FDA 510(k) Premarket Notification

K120053 IPS E.MAX PRESS - ABUTMENT SOLUTIONS

CLEARED

Oct 18, 2012
286 days to decision

K120053 · Product code: **NHA** · Dental
Source: https://www.510kdatabase.net/k120053/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Jan 6, 2012
Decision date	Oct 18, 2012
Days to decision	286 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ivoclar Vivadent, AG**
Location	Amherst, NY, US
Contact	DONNA MARIE HARTNETT
510(k) history	31 submissions · 31 cleared · 2004-2022