

# EC Declaration of Conformity



|                                                                     |                                                                                                                                                                                                                                      |
|---------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>Manufacturer:</b>                                                | Procter & Gamble Technical Centres Limited,<br>The Heights, Brooklands, Weybridge, Surrey,<br>KT13 0XP, United Kingdom                                                                                                               |
| <b>Authorised Representative</b>                                    | Procter & Gamble Service GmbH<br>Sulzbacher Str. 40<br>65824 Schwalbach a. Ts.<br>Germany                                                                                                                                            |
| <b>General product group name:</b>                                  | Desensitization dentifrice                                                                                                                                                                                                           |
| <b>Global Medical Devices Nomenclature System (GMDN) name/code:</b> | GMDN Term Name: Dentifrice<br>GMDN code: 11168                                                                                                                                                                                       |
| <b>Trade (brand) name:</b>                                          | <ul style="list-style-type: none"><li>• Oral B</li></ul>                                                                                                                                                                             |
| <b>Product reference:</b>                                           | Formula Card: <ul style="list-style-type: none"><li>• GCAS: 90509374</li><li>• GCAS: 90520592</li><li>• GCAS: 90545585</li><li>• GCAS: 91510704</li><li>• GCAS: 91530314</li><li>• GCAS: 95926870</li><li>• GCAS: 99922971</li></ul> |
| <b>Class:</b>                                                       | Ila                                                                                                                                                                                                                                  |
| <b>Notified body:</b>                                               | TÜV Rheinland LGA Products GmbH<br>Tillystraße 2<br>90431 Nürnberg<br>Germany                                                                                                                                                        |
| <b>Notified body registration number:</b>                           | 0197                                                                                                                                                                                                                                 |
| <b>EC Certificate Number</b>                                        | HD 60123454 0001                                                                                                                                                                                                                     |

We hereby declare that the above mentioned medical devices comply with all the provisions of EC Directive 93/42/EEC (as amended) which apply, and accordingly, the  mark with Notified Body registration number 0197 will be placed on all production batches placed on the market in the member states of the European Union and EFTA as defined in this declaration under Annex II (excluding point 4) of EC Directive 93/42/EEC (as amended). This Declaration of Conformity is only valid in connection with the release documents for the respective batch of produced devices. All supporting documentation is retained by the manufacturer. The declaration of conformity is issued under the sole responsibility of the manufacturer.

Weybridge

Place

Date

Maneesh Nerurkar

Digitally signed by Maneesh Nerurkar  
Date: 2021.02.08 07:40:22 Z

Signature

Maneesh Nerurkar  
Top Management, Medical Devices  
On behalf of  
Procter & Gamble Technical Centres Ltd.