FDA 510(k) Premarket Notification

K060306 ACCELL CONNEXUS DEMINERALIZED BONE MATRIX PUTTY

CLEARED

Mar 27, 2006
48 days to decision

K060306 · Product code: **NUN** · Dental
Source: https://www.510kdatabase.net/k060306/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bone Grafting Material, Human Source (NUN)
Date received	Feb 7, 2006
Decision date	Mar 27, 2006
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Isotis NV**
Location	North Attleboro, MA, US
Contact	ELIANE SCHUTTE
510(k) history	8 submissions · 8 cleared · 2000-2007