

Full Quality Assurance System
Directive 93/42/EEC on Medical devices, Annex II excluding (4)

CE Certiso Ltd. (NB 2409) certifies that the following manufacturer's quality management system concerning to the listed devices and device categories meets the requirements of the related requirements of the directive.

Name of the manufacturer:

KEROX Ipari és Kereskedelmi Kft.

Headquarters: **2038. Sóskút, Kerox u. 1., Hungary**

Scope:

Dental colouring liquids for zirconia restorations

The certificate covers the following devices:

| Description of the device | Type | Intended use | Model | Risk class |
|---------------------------|-------------------|---|--|------------|
| Colouring Liquid | KEROX Farben | Colouring of dental zirconia restorations | A1, A2, A3, A3,5, A4, B1, B2, B3, B4, C1, C2, C3, C4, D2, D3, D4 | IIa |
| | KEROX Effekten | | Braun, Grey, Pink lite, Pink dark | |

This certificate is valid only in case of successfully conducted annual surveillance audits.

ID number of the related audit report: 180-CE-180912

Issue: 1

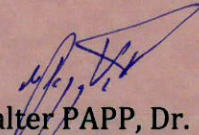
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