

Philips HeartStart FR2+ AED

Philips is recalling roughly 5400 of its HeartStart FR2+ automated external defibrillators (AEDs) over concerns of a memory chip failure that could leave the devices inoperable, the FDA announced on Monday.

Four models, listed in the press release, are affected by the recall. The chip failure occurred in some devices during self-testing, not emergency use.

The company is working with customers to replace the affected AEDs. Customers may also check the Philips website to see whether their device is implicated in the recall.

Link(s):

FDA safety alert (Free)

<http://click.jwatch.org/cts/click?q=227%3B67289980%3BPLEeZdnJbvF1lTP0aD0TB74RC8HeT5BPMblrSmtOqgg%3D>

Manufacturer news release (Free)

<http://click.jwatch.org/cts/click?q=227%3B67289980%3BPLEeZdnJbvF1lTP0aD0TB0nClXdez5HAMblrSmtOqgg%3D>

Manufacturer product recall site (Free)

<http://click.jwatch.org/cts/click?q=227%3B67289980%3BPLEeZdnJbvF1lTP0aD0TB6nBq9c3hFT5MblrSmtOqgg%3D>