FDA 510(k) Premarket Notification

K812011 BLADE GAUGE

CLEARED

Jul 28, 1981
11 days to decision

K812011 · Product code: **HOE** · Ophthalmic
Source: https://www.510kdatabase.net/k812011/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Caliper, Ophthalmic (HOE)
Date received	Jul 17, 1981
Decision date	Jul 28, 1981
Days to decision	11 days
Third-party review	No

APPLICANT

Company	**Koi, Inc.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1981-1986