FDA 510(k) Premarket Notification

K882578 ASTRA DENTAL IMPLANT SYSTEM

CLEARED

Aug 12, 1988
51 days to decision

K882578 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k882578/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jun 22, 1988
Decision date	Aug 12, 1988
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Astra Meditec AB**
Location	Sweden, SE
Contact	ANDERS HOLMEN
510(k) history	3 submissions · 2 cleared · 1988-1993

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k882578/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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