

**K782036 TREMOR MONITOR**Jan 23, 1979  
49 days to decisionK782036 · Product code: **GYD** · Neurology  
Source: <https://www.510kdatabase.net/k782036/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Tremor (GYD)
Date received	Dec 5, 1978
Decision date	Jan 23, 1979
Days to decision	49 days
Third-party review	No

**APPLICANT**

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Company	<b>Columbus Instruments Intl. Corp.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1990

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Device record: <https://www.510kdatabase.net/k782036/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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