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Who gmp guidelines for ayurvedic medicines

Migraine	Cafergot Sumatriptan/Imtrex Verapamil	\$1.20 \$13.56(max) \$0.13	\$7.20 \$162.67 \$0.38	Feverfew extract	\$0.18	\$0.55
Milk Deficiency	Reglan	\$0.73	\$2.91	Fenugreek	\$0.08	\$0.24
Motion Sickness	Transderm Scop Dramamine	\$5.25 \$0.57	\$5.25 \$2.29	Ginger extract	\$0.19	\$0.39
Myalgia	Motrin	\$0.06	\$0.34	Capsicum extract	\$0.13	\$0.40
Narcolepsy	Cyclobenzabrene (generic)/Dexedrine Modafinil/Provigil	\$0.95 \$87/1.49 \$4.08	\$2.85 \$2.60/4.48	Watergreen Ephedra Yohimbe extract	\$4.25/ 330 \$0.12 \$0.31	\$0.25 \$0.63
Nyctalopia	Bright Life Vit. A	\$0.02	\$0.02	Bilberry extract	\$0.53	\$1.07
Obesity	Prozac	\$2.27	\$2.27	Evening Primrose extract	\$0.59	\$0.59
Osteoporosis	Premarin Fosamax	\$0.45 \$1.97	\$0.45 \$1.97	Stinging Nettle extract	\$0.53	\$1.07
PMS	Motrin Paxil Tizanidine	\$0.06 \$2.34 \$0.72	\$0.34 \$2.34 \$0.72	Chasteberry extract Evening Primrose extract	\$0.40 \$0.59	\$0.80 \$0.59
Prostatic Dysuria	Erythromycin Penicillin Hytrin Proscar	\$0.26 \$0.21 \$1.55 \$2.01	\$1.04 \$0.85 \$1.55 \$2.01	Stinging Nettle extract	\$0.53	\$1.07
Retinopathy				Bilberry extract	\$0.53	\$1.07
Rhinitis	Cromolyn Varenese	\$0.31 \$1.84/gram	\$1.84	Echinacea Extract	\$0.53	\$1.60
SAD				Hypericum extract	\$0.17	\$0.50
Stingles	Acyclovir Famvir	\$2.84 \$5.59	\$14.19 \$16.76	Capsicum extract	\$0.13	\$0.40



GMP guidelines

- ◆ GMP as per Schedule "M"
www.cdscop.nic.in
- ◆ GMP as per WHO
www.who.int
- ◆ GMP as per MCA now known as MHRA
www.mca.gov.uk
- ◆ GMP as per TGA
www.tga.gov.au
- ◆ GMP as per US FDA
www.fda.gov
- ◆ GMP as per ICH guidelines
www.ich.org

22/10/2008 Department of Pharmaceutics 19

ACIDBEN
Efficient against Acid Exposure

INDICATIONS:
Acid reflux, heartburn, indigestion, and stomach discomfort.

CONTRAINDICATIONS:
None.

Prevents Acid Reflux, Heartburn, Indigestion

• 100% Natural
• No Side Effects
• No Drowsiness

BENCOLD PLUS
Efficient against cold, cough, and flu

INDICATIONS:
Cold, cough, flu, fever, and sore throat.

CONTRAINDICATIONS:
None.



3. 9) The standards for identity, purity and force depending on the respective pharmacopias of Ayurveda, Siddha and medicinal medicinal systems published by the government of India have been respected. The duration of the record conservation should be the expiry date of the lot. Buildings The buildings used for the factory are such as to allow the production of drugs in hygienic conditions and should be devoid of cobwebs and insects / rodents. Ayurveda sasa shala v / s gmp Atank Rahite deshe ----- samyakvatayanopeta divyechitravichitra 1 (r.sc.6 / 11-16) rasa shala prakurvita ----- NANA UPKARANUPETA PRAKAAREN SUSHOBHITA L (R.R.S. 7 / 1-8) 4. Record of the receipt, tests and approval or rejection and use of the raw material must be maintained. 20. The main requirements for GMP for herbal medicines are common to GMP for pharmaceutical products. 8) registration of the specific method and the preparation procedure, ie "Bhavana", "marrana" and "Putra" and the record of each process carried out by the manufacturer must be maintained. Batch production records Production records are required to provide a report of the list of raw materials and their quantities obtained from the store, tests conducted during the various phases of manufacturing such as taste, color, the physical characteristics and chemical tests that may be necessary or indicated in the approved books of Ayurveda, Siddha and Honed mentioned in the first program of the law on drugs and cosmetics, 1940 (23 of 1940. 3) The production record should be maintained for various processes. 31. Furthermore, moreover Member States have begun to establish its specific national GMP for herbal medicines. Raw material from the animal source 4. Each container used for conservation of raw materials must adequately identified with the label indicating the name of the raw material, the power source and clearly declares the status of Material like test at 5. 2nd List of factory mechanisms, equipment Description of general production requirements 8. 17. Manisha Goyal Assistant Professor Rasa Shastra Dept. Adequate Performance of water To wash the premises must be made 15. 30. Foot area for the quality control section. 25. - Drying space: There should be separate space for drying raw materials, in the process of medicine or medicinal products require drying before packaging. The quality of finished herbal products is also influenced by the quality of the raw materials used. This volume also contains the content page of the content of "Quality Assurance Compendium Vol. 32. In order to consolidate the information and to carry out a user-friendly technical guide, it was proposed to fill out a WHS monograph On GMP for herbal medicines, which combines two series of technical guidelines. There is no doubt that GMP is a fundamental step to ensure the safety and efficacy of herbal medicines. Concentrated / plant extracts and exudates / resins. The floor and walls should not be damp or wet. These equipment must be installed and managed correctly with proper cleaning. Dr. should have an adequate performance of light and ventilation. 7. Records must be maintained. 12. Workspace The production area must provide adequate space for orderly positioning of equipment and material used in any of the operations for which these used to facilitate easy and safe operation and To minimize or eliminate any risk of mixtures -up between different drugs, raw materials and to prevent the possibility of cross-contamination of a drug from another drug that comes Stored or managed at the same locals. The registry must also be available for inspection during any inspection of the premises. Rasa Shala Veda n Vayav Ishan Prakhshan Ras Bhairav

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