FDA 510(k) Premarket Notification

K991256 VIEWPOINT ENT/ORTHOPEDIC OPTION

CLEARED

Oct 8, 1999
178 days to decision

K991256 · Product code: **HAW** · Neurology
Source: https://www.510kdatabase.net/k991256/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Apr 13, 1999
Decision date	Oct 8, 1999
Days to decision	178 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Philips Medical Systems (Cleveland), Inc.**
Location	Mchenry, IL, US
Contact	ELAINE K KEELER
510(k) history	190 submissions · 190 cleared · 1977-2017