

**K013861 MODIFICATION TO LEKSELL SURGIPLAN WITH
ATLASPACE**Dec 6, 2001
15 days to decisionK013861 · Product code: **HAW** · Neurology
Source: <https://www.510kdatabase.net/k013861/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Nov 21, 2001
Decision date	Dec 6, 2001
Days to decision	15 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Elekta Instrument AB
Location	Lake Forest, CA, US
Contact	CAROL PATTERSON
510(k) history	35 submissions · 35 cleared · 1996-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013861/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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