California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Philips North America Llc Microstream Advance Neonatal-Infant Intubated CO2 Filter Line and Related Products

Recall Date	Product Description	Recalling Firm	Recall Reason
05/05/2025	Microstream Advance Neonatal-Infant Intubated CO2 Filter Line marketed as Adt/Pedi Intub CO2 Line STerm , Adt/Pedi Intub CO2 Line STerm Lng , Adt/Pedi Intub CO2 Line High Humidity , Adt/Pedi Intub CO2 Line LTerm Lng , Adt/Pedi Intub CO2 Line LTerm , Neo/Inf Intub CO2 Line LTerm , Neo/Inf Intub CO2 Line High Humidity , Neo/Inf Intub CO2 Line LTerm Lng , VitaLine H Set Adult/Pediatric , VitaLine H Set Infant/Neonatal , FilterLine Set Long Adult/Pediatric , FilterLine H Set Long Adult/Pediatric , FilterLine H Set Long Infant/Neonatal , Trade Compliant: FilterLine H, Adult/Ped , Trade Compliant: FilterLine H, Infant/Neo , FilterLine Set Adult/Pedi , FilterLine H Set Adult/Pedi and FilterLine H Set Infant/Neonatal. Carbon dioxide gas analyzer.	Philips North America Llc	Disconnection of the adapter from the endotracheal tube can potentially be difficult or not possible.

Recall Class	Product Identification	Distribution	Affected Dates
II	Model No. 989803204511, 989803204321, 989803204301, 989803204331, 989803204521,	32,355 Distributed in California	April 2025 and Prior

	989803204531, 989803204311, 989803204341, 989803159571, 989803159581, 989803160241, 989803160251, 989803160261, 989803182921, 989803182931, 989803105531, 989803105541, 989803105561; System Codes 989803105531, 989803105541, 989803105561, 989803159581, 989803160241, 989803160251, 989803160261, 989803182921, 989803182931, 989803204301, 989803204311, 989803204321, 989803204331, 989803204341, 989803204511, 989803204521, 989803204531, 989803159571, 989803159581, 989803160241, 989803160251, 989803160261, 989803182921, 989803182931, 989803204301, 989803204311, 989803204321, 989803204331, 989803204341, 989803204511, 989803204521, 989803204531, M1920A, M1921A, M1923A. UDI: 20884838093137, 20884838093601, 20884838093809, 20884838091836, 20884838091850, 20884838093564, 20884838091867, This issue is not confined to a lot number or serial number.		

For additional information, please visit the FDA Website.