FDA report “Sagent Pharmaceuticals, Inc. announced the voluntary nationwide recall of two lots of Atracurium Besylate Injection, USP, 50mg/5mL single-dose vials (NDC 25021-659-05) and four lots of Atracurium Besylate Injection, USP, 100mg/10mL multi-dose vials (NDC 25021-672-10) manufactured by Emcure Pharmaceuticals Ltd. and distributed by Sagent. Sagent has initiated this voluntary recall of Atracurium Besylate Injection, USP, 50mg/5mL and 100mg/10mL to the user level due to FDA observations pertaining to aseptic and GMP practices at the manufacturer’s site potentially impacting product sterility.”

Recall of Atracurium Besylate injection by Sagent Pharmaceuticals http://ctt.ec/x12nj+ @ivteam #ivteam

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