

**K070542 CARDIUS 1, 2 AND 3 XPO AND 2020TC SPECT IMAGING SYSTEMS**

Mar 23, 2007  
25 days to decision

K070542 · Product code: **KPS** · Radiology  
Source: <https://www.510kdatabase.net/k070542/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Tomography, Computed, Emission (KPS)
Date received	Feb 26, 2007
Decision date	Mar 23, 2007
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Digirad Corp.</b>
Location	Palo Alto, CA, US
Contact	Joel Tuckey
510(k) history	16 submissions · 16 cleared · 1983-2013

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

Device record: <https://www.510kdatabase.net/k070542/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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