

**K913079 02810-01/02810-02 ORTHORANGER II/02810-3 ROM  
360**Sep 25, 1991  
76 days to decisionK913079 · Product code: **KQX** · Neurology  
Source: <https://www.510kdatabase.net/k913079/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Jul 11, 1991
Decision date	Sep 25, 1991
Days to decision	76 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Lucas Sensing Systems, Inc.</b>
Location	Phoenix, AZ, US
Contact	WAYNE SALUTE
510(k) history	1 submissions · 1 cleared · 1991-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k913079/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](https://accessdata.fda.gov). - Generated July 1, 2026