

K103742 ACCELL EVO3 (FORMERLY ACCELL A2I)

Mar 17, 2011
84 days to decision

K103742 · Product code: **MBP** · Orthopedic
Source: <https://www.510kdatabase.net/k103742/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Osteoinduction (w/o Human Growth Factor) (MBP)
Date received	Dec 23, 2010
Decision date	Mar 17, 2011
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Integra Life Sciences
Location	Las Vegas, NV, US
Contact	JENNY FAM
510(k) history	3 submissions · 3 cleared · 2011-2012

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k103742/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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