FDA 510(k) Premarket Notification

CLEARED

K934368 NK PRESSURE-SPECIFIED SENSORY DEVICE, MODEL PSSD-001

Aug 11, 1994
338 days to decision

K934368 · 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