FDA 510(k) Premarket Notification

K013672 BRUGES DISTRACTION - ANCHORING - OSTEOSYNTHESIS SYSTEM

CLEARED

Apr 8, 2002
153 days to decision

K013672 · Product code: **JEY** · Dental
Source: https://www.510kdatabase.net/k013672/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	Nov 6, 2001
Decision date	Apr 8, 2002
Days to decision	153 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Azary Technologies, LLC**
Location	Huntington, CT, US
Contact	JOSEPH M AZARY
510(k) history	1 submissions · 1 cleared · 2002-2002