SPEAK FACTS

VITAMIN K_1 INJECTION
Phytonadione
Injectable Emulsion, USP

A BLACK BOX WARNING IS THE STRICTEST WARNING PUT IN THE LABELING OF PRESCRIPTION DRUGS OR DRUG PRODUCTS BY THE FDA

Aqueous Dispersion of Vitamin K_1

Ampul R_x only

Protect from light. Keep ampuls in tray until time of use.

VISIT OUR WEBSITE FOR FACT BASED LINKS THAT SUPPORT THIS INFO: PROJECTSPEAK.NET

WARNING — INTRAVENOUS AND INTRAMUSCULAR USE
Severe reactions, including fatalities, have occurred during and immediately after INTRAVENOUS injection of phytonadione, even when precautions have been taken to dilute the phytonadione and to avoid rapid infusion. Severe reactions, including fatalities, have also been reported following INTRAMUSCULAR administration. Typically these severe reactions have resembled hypersensitivity or anaphylaxis, including shock and cardiac and/or respiratory arrest. Some patients have exhibited these severe reactions on receiving phytonadione for the first time. Therefore the INTRAVENOUS and INTRAMUSCULAR routes should be restricted to those situations where the subcutaneous route is not feasible and the serious risk involved is considered justified.

VITAMIN K 1

"BASED ON A REVIEW OF THE LITERATURE, USE OF PARENTERAL VITAMIN K1 MAY RESULT IN SEVERE HYPOTENSION, BRADYCARDIA OR TACHYCARDIA, DYSPNEA, BRONCHOSPASM, CARDIAC ARREST, AND DEATH."
- JOURNALS.SAGEPUB.COM

VIEW THE LEAKED VIDEO ON OUR WEBSITE OF PAC, IL COLLUDING WITH DCFS ON HOW THEY CAN ADMINISTER THIS INJECTION TO BABIES WITHOUT THE CONSENT OF THE PARENTS, WHILE IN THEIR CUSTODY.

SPEAK PROJECT