

**K091804 CX50 VERSION 2.0 DIAGNOSTIC ULTRASOUND
SYSTEM AND TRANSDUCERS**Jul 17, 2009
29 days to decisionK091804 · Product code: **ITX** · Radiology
Source: <https://www.510kdatabase.net/k091804/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Ultrasonic, Diagnostic (ITX)
Date received	Jun 18, 2009
Decision date	Jul 17, 2009
Days to decision	29 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Philips Ultrasound, Inc.
Location	Santa Ana, CA, US
Contact	ROB BUTLER
510(k) history	46 submissions · 46 cleared · 1985-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091804/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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