China National Medical Products Administration (NMPA)
国家药品监督管理局

Address: No.1 North Luyuan, Exhibition Road, Xicheng Qu, Beijing 100037 | **Phone:** 86-10-6831-1166 | **Website:** http://www.nmpa.gov.cn

The National Medical Products Administration (NMPA), a vice-ministerial level body under the State Administration for Market Regulation (SAMR), is responsible for creating and supervising the implementation of policies, plans and standards governing the quality and safety of drugs, cosmetics, and medical devices. NMPA oversees standards setting, registration, and quality management of drugs, cosmetics and medical devices, as well as post-market inspection and risk management and registration of licensed pharmacists.

SAMR Leading Party Group Member		Commissioner	
Li Li **李利** **NMPA Deputy Commisioner**		**Jiao Hong** **焦红**	

Deputy Directors		
Xu Jinghe 徐景和 (Deputy Commissioner) Oversees the Department of Polic and Regulations, Departments of Medical Device Registration and Regulation, and Department of Science, Technology and International Cooperation. In charge of the Center for Drug Reevaluation (National Center for ADR Monitoring), Center for Medical Device Evaluation, International Communication Center, Southern Medical Economic Research Institute, Complaints and Reporting Center.	**Chen Shifei 陈时飞** (Deputy Commissioner) Oversees Departments of Drug Registration and Regulation, and Department of Human Resources. In charge of National Institutes for Food and Drug Control, Center for Drug Evaluation, Chinese Pharmacopoeia Commission, Center for Licensed Pharmacists, 146 Warehouse (responsible for the acquisition, allocation and storage of national special drugs), and Center for Drug Examination and Inspection Center.	**Yan Jiangying 颜江瑛** (Deputy Commissioner) Oversees the Department of Comprehensive Affairs, Planning, and Financial Affairs, Department of Cosmetics Regulation, Party Committee, and Bureau of Retired Officials. In charge of Advanced Research Institution, News Center, Media Group, Service Center, Information Center, Chinese Pharmaceutical Association.

Department Responsibilities		
Comprehensive Affairs, Planning, and Financial Affairs (综合和规划财务司) Responsible for administrative, government and financial affairs, as well as media and publicity. Drafts working plans and oversees implementation; responsible for security and information technology; manages state-owned assets and finances of subordinate units; responsible for auditing, archival work, and organizing conferences.	**Policy and Regulations (政策法规司)** Researches major policies, drafts laws, regulations and standards, and provides legal review of relevant regulatory documents. Responsible for publicizing related laws and standards, and overseeing the coordination of administrative and law enforcement. Responsible for the promotion of law and related work involving the World Trade Organization. Undertakes the coordination work related to comprehensively deepening reform. Responsible for the routine office work of the vaccine quality management system (QMS).	**Drug Registration (TCM and Ethno-Medicines Regulation) (药品注册管理司,中药民族药监督管理司)** Drafts regulations and technical guidelines for drug standards and supervises enforcement. Oversees implementation of quality control standards for clinical trials and non-clinical research, and Traditional Chinese Medicines; responsible for R&D facility inspections and investigation of illegal activities; participates in the formulation and implementation of the National Essential Drugs List System.
Drug Regulation (药品监督管理司) Drafts and supervises the implementation of quality management standards for drug production, operation and use. Guides manufacturing facility inspections and conducts quality sampling inspections and reporting; investigates and punishes serious illegal acts; monitors adverse drug reaction (ADR) reporting and response; supervises radioactive drugs, narcotic drugs, toxic drugs, psychotropic drugs and precursor chemicals; guides and supervises the management of batch issuance of biological products.	**Medical Device Registration (医疗器械注册管理司)** Drafts and supervises the implementation of standards for medical device registration. Establishes classification, naming and coding rules; sets quality management standards and technical guidelines for clinical trials; oversees the medical device registration management system. Responsible for R&D facility inspections and investigating and punishing illegal acts.	**Medical Device Regulation (医疗器械监督管理司）** Drafts and supervises the implementation of quality management standards for medical device production, operation and use. Guides manufacturing facility inspections and conducts quality sampling inspections and reporting; investigates and punishes serious illegal acts; monitors ADR reporting and response.
Cosmetics Regulation (化妆品监督管理司) Drafts and supervises the implementation of quality management standards for the registration of cosmetics. Responsible for setting classification rules and technical guidelines; developing a cosmetics inspection system; guiding manufacturing facility inspections and reporting; investigation and punishing serious illegal acts; monitoring ADR reporting and response.	**Science, Technology and International Cooperation (Hong Kong, Macao and Taiwan Office) (科技和国际合作司,港澳台办公室)** Researches scientific tools and methods for the review, supervision and management of drugs, medical devices and cosmetics. Drafts policies and standards for the management and use of new technologies and products; laboratory construction and management; qualifications for inspection and testing institutions and inspection norms. Manages major scientific and technological projects. Manages foreign-related affairs. Coordinates and participates in the formulation of international regulatory rules and standards.	**Human Resources (人事司)** Human resource and talent management; manages qualification process for licensed pharmacists.

Department Responsibilities		
Party Committee (机关党委） Responsible for party building and related work of the NMPA administration and its directly affiliated units in Beijing.	**Bureau of Retired Officials (离退休干部局)** Responsible for the work of retired cadres.	

Information is accurate as of May 2021.
Source: China National Medical Product Administration