FDA 510(k) Premarket Notification

K880982 ORALIX PAN DC III CEPH OR PANORAMIC MODE ONLY

CLEARED

Jun 6, 1988
90 days to decision

K880982 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k880982/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Mar 8, 1988
Decision date	Jun 6, 1988
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Orion Corporation Soredex**
Location	Helsinki, FI
Contact	ENSIO KOSKENNURMI
510(k) history	4 submissions · 4 cleared · 1988-1998