NEWS RELEASE

Impax Provides Update on Epinephrine Injection, USP Auto-Injector

8/30/2016

-- Epinephrine Injection, USP Auto-Injector, 0.15 mg and 0.3 mg is Currently Available in Pharmacies across the United States --

HAYWARD, Calif., Aug. 30, 2016 /PRNewswire/ -- **Impax Laboratories, Inc.** (NASDAQ: IPXL) today provided an update and additional information to patients, physicians and customers on its **epinephrine injection, USP auto-injector**, 0.15 mg and 0.3 mg., the authorized generic of Adrenaclick®.

Epinephrine injection, USP auto-injector (also called epinephrine auto-injector) is an emergency injection (shot) of epinephrine used for the treatment of life-threatening allergic reactions known as anaphylaxis.

"With all the recent news related to epinephrine auto-injection products, we are increasing our mission to inform patients, caregivers and the professional community regarding the availability of this epinephrine auto-injector product," said Fred Wilkinson, President and Chief Executive Officer of Impax. "Our epinephrine auto-injector represents a proven, low-cost treatment for patients in the U.S. who require the use of epinephrine products."

The auto-injector is conveniently packaged with numbered and color-coded instructions and is designed for single-dose use by patients and caregivers in an anaphylactic emergency. It is available in a package of two injectors and trainers are available upon request. For more information about the Impax' epinephrine auto-injector and its $0 co-pay program, please visit: **http://epinephrineautoinject.com/** or call 855-449-4712.

About Epinephrine Injection, USP Auto-Injector

Epinephrine injection, USP auto-injector (also called epinephrine auto-injector) is an emergency injection (shot) of epinephrine used for the treatment of life-threatening allergic reactions known as anaphylaxis. Before you use the epinephrine auto-injector, tell your healthcare provider about all your medical conditions. With numbered and

color-coded instructions, epinephrine auto-injector is designed for single-dose use by patients and caregivers in an anaphylactic emergency and does not take the place of emergency medical care. After using epinephrine auto-injector go to your doctor or emergency room right away for more medical treatment. For full Prescribing Information see **http://epinephrineautoinject.com/pdf/Prescribing-Information.pdf**. For more information about epinephrine injection, USP auto-injector, ask your doctor or call 1-888-894-6528.

"Safe Harbor" statement under the Private Securities Litigation Reform Act of 1995:
To the extent any statements made in this news release contain information that is not historical; these statements are forward-looking in nature and express the beliefs and expectations of management. Such statements are based on current expectations and involve a number of known and unknown risks and uncertainties that could cause the Company's future results, performance, or achievements to differ significantly from the results, performance, or achievements expressed or implied by such forward-looking statements. Such risks and uncertainties include, but are not limited to: fluctuations in revenues and operating income; the Company's ability to successfully develop and commercialize pharmaceutical products in a timely manner; reductions or loss of business with any significant customer; the substantial portion of the Company's total revenues derived from sales of a limited number of products; the impact of consolidation of the Company's customer base; the impact of competition; the Company's ability to sustain profitability and positive cash flows; any delays or unanticipated expenses in connection with the operation of the Company's manufacturing facilities; the effect of foreign economic, political, legal, and other risks on the Company's operations abroad; the uncertainty of patent litigation and other legal proceedings; the increased government scrutiny on the Company's agreements with brand pharmaceutical companies; product development risks and the difficulty of predicting FDA filings and approvals; consumer acceptance and demand for new pharmaceutical products; the impact of market perceptions of the Company and the safety and quality of the Company's products; the Company's determinations to discontinue the manufacture and distribution of certain products; the Company's ability to achieve returns on its investments in research and development activities; changes to FDA approval requirements; the Company's ability to successfully conduct clinical trials; the Company's reliance on third parties to conduct clinical trials and testing; the Company's lack of a license partner for commercialization of NUMIENT™ (IPX066) outside of the United States; impact of illegal distribution and sale by third parties of counterfeits or stolen products; the availability of raw materials and impact of interruptions in the Company's supply chain; the Company's policies regarding returns, allowances and chargebacks; the use of controlled substances in the Company's products; the effect of current economic conditions on the Company's industry, business, results of operations and financial condition; disruptions or failures in the Company's information technology systems and network infrastructure caused by third party breaches or other events; the Company's reliance on alliance and collaboration agreements; the Company's reliance on licenses to proprietary technologies; the Company's dependence on certain employees; the Company's ability to comply with legal and regulatory requirements governing the healthcare industry; the regulatory environment; the effect of certain provisions in the Company's government contracts; the Company's ability to protect its intellectual property; exposure to product liability claims; risks relating to goodwill and intangibles; changes in tax regulations; the Company's ability to manage growth, including through potential acquisitions and investments; the risks related to

the Company's acquisitions of or investments in technologies, products or businesses; the restrictions imposed by the Company's credit facility and indenture; the Company's level of indebtedness and liabilities and the potential impact on cash flow available for operations; uncertainties involved in the preparation of the Company's financial statements; the Company's ability to maintain an effective system of internal control over financial reporting; the effect of terrorist attacks on the Company's business; the location of the Company's manufacturing and research and development facilities near earthquake fault lines; expansion of social media platforms and other risks described in the Company's periodic reports filed with the Securities and Exchange Commission. Forward-looking statements speak only as to the date on which they are made, and the Company undertakes no obligation to update publicly or revise any forward-looking statement, regardless of whether new information becomes available, future developments occur or otherwise.

About Impax Laboratories, Inc.

Impax Laboratories, Inc. (Impax) is a specialty pharmaceutical company applying its formulation expertise and drug delivery technology to the development of controlled-release and specialty generics in addition to the development of central nervous system disorder branded products. Impax markets its generic products through its Impax Generics division and markets its branded products through its Impax Specialty Pharma division. Additionally, where strategically appropriate, Impax develops marketing partnerships to fully leverage its technology platforms and pursues partnership opportunities that offer alternative dosage form technologies, such as injectables, nasal sprays, inhalers, patches, creams and ointments. For more information, please visit the Company's Web site at: **www.impaxlabs.com**.

Company Contacts:

Mark Donohue
Investor Relations and Corporate Communications
(215) 558-4526
www.impaxlabs.com

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