The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

Product Description	**Part Number**	**Lot Number**	**Expiration Date on box**	**Genentech Distribution Date**
Susvimo Ocular Implant and Insertion Tool Assembly	UDI 81004259001 GTIN 00810042590014	3456735 3456737 3477671 3480781 3506526 3506531	10/23/2026 10/29/2026 10/31/2026 12/19/2026 02/25/2027 04/15/2027	11/29/2021 to 10/04/2022
Susvimo drug vial and initial fill needle	NDC 50242-0078-55	3499188 3523071	10/31/20022 6/30/2023	11/29/2021 to 10/04/2022

Genentech has initiated a recall of the Susvimo Ocular Implant and Insertion Tool Assembly, including the lot numbers outlined above for the Susvimo (ranibizumaab) drug vial and initial fill needle, which are sold together. This recall should be carried out to the user/healthcare provider level.

Genentech is conducting this recall because the commercial implants do not meet the filed specification for the intended use. A few patients have experienced an issue with the ocular implants that renders it non-functioning. There is a potential of insufficient ranibizumab concentration in the vitreous over time, resulting in disease progression and potential vision loss if the patient does not receive additional treatment to maintain disease control. Additionally, there is an unknown risk of retaining or removing the non-functioning implant. At this time, there are no issues with the drug itself. A pause in all new implantations is required.