

**K171303 Added Screen Displays For FDA Cleared HeartSee Cardiac P.E.T Processing Software K143664**

Sep 22, 2017  
142 days to decision

K171303 · Product code: **KPS** · Radiology  
Source: <https://www.510kdatabase.net/k171303/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Tomography, Computed, Emission (KPS)
Date received	May 3, 2017
Decision date	Sep 22, 2017
Days to decision	142 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>University of Texas Medical School AT Houston, Texas</b>
Location	Houston, TX, US
Contact	K. Lance Gould
510(k) history	1 submissions · 1 cleared · 2017-2017

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

Device record: <https://www.510kdatabase.net/k171303/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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