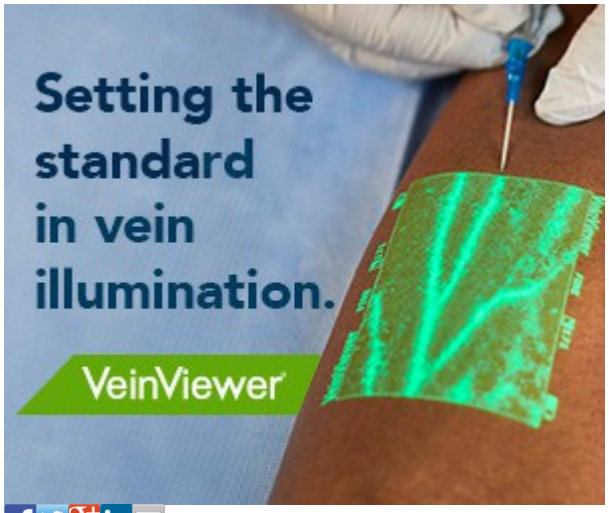

Intravenous products: FDA report “Pentec Health, Inc. initiated a limited, voluntary recall of in-date nutritional prescriptions for renal patients due to lack of sterility assurance associated with one of its laminar flow hoods used in compounding. Pentec Health has received no reports of injury or illness associated with any of the prescriptions subject to this recall. However, because patients are at increased risk of infection in the event a sterile product is compromised, the pharmacy is recalling any unused product whose beyond-use date has not passed.”