FDA 510(k) Premarket Notification

K955946 ACETAL DENTAL/PREFORMED CLASP

CLEARED

Dec 11, 1995
45 days to decision

K955946 · Product code: **EHP** · Dental
Source: https://www.510kdatabase.net/k955946/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clasp, Preformed (EHP)
Date received	Oct 27, 1995
Decision date	Dec 11, 1995
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Pressing DI Monticelli Rag. Stefano**
Location	Cattilica, IT
Contact	STEFANO MONTICELLI
510(k) history	2 submissions · 2 cleared · 1995-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k955946/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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