

K781715 RULER, WECK SCOTTDec 4, 1978
60 days to decisionK781715 · 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

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Device record: <https://www.510kdatabase.net/k781715/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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