FDA 510(k) Premarket Notification

K810538 PLASTICEPH

CLEARED

Apr 29, 1981
62 days to decision

K810538 · Product code: **EAG** · Radiology
Source: https://www.510kdatabase.net/k810538/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cephalometer (EAG)
Date received	Feb 26, 1981
Decision date	Apr 29, 1981
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Morningstar Dental Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981