FDA 510(k) Premarket Notification

K913079 02810-01/02810-02 ORTHORANGER II/02810-3 ROM 360

CLEARED

Sep 25, 1991
76 days to decision

K913079 · Product code: **KQX** · Neurology
Source: https://www.510kdatabase.net/k913079/

SUBMISSION DETAILS

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

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