

K813257 ALVEOGRAF DURAPATITE(18-40 MESH)Jan 5, 1982
47 days to decisionK813257 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k813257/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Nov 19, 1981
Decision date	Jan 5, 1982
Days to decision	47 days
Third-party review	No

APPLICANT

Company	Sterling Drug, Inc.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1978-1990

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Device record: <https://www.510kdatabase.net/k813257/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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