



California Drug Recall Information



Recall Name

Affymax Recalls OMONTYS (Peginesatide) Injection Due to Possible Hypersensitivity

Recall Date	Product Description	Recalling Firm	Recall Reason
2/23/13	OMONTYS (peginesatide) Injection: <ul style="list-style-type: none"> • 10mg Multi-dose Vials – NDC 64764-610-10 • 20mg Multi-dose Vials – NDC 64764-620-20 	Affymax, Inc. Palo Alto, CA	<i>Potential of serious and life threatening hypersensitivity reactions, including anaphylactic shock, and death.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	OMONTYS (peginesatide) Injection: All Lots Recalled Including: <ul style="list-style-type: none"> • 10 mg multi-dose vials – C18685, C18881, C19258 • 20mg multi-dose vials – C18686, C18696 Product Labels	CA , nationwide	All

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm340893.htm>