FDA 510(k) Premarket Notification

K020887 EBI AIS SPINE SYSTEM

CLEARED

May 16, 2002
59 days to decision

K020887 · Product code: **MQP** · Orthopedic
Source: https://www.510kdatabase.net/k020887/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Mar 18, 2002
Decision date	May 16, 2002
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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