FDA 510(k) Premarket Notification

K800223 PATIENT POSITIONING SYSTEM 1D OR 2D

CLEARED

Mar 3, 1980
28 days to decision

K800223 · Product code: **IWE** · Radiology
Source: https://www.510kdatabase.net/k800223/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	Feb 4, 1980
Decision date	Mar 3, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Laboratoire National D'Essais**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980