

**K133535 INFX-8000V, V5.30**Jul 23, 2014  
247 days to decisionK133535 · Product code: **OWB** · Radiology  
Source: <https://www.510kdatabase.net/k133535/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Interventional Fluoroscopic X-ray System (OWB)
Date received	Nov 18, 2013
Decision date	Jul 23, 2014
Days to decision	247 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Toshibamedical Systems Corporation</b>
Location	Tustin, CA, US
Contact	CHARLEMAGNE CHUA
510(k) history	80 submissions · 80 cleared · 2004-2018

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133535/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 16, 2026