EU DECLARATION OF CONFORMITY

Manufacturer:
Air Techniques Inc.
1295 Walt Whitman Road
Melville, New York 11747
USA

Authorized Representative: MDSS GmbH Schiffgraben 41 30175 Hannover Germany	**Notified Body:** N/A **NB Id:** N/A
SRN (Single Registration Number): N/A	**Basic UDI-DI:** ++D67821701100016B
Medical Device Name/Trade Name and Identification Number: VistaScan Ultra View (J1300) REF: 2170110001 / 2170100500	**Device Class:** Class I, Rule 13 **Conformity Assessment Route:** Annex II and Annex III, Article 19 of EU MDR 2017/745

Intended Use: The unit is intended exclusively for use in dental applications for the scanning and processing of image data on an image plate.

This declaration of conformity is issued under the sole responsibility of Air Techniques Inc. We hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices. This declaration is supported by the Quality System approval to ISO 13485 issued by Intertek Medical Notified Body AB.
[We hearby declare that the subject device manufactured by Air Techniques Inc. meets the RoHS 3 requirements outlined under the directive EU 2015/863.]
All supporting documentation is retained at the premises of the manufacturer.

DocuSigned by:
Joseph Latkowski
Signer Name: Joseph Latkowski
Signing Reason: I have reviewed this document
Signing Time: 12/1/2022 | 11:12:03 AM PST
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Joseph Latkowski
Director of Quality and Regulatory Affairs

Melville, NY USA
12/1/2022
Place & Date of Issue