FDA 510(k) Premarket Notification

K951019 INTRAORAL VIDEO SCOPE AND WINDOW TUBE

CLEARED

May 10, 1995
64 days to decision

K951019 · Product code: **EIA** · Dental
Source: https://www.510kdatabase.net/k951019/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Operative Dental (EIA)
Date received	Mar 7, 1995
Decision date	May 10, 1995
Days to decision	64 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Cuda Products Co.**
Location	Jacksonville, FL, US
Contact	KIM REED
510(k) history	17 submissions · 17 cleared · 1988-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k951019/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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