FDA 510(k) Premarket Notification

K990511 X-TIP INTRAOSSEOUS PERFORATOR

CLEARED

Apr 27, 1999
68 days to decision

K990511 · Product code: **DZM** · Dental
Source: https://www.510kdatabase.net/k990511/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Dental (DZM)
Date received	Feb 18, 1999
Decision date	Apr 27, 1999
Days to decision	68 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Schiff & Co.**
Location	West Cadwell, NJ, US
Contact	ROBERT SCHIFF
510(k) history	15 submissions · 15 cleared · 1988-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k990511/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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