

K072292 NEGAVENT PLUS PEGASO COUGH, MODEL: DA-3 PLUSJan 7, 2008
144 days to decisionK072292 · Product code: **NHJ** · Anesthesiology
Source: <https://www.510kdatabase.net/k072292/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Positive Pressure Breathing, Intermittent (NHJ)
Date received	Aug 16, 2007
Decision date	Jan 7, 2008
Days to decision	144 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Dima Italia Srl
Location	Boulder, CO, US
Contact	LEWIS WARD
510(k) history	4 submissions · 4 cleared · 2008-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072292/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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