FDA 510(k) Premarket Notification

K801888 HANDPIECE, DENTAL

CLEARED

Aug 20, 1980
23 days to decision

K801888 · Product code: **DZA** · Dental
Source: https://www.510kdatabase.net/k801888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drill, Dental, Intraoral (DZA)
Date received	Jul 28, 1980
Decision date	Aug 20, 1980
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Aesculap Instruments Corp.**
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1977-1990