FDA 510(k) Premarket Notification

K811825 ORANGE G-6

CLEARED

Jul 10, 1981
11 days to decision

K811825 · Product code: **HZH** · Pathology
Source: https://www.510kdatabase.net/k811825/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orange G (HZH)
Date received	Jun 29, 1981
Decision date	Jul 10, 1981
Days to decision	11 days
Third-party review	No

APPLICANT

Company	**Surgipath**
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1981-1982

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k811825/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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