FDA 510(k) Premarket Notification

K911112 DEVICE PART NUMBERS: - 1400 SERIES

CLEARED

Jun 10, 1991
117 days to decision

K911112 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k911112/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Feb 13, 1991
Decision date	Jun 10, 1991
Days to decision	117 days
Third-party review	No

APPLICANT

Company	**Minimatic Implant Technology**
Location	Deerfield Beach, FL, US
Contact	LEON SHAW
510(k) history	34 submissions · 34 cleared · 1990-1992

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k911112/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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