

B. Pharm. 3 Semester Theory Improvement Examination (2020-2021)

Date: 23/09/2020

Semester :3

Subject: Pharmaceutical Microbiology (BP303T)

Instructions:

All questions are compulsory

Each question carry 1 mark.

Provide Parul Email Id only

*Required

* Required

1. Email address *

2. 1. Which is the correct pore size for HEPA filter? *

Mark only one oval.

- 0.2 Micrometer
 0.3 micrometer
 0.8 micrometer
 100 micrometer

3. 2. Which of the following has given a guideline for the cleanroom? *

Mark only one oval.

- FS 209E
 ISO 14644
 EU GMP
 All of the above

4. 3. Which is controlled in cleanroom? *

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- Temperature
- Number of particles of a particular size
- Both
- vapour pressure

5. 4.FS 209E was replaced by *

Mark only one oval.

- ISO14644
- EU GMP
- BOTH
- None

6. 5. Which is an important particle size from a Pharmaceutical viewpoint for a cleanroom *

Mark only one oval.

- 0.5 micrometer and more
- 5 micrometer and more
- both
- 100 micrometer

7. 6. Which specify guideline for the number of particles at rest and during operation *

Mark only one oval.

- EU GMP
 ISO
 FS209E
 All of the above

8. 7. Which is correct? *

Mark only one oval.

- ISO has 3 more cleanroom classes as compared to FS209E
 ISO classification of cleanroom is exactly same to FS209E
 EU GMP does not provide cleanroom classification
 All of the above

9. 8. Class 100 room should not have *

Mark only one oval.

- more than 100 particles of 0.5 micrometer
 more than 100 particles of 5 micrometer
 more than 100 particles of 50 micrometer
 more than 100 particles of 500 micrometer

10. 9. Which type of Cleanroom is relevant to the Pharma Industry *

Mark only one oval.

- ISO1
 ISO 3
 ISO 2
 ISO 5

11. 10. Source of contamination in cleanroom is *

Mark only one oval.

- Man
- Material
- Process
- All of the above

12. 11. The cleanroom classes are determined by the maximum acceptable numbers of particles (by size) in the air *

Mark only one oval.

- per cubic mm
- per square mm
- per mm
- None of above is correct

13. 12. Which parameters are controlled in cleanroom *

Mark only one oval.

- Air Velocity
- Differential pressure
- Air flow pattern (Turbulent or Streamline)
- All of the above

14. 13. Products that are not terminally sterilized *

Mark only one oval.

- are manufactured under aseptic conditions
- are not manufactured under aseptic condition
- are not manufactured
- are sterilized by autoclaving

15. 14. Production of the sterile product should be carried out *

Mark only one oval.

- in clean room
- any where in facility
- z category lab
- None is correct

16. 15. The preferred surfaces for walls in cleanrooms are *

Mark only one oval.

- Plastic
- Epoxy coated fiber
- Both
- None

17. 16. Which is correct about Doors of a cleanroom? *

Mark only one oval.

- Doors should be flushed with the walls
- Door should be fitted with maintaining positive pressure airflow
- Doors must be self-closing.
- All of the above are correct

18. 17. To minimize contamination in cleanroom *

Mark only one oval.

- Human intervention should be less
- Air should be passed through HEPA filters before entering cleanroom
- Both of above should be done
- More and more humans should be allowed to enter

19. 18. ULPA is *

Mark only one oval.

- Ultra low particulate air filters
- Ultra pure particulate filters
- Ultra large particulate filters
- unapproved particulate air filters

20. 19. Laminar Air Flow Chamber can be *

Mark only one oval.

- Horizontal
- Vertical
- Both horizontal and vertical
- inverted , horizontal and vertical

21. 20. LAF unit contains *

Mark only one oval.

- UV light
- HEPA Filter
- Pre filter
- All of the above

22. 21. For given pharmaceutical products, Microbiological Assay is carried out *

Mark only one oval.

- Tablets and capsules
- Parenteral preparations
- Antibiotics, Vitamins and Hormones
- None of the above

23. 22. One of the following is not an official method for Microbiological Assay *

Mark only one oval.

- Cup Plate Method/Disc Diffusion Method
- Turbidimetric Method
- Membrane Filtration Method
- None of the above

24. 23. Following is the test organism for Microbiological Assay of Streptomycin Sulfate *

Mark only one oval.

- Bacillus subtilis
- S aureus
- E coli
- S citrus

25. 24. If Reidal Walker Coefficient (RWC) is 1.8 it will indicate *

Mark only one oval.

- Test disinfectant is less potent than Phenol
- Test disinfectant is more potent than Phenol
- Test disinfectant is as potent as Phenol
- No conclusion

26. 25. Out of four one is not under suspension test for evaluation of disinfectant *

Mark only one oval.

- Reidal Walker Test
- Chick Martin Test
- Surface Film test
- None of the above

27. 26. The test organism for Microbiological Assay of Kanamycine Sulfate is *

Mark only one oval.

- S aureus
- Bacillus subtilis
- E coli
- None of the above

28. 27. One of the following is not a disinfectant *

Mark only one oval.

- Ethyl alcohol
- Iso Propyl alcohol
- Citric acid
- Bleaching Powder

29. 28. One of the following will not affect disinfectant activity *

Mark only one oval.

- Nature of material
- Time duration of contact
- Temperature
- Source of materials

30. 29. The formula to calculate RWC is *

Mark only one oval.

- ratio of the dilution of the test disinfectant that kills a microorganism to the dilution of phenol that kills the organism in the same time under identical conditions.
- ratio of the dilution of the pheno that kills a microorganism to the dilution of test disinfectant that kills the organism in the same time under identical conditions.
- Average of the dilution of the disinfectant that kills a microorganism to the dilution of phenol that kills the organism in the same time under identical conditions.
- None of the above

31. 30. CMC in Disinfectant theory stands for *

Mark only one oval.

- Critical Micelle Concentration
- Chick Martin coefficient
- Chemistry Manufacturing and Control
- None of the above

32. 31. Mechanism of Ethyl alcohol to kill microorganisms is *

Mark only one oval.

- by protein denaturation
- by inhibiting nucleic acid biosynthesis
- by inhibiting cell wall synthesis
- None of the above

33. 32. Click the best answer with respect to Microbiological Assay *

Mark only one oval.

- In microbiological assay , potency of the drug as well as purity can be found out.
- It can be assayed in very minute quantities
- It is very useful to assay antibiotic preparations, vitamins and hormones etc.
- all of the above

34. 33. One of the following is not correct with respect to Cup Plate method of Microbiological Assay *

Mark only one oval.

- Diameter of Zone of inhibition is measured in this method
- Turbidity of the sample is measured
- The media is specific for the product as per IP
- The test organisms are specific for a specific preparations.

35. 34. The unit of Potency of any antibiotic preparation as per IP is *

Mark only one oval.

- Gram Unit
- International Unit (IU)
- Dalton (D)
- Base pairs (bps)

36. 35. The best method to assay Vitamin B12 sample is *

Mark only one oval.

- Cup Plate Method of Microbiological Assay
- Turbidimetric Method of Microbiological Assay
- Membrane Filtration
- Direct Inoculation