FDA 510(k) Premarket Notification

K052631 SUPER QUICKANCHOR PLUS (WITH ORTHOCORD SUTURE)

CLEARED

Oct 21, 2005
25 days to decision

K052631 · Product code: **JDR** · Orthopedic
Source: https://www.510kdatabase.net/k052631/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Staple, Fixation, Bone (JDR)
Date received	Sep 26, 2005
Decision date	Oct 21, 2005
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Depuy Mitek, A Johnson & Johnson Company**
Location	Norwood, MA, US
Contact	DENISE LUCIANO
510(k) history	58 submissions · 58 cleared · 2004-2015