

**K000513 MODIFICATION TO EBI SPINELINK ANTERIOR
CERVICAL SPINAL SYSTEM**Mar 7, 2000
20 days to decisionK000513 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k000513/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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