

K961061 CODI SYNTHETIC SPONGE(NON-STERILE AND STERILE)

Jun 6, 1996
80 days to decision

K961061 · Product code: **EFQ** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k961061/>

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Gauze/sponge, Internal (EFQ)
Date received	Mar 18, 1996
Decision date	Jun 6, 1996
Days to decision	80 days
Third-party review	No

APPLICANT

Company	Codi International BV
Location	Boston, MA, US
Contact	G. THACHER STORM
510(k) history	2 submissions · 1 cleared · 1996-1996

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k961061/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 5, 2026