

**K032607 POWERPRO PNEUMATIC SYSTEM, MODELS
PRO6150, PRO6175, PRO6185**

Nov 10, 2003
77 days to decision

K032607 · Product code: **DZI** · Dental
Source: <https://www.510kdatabase.net/k032607/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drill, Bone, Powered (DZI)
Date received	Aug 25, 2003
Decision date	Nov 10, 2003
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Linvatec Corp.
Location	Research Triangle Pa, NC, US
Contact	LAURA KREJCI
510(k) history	93 submissions · 87 cleared · 1992-2009

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k032607/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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