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CLINICAL BULLETIN

Bulletin #	Title			Date Issued
72	BodyGuard Infusion Pump Recall			September 22, 2020
Superseded	**Released By**	**Approved By**	**Source**	**For More Information**
	FSTEMS	Dr. Joey Scollan, DO	FDA	FST/EMS (603) 223-4200

We have become aware of a recent updated infusion pump recall regarding the BodyGuard Infusion Pump System from CME America. It identifies a risk of over and under infusion. The FDA has identified this as a Class I recall, the most serious type of recall. Use of these devices may cause serious injury or death.

You can find the FDA link to this recall at: https://www.bd.com/en-us/support/recall-notifications/recall-notification---cme-america-bodyguard-infusion-pump-system

Additionally, you can find more information from CME American's website at: https://www.bd.com/en-us/support/recall-notifications/recall-notification---cme-america-bodyguard-infusion-pump-system