Acella Pharmaceuticals, LLC. Issues a Voluntary Nationwide Recall of Certain Lots of NP Thyroid®(Thyroid Tablets, USP) Due to Super Potency

Acella Pharmaceuticals, LLC is voluntarily recalling a total of 13 lots of 30mg, 60mg, and 90mg NP Thyroid® to the patient level. The products are being recalled because these lots have been found to be super potent by the manufacturer.

The affected lots of NP Thyroid® include:

Product Description	NDC#	Lot#	Expiry Date
NP THYROID 30MG	42192-0329-01	M329M18-2	11/30/20
		M329J18-3	8/31/20
		M329J18-2	8/31/20
		M329J18-1	8/31/20
		M329H18-1	7/31/20
		M329A19-1	12/31/20
NP THYROID 60MG	42192-0330-01	M330J18-3	8/31/20
		M330J18-2A	8/31/20
NP THYROID 90MG	42192-0331-01	M331M18-2	11/30/20
		M331M18-1	11/30/20
		M331J18-2	8/31/20
		M331J18-1	8/31/20
		M331G18-1	6/30/20

What does this mean?

Patients being treated for underactive thyroid, who receive super potent NP Thyroid, may experience signs and symptoms of overactive thyroid which include, but are not limited to, weight loss, heat intolerance, fatigue, muscle weakness, hypertension, chest pain, rapid heart rate, or heart rhythm disturbances. Pregnant

women who take super potent NP Thyroid may also experience negative maternal and fetal outcomes including miscarriage and/or impairment to fetal development.

What you should do:

Patients who are currently taking NP Thyroid from the lots being recalled should not discontinue use without contacting their prescriber for further guidance and/or a replacement prescription.

Patients with questions about the recall can email Acella Pharmaceuticals at recall@acellapharma.com or contact Acella Customer Service at 1-800-541-4802, **Monday – Friday, 9:00am – 5:00 pm, EST**.

Patients should contact their healthcare provider if they have experienced any problems that may be related to taking or using this drug product.

Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax.

- Complete and submit the report Online: www.fda.gov/medwatch/report.htm
- Regular Mail or Fax: Download form or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178

This recall is being conducted with the knowledge of the U.S. Food and Drug Administration.

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