Appendix A

FDA Social Media Posts linking to Safety Communication

FDA

593,831 followers

1y

We are alerting people who use Philips Respironics ventilators, BiPAP, and CPAP machines and their health care providers that Philips Respironics has recently recalled certain devices due to potential health risks. The sound abatement foam may break down potentially entering the device's air pathway, which could release black debris from the foam or certain chemicals that may be inhaled or swallowed by the person using the device.

If you use one of these affected devices, talk to your health care provider to decide on a suitable treatment for your condition and follow the recommendations listed in the safety communication: https://go.usa.gov/x6AZN

Posted June 30, 2021

FDA

593,832 followers

12mo

Today the FDA issued a notification order to Philips Respironics requiring the company to notify patients and others of the company's June 14, 2021, recall of certain Philips Respironics ventilators, CPAP, & BiPAP machines. Ensuring patients and providers have important information regarding the recall of these critical devices is a top priority for the FDA. Find out more: https://go.usa.gov/xzXVZ

Posted March 10, 2022

FDA Medical Devices
@FDADeviceInfo

Today, the @US_FDA provided the latest information about #MedicalDevice reports that are reportedly associated with the breakdown or suspected breakdown of the foam used in the Philips ventilators, BiPAP machines, and CPAP machines. Find out more. fda.gov/medical-device...

Posted November 22, 2022