

**K000108 RAPID RHINO-ANTERIOR 90MM, MODEL RR-A 90,
RAPID RHINO-POSTERIOR 100MM, MODEL RR-P 100**May 23, 2000
130 days to decisionK000108 · Product code: **EMX** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k000108/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Balloon, Epistaxis (EMX)
Date received	Jan 14, 2000
Decision date	May 23, 2000
Days to decision	130 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bhk Holding
Location	New Port Riche, FL, US
Contact	ART WARD
510(k) history	1 submissions · 1 cleared · 2000-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k000108/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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