Scan this code with your smart phone to verify the certificate

Certificate of Compliance

CE

This is to certify that the technical documentation/inspection/test results comply with the requirement of General product safety directive (GPSD) = 2001 / 95 / EC manufactured by:

NAME: **ARK WELLNESS**

ADDRESS: BASEMENT, B-2/10, SARA MATHEW ROAD, EXTENSION 10, NEAR B2 ARYA SAMAJ MANDIR, SAFDARJUNG ENCLAVE, NEW DELHI, SOUTH EAST DELHI, DELHI- 110029, INDIA

PRODUCTS:- **OHC PROTAPER HAND FILES, OHC NITRILE EXAMINATION GLOVES, OHC DIAMOND FG BUR KIT, OHC DIAMOND BURS, OHC LATEX EXAMINATION GLOVES, OHC CROWN PREPARATION BUR KIT, OHC CAVITY PREPARATION BURS KIT, OHC PORTABLE DENTAL CHAIR WITH AIR TURBINE, OHC ORAL IRRIGATOR(WATER FLOSSER), OHC PORTABLE DENTAL UNIT, OHC PORTABLE DENTAL CHAIR, OHC 3 PLY FACE MASK, OHC GUTTA PERCHA CUTTER, OHC COMPRESSOR OIL FREE - 1 HP/ 0.75 HP, OHC DENTAL CHAIR DELUX (FULLY AUTOMATIC), OHC DENTAL CHAIR DIAMOND (FULLY AUTOMATIC), OHC DENTAL CHAIR GOLD (FULLY AUTOMATIC), OHC PROGRESSIVE TAPER GOLD ROTARY HAND FILES, OHC DENTAL X-RAY (FLOOR MOUNTING & WALL MOUNTING!, OHC DENTAL PORTABLE DC RAY,OHC DENTAL PORTABLE AC X RAY, OHC TISSUES**

COMPLIES WITH THE REQUIREMENTS APPLICABLE TO IT

The Certification Body has performed an independent audit of the above product quality system covering the design, manufacture and final inspection of the certified product. The quality system has been assessed, approved and is subject to continuous surveillance according to the General product safety directive (GPSD) = 2001 / 95 / EC

THIS CERTIFICATE IS ISSUED UNDER THE FOLLOWING CONDITIONS:

1. It applies only to the quality system maintained in the manufacture of above referenced models and it does not substitute the design or type-examination procedures, if requested.
2. The certificate remains valid until the manufacturing conditions or the quality systems are changed.
3. The certificate validity is conditioned by positive results or surveillance audits.
4. After fulfilling the relevant EU legislation, the manufacturer shall affix to each device, of the above referenced models.
5. The CE mark as shown above can be used, under the responsibility of the manufacturer, after completion of an EC Declaration of conformity and compliance with all relevant EC Directives. The statement is based on a single evaluation of one sample of above mentioned product. It does not imply an assessment of the whole production.

Certification Calendar:

Client Id: 50194

Certificate No: CD-53277/0425

Initial Registered Date: 30.04.2025 Issuance Date: 30.04.2025

Date of Expiry: 29.04.2028

1st Surv. Due: 29.04.2026 2nd Surv. Due: 29.04.2027

Authorized Signatory