

TP Orthodontics, Inc.

100 Center Plaza La Porte, Indiana 46350-9672 UNITED STATES

REPs Facility ID: 005432299

UL Medical Regulatory Services of UL LLC[®](UL) issues this certificate to the Firm named above, after auditing the Firm's quality management system and finding it in conformance per the defined scope with respect to:

ISO 13485:2016

with additional regulatory requirements listed on final page of this certificate.

The design, manufacture and distribution of non sterile orthodontic brackets, wires, ligatures, adhesives, elastomers and related accessories.



Authorized by

Michael J. Windler, P.E. Manager of Global Regulatory Service Distinguished Member of the Technical Staff UL Life and Health Sciences UL LLC

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Check Certificate Status: <u>here</u>

File Number Certificate Number Initial Issue Date A28804 2756.191031 October 31, 2019 Cycle Start Date Effective Date Expiry Date October 31, 2019 October 31, 2019 October 30, 2022

This quality system registration is included in UL's Directory of Registered Firms and applies to the provision of goods and/or services as specified in the scope of registration from the address(es) shown above. By issuance of this certificate the firm represents that it will maintain its registration in accordance with the applicable requirements. This certificate is not transferable and remains the property of UL Medical and Regulatory Services of UL LLC. Certificates may be verified by visiting the Online Certifications Directory on UL.com.



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UL LLC 333 Pfingsten Road Northbrook, IL 60062-2096 USA

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Additional Regulatory Requirements

Australia:

- Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Brazil:

- RDC ANVISA n. 16/2013
- RDC ANVISA n. 23/2012
- RDC ANVISA n. 67/2009

Canada:

- Medical Devices Regulations - Part 1- SOR 98/282

Japan:

- MHLW Ministerial Ordinance 169, Article 4 to Article 68
- PMD Act (,as applicable)

United States:

- 21 CFR 820
- 21 CFR 803
- 21 CFR 806
- 21 CFR 807 Subparts A to D
- 21 CFR 821 (where applicable)

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