

K001803 ACCELERATOR RECIPROCATING CANNULAAug 4, 2000
51 days to decisionK001803 · Product code: **QPB** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k001803/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Suction, Lipoplasty For Removal (QPB)
Date received	Jun 14, 2000
Decision date	Aug 4, 2000
Days to decision	51 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Byron Medical
Location	Tucson, AZ, US
Contact	REGINA S HARRIS
510(k) history	27 submissions · 24 cleared · 1986-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001803/> 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 July 2, 2026