FDA 510(k) Premarket Notification

K884003 GONIOMETER AND BIOMEDICAL RECORDER SYSTEM

CLEARED

Dec 19, 1988
89 days to decision

K884003 · Product code: **KQX** · Neurology
Source: https://www.510kdatabase.net/k884003/

SUBMISSION DETAILS

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

Company	**Penny & Giles Blackwood , Ltd.**
Location	England, GB
Contact	DR. I THOMAS
510(k) history	1 submissions · 1 cleared · 1988-1988