

**K974355 INTERVENTIONAL MOBILE DIGITAL IMAGING SYSTEM (HEREIN CALLED IMDIS)**Feb 17, 1998  
90 days to decisionK974355 · Product code: **OWB** · Radiology  
Source: <https://www.510kdatabase.net/k974355/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Interventional Fluoroscopic X-ray System (OWB)
Date received	Nov 19, 1997
Decision date	Feb 17, 1998
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ge Dec Medical Systems</b>
Location	Salt Lake City, UT, US
Contact	TED L PARROT
510(k) history	10 submissions · 10 cleared · 1994-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k974355/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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