FDA 510(k) Premarket Notification

K772371 BAYONET SURGICAL INSTRUMENTS

CLEARED

Jan 6, 1978
10 days to decision

K772371 · Product code: **HAO** · Neurology
Source: https://www.510kdatabase.net/k772371/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Non-powered (HAO)
Date received	Dec 27, 1977
Decision date	Jan 6, 1978
Days to decision	10 days
Third-party review	No

APPLICANT

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