

K934519 RADIATION EXIT VERIFICATION SYSTEM/BACK POINTER SYSTEM

Dec 13, 1993
87 days to decision

K934519 · Product code: **IWE** · Radiology
Source: <https://www.510kdatabase.net/k934519/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	Sep 17, 1993
Decision date	Dec 13, 1993
Days to decision	87 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k934519/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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