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| **Domain** | **Reference** | **Description** | **Evidence**(please indicate on which page of you portfolio this evidence can be found) | **Proposer (s) signature** |
| **A. Safe Working Practice** | A1 | Provide evidence that you are competent with a range of generic skills including mandatory training e.g. infection control and basic life support. |   |  |
| A2 | Demonstrates an understanding and application of health & safety and risk management in all aspects of the Clinical Technologists role. |   |  |
| A3 | Demonstrates an understanding of, and works within all relevant legislation to their role including departmental local rules and employers procedures. |   |  |
| A4 | Perform health & safety risk assessments (including radiation risk assessments for ionising radiation) in accordance with standard operating procedures. |   |  |
| A5 | Provide evidence of radiation incident reporting. |   |  |
| A6 | Demonstrates effective communication skills and team working. |   |  |
| A7 | Demonstrates a professional approach to all aspects of the Clinical Technologists role. |   |  |
| A8 | Assists in giving instructions to patients and colleagues regarding radiation hazards, doses and restrictions. |   |  |
| A9 | Demonstrates reflective practice as part of the learning process. |  |  |

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| **Domain** | **Reference** | **Description** | **Evidence**(please indicate on which page of you portfolio this evidence can be found) | **Proposer (s) signature** |
| **B. Equipment Management** | B1 | Assist in the process for the procurement of equipment, accessories or consumables.  |   |  |
| B2 | Demonstrates the use of an equipment inventory system. |   |  |
| B3 | Performs cleaning/decontamination of equipment. |   |  |
| B4 | Performs routine equipment quality control checks and review and interpret results.  |   |  |
| B5 | Performs a range of fault finding and first line user maintenance. |   |  |
| B6 | Demonstrates knowledge of radioactive source management and disposal. |  |  |
| B7 | Demonstrates an understanding of quality management systems. |  |  |
| B8 | Observe and assist with equipment life cycle procedures as an equipment user. |  |  |

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| **Domain** | **Reference** | **Description** | **Evidence**(please indicate on which page of you portfolio this evidence can be found) | **Proposer (s) signature** |
| **C. Nuclear Medicine**  | C1 | Perform all aspects of patient preparation for in vivo/ imaging/therapy treatment; and compliance with legislation. Adhering to standards of professional practice throughout. |   |  |
| C2 | Operate equipment safely across a range of acquisitions and recording techniques to produce high quality results for interpretation. |   |  |
| C3 | Assist with a range of therapy procedures. |  |  |
| C4 | Assist with commissioning checks on a range of systems, and review and interpret results. |  |  |
| C5 | Assist in appointment scheduling. |  |  |
| C6 | Assist with clinical audit. |  |  |
| C7 | Demonstrates accurate recording keeping. |  |  |
| C8 | Performs a range of QC checks including the environment; review and interpret results. |  |  |
| C9 | Performs a range of radioactive manipulations to include activity calculations and measurements. |  |  |
| C10 | Performs a range of tests to demonstrate problems associated with assay. |   |  |

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| **Domain** | **Reference** | **Description** | **Evidence**(please indicate on which page of you portfolio this evidence can be found) | **Proposer (s) signature** |
| **D. Radiopharmaceuticals** | D1 | Complete training courses for venous sampling, the administration of radiopharmaceuticals and the giving of adjunct drugs and perform these tasks. |  |  |
| D2 | Adhere to relevant standards of professional practice as defined in good manufacturing practice. |  |  |
| D3 | Elute the 99mTc generator and reconstitute commercial kits in accordance with written procedures. |  |  |
| D4 | Observe or assist with cell labelling procedures. |  |  |
| **Domain** | **Reference** | **Description** | **Evidence**(please indicate on which page of you portfolio this evidence can be found) | **Proposer (s) signature** |
| **E. Radiation Transport** **and Dosimetry** | E1 | Perform source checks and completes all relevant paper work prior to transport. |  |  |
| E2 | Perform sealed source leak tests, review results and take appropriate action. |   |  |  |
| E3 | Perform contamination checks and maintain appropriate records. |  |  |

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| **Domain** | **Reference** | **Description** | **Evidence**(please indicate on which page of you portfolio this evidence can be found) | **Proposer (s) signature** |
| **F. Good Scientific Practice** | F1 | Adhere to relevant standards of professional practice as defined in Good Scientific Practice. Demonstrate that you have read, understood and comply with this document in all aspects of work. |  |  |

Applicant’s Name (printed): Applicant’s signature: Date: