HIRAY NEWS

Manufacturing License Granted!

Jan 21, 2022

Hiray Pharma Solutions state-of-the-art manufacturing facility in Jingmen, China was granted its license for the manufacture of pharmaceutical products by the Hubei Provincial regulatory authorities on January 19, 2022. Take a virtual tour of our site!

Obtaining our pharmaceutical products manufacturing license at our newest site means that Hiray Pharma Solutions has been recognized by the Chinese government as qualified to produce active pharmaceutical ingredients (APIs) at Jingmen, China, greatly expanding our capacity to serve clients around the world.

In order to reach this milestone, Hiray worked hard to meet and exceed standards in five areas: quality management, personnel, facilities, analytical capability, and quality assurance. Our quality management system has been recognized to meet the requirements of the GMP standard. Hiray Pharma Solutions' quality management team is responsible for the supervision of drug quality in the whole production process, and for regulatory filings of drug production. Our management personnel

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and engineering technicians at all levels have the necessary training and experience to make correct judgments and deal with practical problems. Our facility boasts a clean environment, with cutting edge pollution controls. Air and water quality meet all requirements of drug production. Our production equipment is adapted to the drugs produced and is reasonably arranged to support the required technological process. Equipment, instruments, and meters are managed by specially trained personnel, regularly maintained, and are registered and archived. In terms of hygiene, Hiray Pharma Solutions complies with all hygiene requirements, and the factory area and workshop are kept clean. Production staff have regular physical examinations every year with established health records. Health education for drug production staff is regularly carried out. Our product management team formulates the technological process and operating procedures of each process according to the legal quality standards.

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