Attention-Deficit Hyperactivity Disorder (ADHD)

Comprehensive ADHD API Portfolio of Active Pharmaceutical Ingredients Optimized for Finished Dosage Forms

- High-purity Amphetamines & Dextroamphetamines,
 Methylphenidate Hydrochloride, Dexmethylphenidate
 Hydrochloride, and Lisdexamfetamine Dimesylate.
- Custom API particle sizes for optimal product processability and performance.
- Proven supply chain with multisite API manufacture at cGMP facilities.





Advantages of Partnering with Noramco in ADHD

Noramco helps you advance formulations in line with your commercial requirements. Whether your goal is making ADHD products easy to swallow, or designing novel dosage forms to improve patient compliance and extend the lifecycle of your intellectual property, Noramco partners with you to supply the API properties best suited to your products. We optimize the particle size of your API, and provide the flexibility you require to produce a range of oral dosage forms.

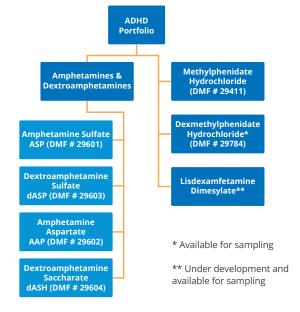
Regulatory filings covering the USA, EU, Canada and other selected countries are available for reference. Noramco also supplies high-purity reference standards for analytical method development.

Technical services include, but are not limited to:

- In-depth API technical package to help support R&D, and filings.
- Regulatory support to help you secure approval of your dosage forms.

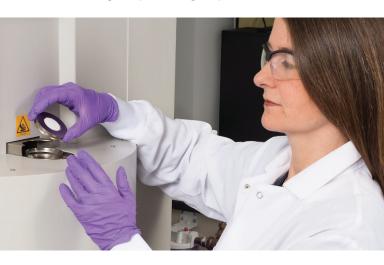
ADHD Portfolio

Noramco supplies all ADHD technologies on a commercial scale along with a guarantee of superior quality and long-term stability. Our ADHD technology offering includes:



Global Supply of ADHD APIs

Noramco has synthesis processes in place at several cGMP production facilities to reliably supply the world market with all ADHD products in a particle size distribution best suited to specific oral dosage forms. A supply chain management system based on multisite production provides the assurance of continuous supply for all ADHD products that best support your performance and manufacturing (i.e., processing) requirements.



A Dedicated Team to help you Manage your Commercial Product Objectives

Are you evaluating a range of oral dosage forms to serve a specific patient population better? Do you require custom API solutions along with regulatory assistance to get to market with novel dosage forms? Ask us how we leverage more than 30 years of expertise in controlled-substance development, licensing and scale up to help you secure your commercial objectives.

- We provide a dedicated team with the skills to provide custom API solutions from concept to commercialization.
- Full support with DEA, FDA & other regulatory bodies, including EDQM, Health Canada.
- International logistics team with technical and commercial presence in Europe & USA.
- Long-term capacity planning with life-cycle commitment to cost reduction.



The Noramco Difference

Find out how a controlled-substance supply chain development program from Noramco helps you go to market faster, with less complexity. Partner with us from the API development phase, to a preformulation and formulation development solution, and obtain the benefits of an integrated team that support best overall outcomes.

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