FDA 510(k) Premarket Notification

K820050 KREMER BLADE GAUGE

CLEARED

Feb 24, 1982
47 days to decision

K820050 · 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