

K002424 3D ANGIOGRAPHIC IMAGING SYSTEM, MODEL XIDF-100A

Oct 27, 2000
80 days to decision

K002424 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k002424/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Aug 8, 2000
Decision date	Oct 27, 2000
Days to decision	80 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Toshiba America Medical Systems, In.C
Location	Tustin, CA, US
Contact	DIANA THORSON
510(k) history	146 submissions · 146 cleared · 1989-2014

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Device record: <https://www.510kdatabase.net/k002424/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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