

**K012380 MODIFICATION TO: STRYKER NAVIGATION SYSTEM,
STRYKER FLUOROSCOPY SYSTEM,VIRTUAL FLUOROSCOPY**

Dec 3, 2001
130 days to decision

K012380 · Product code: **HAW** · Neurology
Source: <https://www.510kdatabase.net/k012380/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Jul 26, 2001
Decision date	Dec 3, 2001
Days to decision	130 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Stryker Leibinger
Location	Kalamazoo, MI, US
Contact	ROBIN L ROWE
510(k) history	12 submissions · 12 cleared · 1999-2005

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k012380/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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