

**K951660 DYNAREX GAUZE SPONGE**May 9, 1995  
29 days to decisionK951660 · Product code: **EFQ** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k951660/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze/sponge, Internal (EFQ)
Date received	Apr 10, 1995
Decision date	May 9, 1995
Days to decision	29 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Dynarex Corp.</b>
Location	Brewster, NY, US
Contact	JOHN G MOULDEN
510(k) history	19 submissions · 18 cleared · 1986-2005

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951660/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 25, 2026