**SARDAR BHAGWAN SINGH P. G INSTITUTE OF BIOMEDICAL SCIENCES & RESEARCH, BALAWALA, DEHRADUN**

**SCHOOL OF PHARMACEUTICAL SCIENCES AND TECHNOLOGY**

**B. PHARM IIIYEAR, VI SEMESTER**

**SUBJECT- HERBAL DRUG TECHNOLOGY (BP 603 T)**

**ASSIGMENT - VI**

**Attempt all questions.**

**I. Fill in the blanks.**

1. The ultimate goal of GMP is …………..
2. The quality control section shall have minimum…………area
3. The premises used for manufacturing, processing, packaging and labelling will be in conformity with the provision of ……………..
4. Procedure of …………… should be adopted for raw materials wherever necessary.
5. The finished goods transferred from the production area after proper packaging shall be stored in the finished goods stored, marked as……………..
6. Recors of sale and distribution of each batch of Ayurveda, Siddha and Unani drugs shall be maintained up to ………….. of the exhausting of stock.
7. The manufacturer shall submit the record related to complaints to the liscensing Authority, once in a period of …………..
8. Each batch record should be maintained irrespective of …………….
9. Raw materials used in the manufacture of drugs are ………..
10. Water needed for manufacturing should be ……………..

**II. Short Answer Type Questions.**

Q1. Describe importance of GMP for manufacturing ASU drugs.

Q2. What are the principle and objectives of GMP?

Q3. Describe facility required for quality control section.

**III. Short Answer Type Questions.**

Q1. Write a detail note on documentation and records maintenance during manufacturing of herbal drugs.

Q2. Describe different components of GMP in detail.