

# Indian Bio-Medical Skills Consortium

## IBSC Modules

SI No	Subject
1	Anatomy and Physiology
2	Fundamentals of Electricity and Electronics
3	Healthcare Technology Function and Operation
4	Healthcare Technology Problem Solving and Troubleshooting
5	Healthcare Information Technology
6	Healthcare Technology Management
7	Healthcare Safety & Standards
8	NABH & NABL Accreditation
9	Product Development, Testing, Evaluation, & Modification
10	Medical Terminology for Engineers
11	Risk Management / Safety
12	Radiation Safety
13	Medical Device Regulatory
14	Facilities / General Management
15	Service Delivery Management

<b>Detail Course Content</b>	
<p><b>1. Anatomy &amp; Physiology</b></p> <ol style="list-style-type: none"> <li>1. The Human Body: An Orientation</li> <li>2. Cells and Tissues</li> <li>3. Skin and Body Membranes</li> <li>4. Hematological System</li> <li>5. Eye, Ear and Endocrine System</li> <li>6. Skeletal and Muscular System</li> <li>7. Nervous System</li> <li>8. Cardiovascular System</li> <li>9. Respiratory System</li> <li>10. Digestive, Excretory &amp; Reproductive System</li> </ol>	<p><b>2. Fundamentals of Electricity and Electronics</b></p> <ol style="list-style-type: none"> <li>1. Transducers</li> <li>2. Calculations and Conversions</li> <li>3. Active &amp; Passive Devices, Solid State Devices, (Analog &amp; Digital)</li> <li>4. Circuits &amp; Components</li> <li>5. Oscillators</li> <li>6. CRTs, X-Ray tubes, Photomultipliers</li> <li>7. AC Power</li> <li>8. Display devices.</li> <li>9. Test Equipment</li> <li>10. Power Distribution &amp; Storage Systems</li> </ol>
<p><b>3. Healthcare Technology and Function</b></p> <ol style="list-style-type: none"> <li>1. Understand physiological concepts as applicable to healthcare technology (e.g., PEEP sphygmomanometer, manometer, Korotkoff sounds, Einthoven's triangle, 10-20-10 EEG pattern).</li> </ol>	<p><b>4. Healthcare Technology Problem Solving &amp; Troubleshooting</b></p> <ol style="list-style-type: none"> <li>1. Identify and resolve fault conditions of modules/subsystems including power supplies.</li> <li>2. Prioritize repairs of medical devices based on level of risk and/or urgency.</li> </ol>

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<ol style="list-style-type: none"> <li>2. Understand normal function, use, and underlying technology of test equipment (electrical safety analyzer, defibrillator analyzer, electro surgical analyzer, physiologic simulators, DVM, meters).</li> <li>3. Understand normal function and underlying technology of monitoring systems (e.g., EtCO<sub>2</sub>, ECG, EEG, non-invasive blood pressure, invasive blood pressure, pulse oximetry, fetal monitor, respiration).</li> <li>4. Understand normal function and underlying technology of laboratory equipment (e.g., centrifuges, water baths, analyzers, cryostats, microtomes).</li> <li>5. Understand normal function and underlying technology of imaging devices (e.g., Ultrasound, Radiographic / Fluoroscopy).</li> <li>6. Understand normal function and underlying technology of diagnostic equipment (e.g., otoscope, ophthalmoscope, audiometer, uroflow meter).</li> <li>7. Understand normal function and underlying technology of infusion equipment (e.g., feeding pumps, infusion devices, syringe pumps, PCA pumps).</li> <li>8. Understand normal function and underlying technology of life support equipment (e.g., defibrillators, anesthesia machines, ventilators, balloon pumps, external pacemakers).</li> <li>9. Understand normal function and underlying technology of therapeutic equipment (e.g., infant warmers, ultrasound therapy, hypo/hyperthermia, aspirators, SCD, Bilirubin light).</li> </ol>	<ol style="list-style-type: none"> <li>3. Differentiate between a device error and a use error (User Training, Applications) to determine appropriate action.</li> <li>4. Differentiate between an issue with a localized monitoring device on a network and a system-wide problem.</li> <li>5. Identify the fault conditions and apply appropriate corrective action for monitoring systems (EtCO<sub>2</sub>, ECG, EEG, non-invasive blood pressure, invasive blood pressure, pulse oximetry, fetal monitor, respiration).</li> <li>6. Identify the fault conditions and apply appropriate corrective action for laboratory equipment (Centrifuges, Water Baths, Analyzers, cryostats, microtomes).</li> <li>7. Identify the fault conditions and apply appropriate corrective action for diagnostic equipment (otoscope, ophthalmoscope, audiometer, uroflow meter).</li> <li>8. Identify the fault conditions and apply appropriate corrective action for infusion equipment (feeding pumps, infusion devices, syringe pumps, PCA pumps).</li> <li>9. Identify the fault conditions and apply appropriate corrective action for therapeutic equipment (infant warmers, ultrasound therapy, hypo/hyperthermia, aspirators, SCD, Bilirubin light, defibrillators, external pacemakers).</li> <li>10. Identify the fault conditions and apply appropriate corrective action for operating room equipment (electro surgical generators, video equipment, tourniquets, sterilizers, warmers).</li> </ol>
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<p>10. Understand normal function and underlying technology of operating room equipment (e.g., electro surgical generators, video equipment, lasers, tourniquets, sterilizers, warmers).</p>	
<b>5. Healthcare Information Technology</b>	<b>6. Healthcare Technology Management</b>
<p><b>A. Foundations</b></p> <ol style="list-style-type: none"> <li>1. Hardware             <ol style="list-style-type: none"> <li>a. Topology</li> <li>b. PCs / Laptops / Servers</li> <li>c. Wiring / Structured Cabling / Connectors</li> <li>d. Switches / Hubs / Routers</li> <li>e. Wireless Communications</li> </ol> </li> <li>2. Software / Middleware / Applications             <ol style="list-style-type: none"> <li>a. EMR/EHR</li> <li>b. Healthcare Information Systems (PACs, LIS, RIS)</li> <li>c. Network Protocols (IP, CCP, UDP)</li> <li>d. Operating Systems</li> </ol> </li> </ol> <p><b>B. Function and Operation</b></p> <ol style="list-style-type: none"> <li>1. Hardware             <ol style="list-style-type: none"> <li>a. PCs, Switches, Patch Panels</li> <li>b. Networks, Topology</li> <li>c. Peripherals</li> </ol> </li> <li>2. Integration             <ol style="list-style-type: none"> <li>a. Bedside Medical Device Integration (BMDI)</li> <li>b. Medical Device Integration (MDI) (Labs, Printers, etc.)</li> <li>c. Mobile Devices (Handhelds, Smart Phones, Tablets, etc.)</li> </ol> </li> <li>3. Test Equipment             <ol style="list-style-type: none"> <li>a. Cable Test Devices (Copper, Fiber)</li> <li>b. Network Test Devices</li> </ol> </li> <li>4. Security</li> </ol> <p><b>C. Problem Solving</b></p> <ol style="list-style-type: none"> <li>1. Computer Networks</li> <li>2. Integration</li> <li>3. PCs, Switches, Hubs</li> </ol>	<ol style="list-style-type: none"> <li>1. Product Selection / Vendor Selection             <ol style="list-style-type: none"> <li>a. Technology Assessment</li> <li>b. Healthcare Technology Strategic Planning</li> </ol> </li> <li>2. Project Management             <ol style="list-style-type: none"> <li>a. Capital Planning</li> <li>b. Return on Investment (ROI) Analysis</li> </ol> </li> <li>3. Life Cycle Analysis</li> <li>4. Usability/Compatibility Assessment             <ol style="list-style-type: none"> <li>a. Device/System Upgrade Planning</li> <li>b. Clinical Device Use and/or Application</li> </ol> </li> <li>5. Device Integration Planning             <ol style="list-style-type: none"> <li>a. Clinical Systems Networking</li> </ol> </li> <li>6. Coordinating Device Interoperability / Interfacing</li> <li>7. EMI/RFI Management             <ol style="list-style-type: none"> <li>a. Quality Management</li> <li>b. Pre-clinical Procedure Set-up / Testing</li> <li>c. Clinical Trials Management (Non-investigational)</li> <li>d. Participation in Clinical Procedures (e.g., surgery)</li> </ol> </li> <li>8. Interpretation of Codes and Standards</li> <li>9. Other Technology Management Responsibilities</li> </ol>

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<p><b>7. Healthcare Safety &amp; Standards</b></p> <ol style="list-style-type: none"> <li>1. Electrical – Micro / Marco-shock, Electrical Safety Testing</li> <li>2. Chemical - Material Safety Data Sheet</li> <li>3. Biological - Universal Precautions</li> <li>4. Fire (Class, Fire Extinguishers)</li> <li>5. Regulations, Codes and Standards (CSA Standards, Electromedical, Laser Safety, Low Pressure Connecting Assemblies (Medical Gases), Stability and Transport)</li> <li>6. Biomedical Waste Management             <ol style="list-style-type: none"> <li>i) Non-Hazardous</li> <li>ii) Hazardous (Radioactive waste, Discarded Glass, pressurized Containers, Chemical Waste, Cytotoxic Waste, Plastic Disposables, Liquid Wastes)</li> </ol> </li> </ol>	<p><b>8. NABH &amp; NABL Accreditation</b></p> <ol style="list-style-type: none"> <li>1. Access, Assessment &amp; Continuity of Care (AAC), &amp; Management of Medication (MOM) For NABH</li> <li>2. Care of Patients (COP) For NABH</li> <li>3. Patient Rights &amp; Education (PRE), Responsibilities Management (ROM), Human Resource Management (HRM) for NABH</li> <li>4. Hospital Infection Control (HIC), Continuous Quality Improvement (CQI) for NABH</li> <li>5. Facility management &amp; Safety (FMS), Information Management System (IMS) For HABH.</li> <li>6. Documentation Requirements &amp; Implementation Guidelines for NABH &amp; NABL</li> <li>7. Accreditation Process of NABH &amp; NABL</li> <li>8. Accreditation criteria &amp; their interpretations for NABH</li> <li>9. Accreditation and International perspective of NABL</li> <li>10. Final Assessment of NABL &amp; NABH</li> </ol>
<p><b>9. Product Development, Testing, Evaluation, &amp; Regulatory Compliance</b></p> <ol style="list-style-type: none"> <li>1. Regulatory Compliance Activities</li> <li>2. New Product Testing &amp; Evaluation</li> <li>3. Documentation Development / Management.</li> <li>4. Human Factors Engineering</li> <li>5. Product / Systems Quality Management</li> <li>6. Device Modifications</li> <li>7. Medical Device Design</li> <li>8. Product Research and Development</li> <li>9. Medical Device Concept Development / Invention</li> <li>10. Other Product Development Responsibilities</li> <li>11. Product Sales / Sales Support</li> </ol>	<p><b>10. Medical Terminology for Engineers</b></p> <ol style="list-style-type: none"> <li>1. Disease and Treatment</li> <li>2. Circulation, Blood and Immunity</li> <li>3. Respiration and Digestion</li> <li>4. Urinary and Male Reproductive System</li> <li>5. Female Reproductive System, Pregnancy and Birth</li> <li>6. Endocrine and Nervous Systems, Behavioral Disorders</li> <li>7. The Senses</li> <li>8. The Skeleton and Muscular Systems</li> <li>9. The Skin</li> </ol>
<p><b>11. Risk Management / Safety</b></p>	<p><b>12. Radiation Safety</b></p>

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<ol style="list-style-type: none"> <li>1. Patient Safety</li> <li>2. Product Safety / Hazard Alerts / Recalls</li> <li>3. Incident / Untoward Event Investigation</li> <li>4. Engineering Assessment of Medical Device Failures</li> <li>5. Root Cause Analysis</li> <li>6. Medical Device Incident Reporting</li> <li>7. Infection Control</li> <li>8. Failure Mode and Effect Analysis</li> <li>9. Hazardous Materials</li> <li>10. Other Risk Management / Safety Responsibilities</li> </ol>	<ol style="list-style-type: none"> <li>12. X-Ray Equipment and Production</li> <li>13. Radiation Units for Measurement of Radioisotopes</li> <li>14. Radiation Units for Measurement of Ionizing Radiation</li> <li>15. Personal Maximum Permissible Doses</li> <li>16. Proposed I.C.R.P. New Maximum Permissible Doses</li> <li>17. Radiation Detection and Monitoring</li> <li>18. Basic Principles of Radiation Protection</li> <li>19. Leak Test and Storage of Radiographic Exposure Devices</li> <li>20. Biological Effects of Ionizing Radiation</li> <li>21. Nuclear Regulatory Commission Regulations</li> <li>22. Documentation and Record Keeping</li> <li>23. Transportation of Radioactive Material</li> </ol>
<b>13. Medical Device Regulatory</b>	<b>14. Facilities / General Management</b>
<ol style="list-style-type: none"> <li>1. Medical Devices, In –vitro devices, Biologics &amp; Combination Products: Introductory Module</li> <li>2. FDA Regulations and Guidelines on Medical Devices</li> <li>3. European Union Regulatory Guidelines on Medical Devices</li> <li>4. Medical Device Regulations from Indian perspective</li> <li>5. Management of the risks associated with Medical devices</li> <li>6. Biocompatibility Studies and Medical Devices</li> <li>7. Clinical Trials: Medical Devices</li> <li>8. Overview of In – Vitro Device Regulation</li> <li>9. Overview of Combination Products Regulation</li> <li>10. Dossier preparation in CTD format, eCTD submissions</li> </ol>	<ol style="list-style-type: none"> <li>1. Hospital Building plan Review and Hospital building design</li> <li>2. Emergency Electrical Power and Medical Gas system testing</li> <li>3. Facility emergency preparedness activities</li> <li>4. Supervise Manage / Direct Facilities Management and facility / Utility Remediation planning, other facility Management Responsibilities.</li> <li>5. Budget develop / Execution, Business, / Operation, Plan development, / Management. Committee Management, Performance Improvement.</li> <li>6. Personnel Management / Supervision, Staffing, Staff skills, Competency assessment, Policy, / Procedure management, / Development.</li> </ol>
<b>15. Service Delivery Management</b>	
<ol style="list-style-type: none"> <li>1. Technician / Service Supervision</li> <li>2. Equipment Repair and Maintenance</li> </ol>	

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<ol style="list-style-type: none"><li>3. Equipment Acceptance</li><li>4. Service Contract Management</li><li>5. Equipment Performance Testing</li><li>6. Maintenance                      Software Administration</li><li>7. Develop Test/Calibration/Maintenance Procedures</li><li>8. Parts/Supplies    Purchase    and/or Inventory Management</li><li>9. Other            Service            Delivery Responsibilities.</li><li>10. Technical Library / Service Manuals Management</li></ol>	
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