



Mitsubishi Tanabe Pharma America, Inc.  
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June 8<sup>th</sup>, 2021  
Vermont Attorney General’s Office  
Via Email ([AGO.highcostprescriptiondrugs@vermont.gov](mailto:AGO.highcostprescriptiondrugs@vermont.gov))

**RE: New Prescription Drug – 30 Day Report**

In accordance with 18 V.S.A. § 4637, Mitsubishi Tanabe Pharma America, Inc. (MTPA) is reporting information regarding the commercial introduction of the following new NDC at a wholesale acquisition cost that exceeds the threshold set for a specialty drug under the Medicare Part D program. Per 18 V.S.A. § 4637(d), MTPA has limited the information reported to that which is otherwise in the public domain or publicly available.

NDC	PRODUCT
70510-2201-02	Exservan (riluzole) oral film, 50 mg

**(1) A description of the marketing and pricing plans used in the launch of the new drug in the United States and internationally:**

This information is not otherwise in the public domain or publicly available.

**(2) The estimated volume of patients who may be prescribed the drug:**

This information is not otherwise in the public domain or publicly available.

**(3) Whether the drug was granted breakthrough therapy designation or priority review by the FDA prior to final approval:**

The drug was not granted breakthrough therapy designation or priority review by the FDA prior to final approval.

**(4) The date and price of acquisition if the drug was not developed by the manufacturer:**

January 21, 2021

Please do not hesitate to contact us if you have any questions.

Best Regards,

**Ekaterina Miteiko**

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