

K820392 ORTHO* BI-LEVEL ASSAYED ALCOHOL DILUENMar 4, 1982
22 days to decisionK820392 · Product code: **LAS** · Chemistry
Source: <https://www.510kdatabase.net/k820392/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Specific Control Materials (LAS)
Date received	Feb 10, 1982
Decision date	Mar 4, 1982
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Ortho Diagnostic Systems, Inc.
Location	Carpinteria, CA, US
510(k) history	126 submissions · 126 cleared · 1981-1997

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Device record: <https://www.510kdatabase.net/k820392/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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