

K803049 DIGITAL RADIOGRAPHY CHEST DEVICEMar 2, 1981
91 days to decisionK803049 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k803049/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 16, 2026