FDA 510(k) Premarket Notification

K811113 RIESS-LUHR

CLEARED

Jun 16, 1981
54 days to decision

K811113 · Product code: **DZA** · Dental
Source: https://www.510kdatabase.net/k811113/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drill, Dental, Intraoral (DZA)
Date received	Apr 23, 1981
Decision date	Jun 16, 1981
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Allomedic**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k811113/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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