

The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

Description	Lot #	UPC
ADM 20 DRP INTEGRAT CLAVE 3011	4794307; 4806855; 4849220; 4861176; 4734249; 4877472	
CHEMOLOCK 20 DR CH CL3020ICU50	4763384; 4800367; 4866943; 4908482	88770907292
CHEMOLOCK 30 SET DRIP 3011 50	4868079; 4868080; 4808098; 4803047; 4735699; 4720479; 4724743; 4775051	
CHEMOLOCK ADMIN SET	4808060; 4724700; 4848204; 4787228; 4775030; 4758414; 4742639	88770907518
DILUENT SET 47" 50 ML CH4009	4853711; 4894570	84061908276
ONCOKIT SP SPR/CL CL3941ICU50	4790865	88770907918
ONCOKIT W/VNT CAP CL3946ICU50	4848213; 4735669; 4742630; 4758441; 4764898; 4787232; 4867995	88770908029
ONCOLOGY CLAVE UNI VL SPK 3546	4749463	
ONCOLOGY KIT W/ CLAVE CH3507	4749895; 4800302	
ONCOLOGY KT W/5 CAP 3538 50	4724728; 4765922; 4867983	
ONCOLOGY KT W/CAP CL3948ICU50	4794503; 4743901; 4765927; 4775070; 4777973	88770908126
SPIN SPIR CLS M LUER 2000S 100	4895432; 4800185; 4726692; 4756786	
SPIN SPIR MALE LUER 2000S-C 50	4757618; 4858381; 4849195; 4713514; 4806838; 4806832; 4763178; 4750952; 4750950; 4750947; 4749755; 4713516; 4749750; 4895446	
SPIRS MALE LR 2000SC-10 10UN	4858387; 4726709; 4756789; 4761899; 4783763; 4962299	

On 31 August 2020, ICU Medical Inc. issued an Urgent Medical Device Recall letter informing customers of a recall of the potential for leaks to occur with the Spinning Spiros Male Luer in certain lots. As a result of continued evaluation, ICU Medical is expanding the scope of the

previously distributed Urgent Medical Device Recall to include additional list and lot numbers that may be affected by this issue.

This recall is to the retail level. Affected product started shipping January 2020.

NOTE: Additional products not listed here are impacted by this recall. For information, contact ICU Medical:

ICU Medical Contact	Contact Information	Areas of Support
Global Complaint Management	1-844-654-7780 or ProductComplaintsPP@icumed.com .	To report adverse events or product complaints
Customer Service	1-866-829-9025, option 8; or customerservice@icumed.com	Additional information or assistance
	(M-F, 8am-6pm CT)	