The US Food and Drug Administration report on the Manufacturer and User Facility Device Experience – MAUDE

The MAUDE database houses medical device reports submitted to the FDA by mandatory reporters (manufacturers, importers and device user facilities) and voluntary reporters such as health care professionals, patients and consumers.

https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfmaude/search.cfm

Event Type = malfunction
Manufacturer comparators = Zoll, Physio
Product comparators = X Series, LP 15

	2013	2014	2015	2016	2017	Annual Average
Zoll Medical	125	239	311	408	500	**316.6**
Physio Control	1	1	1	3	3	**1.8**