

**K083351 PROABLE PULSE OXIMETER, MODEL MD-600P,  
TRAVELER OXIMETER, MODEL MD-650P AND PALM PULSE  
OXIMETER, MODEL MD-680P**Apr 24, 2009  
162 days to decisionK083351 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k083351/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Nov 13, 2008
Decision date	Apr 24, 2009
Days to decision	162 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Comdek Industrial Corp.</b>
Location	Taipei, TW
Contact	Y.S. KUO
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083351/> 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 24, 2026