FDA 510(k) Premarket Notification

K760316 REGULATORS, LOW SUCTION

CLEARED

Aug 4, 1976
9 days to decision

K760316 · Product code: **KDP** · General Hospital
Source: https://www.510kdatabase.net/k760316/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Vacuum (KDP)
Date received	Jul 26, 1976
Decision date	Aug 4, 1976
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**Harris-Lake, Inc.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1976-1977