

**K880982 ORALIX PAN DC III CEPH OR PANORAMIC MODE ONLY**Jun 6, 1988  
90 days to decisionK880982 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k880982/>**SUBMISSION DETAILS**

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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**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880982/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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