

K820050 KREMER BLADE GAUGEFeb 24, 1982
47 days to decisionK820050 · Product code: **HOE** · Ophthalmic
Source: <https://www.510kdatabase.net/k820050/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Caliper, Ophthalmic (HOE)
Date received	Jan 8, 1982
Decision date	Feb 24, 1982
Days to decision	47 days
Third-party review	No

APPLICANT

Company	Accutome, Inc.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1981-2015

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Device record: <https://www.510kdatabase.net/k820050/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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