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To
PAUL HARTMANN AG
89522 Heidenheim
Paul-Hartmann-Strasse 12
Germany

15 October 2009

We, **Supermax latex Products**, hereby confirm that we will comply with the European Medical Device Directive (MDD 93/42/EEC) that has been amended in September 2007 by EU-Directive 2007/47/EC.

We declare that this compliance will be effective by October 01st, 2009, and will cover all deliveries of the following medical devices to PAUL HARTMANN AG from that date on (except labelling requirements):

- Peha soft (pwd and pf)

Yap Peak Geeh



Name / signature / stamp

Manager, Quality Assurance

Function (Regulatory / Quality dept. only)