FDA 510(k) Premarket Notification

K982929 MAXAIR NASAL DILATOR SYSTEM

CLEARED

Sep 9, 1998
20 days to decision

K982929 · Product code: **LWF** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k982929/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Nasal (LWF)
Date received	Aug 20, 1998
Decision date	Sep 9, 1998
Days to decision	20 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Hnl Technologies**
Location	San Mateo, CA, US
Contact	HANFORD N LOCKWOOD, JR.
510(k) history	1 submissions · 1 cleared · 1998-1998