FDA 510(k) Premarket Notification

K953810 FLUFF GAUZE BANDAGE, STERILE & NON-STERILE

CLEARED

Sep 14, 1995
37 days to decision

K953810 · Product code: **EFQ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k953810/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze/sponge, Internal (EFQ)
Date received	Aug 8, 1995
Decision date	Sep 14, 1995
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**K. W. Griffen Co.**
Location	Stamford, CT, US
Contact	DAVID M PIERATOS
510(k) history	4 submissions · 3 cleared · 1987-1996