

K820568 MUSCLE DYSFUNCTION MONITORApr 9, 1982
38 days to decisionK820568 · Product code: **DZM** · Dental
Source: <https://www.510kdatabase.net/k820568/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Dental (DZM)
Date received	Mar 2, 1982
Decision date	Apr 9, 1982
Days to decision	38 days
Third-party review	No

APPLICANT

Company	A Company, Inc.
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1982-1990

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Device record: <https://www.510kdatabase.net/k820568/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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