

K801888 HANDPIECE, DENTALAug 20, 1980
23 days to decisionK801888 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k801888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026