MEDES limited
AUTHORIZED REPRESENTATIVE

CERTIFICATE OF CONFORMANCE

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

The following products listed hereunder or in the "Products Schedule" have been registered with the European Competent Authority in the UK (MHRA) on 5 of May 2009 Ref. No. CA010953 As Class I.

Products group

A Denture Base Polymer

By submitting the information to the MHRA, the legal requirements for registration have been met. **Perflex Ltd.** complies with the requirements of the Medical Device Directive 93/42 EEC amended by 2007/47 and the above products can be CE marked according to the above directive.

Product Schedule

Sr. No.	Product Name/Family
1	Perflex FN - Flexi Nylon
2	Perflex TC - T-Crystal
3	Perflex BS - BioSens
4	Perflex AC - Acetal
5	Perflex AF - Acry Free
6	Perflex TF - Thermofix
7	Perflex PPP - Perflex Pure Permanent
8	Perflex EC - Easy Clasp

For Medes Ltd.

Benny Arazy
General Manager
Medes Ltd.

MEDES LTD.

Date: 01.01.18

CE

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PERFLEX®
Innovative Dental Technologies

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New Industrial zone Ramat Poleg, Netanya.
Zip code: 42504, POB: 8123
TEL: 972-9-8825473 FAX:972-9-8614474

SMART 101-400

The machine, smart 101 manufactured by the company Perflex ltd.

EC CONFORMITY DECLARATION

98/37 CE - EC -- 73/23 CEE - EEC -- 89/336 CEE - EEC

Manufacturer: PERFLEX LTD
Located at:
7 Giborey Israel st. Ind. Area Poleg Netanya 4250407 Israel

Declares under our own sole responsibility that the product:

Model SMART 101-400
To which this declaration refers complies with the following norms:
EN 292/1 – EN 292/2 – EN 60335/1 – EN 50082/1 EN 50081/1

Issued At:
Perflex Ltd.
7, Giborey Israel st.
Netanya 4250407
ISRAEL

Signed by:
Mr Vidal Ben Simon – President

PERFLEX LTD
VAT.NO 514013655