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SCHILLER
SWITZERLAND

DECLARATION OF CONFORMITY

Monitoring System: **Argus TM – 7**

Serial numbers starting with: 710.

We, the undersigned, hereby declare that the medical device (classe II b) specified above conforms with the Essential Requirements listed in Annex I, of EC Directive 93/42/EEC

This declaration is supported by:

TÜV Product Service GmbH, Management Service, D – 80339 Munich

Certificate of approval No:

12 100 13897 DIN EN ISO 9001:2000

Q1Z 01 03 41505 002 DIN EN ISO 9001:1994 / DIN EN 46001:1996

G1 01 03 41505 001 Annex II, Section 3 of the Directive 93/42/EEC Medical Devices

Valid date 02/2004.

CE 0123

Baar (Switzerland), 26.10.2001



Markus Bütler
Quality Assurance Manager



THE ART OF DIAGNOSTICS