FDA 510(k) Premarket Notification

K020265 SERRALENE, MODEL CATALOG NO 1S

CLEARED

Mar 8, 2002
42 days to decision

K020265 · Product code: **GAW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k020265/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Nonabsorbable, Synthetic, Polypropylene (GAW)
Date received	Jan 25, 2002
Decision date	Mar 8, 2002
Days to decision	42 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Serral, S.A. DE C.V.**
Location	Williamsburg, VA, US
Contact	SCOTT HENDERSON
510(k) history	6 submissions · 6 cleared · 2002-2004