



EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

No. CE 685475

Issued To: Atrasorb Industria

de Produtos Hospitalares Ltda

Av. Piracicaba, 351 Vila Nova São Roque

São Roque São Paulo 18131-230 Brasil

In respect of:

Design and manufacture of carbon dioxide absorbers for anaesthesia devices.

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):

Gary E Slack, Senior Vice President Medical Devices

Gay C Stade

First Issued: **2019-07-31** Date: **2019-07-31** Expiry Date: **2024-05-26**

...making excellence a habit."

Page 1 of 2

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.





EC Certificate - Full Quality Assurance System

Supplementary Information to CE 685475

Issued To: Atrasorb Industria

de Produtos Hospitalares Ltda

Av. Piracicaba, 351 Vila Nova São Roque

São Roque São Paulo 18131-230 Brasil

Number	Device Name	Intended purpose per IFU		
Class IIa				
MD 0101	Atrasorb Pharma			
MD 0101	Atrasorb Pharma Free			
MD 0101	Astrasorb Pharma FIX			

First Issued: **2019-07-31** Date: **2019-07-31** Expiry Date: **2024-05-26**

...making excellence a habit.

Page 2 of 2

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.





EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No: **CE 685475**Date: **2019-07-31**

Issued To: Atrasorb Industria

de Produtos Hospitalares Ltda

Av. Piracicaba, 351 Vila Nova São Roque

São Roque São Paulo 18131-230 Brasil

Subcontractor:

Service(s) supplied

CINTERQUAL – Soluções de Comércio Internacional Ltda.

Rua Fran Pacheco Nr. 220, 2nd floor Setúbal 2900-374

2900-374 Portugal **EU Representative**

...making excellence a habit."





EC Certificate - Full Quality Assurance System Certificate History

Certificate No: **CE 685475**Date: **2019-07-31**

Issued To: Atrasorb Industria

de Produtos Hospitalares Ltda

Av. Piracicaba, 351 Vila Nova São Roque

São Roque São Paulo 18131-230 Brasil

Date	Reference Number		Action
Current	8866857	First Issue.	AND THE RESERVE

...making excellence a habit.

Page 1 of 1

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.