MEDES LIMITED
Regulatory Authorized Representative

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TO WHOME IT MAY CONCERN Date: 22/05/11

Acting as Regulatory Authorized Representative in Europe for **Perflex Lrd.** from **Israel**, We hereby declare that:

The Following products listed hereunder, or in the attached "Product Schedule" have been registered with the European Competent authority in the UK (MHRA) on 20 of April 2009 as Class I devices.

Products:

- **A Denture Base Polymer**

Perflex Ltd. complies with the requirements of the Medical Device Directive 93/42 EEC and the above products can be CE marked according to the above directive.

Yours Sincerely

Benny Arazy
General Manager
Medes Limited.

Product Schedule

CE

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#	Product Name/Family	Class
1	A Denture Base Polymer	I
1.1	Perflex FN - Flexi Nylon	I
1.2	Perflex T- Crystal	I
1.3	Perflex AC- Acetal	I
1.4	Perflex AF- Acry Free	I