

**K123263 HDRPLUS, ADD IN PROSTATE MODULE FOR HDRPLUS**

Apr 4, 2013  
168 days to decision

K123263 · Product code: **MUJ** · Radiology  
Source: <https://www.510kdatabase.net/k123263/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	System, Planning, Radiation Therapy Treatment (MUJ)
Date received	Oct 18, 2012
Decision date	Apr 4, 2013
Days to decision	168 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Eckert &amp; Ziegler Bebig GmbH</b>
Location	Veenendaal, NL
Contact	HUB VAN DE BERGH
510(k) history	4 submissions · 4 cleared · 2013-2015

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

Device record: <https://www.510kdatabase.net/k123263/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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