

Certificate

Certificate No.: MD 234172058-1
Manufacturer: **Whip Mix Corporation**
361 Farmington Ave
Louisville KY 40217
USA

REPs Facility ID: F004250
Certification criteria: ISO 13485:2016
Canada Medical Devices Regulations – Part 1 – SOR 98/282
United States 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 –
Subparts A to D

Scope: Design and development, manufacture, distribution and service of articulating paper, dental forceps, zirconia discs, PMMA discs, Veriguide OS and VeriSplint OS resins. Design and development, manufacture, distribution and service of facebows and articulators

TUV Rheinland of North America, Inc., an MDSAP recognized Auditing Organization, certifies that the quality management system of the Manufacturer has been audited against and found to conform the Certification criteria for the Scope contained in this certificate. The quality management system is subject to annual surveillance audit(s).

Project No.: 234172058-40
Issue Date: 2021-10-20
Effective Date: 2021-08-26
Expiry Date: 2024-07-22



Certification officer: Sandor Juhasz
TUV Rheinland of North America, Inc.

The validity of the certificate can be verified on https://www.certipedia.com/quality_marks/9000013541?locale=en
or calling 1-888-743-4652.

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The scope of certification includes the following additional sites:

No.	Location	Scope
/01	Whip Mix Corporation 110 Central Avenue Louisville KY 40217 USA	Design and development, manufacture, distribution and service of articulating paper, dental forceps, zirconia discs, PMMA discs, Veriguide OS and VeriSplint OS resins
/02	Whip Mix Corporation-West 1730 E. Prospect Road Fort Collins CO 80525 USA	Design and development, manufacture, distribution and service of facebows and articulators



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