

**K934368 NK PRESSURE-SPECIFIED SENSORY DEVICE,
MODEL PSSD-001**Aug 11, 1994
338 days to decisionK934368 · Product code: LLN · Neurology
Source: <https://www.510kdatabase.net/k934368/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vibration Threshold Measurement (LLN)
Date received	Sep 7, 1993
Decision date	Aug 11, 1994
Days to decision	338 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Nk Biotechnical Engineering Co.
Location	Minneapolis, MN, US
Contact	KAREN GOTFREDSON
510(k) history	15 submissions · 15 cleared · 1991-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k934368/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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