FDA 510(k) Premarket Notification

K834364 CRYSTALLINE BONE SCREW ENDO-OSSEOUS

CLEARED

Mar 6, 1984
131 days to decision

K834364 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k834364/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Oct 27, 1983
Decision date	Mar 6, 1984
Days to decision	131 days
Third-party review	No

APPLICANT

Company	**C.B.S. Biotechnics, Inc.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1984-1984

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k834364/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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