FDA 510(k) Premarket Notification

K770944 INSTATEMP

CLEARED

Jul 27, 1977
63 days to decision

K770944 · Product code: **KDP** · General Hospital
Source: https://www.510kdatabase.net/k770944/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Vacuum (KDP)
Date received	May 25, 1977
Decision date	Jul 27, 1977
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**H.I. Silverman, D.Sc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977