

K013884 BOUSSIGNAC C.P.A.P. DEVICEJan 21, 2003
424 days to decisionK013884 · Product code: **BYE** · Anesthesiology
Source: <https://www.510kdatabase.net/k013884/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Attachment, Breathing, Positive End Expiratory Pressure (BYE)
Date received	Nov 23, 2001
Decision date	Jan 21, 2003
Days to decision	424 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vygon SA
Location	Temecula, CA, US
Contact	LAETITIA BERNARD
510(k) history	2 submissions · 2 cleared · 2003-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013884/> 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 25, 2026