FDA 510(k) Premarket Notification

K951808 MED-TEC REDI-FOAM

CLEARED

Jul 6, 1995
78 days to decision

K951808 · Product code: **IYE** · Radiology
Source: https://www.510kdatabase.net/k951808/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Apr 19, 1995
Decision date	Jul 6, 1995
Days to decision	78 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medtec, Inc.**
Location	Orange City, IA, US
Contact	DONALD RIIBE
510(k) history	41 submissions · 41 cleared · 1993-2014