FDA 510(k) Premarket Notification

CLEARED

K000108 RAPID RHINO-ANTERIOR 90MM, MODEL RR-A 90, RAPID RHINO-POSTERIOR 100MM, MODEL RR-P 100

May 23, 2000
130 days to decision

K000108 · 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