FDA 510(k) Premarket Notification

K803049 DIGITAL RADIOGRAPHY CHEST DEVICE

CLEARED

Mar 2, 1981
91 days to decision

K803049 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k803049/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Dec 1, 1980
Decision date	Mar 2, 1981
Days to decision	91 days
Third-party review	No

APPLICANT

Company	**Philips Medical Systems (Cleveland), Inc.**
Location	Mchenry, IL, US
510(k) history	190 submissions · 190 cleared · 1977-2017