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

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The National Medical Product 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 Party Secretary		Director	
Li Li **李利** **NMPA Deputy Director**		**Jiao Hong** **焦红**	

Deputy Directors		
Xu Jinghe 徐景和	**Chen Shifei 陈时飞**	**Yan Jiangying 颜江瑛**

Department Responsibilities		
Comprehensive and Financial Planning (综合规划和财务司) 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, Law and Regulation (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.	**Drug Registration (TCM and Ethno-Medicine Supervision) (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 Supervision and Administration (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.	**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 Supervision and Administration (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 Supervision and Administration (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.	**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; manages foreign-related affairs and participates in the formulation of rules and standards for international supervision. 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.	**Personnel (人事司)** Human resource and talent management; manages qualification process for licensed pharmacists.

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

Information is accurate as of September 2019.
Source: China National Medical Product Administration