January 7, 2019

COM-2019-003

Recall: Temozolomide Capsules recalled

Dear provider of pharmaceutical services,

At PharmPix we are committed to the health and well-being of patients. It is for this reason that we inform you that on January 4, 2019 Amerigen Pharmaceuticals Inc. issued a recalling of 3552 bottles of temozolomide capsules, 20mg. The recall was made because samples failed to meet dissolution specifications according to Food and Drug Administration (FDA) Enforcement Report. The products affected by the recall are detailed on Table 1.

Table 1. Affected products

Product	Affected Lot	Expiration date	NDC
Temozolomide capsules, 20 mg	18B005 A	02/2020	43975-253-05
Temozolomide capsules, 20 mg	18B005 B	02/2020	43975-253-14

The Pharmacy must:

1. Identify if they have the product in inventory and immediately stop using and dispensing the product.
2. Contact all members that in the previous 90 days received the recalled medication and advised them to talk to their doctor.
3. For questions regarding this recall please visit Managed Health Care Connect at: https://www.managedhealthcareconnect.com/content/cancer-drug-recalled-2?page=4,0

For additional information visit:

https://www.managedhealthcareconnect.com/content/cancer-drug-recalled-2?page=3,0

Department of Clinical Pharmacy

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