

**K884003 GONIOMETER AND BIOMEDICAL RECORDER SYSTEM**Dec 19, 1988  
89 days to decisionK884003 · Product code: **KQX** · Neurology  
Source: <https://www.510kdatabase.net/k884003/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Sep 21, 1988
Decision date	Dec 19, 1988
Days to decision	89 days
Third-party review	No

**APPLICANT**

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Company	<b>Penny &amp; Giles Blackwood , Ltd.</b>
Location	England, GB
Contact	DR. I THOMAS
510(k) history	1 submissions · 1 cleared · 1988-1988

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884003/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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