FDA 510(k) Premarket Notification

K000513 MODIFICATION TO EBI SPINELINK ANTERIOR CERVICAL SPINAL SYSTEM

CLEARED

Mar 7, 2000
20 days to decision

K000513 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k000513/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Feb 16, 2000
Decision date	Mar 7, 2000
Days to decision	20 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ebi, L.P.**
Location	Parsippany, NJ, US
Contact	JON CAPAROTTA
510(k) history	95 submissions · 94 cleared · 1997-2010