Nexus Pharmaceuticals Receives FDA Approval for Isoproterenol Injection

Vernon Hills, Ill., August 3rd, 2017 – Nexus Pharmaceuticals announced today that it has received approval from the U.S. Food and Drug Administration (FDA) for its Abbreviated New Drug Application (ANDA) for Isoproterenol Hydrochloride Injection. Nexus Pharmaceuticals' Isoproterenol Hydrochloride Injection is available as a single dose vial containing 0.2 mg per 1mL or 1 mg/5 mL and is an AP Rated generic equivalent of Isuprel®

Isoproternol HCl Injection is indicated for mild or transient episodes of heart block that do not require electric shock or pacemaker therapy, for serious episodes of heart block and Adams-Stokes attacks (except when caused by ventricular tachycardia or fibrillation), for use in cardiac arrest until electric shock or pacemaker therapy, the treatments of choice, is available, for bronchospasm occurring during anesthesia and as an adjunct to fluid and electrolyte replacement therapy and the use of other drugs and procedures in the treatment of hypovolemic and septic shock, low cardiac output (hypoperfusion) states, congestive heart failure, and cardiogenic shock.

About Nexus Pharmaceuticals Inc.

Nexus Pharmaceuticals (www.nexuspharma.net) is a U.S. based healthcare company that specializes in developing and marketing generic sterile injectable products. Through our high quality generic products, Nexus Pharmaceuticals is committed to providing patients with affordable prescription medicines that lower healthcare costs and provide a better quality of life. The company's headquarters is in Vernon Hills, Illinois.

Isuprel® is a registered trademark of Hospira Inc.

Media Contact
Nexus Pharmaceuticals Inc.
Ayesha Ahmed
847-996-3790
aahmed@nexuspharma.net