

## Supplementary Information to CE 504480

Issued to:  
**Viamed Ltd**

**Keighley**

For the design and manufacturing operations currently certified by AMTAC MEDIQA (Notified Body 0473, certificate 485CE, Medical Device Directive 93/42/EEC, Annex II, Section 3) at:

Analytical Instruments Inc.  
2855 Metropolitan Place  
Pomona  
California 91767  
USA

First Issued: **31 Mar 2006**

Date: **31 Mar 2006**

Expiration Date: **30 Mar 2011**

Page: 2 of 2