FDA 510(k) Premarket Notification

K791983 SPATULA'S #'S 7618,7248,7240,7242,5108

CLEARED

Oct 4, 1979
9 days to decision

K791983 · Product code: **HND** · Ophthalmic
Source: https://www.510kdatabase.net/k791983/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spatula, Ophthalmic (HND)
Date received	Sep 25, 1979
Decision date	Oct 4, 1979
Days to decision	9 days
Third-party review	No

APPLICANT

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