FDA 510(k) Premarket Notification

K770477 CITADEL REFERENCE ELECTRODE

CLEARED

Mar 30, 1977
19 days to decision

K770477 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k770477/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Mar 11, 1977
Decision date	Mar 30, 1977
Days to decision	19 days
Third-party review	No

APPLICANT

Company	**Sherwood Medical Industries**
Location	Mchenry, IL, US
Website	https://www.bd.com
510(k) history	62 submissions · 62 cleared · 1976-1980

Sherwood Medical Industries is a medical device manufacturer based in McHenry, US. The company specialized in diagnostic and surgical devices across multiple therapeutic areas. Sherwood Medical Industries received FDA 510(k) clearances from total submissions between 1976 and 1980. The company's cleared devices spanned anesthesiology, chemistry, immunology, obstetrics and gynecology, pathology, ophthalmic, and dental specialties. Notable products included the Argyle Esophageal Stethoscope and Monoject Arterial Blood Sampling Device. This company is inactive and represents ...