

K840108 NEUROMUSCULAR TRANSMISSION MONITORFeb 17, 1984
38 days to decisionK840108 · Product code: **KOI** · Anesthesiology
Source: <https://www.510kdatabase.net/k840108/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Peripheral, Electric (KOI)
Date received	Jan 10, 1984
Decision date	Feb 17, 1984
Days to decision	38 days
Third-party review	No

APPLICANT

Company	Datex Division Instrumentarium Corp.
Location	Walker, MI, US
510(k) history	55 submissions · 55 cleared · 1982-1997

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Device record: <https://www.510kdatabase.net/k840108/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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