

**K812908 ELECTRO-GONIOMETER**Nov 24, 1981  
36 days to decisionK812908 · Product code: **KQX** · Neurology  
Source: <https://www.510kdatabase.net/k812908/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Oct 19, 1981
Decision date	Nov 24, 1981
Days to decision	36 days
Third-party review	No

**APPLICANT**

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Company	<b>Chattanooga Group, Inc.</b>
Location	Mchenry, IL, US
510(k) history	70 submissions · 68 cleared · 1980-2001

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Device record: <https://www.510kdatabase.net/k812908/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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