

**K925705 ACCURAD-200 SYSTEM & ACCURAD-300 PLUS SYSTEM**

Dec 28, 1992  
46 days to decision

K925705 · Product code: **IWY** · Radiology  
Source: <https://www.510kdatabase.net/k925705/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Holder, Head, Radiographic (IWY)
Date received	Nov 12, 1992
Decision date	Dec 28, 1992
Days to decision	46 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Denar Corp.</b>
Location	Anaheim, CA, US
Contact	THOMAS J MULDOON
510(k) history	5 submissions · 5 cleared · 1985-1992

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

Device record: <https://www.510kdatabase.net/k925705/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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