FDA 510(k) Premarket Notification

K072219 CYTOPHIL TISSUE MARKER

CLEARED

Jan 18, 2008
161 days to decision

K072219 · Product code: **NEU** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k072219/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Marker, Radiographic, Implantable (NEU)
Date received	Aug 10, 2007
Decision date	Jan 18, 2008
Days to decision	161 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cytophil, Inc.**
Location	Whitefish Bay, WI, US
Contact	Greg Johnson
510(k) history	8 submissions · 8 cleared · 2008-2015