FDA 510(k) Premarket Notification

K080072 MRI PATIENT POSITIONING DEVICES

CLEARED

Jun 4, 2008
145 days to decision

K080072 · Product code: **IYE** · Radiology
Source: https://www.510kdatabase.net/k080072/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jan 11, 2008
Decision date	Jun 4, 2008
Days to decision	145 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**CIVCO Medical Instruments Co., Inc.**
Location	Walker, MI, US
Contact	ARTHUR WARD
510(k) history	29 submissions · 29 cleared · 1982-2023