FDA 510(k) Premarket Notification

K781715 RULER, WECK SCOTT

CLEARED

Dec 4, 1978
60 days to decision

K781715 · Product code: **HOE** · Ophthalmic
Source: https://www.510kdatabase.net/k781715/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Caliper, Ophthalmic (HOE)
Date received	Oct 5, 1978
Decision date	Dec 4, 1978
Days to decision	60 days
Third-party review	No

APPLICANT

Company	**Edward Weck, Inc.**
Location	Mchenry, IL, US
510(k) history	140 submissions · 140 cleared · 1976-1992