No. of Printed Pages: 3

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MVP-004

POST GRADUATE DIPLOMA IN FOOD SAFETY AND QUALITY MANAGEMENT (PGDFSQM)

Term-End Examination December, 2014

| MVP-004 : FOOD SAFETY AND QUALITY MANAGEMENT SYSTEMS | | | | | | |
|--|----------|---|--|--|--|--|
| Tir | ne : 3 l | hours Maximum Marks: 100 | | | | |
| No | | ttempt any five questions. All questions carry qual marks. | | | | |
| 1. | (a) | As per the Clause 4 of ISO 1401, what are the principles of auditing? | | | | |
| | (b) | List the personal attributes and knowledge and skills of an auditor. $5+5=10$ | | | | |
| | (c) | Explain the 2-tier structure of QCI. 5 | | | | |
| 2. | (a) | What are the five conditions that must be considered by the top management regarding quality policy? 5 | | | | |
| | (b) | Briefly describe the hierarchical levels of documentation structure of ISO 9001 : 2000/2008. | | | | |
| | (c) | List three standards under ISO 9000 family with functions of each. 5 | | | | |

| 3. | (a) | What does ISO 22000 : 2005 bring to HACCP method? | 3 |
|----|--------------|--|-----|
| | (b) | Name 5 types of organizations with examples which can use ISO 22000 : 2005. | 5 |
| | (c) | Explain the similarity and difference between OPRP and HACCP. | 5 |
| | (d) | List the elements of structure of FSMS documentation. Explain any two elements. | 7 |
| 4. | (a) | What are the factors that affect the Test Result? | 3 |
| | (b) | Discuss the factors which will enhance a laboratory's capability to do the analysis on a consistent basis. | 4 |
| | (c) | Write a short note on Laboratory Accreditation. | 3 |
| | (d) | What is the importance of Method Validation and Certified Reference Material (CRM) for a testing laboratory? Discuss when should a method be validated and in what activities the CRM is used. 2+4+4= | =10 |
| 5. | (a) | Describe the procedure for certification of India Gap. | 10 |
| | (b) | Discuss the benefits, principles and scope of | |
| | / | BRC Global. 3+4+3: | =10 |

| 6. | Defi | ne any <i>ten</i> of the following terms : | 10×2=20 |
|----|--------------|---|--------------|
| | (a) | Threat | |
| | (b) | Critical Limit | |
| | (c) | Quality | |
| | (d) | Validation | |
| | (e) | Non-conforming Product | |
| | (f) | OPRP | |
| | (g) | Traceability | |
| | (h) | Uncertainty of measurement | |
| | (i) | Proficiency Testing | |
| | (j) | PRP | |
| | (k) | Limit of Detection | |
| | (1) | Limit of Quantitation | |
| 7. | Writ | e short notes on any <i>five</i> of the following | ;: 5×4=20 |
| | (a) | Internal quality control in a laboratory | |
| | (b) | Biosafety Cabinet | |
| | (c) | SOP | |
| | (d) | SQF-1000 | |
| | (e) | IFS | |
| | (f) | Quality Manual | * |
| | (g) | OHSAS | |
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