FDA 510(k) Premarket Notification

K060275 PIEZOTOME

CLEARED

May 3, 2006
90 days to decision

K060275 · Product code: **ELC** · Dental
Source: https://www.510kdatabase.net/k060275/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Scaler, Ultrasonic (ELC)
Date received	Feb 2, 2006
Decision date	May 3, 2006
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Satelec**
Location	Merignac, FR
Contact	STEVE SALESKY
510(k) history	24 submissions · 24 cleared · 1990-2011

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k060275/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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