FDA 510(k) Premarket Notification

K842609 R.L. MEDICAL TWIN LIGHT SOURCE

CLEARED

Aug 9, 1984
35 days to decision

K842609 · Product code: **FCW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k842609/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Source, Fiberoptic, Routine (FCW)
Date received	Jul 5, 1984
Decision date	Aug 9, 1984
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Tridak Division of Indicon, Inc.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1984-1990