

**To:** Office of Attorney General  
[AGO.highcostprescriptiondrugs@vermont.gov](mailto:AGO.highcostprescriptiondrugs@vermont.gov)

**From:** Mylan Pharmaceuticals Inc.  
781 Chestnut Ridge Road  
Morgantown, West Virginia 26505

**Date:** September 5, 2018

**Re:** 18 V.S.A § 4637

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In compliance with 18 V.S.A. § 4637, on August 10, 2018 Mylan Pharmaceuticals Inc. (“Mylan”) provided written notice to the Office of the Attorney General that it introduced a new prescription drug, Tadalafil Tablets USP, 20mg (“the Product”), to the commercial market on August 9, 2018 at a wholesale acquisition cost that exceeds the threshold set for a specialty drug under the Medicare Part D program.

This letter provides the additional required information by 18 V.S.A. § 4637(c) regarding the Product. Mylan notes that the Office of the Attorney General has not yet prescribed a format for submissions under this section. Further, as authorized by 18 V.S.A. § 4637(d), Mylan has limited the information reported to that which is in the public domain or publicly available.

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

*The Wholesale Acquisition Cost (WAC) for the product in the United States is below:*

Product	Package Size	NDC	WAC <sup>[1]</sup>
Tadalafil Tablets USP, 20mg	30	0378-6976-93	\$1,827.65

*Discounted prices negotiated with customers are confidential. There are no marketing or pricing plans in the United States in the public domain or publicly available. The Product has not yet launched in any international jurisdiction.*

- (2) the estimated volume of patients who may be prescribed the drug;

*No information responsive to this request is 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 Product was not granted breakthrough therapy designation or priority review by the FDA.*

- (4) the date and price of acquisition if the drug was not developed by the manufacturer.

*The Product was not a result of an acquisition.*