K193401 Altatec GmbH CAMLOG/CONELOG PROGRESSIVE-LINE Implants

CLEARED

Sep 29, 2020
298 days to decision

K193401 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k193401/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Dec 6, 2019
Decision date	Sep 29, 2020
Days to decision	298 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Altatec GmbH**
Location	Reno, NV, US
Contact	Reto Pusterla
510(k) history	14 submissions · 14 cleared · 2006-2020

REGULATORY CONSULTANT

Consulting firm	**BioHorizons Implant Systems, Inc.**
Contact	Bill Hornbuckle

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov