

**K013273 FUKUDA DENSHI MODEL HG-500, PULSE OXIMETER  
MODULE**Dec 20, 2001  
80 days to decisionK013273 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k013273/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Oct 1, 2001
Decision date	Dec 20, 2001
Days to decision	80 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Fukuda Denshi USA, Inc.</b>
Location	Mchenry, IL, US
Contact	LARRY WALKER
510(k) history	68 submissions · 68 cleared · 1984-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013273/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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