From:	General Board of Pharmacy Subscriber List on behalf of Board of Pharmacy
To:	PHARM-GENERAL@DCALISTS.CA.GOV
Subject:	Product Recall - Par Pharmaceuticals
Date:	Thursday, September 29, 2016 10:36:22 AM

Par Pharmaceuticals, Inc. is conducting a drug recall of the drug product lots listed below. The impacted lots of drug products have exhibited a reduction in the level of Ethinyl Estradiol. This recall is to the pharmacy level. Product shipped between December 30, 2014 and July 21, 2016.

Gildess® 1.5/30, NDC 0603-7606-48, 3 blisters x 21 tablets

Gildess® 1.5/30, NDC 0603-7606-15, 6 blisters x 21 tablets

Gildess® 1/20, NDC 0603-7607-48, 3 blisters x 21 tablets

Gildess® 1/20, NDC 0603-7607-15, 6 blisters x 21 tablets

Gildess® FE 1/20, NDC 0603-7609-17, 6 blisters x 28 tablets

Gildess® FE 1.5/30, NDC 0603-7608-17, 6 blisters x 28 tablets

Gildess® 24 FE, NDC 0603-7610-49, 3 blisters x 28 tablets

Gildess® 24 FE, NDC 0603-7610-17, 6 blisters x 28 tablets

Please immediately check your inventory, quarantine, discontinue distribution of the affected product, and contact your wholesaler for directions.

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