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(NMPA) and the National Health Commission of the People ' s Republic of China (NHC) released an announcement on promulgating the 2020 edition of the Pharmacopoeia of the People ' s Republic of China (hereinafter referred to as Chinese

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Pharmacopoeia). This is the 11th edition of Chinese Pharmacopoeia and will be implemented officially from December 30, 2020.

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The 2020 edition of the Chinese

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Page 38/78

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dition knowledge of the matter said. Financial details of the possible transaction have not been firmed up and Ant, the Alibaba-backed payments-to-consumer ...

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revised; Volume III contains
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pharmaceutical companies
producing medicines for the
Chinese market.

The is the 2005 English edition of
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compiled by the State
Pharmacopoeia Commission of the

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Ministry of Public Health. The data is effective from July 2005 and includes information on standards of purity, descriptions and dosage, with a total of over 3200 monographs of drugs with 525 new admissions. The publication is in three volumes: volume one

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Edition 2 contains 1146 monographs of drugs including Chinese materia medica and traditional patent medicines; volume two contains 1967 monographs relating to chemical drugs, antibiotics, biochemicals, radiopharmaceuticals and and excipients for

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pharmaceutical use; and volume three has 101 monographs on biological products.

This book is the first volume of a comprehensive 2-volume book covering modern pharmacological and clinical studies of the most

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Edition 2 of herb. The description on each drug includes an introduction (source, character and taste, actions and indications according to traditional Chinese medicine etc.), chemical composition, pharmacology, clinical studies, adverse effects and references.

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The translation of the book into English was done by qualified professionals in the field and the terms used are consistent with those used in Index Medicus, Chemical Abstracts and Botanical journals.

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determination of residual solvents
in pharmaceutical packaging
materials. This method is based on
the gas - solid balance. Take a

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certain area of the sample. Place into a sealed container. Under a certain temperature and time conditions, the organic solvent resided in the sample is heated and evaporated. After the equilibrium is reached, take headspace to quantitative inject into the gas

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Chromatograph for analysis. Use the maintaining time to perform qualification. Use the peak area to perform quantitation. Determine according to the determination of residual organic solvents (Part 2, Chinese Pharmacopoeia 2010 edition).

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Application of Western analytical methods. It continues with an examination of the concept of synergistic, multi-targeted action among components, followed by an analysis of how analytical chemistry and other methods are used to assure quality and

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Uniformity of natural products within tolerable limits. It then addresses the complexity introduced by absorption, distribution, and metabolism of an already complex drug. The last chapter advocates a system biological approach as the way

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and promulgated earlier heterogeneous medical manuscripts belonging to various traditions the government established unique imperially sponsored medical institutions to handle public health and other aspects of medicine. As the first

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medicine underwent during the
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