foc pot fie liberate:

Dioxid sau monoxid de carbon

Gaze iritante/corozive, inflamabile si toxice

#### Echipament special de protectie pentru stingerea focului

In caz de foc purtati costume prevazute cu sistem de respiratie

#### Alte informatii

Recipientele reci supuse riscului, vor fi stropite cu apa

Vaporii sunt mai grei decat aerul si pot forma amestecuri explosive cu aerul, chiar si in recipienti goi, care nu au fost curatati

# Zhermack S.p.a

## C810060-C810061-ZETA 3 SERVETELE

Revizia nr. 1

Data 02/09/08

Tiparit 11/11/2008

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### 6. Masuri contra accidentelor

(pentru informatii suplimentare cititi capitolul 16: Alte informatii)

#### Precautii personale

Asigurati o ventilatie adecvata. In cazul formarii de vapori (concentratie ridicata) utilizati masca de gaze.

Evitati contactul cu pielea, ochii si imbracamintea.

#### Precautii pentru mediul inconjurator

Nu aruncati in apele de suprafata sau in sistemul de scurgere.

#### Metode de curatare

Folositi un material absorbant (ex: nisip, silica gel, etc)

Aruncati materialul absorbant imbibat in containere corespunzatoare

### 7. Manipulare si depozitare

(pentru informatii suplimentare cititi capitolul 16: Alte informatii)

#### Manipularea

##### Sfaturi pentru manipularea in siguranta

Evitati contactul cu ochii.

Asigurati o aerisire adecvata, mai ales in incaperi inchise.

##### Masuri de precautie impotriva focului si exploziilor

Pastrati recipientul departe de foc, surse de aprindere sau suprafete incinse – nu fumati

#### Depozitarea

##### Cerinte pentru camerele si vasele de depozitare

Pastrati containerele bine inchise intr-un loc uscat si bine aerisit. Protejati impotriva razelor solare directe.

##### Asamblul de depozitare

Incompatibil cu agentii oxidanti.

##### Alte informatii legate de conditiile de depozitare

Tineti departe de mancare, bauturi sau mancarea animalelor.

### 8. Controlul expunerii / protectia personala.

#### 8.1 Valorile limita ale expunerii

|              | Tip       | Tara | TWA/8h |     | STEL/15 min |     |       |
|--------------|-----------|------|--------|-----|-------------|-----|-------|
|              |           |      | mg/m3  | ppm | mg/m3       | ppm |       |
| PROPAN-2-OLO | TLV-ACGIH |      | 491    |     | 982         |     | Piele |
|              | OEL       | IRL  |        | 400 |             | 500 | Piele |
|              | WEL       | UK   |        | 400 |             | 500 | Piele |

TLV al amestecului solventului: 491 mg/m3

#### 8.2 Controlul expunerii

Pentru a minimaliza expunerea pe cat de mult posibil, se recomanda masuri adecvate, individuale de protectie: masca, ochelari, manusi. Nu mancati, nu beti si nu fumati atunci cand lucrati cu acest produs; spalati-va pe maini cu sapun si apa inainte de masa si dupa servicii; se recomanda un dus.

### 9. Proprietati fizice si chimice

#### Aspect

##### Statut fizic: culoare, miros

Solid, servetele albe cu alcool

##### Date legate de siguranta (a solutiei)

##### Valori unitare

Punct de fierbere 85° C

Punct de topire < -10° C

Punct de aprindere 25° C

Inflamabil (Solid/faz) n.a. °C

#### Aspect

##### Statut fizic culoare, miros

Solid, servetele albe cu alcool

Temperatura de aprindere > 425° C

Temperatura de autoaprindere n.a. °C

Gravitate specifica (20° C) aprox. 0.94 g/ml

Valorile minime de expunere 3.5 vol -%

Valorile maxime de expunere 19 vol -%

Solubilitatea in apa (20° C)

pH – valoare 6-8

### 10. Stabilitate si reactivitate

(pentru informatii suplimentare cititi capitolul 16: Alte informatii)

#### Conditii de evitat

La caldura pot fi eliberati vapori inflamabili

**Zhermack S.p.a**  
**C810060-C810061-ZETA 3 SERVETELE**

Revizia nr. 1

Data 02/09/08

Tiparit 11/11/2008

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**Materiale ce trebuie evitate**

Agenti oxidanti

**Produce descompuse periculoase**

Focul poate produce:

Dioxid si monoxid de carbon

Gaze iritante/corozive, inflamabile si toxice

**Alte informatii**

Nu vor aparea descompuneri daca modul de depozitare este corect

**11. Informatii toxicologice**

(pentru informatii suplimentare cititi capitolul 16: Alte informatii)

**Efecte asupra oamenilor**

Contactul cu ochii poate cauza iritatie

Poate provoca iritarea membranelor mucoase

**Alte informatii**

Inhalarea unei concentratii mari de vapori poate cauza efecte narcotice.

Daca este utilizat corespunzator si in conformitate cu regulile generale de igiena, nu este daunator.

**12. Informatii legate de ecologie**

(pentru informatii suplimentare cititi capitolul 16: Alte informatii)

**Informatii generale**

Nu aruncati produsul in apele de suprafata

Pericol minim pentru apa.

**13. Aruncarea produsului**

Codul/denumirea

07 06 99 fara specificatii

**Produsul**

**Recomandare**

Reciclarea este preferabila, atunci cand este posibila.

Poate fi incinerat conform legilor in vigoare.

**Pachet contaminat**

Recomandare

Containerele trebuie golite in mod optim, dupa o spalare adecvata poate fi dat la reciclare. Pachetele care nu pot fi curatate trebuie aruncate ca si produsul in sine.

**14. Transportarea**

**Transportul** (ADR/RID/GGVS/GGVE/ADNR)

**Identificare** 3175 SOLIDE CARE CONTIN LICHIDE INFLAMABILE, N.O.S. (contine etanol)

**Clasa** 4.1

**UN-No** 3175

**Poluant marin no**

**PG II**

**EMS-No** F-A, S-I

**Remarci** Cantitati limitate (capitolul 3.4): pachet combinat: 1.00 kg/30 kg (masa totala bruta)

**Transport aerian** ICAO/IATA

**Denumirea corespunzatoare de transport** SOLIDE CARE CONTIN LICHIDE IMFLAMABILE, N.O.S. (contine etanol)

**Clasa** 4.1

**UN/ID No.** 3175

**PG II**

**Remarci** PAC Y415: 0.5 kg/5

**15. Reguli**

**Clasificare**

In conformitate cu reglementarea CE privind substantele periculoase, acest produs trebuie marcat dupa cum urmeaza:

F Inflamabil puternic

R11 FOARTE IMFLAMABIL

S7 Pastrati containerul bine inchis

S16 Tineti departe de surse de aprindere – nu fumati

**16. Alte informatii**

Datele din paragrafele 4-8 si 10-12, partial nu se refera la utilizarea si domeniul de utilizare al produsului (in acest sens consultati informatiile referitoare la produs si la modul de utilizare). Aceste paragrafe se refera la masuri impotriva accidentelor si neregularitatilor.

(n.a. nu se aplica, n.d-nedeterminat)