FDA report “Hospira notified the public of a nationwide recall of seven lots of Propofol Injectable Emulsion, 1%, 200 mg/20 mL (10 mg/mL) to the user level due to a glass defect located on the interior neck of the vial. The defect was identified during a sample inspection where the glass vial contained visible embedded metal particulate. Free-floating metal particulates were also identified in vials upon further analysis.”

Other intravenous and vascular access resources that may be of interest (External links – IVTEAM has no responsibility for content).

Guide for intravenous chemotherapy and associated vascular access devices from Macmillan.
CancerUK IV chemotherapy information.