FDA 510(k) Premarket Notification

K103577 TRINON Q & Q3 IMPLANT SYSTEM

CLEARED

Jul 3, 2012
575 days to decision

K103577 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k103577/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Dec 6, 2010
Decision date	Jul 3, 2012
Days to decision	575 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Trinon Titanium GmbH**
Location	Tuttlingen, DE
Contact	MARKUS DENK
510(k) history	2 submissions · 2 cleared · 2012-2017

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k103577/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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