FDA 510(k) Premarket Notification

K953173 NONPOWERED NEUROSURGICAL INSTRUMENT

CLEARED

Oct 3, 1995
88 days to decision

K953173 · Product code: **GWG** · Neurology
Source: https://www.510kdatabase.net/k953173/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscope, Neurological (GWG)
Date received	Jul 7, 1995
Decision date	Oct 3, 1995
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**United States Surgical, A Division of Tyco Healthc**
Location	Mchenry, IL, US
Contact	JANET G JOHNSON
510(k) history	218 submissions · 200 cleared · 1977-2007