

**K912517 FLOOR MOUNT TUBESTAND PERFORMER,  
6610.101/S-9109**

Sep 19, 1991  
105 days to decision

K912517 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k912517/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Jun 6, 1991
Decision date	Sep 19, 1991
Days to decision	105 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Continental X-Ray Corp.</b>
Location	Chicago, IL, US
Contact	OSCAR KHUTORYANSKY
510(k) history	27 submissions · 27 cleared · 1988-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k912517/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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