

**APPROVAL**  
**EC Directive 93/42/EEC Annex II, Article 3**  
**Full Quality Assurance System**  
**Medical Devices**

**Registration No.:** HD 60022552 0001

**Report No.:** 21137970 005

**Manufacturer:** ASSKEA GmbH  
Haßlocher Str. 9  
99189 Gebesee  
Deutschland

**Scope:** Design/development, manufacture and final inspection  
of medical suction equipment


Replaces Approval, Registration No.: HD 60011810 0001

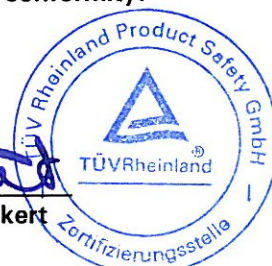
**Date of Expiry:** 14.01.2014

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex II, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex II, Article 5 of the aforementioned EC Directive, and can be used by the company with the manufacturer's declaration of conformity.

Notified Body

Cologne, 07.05.2009

  
Dipl.-Ing. U. Frenkert



**TÜV Rheinland Product Safety GmbH - Am Grauen Stein - D-51105 Köln**  
Accredited by Zentralstelle der Länder für Sicherheitstechnik (ZLS) and  
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten (ZLG).

Notified under No. **0197** to the EC Commission.

CE The CE marking may be used if all relevant and effective EC Directives are complied with.

CE