

K883229 OHMEDA CONTINUOUS SUCTION REGULATOROct 19, 1988
79 days to decisionK883229 · Product code: **KDP** · General Hospital
Source: <https://www.510kdatabase.net/k883229/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Vacuum (KDP)
Date received	Aug 1, 1988
Decision date	Oct 19, 1988
Days to decision	79 days
Third-party review	No

APPLICANT

Company	Ohmeda Medical
Location	Madison, WI, US
Contact	ALBERTO F PROFUMO
510(k) history	120 submissions · 118 cleared · 1984-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k883229/>; Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026