

**K233230 See Factor CT3™**Jun 21, 2024  
267 days to decisionK233230 · Product code: **OWB** · Radiology  
Source: <https://www.510kdatabase.net/k233230/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Interventional Fluoroscopic X-ray System (OWB)
Date received	Sep 28, 2023
Decision date	Jun 21, 2024
Days to decision	267 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Epica International, Inc.</b>
Location	San Clemente, CA, US
Contact	Craig Glaiberman
510(k) history	2 submissions · 2 cleared · 2019-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k233230/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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