

K063067 CONFIDENCE FENESTRATED INTRODUCER NEEDLE

Nov 3, 2006
28 days to decision

K063067 · Product code: **NDN** · Orthopedic
Source: <https://www.510kdatabase.net/k063067/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Cement, Bone, Vertebroplasty (NDN)
Date received	Oct 6, 2006
Decision date	Nov 3, 2006
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Disc-O-Tech Medical Technologies, Ltd.
Location	Washington, Dc, DC, US
Contact	JONATHAN S KAHAN
510(k) history	16 submissions · 16 cleared · 2001-2007

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k063067/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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