

**K141979 ALLURA XPER FD SERIES / ALLURA XPER OR TABLE SERIES**

Aug 19, 2014  
29 days to decision

K141979 · Product code: **OWB** · Radiology  
Source: <https://www.510kdatabase.net/k141979/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Interventional Fluoroscopic X-ray System (OWB)
Date received	Jul 21, 2014
Decision date	Aug 19, 2014
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Medical Systems Netherland BV</b>
Location	Best, NL
Contact	JEANETTE BECKER
510(k) history	1 submissions · 1 cleared · 2014-2014

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

Device record: <https://www.510kdatabase.net/k141979/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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