

**K072191 CLEARVISION NUCLEAR MEDICINE IMAGING SYSTEM**

Oct 25, 2007  
80 days to decision

K072191 · Product code: **KPS** · Radiology  
Source: <https://www.510kdatabase.net/k072191/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Tomography, Computed, Emission (KPS)
Date received	Aug 6, 2007
Decision date	Oct 25, 2007
Days to decision	80 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Gvi Technology Partners</b>
Location	Twinsburg, OH, US
Contact	KEVIN M MURROCK
510(k) history	3 submissions · 3 cleared · 2003-2007

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

Device record: <https://www.510kdatabase.net/k072191/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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