

**SPECIALTY TRAINING CURRICULUM**

**FOR**

**NUCLEAR MEDICINE**

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**Joint Royal Colleges of Physicians Training Board**

**5 St Andrews Place  
Regent's Park  
London NW1 4LB**

**Telephone: (020) 79351174**

**Facsimile: (020)7486 4160**

**Email: [ptb@jrcptb.org.uk](mailto:ptb@jrcptb.org.uk)**

**Website: [www.jrcptb.org.uk](http://www.jrcptb.org.uk)**

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## **1 Introduction**

Nuclear Medicine is the specialty responsible for the administration of unsealed radioactive substances to patients for the purposes of diagnosis, therapy or research (in contrast to radionuclide radiology which does not include the use of radionuclides for therapy or for non-imaging diagnostic studies). Nuclear medicine trainees will be expected to combine their skills as a physician with that of a physiological imager to solve diagnostic problems. They will provide a unique insight into the pathophysiology of disease and where appropriate offer a radionuclide therapeutic option for treatment. Trainees will require appropriate instruction in the clinical, scientific and legal aspects of the specialty. Specialists in Nuclear Medicine have ultimate responsibility for Nuclear Medicine services and must hold the appropriate certificate from Health Ministers to administer radioactive substances.

The trainee in Nuclear Medicine needs to gain a broad view of the needs of the community he or she serves. This requires not only the acquisition of certain knowledge and skills but also the development of appropriate attitudes enabling the trainee to look after the interests of patients, to work with other relevant health care professionals, to keep up with developments in the field and to bring these developments into the clinical arena. The trainee will have to demonstrate a good understanding of the pathophysiology of the diseases they are imaging or treating. They will need to maintain skills in taking competent histories, relevant clinical examination and the care of both in-patients and out-patients. They will need to learn how they, as medical practitioners, should interact with other clinicians and non-medically trained professional groups. They will need to develop the confidence to present their opinion on patient management as necessary.

## **2 Rationale**

### **2.1 Purpose of the Curriculum**

The purpose of this curriculum is to train a specialist in Nuclear Medicine. The curriculum describes the competencies required for the award of a Certificate of Completion of Training (CCT) and to be included on the Specialist Register in Nuclear Medicine. The CCT specialist will be able to work as a consultant specialist within the National Health Service and will have the knowledge, skills and attitudes required to do this. It is expected that the trainee at the time of their CCT will be competent in the understanding of the scientific knowledge base of nuclear medicine and in the practice of diagnostic and therapeutic nuclear medicine.

The curriculum covers training for all four nations of the UK.

### **2.2 Development**

This curriculum was developed by a sub committee of the Specialty Advisory Committee for Nuclear Medicine under the direction of the Joint Royal Colleges of Physicians Training Board (JRCPTB). It replaces the previous version of the curriculum dated May 2007, with changes to ensure the curriculum meets GMC's standards for Curricula and Assessment, and to incorporate revisions to the content and delivery of the training programme. Major changes from the previous curriculum include the incorporation of leadership, health inequalities and common competencies.

The majority of the SAC members are teachers, trainers and trainees in the specialty, as well as representatives from the Royal College of Radiologists.

### **2.3 Entry Requirements and Training Pathway**

Specialty training in Nuclear Medicine consists of core and higher speciality training. Core training provides physicians with the ability to investigate, treat and diagnose patients with acute and chronic medical symptoms, and with high quality review skills for managing inpatients and outpatients. Higher speciality training then builds on these core skills to develop the specific competencies required to practise independently as a consultant in Nuclear medicine.

Core Medical training programmes are designed to deliver core training for specialty training by acquisition of knowledge and skills as assessed by the workplace based assessments and the MRCP. Programmes are usually for two years and are broad based consisting of four to six placements in medical specialties. These placements over the two years must include direct involvement in the acute medical take. Trainees are asked to document their record of workplace based assessments in an ePortfolio which will then be continued to document assessments in specialty training. Trainees completing core training will have a solid platform of common knowledge and skills from which to continue into Specialty Training at ST3, where these skills will be developed and combined with specialty knowledge and skills in order to award the trainee with a certificate of completion of training (CCT).

There are common competencies that should be acquired by all physicians during their training period starting within the undergraduate career and developed throughout the postgraduate career. These are initially defined for CMT and then developed further in the specialty. This part of the curriculum supports the spiral nature of learning that underpins a trainee's continual development. It recognises that for many of the competences outlined there is a maturation process whereby practitioners become more adept and skilled as their career and experience progresses. It is intended that doctors should recognise that the acquisition of basic competences is often followed by an increasing sophistication and complexity of that competence throughout their career. This is reflected by increasing expertise in their chosen career pathway.

In view of the multi-disciplinary nature of Nuclear Medicine, the specialty is considered to be strengthened by inclusion of practitioners from a variety of clinical backgrounds. Thus, this curriculum allows for entry into specialty training not only from a background in clinical medicine but also from radionuclide radiology and other specialties such as surgery and paediatrics.

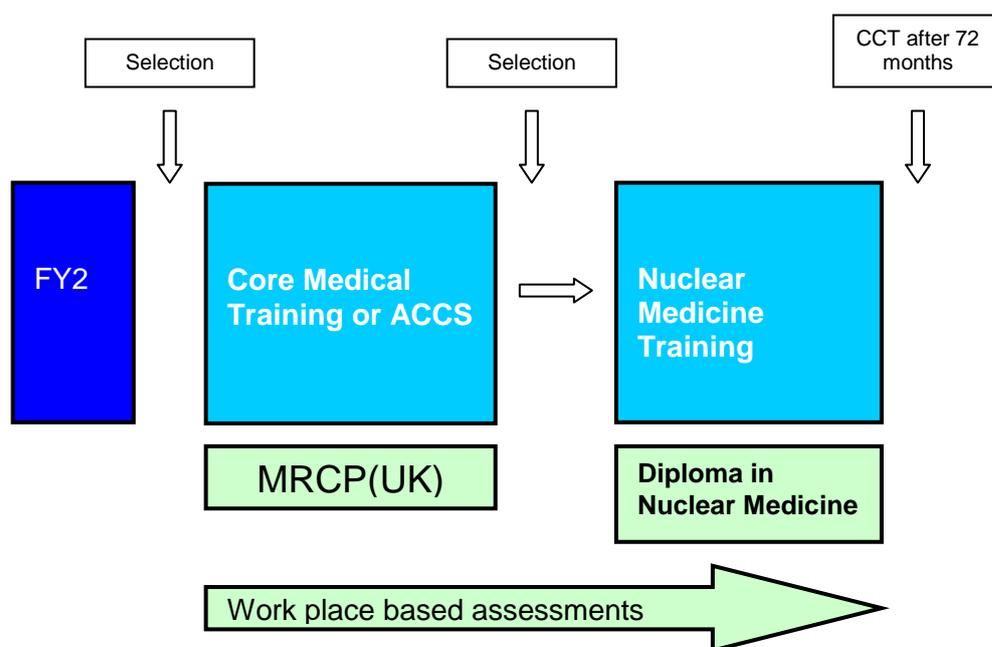
#### **Entry from Clinical Medicine:**

Applicants for Specialty Training year 3 should have successfully completed Foundation training and either a) successfully completed approved core medical training (ST1 and ST2) or b) provide other evidence of achievement of core medical competencies. They must hold the full MRCP (UK) (after 2011)

Core training may be completed in either a Core Medical Training (CMT) or Acute Care Common Stem (ACCS) programme. The full curriculum for specialty training in Nuclear Medicine for trainees entering the specialty through core training therefore consists of the curriculum for either CMT or ACCS plus this specialty training curriculum for Nuclear Medicine.

The approved curriculum for CMT is a sub-set of the Curriculum for General Internal Medicine (GIM). A “Framework for CMT” has been created for the convenience of trainees, supervisors, tutors and programme directors. The body of the Framework document has been extracted from the approved curriculum but only includes the syllabus requirements for CMT and not the further requirements for acquiring a CCT in GIM.

**Figure 1.0 shows the training pathway for a trainee entering through Clinical Medicine.**



#### **Entry from Radionuclide Radiology.**

Applicants for a CCT in Nuclear Medicine in addition to a CCT in radiology should have FRCR and undergone special interest training in Radionuclide radiology. Applicants should have successfully completed Foundation training and either a) successfully completed approved core medical training or b) provide other evidence of achievement of core medical competencies. This should be provided by the candidate’s record of training and will be assessed on an individual basis by the STC. If there is insufficient evidence of core medical competencies, applicants may be admitted to Nuclear Medicine training only if the training period is extended to enable the provision of additional core medical training. Due to the commonality between the nuclear medicine curriculum and the special interest training curriculum for radionuclide radiology it would be expected that all candidates entering from radionuclide radiology will have obtained all level 1 and almost all level 2 competencies as outlined in the nuclear medicine syllabus so that entry from radionuclide radiology will be assessed on an individual basis and based on the competencies already gained the trainee will be placed at the appropriate stage in the training scheme. This is possible because there are a maximum of 6 trainees per year and we work with a single post-graduate dean for the whole UK.

#### **Entry from other Clinical Backgrounds:**

Applicants without the full MRCP (UK) or FRCR who compete for specialty training year 3 posts must provide evidence of appropriate knowledge, training and experience. As a minimum they must have evidence of completion of foundation competencies. If there is insufficient evidence of core medical competencies,

applicants may be admitted to Nuclear Medicine training only if the training period is extended to enable the provision of additional core medical training. Overseas graduates must also provide evidence of satisfactory completion of appropriately supervised general professional training.

The Nuclear Medicine Specialty Curriculum builds on the general competencies delivered in core medical training and other training pathways. Nuclear medicine trainees are expected to be involved in a range of clinical activities. They must also show that they can perform as physicians of the highest clinical and ethical standard. They should show knowledge of how society shapes disease and the role of nuclear medicine within that disease. They must show they can work within a multi-disciplinary team but be able to take a clinical lead role within that team. They should recognise an understanding of the concerns and fears of their patients including the special requirements of children, the vulnerable and those from different ethnic backgrounds. They must demonstrate, through participation, that they know the importance of audit and research.

## **2.4 Enrolment with JRCPTB**

Trainees are required to register for specialist training with JRCPTB at the start of their training programmes. Enrolment with JRCPTB, including the complete payment of enrolment fees, is required before JRCPTB will be able to recommend trainees for a CCT. Trainees can enrol online at [www.jrcptb.org.uk](http://www.jrcptb.org.uk).

## **2.5 Duration of Training**

Although this curriculum is competency based, the duration of training must meet the European minimum of 4 years for full time specialty training adjusted accordingly for flexible training (EU directive 2005/36/EC). The SAC has advised that training from ST1 will usually be completed in 6 years in full time training (2 years core plus 4 years specialty training).

## **2.6 Less Than Full Time Training (LTFT)**

Trainees who are unable to work full-time are entitled to opt for less than full time training programmes. EC Directive 2005/36/EC requires that:

- LTFT shall meet the same requirements as full-time training, from which it will differ only in the possibility of limiting participation in medical activities.
- The competent authorities shall ensure that the competencies achieved and the quality of part-time training are not less than those of full-time trainees.

The above provisions must be adhered to. LTFT trainees should undertake a pro rata share of the out-of-hours duties (including on-call and other out-of-hours commitments) required of their full-time colleagues in the same programme and at the equivalent stage.

EC Directive 2005/36/EC states that there is no longer a minimum time requirement on training for LTFT trainees. In the past, less than full time trainees were required to work a minimum of 50% of full time. With competence-based training, in order to retain competence, in addition to acquiring new skills, less than full time trainees would still normally be expected to work a minimum of 50% of full time. If you are returning or converting to training at less than full time please complete the LTFT application form on the JRCPTB website [www.jrcptb.org.uk](http://www.jrcptb.org.uk).

Funding for LTFT is from deaneries and these posts are not supernumerary. Ideally therefore 2 LTFT trainees should share one post to provide appropriate service cover.

Less than full time trainees should assume that their clinical training will be of a duration pro-rata with the time indicated/recommended, but this should be reviewed during annual appraisal by their TPD and chair of STC and Deanery Associate Dean for LTFT training. As long as the statutory European Minimum Training Time (if relevant), has been exceeded, then indicative training times as stated in curricula may be adjusted in line with the achievement of all stated competencies.

## **2.7 Dual CCT (GIM and Nuclear Medicine)**

Trainees who wish to achieve a CCT in General Internal Medicine (GIM) as well as nuclear medicine must have applied for and successfully entered a training programme which was advertised openly as a dual training programme. Trainees will need to achieve the competencies, with assessment evidence, as described in both the Nuclear Medicine and GIM curricula. Individual assessments may provide evidence towards competencies from both curricula. Postgraduate Deans wishing to advertise such programmes should ensure that they meet the requirements of both SACs.

## **3 Content of Learning**

Nuclear Medicine trainees will be expected to maintain and extend the clinical skills required in obtaining relevant clinical assessment of patients. As nuclear cardiology may represent a significant workload the trainee will need to develop competency in forms of safe cardiac stressing and maintain ALS skills throughout their training and beyond. The trainee will also have significant exposure to patients with chronic and life threatening illness and they will be expected to manage these patients in an empathetic and professional way. Unlike many physicians they will also need to interact with children and will need to develop the requisite skills in working with children as well as being aware of the legal framework for the care of children in the NHS.

Nuclear medicine is unusual in that the nuclear medicine physician is often part of a highly professional and educated team which may involve senior scientists and technical staff. They will need to develop the skills to work as part of a multi-disciplinary team but learn how to provide clinical leadership within that group. Nuclear medicine interacts with a large number of clinicians including surgeons, paediatricians, psychiatrists etc. The trainee therefore should retain and develop an interest in a wide range of medical conditions, their presentation, complications and treatment. They need to develop the confidence in their abilities within the multi-disciplinary team setting.

Nuclear medicine does not exist in isolation from society and as physicians we should be aware of opportunities of providing appropriate health advice to our patients. This could include smoking cessation advice to a patient having a cardiac stress test and life style advice to a patient with osteoporosis. The trainee should also be aware of the cultural diversity of patients and fellow staff and be aware of how and when this may conflict with the practice of nuclear medicine and determine solutions that allow the dignity of colleagues and staff to be maintained.

The detailed syllabus is included below in section 10 of this document.

### 3.1 Specific Skills to be Acquired during Speciality Training

**1. Basic radiation safety:**

The trainee will be able to ensure the safe handling of radiopharmaceuticals both as administered to patients, to him/herself, other staff members and the patient's family and others in whom they are in close contact. Special note will be taken of women who may be or who are pregnant and lactating mothers. The trainee will learn and apply the principles of ALARP (as low as reasonably practical) as defined as lowest radiation dose to the patient to achieve a diagnostic image or therapeutic response. Competency should be obtained by the end of year 1 with consolidation over the training period.

**2. Understanding of the legal requirements for safe handling of radioisotopes:**

The trainee will be taught the legal framework for the safe administration of radiopharmaceuticals including the general instructions for ionising radiation (IR(ME)R 2000) and those specific to the practice of nuclear medicine (MARS and ARSAC regulation) Competence will be obtained by the time of their CCT

**3. Basic science underpinning safe practice of nuclear medicine:**

The trainees should acquire an understanding of the different forms of radioactive decay, their effects on human tissue, how basic nuclear medicine imaging devices work and the factors which effect image quality. This should be achieved within the first year of the trainee's appointment with further in depth knowledge gained before CCT

**4. Assessment of patient's condition and appropriateness of diagnostic test:**

An understanding of why nuclear medicine tests are required and how the patient's condition can affect the interpretation of the diagnostic image. These skills will be acquired throughout the course

**5. An understanding of how to conduct nuclear medicine tests and the skills to report those tests accurately and understand how these results fit into the patient's ongoing management:**

These will include interaction with referring clinicians both informally and through formal MDMs.

Training in these areas will be delivered in a method that shows progression from the simplest procedures defined as level 1 in the first two years of training to the most complex studies and therapies performed in the last 2 years of training (level 2 and 3) studies. However these are not isolated and the competencies gained in performing level 1 procedures will be essential for progression to level 2 and 3 competencies

**6. To understand the appropriate and safe administration of radionuclide therapy and relevant patient aftercare for the patient and their families:**

This will include the indications for radionuclide therapy, patient preparation, radiation protection for both nuclear medicine and other hospital staff and the patient's family as well as the legal framework for the safe administration of radionuclide therapy, the mechanisms required for administration, expected side effects and effective follow-up following therapy. Training in these aspects will be delivered throughout the 4 years but will be a main focus of

year 4 to allow those with radionuclide radiology to gain a CCT in Nuclear Medicine

**7. Communication with patients and other members of the nuclear medicine team:**

These skills will also be strengthened through the generic curriculum but with special emphasis on the uses of radionuclides for diagnosis and therapy

**8. Understanding the inter-relationship of nuclear medicine studies and other diagnostic tests:**

Training in these aspects will occur both throughout the course but also with special reference to cross-sectional radiology as a specific rotation.

**9. Building skills in communicating results of investigations with clinicians:**

This will be occurring throughout the course aided by the generic training and skills learnt in Foundation years and core medical training or equivalent.

**10. Safe and appropriate uses of interventions such as cardiac testing:**

This will include both physical and pharmacological stress and maintaining skills in cardiac resuscitation again building on skills gained in Foundation and Core Medical Training.

**11. Understanding the role of the Nuclear Medicine Physician as a medical profession in the health service**

**12. Promoting personal and professional development.**

### **3.2 Levels of Competence**

As Nuclear Medicine contains discrete quanta of knowledge and competency, a trainee cannot be 'half' competent in reading a scan. All nuclear medicine procedures in the syllabus below have been divided into 3 levels of competencies with the trainees making a step wise progression from the simplest (level 1) to the most complex (level 3).

### **3.3 Good Medical Practice**

In preparation for the introduction of licensing and revalidation, the General Medical Council has translated Good Medical Practice into a Framework for Appraisal and Assessment which provides a foundation for the development of the appraisal and assessment system for revalidation. The Framework can be accessed at [http://www.gmc-uk.org/Framework\\_4\\_3.pdf\\_25396256.pdf](http://www.gmc-uk.org/Framework_4_3.pdf_25396256.pdf)

The Framework for Appraisal and Assessment covers the following domains:

Domain 1 – Knowledge, Skills and Performance

Domain 2 – Safety and Quality

Domain 3 – Communication, Partnership and Teamwork

Domain 4 – Maintaining Trust

The "GMP" column in the syllabus defines which of the 4 domains of the Good Medical Practice Framework for Appraisal and Assessment are addressed by each competency. Most parts of the syllabus relate to "Knowledge, Skills and Performance" but some parts will also relate to other domains.

## 4 Learning and Teaching

### 4.1 The Training Programme

The organisation and delivery of postgraduate training is the statutory responsibility of the General Medical Council (GMC) which devolves responsibility for the local organisation and delivery of training to the deaneries. Each deanery oversees a "School of Medicine" which is comprised of the regional Specialty Training Committees (STCs) in each medical specialty. Responsibility for the organisation and delivery of specialty training in Nuclear Medicine in each deanery is, therefore, the remit of the regional Nuclear Medicine STC. Each STC has a Training Programme Director who coordinates the training programme in the specialty.

The sequence of training should ensure appropriate progression in experience and responsibility. The training to be provided at each training site is defined to ensure that, during the programme, the entire curriculum is covered and also that unnecessary duplication and educationally unrewarding experiences are avoided. However, the sequence of training should ideally be flexible enough to allow the trainee to develop a special interest.

#### Acting up as a consultant (AUC)

"Acting up" provides doctors in training coming towards the end of their training with the experience of navigating the transition from junior doctor to consultant while maintaining an element of supervision.

Although acting up often fulfills a genuine service requirement, it is not the same as being a locum consultant. Doctors in training acting up will be carrying out a consultant's tasks but with the understanding that they will have a named supervisor at the hosting hospital and that the designated supervisor will always be available for support, including out of hours or during on-call work. Doctors in training will need to follow the rules laid down by the Deanery / LETB within which they work and also follow the JRCPTB rules which can be found at [www.jrcptb.org.uk/trainingandcert/Pages/Out-of-Programme](http://www.jrcptb.org.uk/trainingandcert/Pages/Out-of-Programme).

### 4.2 Teaching and Learning Methods

The curriculum will be delivered through a variety of learning experiences. Trainees will learn from practice, clinical skills appropriate to their level of training and to their attachment within the department.

Trainees will achieve the competencies described in the curriculum through a variety of learning methods. There will be a balance of different modes of learning from formal teaching programmes to experiential learning 'on the job'. The proportion of time allocated to different learning methods may vary depending on the nature of the attachment within a rotation.

This section identifies the types of situations in which a trainee will learn.

**Learning with Peers** - There are many opportunities for trainees to learn with their peers. Local postgraduate teaching opportunities allow trainees of varied levels of experience to come together for small group sessions. The taught programme encourages group learning. Examination preparation encourages the formation of self-help groups and learning sets.

**Work-based Experiential Learning** - The majority of the curriculum is suited to delivery by work-based experiential learning and on-the-job supervision. Where it is clear from trainees' experience that parts of the curriculum are not being delivered

within their work place, appropriate off-the job education or rotations to other work places will be arranged. The key will be regular workplace-based assessment by educational supervisors who will be able to assess, with the trainee, their on-going progress and whether parts of the curriculum are not being delivered within their present work place. These will show a progression of skills from the most simple (level 1) to the most complex (level 3).

The content of work-based experiential learning is decided by the local faculty for education but includes active participation in the following, remembering that nuclear medicine has imaging as its primary role and trainees will not be involved in general out-patients or acute medical 'takes'. As almost all procedures are done as out patients the traditional model of learning including from acute assessment, admission and management of patients is not relevant.

Therefore the main work based teaching experiences will be:

- The majority of work based learning will take place in the nuclear medicine department where patients will be assessed to determine if the correct scan has been requested and if they have any co-morbidities or are on medication that will affect the outcome of the scan. This will initially be under direct supervision but the degree of autonomy will increase with the trainees' competence.
- The trainee will learn by first observing image interpretation and reporting skills of a specialist in nuclear medicine but as confidence and competence increases will be expected to report scans under supervision and then with more autonomy. It would be expected that the trainee will gain competence in less complex scan reading (level 1) by the end of ST4, medium complexity (level 2) studies by the end of ST5 and complex (level 3) studies by the end of the training course
- Specialist out-patient clinics such as thyroid and neuroendocrine clinics. After initial induction, trainees will review patients in such clinics, under direct supervision. The degree of responsibility taken by the trainee will increase as competency increases. As experience and clinical competence increase trainees will assess 'new' and 'review' patients and present their findings to their clinical supervisor.
- Personal ward rounds if there are any in-patients and provision of ongoing clinical care on specialist medical wards. The only patients that will be seen as in-patients are those receiving radionuclide therapy. Every patient seen, on the ward or in out-patients, provides a learning opportunity, which will be enhanced by following the patient through the course of their treatment and possible side effects. Also there should be a proper understanding of the information required by the patient's referring clinician to ensure continuing care. Patients seen should provide the basis for critical reading and reflection of clinical problems.
- Multi-disciplinary team meetings. There are many situations where clinical problems are discussed with clinicians in other disciplines. These provide excellent opportunities for observation of clinical reasoning.
- Attachments to other training departments will be organised to supplement the learning experiences as required. Some centres have local arrangements for rotations of trainees between departments or to other departments for specific parts of the training scheme for example paediatrics, therapy and PET-CT. However it is expected that most trainees will spend their first year within a single department

**Formal Postgraduate Teaching** – The content of these sessions are determined by the local faculty of medical education and will be based on the curriculum. There are many opportunities throughout the year for formal teaching in the local postgraduate

teaching sessions and at regional, national and international meetings. Many of these are organised by the Royal Colleges of Physicians.

Suggested activities include:

• The most important teaching will include formal courses of training - see below  
In addition the trainees should take part in other learning activities provided including:

- A programme of formal bleep-free regular teaching sessions to cohorts of trainees (e.g. a weekly core training hour of teaching within a Trust)
- Case presentations
- Journal clubs
- Research and audit projects
- Lectures and small group teaching
- Grand Rounds
- Clinical skills demonstrations and teaching
- Critical appraisal and evidence based medicine and journal clubs
- Joint specialty meetings
- Attendance at training programmes organised on a deanery or regional basis, which are designed to cover aspects of the training programme outlined in this curriculum.

**Independent Self-Directed Learning** -Trainees will use this time in a variety of ways depending upon their stage of learning. These methods will supplement the knowledge based learning. Suggested activities include:

- Reading, including web-based material
- Maintenance of personal portfolio (self-assessment, reflective learning, personal development plan)
- Audit and research projects
- Reading journals
- Achieving personal learning goals beyond the essential, core curriculum

**Formal Study Courses** – Making time available for formal courses is encouraged, subject to local conditions of service. Examples include management courses and communication courses.

**Externally Delivered Education** – As stated above this will supplement the locally delivered knowledge focused training, usually occurring in the ST 4 and 5. This is delivered on a national level by Kings College London with 38 days of face to face learning and some distance learning, It is planned that the majority of teaching will be using e-learning methods supplemented by tutorials with local trainees by 2012 but there will be at least 10 days of face to face learning.

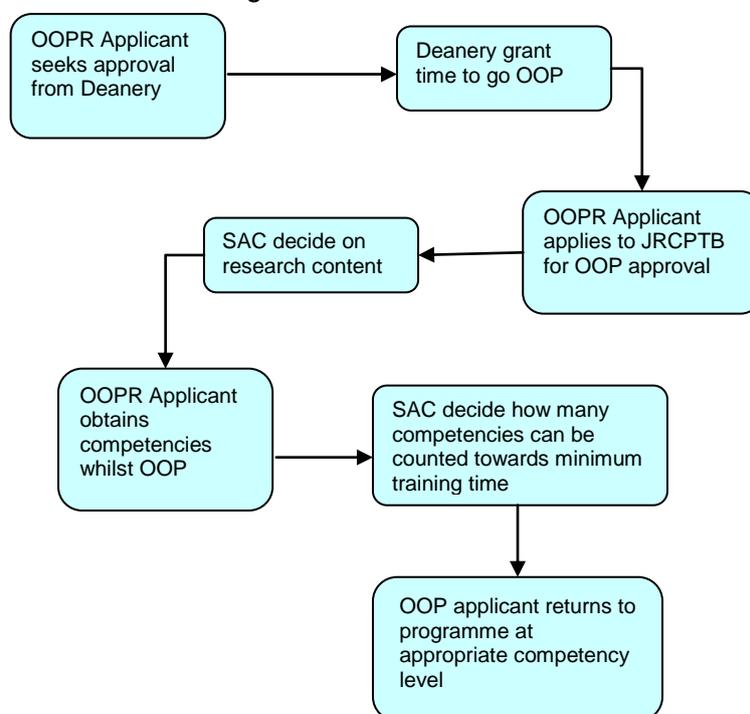
### 4.3 Research

Trainees who wish to acquire research competencies, in addition to those specified in their specialty curriculum, may undertake a research project as an ideal way of obtaining those competencies. For those in specialty training, one option to be considered is that of taking time out of programme to complete a specified project or research degree. Applications to research bodies, the deanery (via an OOPR form) and the JRCPTB (via a Research Application Form) are necessary steps, which are the responsibility of the trainee. The JRCPTB Research Application Form can be accessed via the JRCPTB website. It requires an estimate of the competencies that will be achieved and, once completed, it should be returned to JRCPTB together with a job description and an up to date CV. The JRCPTB will submit applications to the

relevant SACs for review of the research content including an indicative assessment of the amount of clinical credit (competence acquisition) which might be achieved. This is likely to be influenced by the nature of the research (eg entirely laboratory-based or strong clinical commitment), as well as duration (eg 12 month Masters, 2-year MD, 3-Year PhD). On approval by the SAC, the JRCPTB will advise the trainee and the deanery of the decision. The deanery will make an application to the GMC for approval of the out of programme research. All applications for out of programme research must be prospectively approved.

Upon completion of the research period the competencies achieved will be agreed by the OOP Supervisor, Educational Supervisor and communicated to the SAC, accessing the facilities available on the JRCPTB ePortfolio. The competencies achieved will determine the trainee's position on return to programme; for example if an ST3 trainee obtains all ST4 competencies then 12 months will be recognised towards the minimum training time and the trainee will return to the programme at ST5. This would be corroborated by the subsequent ARCP.

This process is shown in the diagram below:



Funding will need to be identified for the duration of the research period. Trainees need not count research experience or its clinical component towards a CCT programme but must decide whether or not they wish it to be counted on application to the deanery and the JRCPTB.

Whilst not obligatory candidates will be encouraged to extend the Post Graduate Diploma by completion of a 5<sup>th</sup> taught module and submission of a 10,000 word research project in the 3rd or 4th year (ST5, ST6) leading to an MSc in Nuclear Medicine.

Time out of programme to take a Clinical Fellowship position is encouraged for good candidates and would be best taken after the candidate has obtained their MSc as many Universities make that a pre-requisite for admission to MRes/MD/PhD programme.

A maximum period of 3 years out of programme is allowed and the SACs will recognise up to 12 months towards the minimum training times.

#### **4.4 Academic Training**

Training centres may be able to offer an Academic Clinical Lectureship programme and that programme can only be offered after completion of an MD/PhD.

For those contemplating an academic career path, there are now well-defined posts at all levels in the Integrated Academic Training Pathway (IATP) involving the National Institute for Health Research (NIHR) and the Academy of Medical Sciences (AMS). For full details see <http://www.nccrcd.nhs.uk/intetacatrain> and <http://www.academicmedicine.ac.uk/uploads/A-pocket-guide.pdf>. Academic trainees may wish to focus on education or research and are united by the target of a consultant-level post in a university and/or teaching hospital, typically starting as a senior lecturer and aiming to progress to readership and professor. A postgraduate degree will usually be essential (see “out of programme experience”) and academic mentorship is advised (see section 6.1). Academic competencies have been defined by the JRCPTB in association with AMS and the Colleges and modes of assessment have been incorporated in the latest edition of the Gold Guide (section 7, see <http://www.jrcptb.org.uk/forms/Documents/GoldGuide2009.pdf>).

Academic integrated pathways to CCT are a) considered fulltime CCTs as the default position and b) are run through in nature. The academic programmes are CCT programmes and the indicative time set for academic trainees to achieve the CCT is the same as the time set for non-academic trainees. If a trainee fails to achieve all the required competencies within the notional time period for the programme, this would be considered at the ARCP, and recommendations to allow completion of clinical training would be made (assuming other progress to be satisfactory). An academic trainee working in an entirely laboratory-based project would be likely to require additional clinical training, whereas a trainee whose project is strongly clinically oriented may complete within the “normal” time (see the guidelines for monitoring training and progress) <http://www.academicmedicine.ac.uk/careersacademicmedicine.aspx>. Extension of a CCT date will be in proportion depending upon the nature of the research and will ensure full capture of the specialty outcomes set down by the Royal College and approved by GMC.

All applications for research must be prospectively approved by the SAC and the regulator, see [www.jrcptb.org.uk](http://www.jrcptb.org.uk) for details of the process.

## **5 Assessment**

### **5.1 The Assessment System**

The purpose of the assessment system is to:

- Enhance learning by providing formative assessment, enabling trainees to receive immediate feedback, measure their own performance and identify areas for development;
- Drive learning and enhance the training process by making clear what is required of trainees and motivating them to ensure they receive suitable training and experience;
- Provide robust, summative evidence that trainees are meeting the curriculum standards during the training programme;

- Ensure trainees are acquiring competencies within the domains of Good Medical Practice;
- Assess trainees' actual performance in the workplace;
- Ensure that trainees possess the essential underlying knowledge required for their specialty;
- Inform the Annual Review of Competence Progression (ARCP), identifying any requirements for targeted or additional training where necessary and facilitating decisions regarding progression through the training programme;
- Identify trainees who should be advised to consider changes of career direction.

The integrated assessment system comprises workplace-based assessments and knowledge – base assessments. Individual assessment methods are described in more detail below.

Because trainees will achieve competencies at different rates, it is not possible to stipulate the numbers of nuclear medicine procedures that should comprise the work-based experiential learning.

The curriculum is blueprinted so that key competencies will be delivered, and the various assessments of knowledge, skills, behaviours and attitudes will be fit for purpose and give coverage across the domains of the curriculum by a process of sampling. All assessments will be appropriate to the training level of the trainee and will be valid, reliable, systematically collected, judged against pre-determined criteria and appropriately weighted. Feedback will be given confidentially to each trainee with suggestions for improvements where appropriate.

Workplace-based assessments will take place throughout the training programme to allow trainees to continually gather evidence of learning and to provide trainees with formative feedback. They are not individually summative but overall outcomes from a number of such assessments provide evidence for summative decision making. The number and range of these will ensure a reliable assessment of the training relevant to their stage of training and achieve coverage of the curriculum.

## 5.2 Assessment Blueprint

In the syllabus (0) the “Assessment Methods” shown are those that are appropriate as **possible** methods that could be used to assess each competency. It is not expected that all competencies will be assessed and that where they are assessed not every method will be used.

## 5.3 Assessment Methods

The following assessment methods are used in the integrated assessment system:

### Examinations and Certificates

- The Post Graduate Diploma in Nuclear Medicine (PGD)
- Advanced Life Support Certificate (ALS)

All trainees must complete the Post Graduate Diploma in Nuclear Medicine as a requirement for achieving a CCT. The Diploma is run by the post graduate education department of Kings College London, which ensures appropriate external review and quality assurance.

Information about the Diploma, including guidance for candidates, is available on the Kings College website:

<http://www.kcl.ac.uk/prospectus/graduate/Nuclear%20Medicine:%20Science%20&%20Practice>

The PGD consists of 4 modules, the first 3 modules and the requisite teaching is undertaken in year 2 (ST4) of the training scheme and consists of 2 compulsory modules Basic science and regulatory and Clinical Nuclear Medicine; with one of two other modules Radiopharmacy or Diagnostic and Therapeutic Nuclear Oncology. Assessment is by locally marked essay (20%) and a formal written examination (80%). A practical module is taken normally in year 3 (ST5) which involves the provision of a log book including expanded CbD, report of basic science experiments performed, presentation of an audit and a clinical film reading assessment.

Marking is performed by a University Appointed Examination Committee with 2 external examiners. Retakes are allowed and there is an appeal mechanism for candidates regulated by Kings College London and the University of London.

### **Workplace-Based Assessments (WPBAs)**

- mini-Imaging Interpretation Exercise (mini-IPX)
- mini-Clinical Evaluation Exercise (mini-CEX)
- Direct Observation of Procedural Skills (DOPS)
- Multi-Source Feedback (MSF)
- Case-Based Discussion (CbD)
- Patient Survey (PS)
- Audit Assessment (AA)
- Teaching Observation (TO)

These methods are described briefly below. More information about these methods including guidance for trainees and assessors is available in the ePortfolio and on the JRCPTB website [www.jrcptb.org.uk](http://www.jrcptb.org.uk). Workplace-based assessments should be recorded in the trainee's ePortfolio. The workplace-based assessment methods include feedback opportunities as an integral part of the assessment process, this is explained in the guidance notes provided for the techniques.

### **mini-Imaging Interpretation Exercise (mini-IPX)**

Because of the imaging nature of about 70% of nuclear medicine practice, mini-IPX is the most common form of assessment. This method of assessment has been developed by the Royal College of Radiologists and is designed to assess a trainee's skills in interpreting an image and to provide rapid and prompt feedback to a trainee in a particular area of diagnostic imaging. More information concerning how to use this assessment is available on [www.rcr.ac.uk/docs/radiology/pdf/RCRmini-IPXGuidanceforAssessors.pdf](http://www.rcr.ac.uk/docs/radiology/pdf/RCRmini-IPXGuidanceforAssessors.pdf).

### **Multisource Feedback (MSF)**

This tool is a method of assessing generic skills such as communication, leadership, team working, reliability etc, across the domains of Good Medical Practice. This provides objective systematic collection and feedback of performance data on a trainee, derived from a number of colleagues. 'Raters' are individuals with whom the trainee works, and includes doctors, administration staff, and other allied professionals. The trainee will not see the individual responses by raters; feedback is given to the trainee by the Educational Supervisor.

### **mini-Clinical Evaluation Exercise (mini-CEX)**

This tool evaluates a clinical encounter with a patient to provide an indication of competence in skills essential for good clinical care such as history taking,

examination and clinical reasoning. The trainee receives immediate feedback to aid learning. The mini-CEX can be used at any time and in any setting when there is a trainee and patient interaction and an assessor is available.

#### **Direct Observation of Procedural Skills (DOPS)**

A DOPS is an assessment tool designed to assess the performance of a trainee in undertaking a practical procedure, against a structured checklist. The trainee receives immediate feedback to identify strengths and areas for development.

#### **Case based Discussion (CbD)**

The CbD assesses the performance of a trainee in their management of a patient to provide an indication of competence in areas such as clinical reasoning, decision-making and application of medical knowledge in relation to patient care. It also serves as a method to document conversations about, and presentations of, cases by trainees. The CbD should include discussion about a written record (such as written case notes, out-patient letter, discharge summary). A typical encounter might be when presenting newly referred patients in the out-patient department.

#### **Patient Survey (PS)**

Patient Survey address issues, including behaviour of the doctor and effectiveness of the consultation, which are important to patients. It is intended to assess the trainee's performance in areas such as interpersonal skills, communication skills and professionalism by concentrating solely on their performance during one consultation.

#### **Audit Assessment Tool (AA)**

The Audit Assessment Tool is designed to assess a trainee's competence in completing an audit. The Audit Assessment can be based on review of audit documentation OR on a presentation of the audit at a meeting. If possible the trainee should be assessed on the same audit by more than one assessor.

#### **Teaching Observation (TO)**

The Teaching Observation form is designed to provide structured, formative feedback to trainees on their competence at teaching. The Teaching Observation can be based on any instance of formalised teaching by the trainee who has been observed by the assessor. The process should be trainee-led (identifying appropriate teaching sessions and assessors).

### **5.4 Decisions on Progress (ARCP)**

The Annual Review of Competence Progression (ARCP) is the formal method by which a trainee's progression through her/his training programme is monitored and recorded. ARCP is not an assessment – it is the review of evidence of training and assessment. The ARCP process is described in A Reference Guide for Postgraduate Specialty Training in the UK (the "Gold Guide" – available from [www.mmc.nhs.uk](http://www.mmc.nhs.uk)). Deaneries are responsible for organising and conducting ARCPs. The evidence to be reviewed by ARCP panels should be collected in the trainee's ePortfolio.

The ARCP Decision Aid is included in section 5.5, giving details of the evidence required of trainees for submission to the ARCP panels.

## 5.5 ARCP Decision Aid

	ST3	ST4	ST5	ST6
Examination			Post Graduate Diploma in Nuclear Medicine attempted	Post Graduate Diploma in Nuclear Medicine passed to obtain CCT
mini-IPX	Trainees should complete sufficient mini-IPX to show that they have completed 25% of level 1 competencies (4 minimum)	Trainees should demonstrate they have completed all of level 1 competencies (4 minimum)	Trainees should demonstrate they have completed 75% of level 2 competencies (8 minimum)	Trainees should demonstrate they have completed all of level 2 and 3 competencies (8 minimum)
mini-CEX	Trainees should complete 50% of all level 1 competencies (1 minimum)	Trainees should complete 100 % of all level 1 competencies (1 minimum)	Trainees should complete 75% of level 2 competencies (1 minimum)	Trainees should complete all level 2 and 3 competencies (1 minimum)
DOPS	Trainees must complete all level 1 competences as these are related to safe practice within a nuclear medicine department (2 minimum)	Trainees must complete all level 1 competences (2 minimum)	Trainees must complete 75% of level 2 competences (1 minimum)	Trainees should complete all level 2 and 3 competencies (1 minimum)
CbD	Trainees should complete a minimum of 4 CbDs related to less complex studies level 1	Trainees should complete a minimum of 4 CbDs related to less and medium complex studies (level 2). At least 2 of these CbDs must be at level 2	Trainees should complete a minimum of 4 CbDs related to medium complex studies level 2	Trainees should complete a minimum of 4 CbDs related to the most complex studies level 3
MSF	Satisfactory		satisfactory	
ALS	Valid	valid	valid	valid

Audit Assessment	The trainees should complete and present at a local audit meeting one audit		The trainees should complete and present at a local audit meeting one audit	
Patient survey		satisfactory		satisfactory
Research		Should show awareness of research methods as taught in PGD course	Should undertake at least one research project encouraged to submit for MSc	By this time should have presented a research paper at national meeting
Attendance at meeting		Encouraged to attend British Nuclear Medicine Society meeting	Encouraged to attend British Nuclear Medicine Society meeting especially if not attended in ST4	Encouraged to attend an International nuclear medicine meeting
Radiation and society	Must show by DOPS, CbD and direct conversation knowledge and application of rules governing safe handling and use of radioisotopes	Must show by DOPS, CbD and direct conversation knowledge and application of rules governing safe handling and use of radioisotopes	Must show by DOPS, CbD and direct conversation knowledge and application of rules governing safe handling and use of radioisotopes	Must show by DOPS, CbD and direct conversation knowledge and application of rules governing safe handling and use of radioisotopes
Passage to next stage	Completion of sufficient level 1 competencies as assessed by ARCP panel	Completion of all level 1 and some level 2 competencies as assessed by ARCP panel	Completion of 75% of all level 2 competencies as assessed by ARCP panel	Completion of all competencies as assessed by ARCP panel leading to CCT

Note: Nuclear medicine contains discrete quanta of knowledge and competency a trainee cannot be 50% competent in reading a scan therefore all nuclear medicine procedures have been divided in 3 levels of competencies level 1,2 and 3 (appendix 1) with the trainees making a step wise progression from the simplest level 1 to the most complex (level 3). Some leeway has to be given in passing these milestones as the number of training centres is very limited for specialist training in PET-CT, therapy and paediatric nuclear medicine so some flexibility on the timing of mile stones is needed

## 5.6 Penultimate Year Assessment (PYA)

The penultimate ARCP prior to the anticipated CCT date will include an external assessor from outside the training programme. JRCPTB and the deanery will coordinate the appointment of this assessor. This is known as "PYA". Whilst the ARCP will be a review of evidence, the PYA will include a face to face component.

## 5.7 Complaints and Appeals

Kings College London has complaints procedures and appeals regulations documented in its website which apply to the Postgraduate Diploma in Nuclear Medicine.

All workplace-based assessment methods incorporate direct feedback from the assessor to the trainee and the opportunity to discuss the outcome. If a trainee has a complaint about the outcome from a specific assessment this is their first opportunity to raise it.

Appeals against decisions concerning in-year assessments will be handled at deanery level and deaneries are responsible for setting up and reviewing suitable processes. If a formal complaint about assessment is to be pursued this should be referred in the first instance to the chair of the Specialty Training Committee who is accountable to the regional deanery. Continuing concerns should be referred to the Associate Dean.

# 6 Supervision and Feedback

## 6.1 Supervision

All elements of work in training posts must be supervised with the level of supervision varying depending on the experience of the trainee and the clinical exposure and case mix undertaken. Outpatient and referral supervision must routinely include the opportunity to personally discuss all cases if required. As training progresses the trainee should have the opportunity for increasing autonomy, consistent with safe and effective care for the patient.

Trainees will at all times have a named Educational Supervisor and Clinical Supervisor, responsible for overseeing their education. Depending on local arrangements these roles may be combined into a single role of Educational Supervisor.

The responsibilities of supervisors have been defined by GMC in the document "Operational Guide for the PMETB Quality Framework". These definitions have been agreed with the National Association of Clinical Tutors, the Academy of Medical Royal Colleges and the Gold Guide team at MMC, and are reproduced below:

### ***Educational Supervisor***

*A trainer who is selected and appropriately trained to be responsible for the overall supervision and management of a specified trainee's educational progress during a training placement or series of placements. The Educational Supervisor is responsible for the trainee's Educational Agreement.*

### ***Clinical Supervisor***

*A trainer who is selected and appropriately trained to be responsible for overseeing a specified trainee's clinical work and providing constructive feedback during a training placement. Some training schemes appoint an Educational Supervisor for each placement. The roles of Clinical and Educational Supervisor may then be merged.*

The Educational Supervisor, when meeting with the trainee, should discuss issues of clinical governance, risk management and any report of any untoward clinical incidents involving the trainee. The Educational Supervisor should be part of the clinical specialty team. Thus if the clinical directorate (clinical director) have any concerns about the performance of the trainee, or there were issues of doctor or patient safety, these would be discussed with the Educational Supervisor. These processes, which are integral to trainee development, must not detract from the statutory duty of the Trust to deliver effective clinical governance through its management systems.

Academic trainees are encouraged to identify an academic mentor, who will not usually be their research supervisor and will often be from outside their geographical area. The Academy of Medical Sciences organises one such scheme (see <http://www.acmedsci.ac.uk/index.php?pid=91>) but there are others and inclusion in an organised scheme is not a pre-requisite. The Medical Research Society organises annual meetings for clinician scientists in training (see [http://www.medres.org.uk/j/index.php?option=com\\_content&task=view&id=54&Itemid=1](http://www.medres.org.uk/j/index.php?option=com_content&task=view&id=54&Itemid=1)) and this type of meeting provides an excellent setting for trainees to meet colleagues and share experiences.

Opportunities for feedback to trainees about their performance will arise through the use of the workplace-based assessments, regular appraisal meetings with supervisors, other meetings and discussions with supervisors and colleagues, and feedback from ARCP.

## **6.2 Appraisal**

A formal process of appraisals and reviews underpins training. This process ensures adequate supervision during training, provides continuity between posts and different supervisors and is one of the main ways of providing feedback to trainees. All appraisals should be recorded in the ePortfolio

### **Induction Appraisal**

The trainee and educational supervisor should have an appraisal meeting at the beginning of each post to review the trainee's progress so far, agree learning objectives for the post ahead and identify the learning opportunities presented by the post. Reviewing progress through the curriculum will help trainees to compile an effective Personal Development Plan (PDP) of objectives for the upcoming post. This PDP should be agreed during the Induction Appraisal. The trainee and supervisor should also both sign the educational agreement in the e-portfolio at this time, recording their commitment to the training process.

### **Mid-Point Review**

This meeting between trainee and educational supervisor is mandatory (except when an attachment is shorter than 6 months), but is encouraged particularly if either the trainee or educational or clinical supervisor has training concerns or the trainee has been set specific targeted training objectives at their ARCP. At this meeting trainees should review their PDP with their supervisor using evidence from the e-portfolio. Workplace-based assessments and progress through the curriculum can be

reviewed to ensure trainees are progressing satisfactorily, and attendance at educational events should also be reviewed. The PDP can be amended at this review.

### **End of Attachment Appraisal**

Trainees should review the PDP and curriculum progress with their educational supervisor using evidence from the e-portfolio. Specific concerns may be highlighted from this appraisal. The end of attachment appraisal form should record the areas where further work is required to overcome any shortcomings. Further evidence of competence in certain areas may be needed, such as planned workplace-based assessments, and this should be recorded. If there are significant concerns following the end of attachment appraisal then the programme director should be informed.

## **7 Managing Curriculum Implementation**

### **7.1 Intended Use of Curriculum by Trainers and Trainees**

This curriculum and ePortfolio are web-based documents which are available from the Joint Royal Colleges of Physicians Training Board (JRCPTB) website [www.jrcptb.org.uk](http://www.jrcptb.org.uk).

The educational supervisors and trainers can access the up-to-date curriculum from the JRCPTB website and will be expected to use this as the basis of their discussion with trainees. Both trainers and trainees are expected to have a good knowledge of the curriculum and should use it as a guide for their training programme.

Each trainee will engage with the curriculum by maintaining a portfolio. The trainee will use the curriculum to develop learning objectives and reflect on learning experiences.

### **7.2 Recording Progress**

On enrolling with JRCPTB trainees will be given access to the ePortfolio for Nuclear Medicine. The ePortfolio allows evidence to be built up to inform decisions on a trainee's progress and provides tools to support trainees' education and development.

The trainee's main responsibilities are to ensure the ePortfolio is kept up to date, arrange assessments and ensure they are recorded, prepare drafts of appraisal forms, maintain their personal development plan, record their reflections on learning and record their progress through the curriculum.

The supervisor's main responsibilities are to use ePortfolio evidence such as outcomes of assessments, reflections and personal development plans to inform appraisal meetings. They are also expected to update the trainee's record of progress through the curriculum, write end-of-attachment appraisals and supervisor's reports.

Deaneries, training programme directors, college tutors and ARCP panels may use the ePortfolio to monitor the progress of trainees for whom they are responsible.

JRCPTB will use summarised, anonymous ePortfolio data to support its work in quality assurance.

All appraisal meetings, personal development plans and workplace based assessments (including MSF) should be recorded in the ePortfolio. Trainees and supervisors should electronically sign the educational agreement. Trainees are encouraged to reflect on their learning experiences and to record these in the ePortfolio. Reflections can be kept private or shared with supervisors.

Reflections, assessments and other ePortfolio content should be linked to curriculum competencies in order to provide evidence towards acquisition of these competencies. Trainees can add their own self-assessment ratings to record their view of their progress. The aims of the self-assessment are:

- To provide the means for reflection and evaluation of current practice
- To inform discussions with supervisors to help both gain insight and assists in developing personal development plans.
- To identify shortcomings between experience, competency and areas defined in the curriculum so as to guide future clinical exposure and learning.

Supervisors can sign-off and comment on curriculum competencies to build up a picture of progression and to inform ARCP panels.

## **8 Curriculum Review and Updating**

The specialty curriculum will be reviewed and updated with minor changes on an annual basis. The curriculum should be regarded as a fluid, living document and the SAC will ensure it responds swiftly to new clinical and service developments. In addition, the curriculum will be subject to three-yearly formal review within the SAC. This will be informed by curriculum evaluation and monitoring. The SAC will have available:

- The trainees' survey, which will include questions pertaining to their specialty (GMC to provide)
- Specialty-specific questionnaires ( if applicable)
- Reports from other sources such as educational supervisors, programme directors, specialty deans, service providers and patients.
- Trainee representation on the Deanery STC and the SAC of the JRCPTB
- Informal trainee feedback during appraisal.

Evaluation will address:

- The relevance of the learning outcomes to clinical practice
- The balance of work-based and off-the-job learning
- Quality of training in individual posts
- Feasibility and appropriateness of on-the-job assessments in the course of training programmes
- Availability and quality of research opportunities
- Current training affecting the service

Evaluation will be the responsibility of the JRCPTB and GMC. These bodies must approve any significant changes to the curriculum.

Interaction with the NHS will be particularly important to understand the performance of specialists within the NHS and feedback will be required as to the continuing needs for that specialty as defined by the curriculum. It is likely that the NHS will have a view as to the balance between generalist and specialist skills, the development of generic competencies and, looking to the future, the need for

additional specialist competencies and curricula. In establishing specialty issues which could have implications for training, the SAC will produce a summary report to discuss with the NHS employers and ensure that conclusions are reflected in curriculum reviews.

Trainee contribution to curriculum review will be facilitated through the involvement of trainees in local faculties of education and through informal feedback during appraisal and College meetings.

The SAC will respond rapidly to changes in service delivery. Regular review will ensure the coming together of all the stakeholders needed to deliver an up-to-date, modern specialty curriculum. The curriculum will indicate the last date of formal review monitoring and document revision.

## **9 Equality and Diversity**

The Royal Colleges of Physicians will comply, and ensure compliance, with the requirements of equality and diversity legislation, such as the:

- Race Relations (Amendment) Act 2000
- Disability Discrimination Act 1995
- Human Rights Act 1998
- Employment Equality (Age) Regulation 2006
- Special Educational Needs and Disabilities Act 2001
- Data Protection Acts 1984 and 1998

The Federation of the Royal Colleges of Physicians believes that equality of opportunity is fundamental to the many and varied ways in which individuals become involved with the Colleges, either as members of staff and Officers; as advisers from the medical profession; as members of the Colleges' professional bodies or as doctors in training and examination candidates. Accordingly, it warmly welcomes contributors and applicants from as diverse a population as possible, and actively seeks to recruit people to all its activities regardless of race, religion, ethnic origin, disability, age, gender or sexual orientation.

Deanery quality assurance will ensure that each training programme complies with the equality and diversity standards in postgraduate medical training as set by GMC.

Compliance with anti-discriminatory practice will be assured through:

- monitoring of recruitment processes;
- ensuring all College representatives and Programme Directors have attended appropriate training sessions prior to appointment or within 12 months of taking up post;
- Deaneries must ensure that educational supervisors have had equality and diversity training (at least as an e learning module) every 3 years
- Deaneries must ensure that any specialist participating in trainee interview/appointments committees or processes has had equality and diversity training (at least as an e module) every 3 years.
- ensuring trainees have an appropriate, confidential and supportive route to report examples of inappropriate behaviour of a discriminatory nature. Deaneries and Programme Directors must ensure that on appointment trainees are made aware of the route in which inappropriate or discriminatory behaviour can be reported and supplied with contact names and numbers. Deaneries must also ensure contingency mechanisms are in place if trainees feel unhappy with the response or uncomfortable with the contact individual.

- monitoring of College Examinations;
- ensuring all assessments discriminate on objective and appropriate criteria and do not unfairly disadvantage trainees because of gender, ethnicity, sexual orientation or disability (other than that which would make it impossible to practise safely as a physician). All efforts shall be made to ensure the participation of people with a disability in training.

## 10 Syllabus

In the tables below, the “Assessment Methods” shown are those that are appropriate as **possible** methods that could be used to assess each competency. It is not expected that all competencies will be assessed and that where they are assessed not every method will be used. See section 5.2 for more details.

“GMP” defines which of the 4 domains of the Good Medical Practice Framework for Appraisal and Assessment are addressed by each competency. See section 3.2 for more details.

The syllabus itself covers 4 distinct areas these are:

The good Nuclear Medicine Physician

Basic Science and Regulations

Diagnostic skills

Therapy with radioisotopes

Within each of these 4 strands there will be 3 levels of competency from simple (level 1) which should be obtained at the start of training to the most complex (level 3) which should be obtained by the end of the training period.

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# The Good Nuclear Medicine Physician

## Level 1 Competencies

<b>The trainee will understand the role of the physician within nuclear medicine. They will understand the skills of other craft groups and their role in clinical leadership</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Definition of Nuclear Medicine	CbD, PGD	1,2,3,4
Understand the skills and knowledge required to perform the role of clinical lead in Nuclear Medicine	CbD, PGD	1,2,3,4
To understand the role of the radiation protection supervisor and subsequent legal relationship	CbD, PGD, MSF	1,2,3,4
Be aware of the need to work in a safe way understanding the concerns of the colleagues within the department	CbD, PGD, MSF	1,2,3,4
Understand the need for close working relationships with non-medically qualified staff	CbD, PGD, MSF	1,2,3,4
<b>Skills</b>		
Requirement to learn the legislative framework under which nuclear medicine operates	PGD	2,3
Ability to both learn from and teach non-medically qualified staff	PGD, TO	2,3
<b>Behaviours</b>		
Understand and accept the roles and limitations of the team delivering nuclear medicine services	MSF	3

<b>The trainee will understand the role that radioisotopes play in society:How they are produced and delivered to the patient</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Definition of Radioactivity	PGD, MSF	1,2
Understand the constraints when using radioisotopes such as half life	PGD	1,2
Understand that the general public may be fearful of the use of radioisotopes. Have sufficient knowledge to discuss and refute those fears	PGD, MSF	1,2
Understand the factors that may restrict the access of radio-isotopes to patients	PGD, MSF	1,2
Understand the security requirements in the holding and disposal of radio-isotopes including the possibility of terrorist mis-use of radioisotopes	PGD, MSF	1,2
Ability to understand the physical basis of radiation and radioisotopes	MSF	1,2
<b>Skills</b>		
Ability to understand the physical basis of radiation and radioisotopes	PGD	1,2
Establish a rapport with the patient and any relevant others (eg carers)	ACAT, CbD, Mini-CEX, Patient Survey	1, 3

Utilise open and closed questioning appropriately		
Listen actively and question sensitively to guide the patient and to clarify information	ACAT, Mini-CEX, PS	1, 3
Identify and manage communication barriers, tailoring language to the individual patient and others and using interpreters when indicated	ACAT, CbD, Mini-CEX, PS	1, 3
Establish a rapport with the patient and any relevant others (eg carers)	ACAT, CbD, Mini-CEX, PS	1, 3
<b>Behaviours</b>		
Good communication skills with patient	MSF, PS	1,2
Maintains a high level of safety awareness and consciousness at all times	CbD, mini-CEX	2
Be willing to provide patients with a second opinion	CbD, mini-CEX, MSF, PS	1, 3
Be confident and positive in one's own values	CbD, mini-CEX	1, 3

<b>The trainee shall be able to take an appropriate clinical history</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Recognises the importance of different elements of history	CbD, mini-CEX, MSF	1,2
Recognises that patients do not present history in structured fashion	CbD, mini-CEX, MSF	1,2
Knows likely causes and risk factors for conditions relevant to mode of presentation	CbD, mini-CEX, MSF	1,2
Recognises that history should inform examination and investigation and advice given to the referring clinician concerning the patients management plan	CbD, mini-CEX, MSF	1,2
<b>Skills</b>		
Focuses on relevant aspects of history	CbD, mini-CEX, MSF	1,2,3
Identifies and overcomes possible barriers to effective communication	CbD, mini-CEX, MSF	1,2,3
Assimilates history from the available information from patient and other sources	CbD, mini-CEX, MSF	1,2,3
Recognises and interprets the use of non verbal communication from patients and carers	CbD, mini-CEX, MSF	1,2,3
Manages alternative and conflicting views from family, carers and friends	CbD, mini-CEX, MSF	1,2,3
Supplements history with standardised instruments or questionnaires when relevant	CbD, mini-CEX, MSF	1,2,3
Manages time and draws consultation to a close appropriately	CbD, mini-CEX, MSF	1,2,3
<b>Behaviours</b>		
Shows respect and behaves in accordance with Good Medical Practice	mini-CEX, MSF	3

<b>The trainee shall be able to perform a relevant clinical examination of the patient</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understands the need for a valid clinical examination	CbD, mini-CEX, MSF	1,2
Understands the basis for clinical signs and the relevance of positive and negative physical signs	CbD, mini-CEX, MSF	1,2
Recognises constraints to performing physical examination and strategies that may be used to overcome them	CbD, mini-CEX, MSF	1,2
<b>Skills</b>		
Performs an examination relevant to the presentation and risk factors that is valid, targeted and time efficient	CbD, mini-CEX, MSF	1,2,3
Recognises the possibility of deliberate harm in vulnerable patients and reports to appropriate agencies	CbD, mini-CEX, MSF	1,2,3
Actively elicits and records important clinical findings	CbD, mini-CEX, MSF	1,2,3
<b>Behaviours</b>		
Shows respect and behaves in accordance with Good Medical Practice	mini-CEX, MSF	3

<b>The trainee shall learn to deal with the conflicting calls upon their time and learn how to prioritise work, especially being aware of the constraints caused by isotope decay and camera time</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand that organisation is key to time management	MSF	2
Understand that some tasks are more urgent or more important than others	MSF	2
Understand the need to prioritise work according to urgency and importance	MSF	2
Understand that some tasks may have to wait or be delegated to others	MSF	2
Outline techniques for improving time management	CbD	2
Understand the importance of delivering promptly the results of nuclear medicine investigations in disease management	MSF	2
<b>Skills</b>		
Estimate the time likely to be required for essential tasks and plan accordingly	MSF	2,3
Group together tasks when this will be the most effective way of working	MSF	2,3
Recognise the most urgent / important tasks and ensure that they are managed expediently	MSF	2,3
Regularly review and re-prioritise personal and team work load	MSF	2,3
Organise and manage workload effectively	MSF	2,3
<b>Behaviours</b>		

Ability to work flexibly and deal with tasks in an effective fashion	MSF	3
Recognise when you or others are falling behind and take steps to rectify the situation	MSF	3
Communicate changes in priority to others	MSF	3
Remain calm in stressful or high pressure situations and adopt a timely, rational approach	MSF	3

**The trainee shall show a correct approach to looking after patients and their families and friends in the context of nuclear medicine**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the concerns of patients with potentially serious and life threatening disease	PGD, MSF	2,3,4
Understand patient's fears about their disease	PGD, MSF	2,3,4
Understand why patients may refuse a test including fear of radiation and be able to explain the science behind why these tests are needed and the safety issues involved	PGD, MSF	2,3,4
Understand the special requirements and legal framework for dealing with the young, the old and the vulnerable and their guardians/parents/carers	PGD, MSF	2,3,4
<b>Skills</b>		
Demonstrate ability to communicate with patients	MSF, PS	2,3
Ability to look at the patient holistically and see their disease within a wider context for their individual health	MSF, PS	2,3
Show an ability to explain the dangers and benefits of radiation to the patient and society as a whole as appropriate	MSF, PS	2,3
<b>Behaviours</b>		
Show that the trainee is able to listen to and comprehend the fears and worries of patients and their families/friends/carers	MSF	3

**The trainee shall be able to give injections of radiopharmaceutical in a way which is safe for themselves, their patients and staff colleagues**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the concepts of sterile injection	PGD, DOPS, MSF	1,2
Understand the concepts of maintaining both sterility and radiation safety especially where this results in variance from normal clinical practice	PGD, DOPS, MSF	1,2
Able to communicate to the patient the reason for the test and obtain the required written and/or verbal consent	PGD, MSF	1,2
Able to explain to the patient the risks and benefits of radiation for their particular situation in particular reference to IR(ME)R	PGD, MSF	1,2
Understand the importance of avoiding radiation during pregnancy and the exceptions to this rule for the diagnosis of pulmonary embolism	PGD, DOPS, MSF	1,2
To understand the special requirements and constraints when administering a radiopharmaceutical including: <ul style="list-style-type: none"> <li>• Via a central venous catheter</li> <li>• In a patient with lymphoedema/vascular fistula</li> <li>• Tc-99m MAA for lung scanning</li> </ul>	PGD, DOPS, MSF	1,2

- TI-201
- Biological which may cause an allergic reaction

### Skills

Able to communicate to the patient the reason for the test and obtain the required written and/or verbal consent	PGD, DOPS, mini-CEX, MSF	2,3
Deal with the concerns of the patient and their friends and family in relation to the administration of radioactivity	PGD, DOPS, mini-CEX, MSF	2,3

### Behaviours

Demonstrate willingness to take advice from non-clinical staff	MSF	3
To empathise with the concerns of the patient and their family and friends	MSF, PS	3

## The trainee shall learn how to deal with a complaint, either against themselves, a colleague or the department

Knowledge	Assessment Methods	GMP
Define the local complaints procedure	CbD	3,4
Recognise factors likely to lead to complaints (poor communication, dishonesty etc)	CbD	3,4
Dealing with dissatisfied patients or relatives	MSF	3,4
Outline the principles of an effective apology	MSF	3,4
Identify sources of help and support when a complaint is made about yourself or a colleague	MSF	3,4
Skills		
Contribute to processes whereby complaints are reviewed and learned from	MSF	1,2,3
Explain comprehensibly to the patient the events leading up to a medical error	MSF	1,2,3
Recognise when something has gone wrong and identify appropriate staff to communicate this with	MSF	3,4
Act with honesty and sensitivity in a non-confrontational manner	MSF	3,4
Deliver an appropriate apology	MSF	1,2,3
Distinguish between system and individual errors	MSF	1,2,3
Show an ability to learn from previous errors	MSF	1,2,3
Behaviours		
Take leadership over complaint issues (If appropriate or legal)	MSF	2,3,4
Recognise the impact of complaints and medical error on staff, patients, and the National Health Service	MSF	2,3,4
Adopt behaviour likely to prevent complaints	MSF	3,4
Contribute to a fair and transparent culture around complaints and errors	MSF	2,3,4
Recognise the rights of patients, family members and carers to make a complaint	MSF	2,3,4

## Level 2 Competencies

**The trainee shall recognise the desirability of monitoring performance, learning from mistakes and adopting no blame culture in order to ensure high standards of care and optimise patient safety**

Knowledge	Assessment Methods	GMP
Understand the elements of clinical governance	CbD, mini-CEX, MSF	1,2
Recognise that governance safeguards high standards of care and facilitates the development of improved clinical services	CbD, mini-CEX, MSF	1,2
Define local and national significant event reporting systems relevant to specialty	CbD, mini-CEX,, MSF	1,2
Recognise importance of evidence-based practice in relation to clinical effectiveness	CbD, mini-CEX, MSF	1,2
Outline local health and safety protocols (fire, manual handling etc)	CbD, mini-CEX, MSF	1,2
Understand risk associated with the trainee's specialty work including biohazards and mechanisms to reduce risk	CbD, mini-CEX, MSF	1,2
Outline the use of patient early warning systems to detect clinical deterioration where relevant to the trainees clinical specialty	CbD, mini-CEX, MSF	1,2
Keep abreast of national patient safety initiatives including National Patient Safety Agency <sup>1</sup> , NCEPOD reports, NICE guidelines etc	CbD, mini-CEX, MSF	1,2
Skills		
Contribute to quality improvement processes e.g.: <ul style="list-style-type: none"> <li>• Audit of personal and departmental performance</li> <li>• Errors / discrepancy meetings</li> <li>• Critical incident reporting</li> <li>• Unit morbidity and mortality meetings</li> <li>• Local and national databases</li> </ul>	CbD, mini-CEX, MSF	1,2,3
Maintain a folder of information and evidence, drawn from your medical practice	CbD, mini-CEX, MSF	1,2,3
Reflect regularly on your standards of medical practice in accordance with GMC guidance on licensing and revalidation	CbD, mini-CEX, MSF	1,2,3
Behaviours		
Show willingness to participate in safety improvement strategies such as critical incident reporting	CbD, mini-CEX, mini-IPX, MSF	2,3,4
Engage with an open no blame culture	CbD, mini-CEX, mini-IPX, MSF	
Respond positively to outcomes of audit and quality improvement	CbD, mini-CEX, mini-IPX, MSF	
Co-operate with changes necessary to improve service quality and safety	CbD, mini-CEX, mini-IPX, MSF	

**The trainee shall know, understand and apply appropriately the principles, guidance and laws regarding medical ethics and confidentiality**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Outline and follow the guidance given by the GMC on confidentiality	CbD, mini-CEX, mini-IPX, MSF	1,2
Define the provisions of the Data Protection Act and Freedom of Information Act	CbD, mini-CEX, mini-IPX, MSF	1,2
Define the role of the Caldicott Guardian within an institution, and outline the process of attaining Caldicott approval for audit or research	CbD, mini-CEX, MSF	1,2
Outline the procedures for seeking a patient's consent for disclosure of identifiable information	CbD, mini-CEX, mini-IPX, MSF	1,2
Outline situations where patient consent, while desirable, is not required for disclosure e.g. communicable diseases, public interest	CbD, mini-CEX, mini-IPX, MSF	1,2
Recall the obligations for confidentiality following a patient's death	CbD, mini-CEX, mini-IPX, MSF	1,2
Recognise the problems posed by disclosure in the public interest, without patient's consent	CbD, mini-CEX, mini-IPX, MSF	1,2
Recognise the factors influencing ethical decision making: religion, moral beliefs, cultural practices	CbD, mini-CEX, mini-IPX, MSF	1,2
Do not resuscitate: Define the standards of practice defined by the GMC when deciding to withhold or withdraw life-prolonging treatment	CbD, mini-CEX, MSF	1,2
Outline the principles of the Mental Capacity Act	CbD, mini-CEX, mini-IPX, MSF	1,2
Be aware of the need to control access to data from various IT systems	CbD, mini-CEX, mini-IPX, MSF	1,2
Be aware of the use of pseudonym based data	CbD, mini-CEX, mini-IPX, MSF	1,2
<b>Skills</b>		
Use and share information with the highest regard for confidentiality, and encourage such behaviour in other members of the team	MSF	1,2,3
Use and promote strategies to ensure confidentiality is maintained e.g. anonymisation	MSF	1,2,3
Counsel patients on the need for information distribution within members of the immediate healthcare team	MSF	1,2,3
Counsel patients, family, carers and advocates tactfully and effectively when making decisions about resuscitation status, and withholding or withdrawing treatment	MSF	1,2,3
<b>Behaviours</b>		
Encourage ethical reflection in others	MSF	2,3,4
Show willingness to seek advice of peers, legal bodies, and the GMC in the event of ethical dilemmas over disclosure and confidentiality	MSF	2,3,4
Respect patient's requests for information not to be shared, unless this puts the patient, or others, at risk of harm	MSF	2,3,4

<b>The trainee will understand what is required to authorize a nuclear medicine request</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Demonstrate a wide knowledge of relevant medical knowledge	CbD, PGD, mini-IPX	1,2
Know the legal framework of IR(ME)R 2000	CbD, PGD, mini-IPX	1,2
Understand the relevancy and importance of the request in that patients clinical pathway	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Know how to discuss requests with referrer	CbD, MSF	2,3
Combine knowledge base of nuclear medicine and clinical medicine	CbD, MSF	2,3
Have confidence in that knowledge	CbD, MSF	2,3
<b>Behaviours</b>		
Ability to discuss and where necessary to contravene the requests of senior colleagues	MSF	3

<b>The trainee will understand how to produce a nuclear medicine report</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Know the role of the physician within the nuclear medicine department	mini-IPX	2,3
Recognise those artefacts which could influence a nuclear medicine report and where possible ensure a patient is re-imaged or if not possible know how this could influence the report	mini-IPX	2,3
Know how to structure a report into the following 4 phases: <ul style="list-style-type: none"> <li>• Indication</li> <li>• Description</li> <li>• Interpretation</li> <li>• Recommendation</li> </ul>	mini-IPX	2,3
Know which results must be communicated rapidly to the referring clinician and how to ensure this happens	mini-IPX	2,3
<b>Skills</b>		
Clear use of English	mini-IPX	2,3
Logical progression of report		
<b>Behaviours</b>		
Be open to comments and criticism of others	mini-IPX, MSF	3
Be confident that final report is correct	mini-IPX, MSF	3

## Level 3 Competencies

<b>The trainee will be able to present results of nuclear medicine in MDM/Ts</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the role of the MDM/T and in the case of cancer the legal framework in which it works	DOPS, MSF	2,3,4
Understand the role of a member of an MDM/T and its role in corporate decision making	DOPS, MSF	2,3,4
A good level of knowledge of both the nuclear medicine and clinical medicine in the topic covered within an MDM/T	DOPS, MSF	2,3,4
<b>Skills</b>		
Know how to work within a team	MSF	2,3,4
Have confidence in own skills and experience	MSF	2,3,4
Know your limitations within MDT/M	MSF	2,3,4
<b>Behaviours</b>		
Work with colleagues from different specialties to enhance patient care	MSF	3,4

<b>The trainee will undertake appropriate audit to measure their practice and their departments against known standards</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Know how to organize an audit	AA	2,4
Understand the idea of an audit standard	AA	2,4
Know how to identify the relevant guidelines available from the National Institute of Clinical Research, the British Society of Nuclear Medicine and the European Association of Nuclear Medicine	AA	2,4
<b>Skills</b>		
Be able to design and undertake one audit per year of training based on clinical practice of nuclear medicine	AA, MSF	2,4
Ability to communicate the results of such an audit	AA, MSF	2,4
Be able to implement changes in practice secondary to audit	AA, MSF	2,4
<b>Behaviours</b>		
Have a positive view of audit and be able to deal with fears induced in fellow workers positively	MSF	3

<b>The trainee will be able to critically assess research in the field of nuclear medicine</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
The trainee shall have the basic skills to be able to understand the following parameters used in assessing research and apply them critically to any relevant scientific paper: <ul style="list-style-type: none"> <li>• ethical basis of research</li> <li>• prospective or retrospective</li> <li>• sample size</li> <li>• appropriate methodology</li> <li>• how the data is assessed</li> <li>• appropriate use of statistics and their meaning</li> <li>• the use of the terms phase 1, phase 2 and phase 3 trials</li> <li>• understanding the requirements and limitations of randomized controlled trials</li> </ul>	CbD, PGD	1
How results may affect practice or determine the need for further research	CbD, PGD	1
The importance of looking at levels of evidence such as the Cochcrane method	CbD, PGD	1
<b>Skills</b>		
Be able to understand both strengths and weaknesses of research	PGD, TO	2
Understand the particular limitations which occur in research in imaging	PGD, TO	2
<b>Behaviours</b>		
An open and enquiring mind	MSF	3,4
Not accepting the status quo without supporting evidence	MSF	3,4

<b>The trainee will understand how to undertake a research project in diagnostic or therapeutic nuclear medicine</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Know how to write a research protocol	CbD	1,2
Know how to obtain any required funding	CbD	1,2
Work within the current ethical guidelines as stated within the European Clinical Trials Directive, national and local research ethics committees	CDB, PGD	1,2
Know how to determine the required sample size and how this can be obtained	PGD	1,2
Know the requirements for undertaking research within the trainee's hospital	CbD, MSF	1
Know the special requirements for research with radioisotopes including the role of ARSAC	PGD	1

<b>Skills</b>		
Have access to required statistical knowledge	PGD	1
Be able to write clearly any grant application forms, EUDRACT and ethics forms	PGD	1
Write a patient information sheet	MSF	1,2
Be able to recruit subjects	MSF	1,2
Be able to obtain consent correctly	MSF	1,2
<b>Behaviours</b>		
Be able to take on suggestions from co-researchers	MSF	3,4
Be positive in the face of adversity and setbacks	MSF	3,4

<b>The trainee will show that they can work with children undergoing nuclear medicine studies or therapy</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Have a working knowledge of how children develop and understand how they see the world at various stages in their development	CbD, PGD, MSF	1,2
Know the current law concerning consent in children and the rights and roles of the parent/guardian	CbD, PGD, MSF	1,2
Know the meaning of Ghillick competency and its application to nuclear medicine	CbD, PGD, MSF	1,2
Be aware of the need to ensure compliance to the principle of ALARA in children whilst ensuring a diagnostic study is performed	CbD, PGD, MSF	1,2
Whether or not any given radiopharmaceutical is licensed in children and if not the legal position	CbD, PGD, MSF	1,2
Be aware of the advantages and disadvantages of sedation in children	CbD, PGD, MSF	1,2
Know when children may need to be anaesthetised for a scan and how this is safely achieved	CbD, PGD, MSF	1,2
<b>Skills</b>		
Be able to communicate what needs to be done with the child	CbD, PGD, MSF	1,2,3
Gain the child's trust and co-operation	CbD, PGD, MSF	1,2,3
Be able to perform intravenous injection or if required bladder catheterisation in a way that discomforts the child the least	CbD, PGD, MSF	1,2,3
Be able to engage the child and maintain co-operation throughout the study	CbD, PGD, MSF	1,2,3
Be aware of the special requirements for resuscitation in children	CbD, PGD, MSF	1,2,3
Know the present guideline from ARSAC vis-à-vis recommended adjustments to activity for children for any given radiopharmaceutical	CbD, PGD, MSF	1,2,3
Know similar dose reductions for Frusemide and sodium pyrophosphate	CbD, PGD, MSF	1,2,3
Be able recruit other team members including play therapists and parents/guardians into the process of scanning the child	CbD, PGD, MSF	1,2,3

<b>Behaviours</b>		
Be open and honest	MSF	3,4
Be aware of the fears and hopes of the child and their parent guardian	MSF	3,4
Involve the parent/guardian in the procedure and be willing to answer their questions	MSF	3,4
Be open to the fears of the child and their parents/guardians about illness especially when terminal or potentially fatal	MSF	3,4

**The trainee will be able to apply for a license from the Administration of Radioactive Substances Advisory Committee (ARSAC) N.B. Though this knowledge must be obtained whilst the candidate is a trainee the law states that an application cannot be made until the candidate has a CCT and has been appointed to a specific Consultant post**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the role of ARSAC	PGD	1,2
Understand the different licences available	PGD	1,2
Know how to fill in an ARSAC application form	PGD	1,2
Provide required evidence of competence	PGD	1,2
<b>Skills</b>		
Able to provide the correct data on forms	PGD	3
Ability to both learn from and teach non-medically qualified staff	PGD	3
<b>Behaviours</b>		
Open to help from colleagues	MSF	3,4

**The trainee shall understand the structure of the NHS and the management of local healthcare systems in order to be able to participate fully in managing healthcare provision**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the guidance given on management and doctors by the GMC	CbD, PGD	1,2,3,4
Understand the local structure of NHS systems in your locality recognising the potential differences between the four countries of the UK	CbD, PGD	1,2,3,4
Understand the structure and function of healthcare systems as they apply to your specialty	CbD, PGD	1,2,3,4
Understand the principles of: <ul style="list-style-type: none"> <li>• Clinical coding</li> <li>• European Working Time Regulations</li> <li>• National Service Frameworks</li> <li>• Health regulatory agencies (e.g. CHI, NICE, Scottish Government)</li> <li>• NHS Structure and relationships</li> </ul>	CbD, PGD	1,2,3,4

<ul style="list-style-type: none"> <li>• NHS finance and budgeting</li> <li>• Consultant contract and the contracting process</li> <li>• Resource allocation</li> <li>• The role of the Independent sector as providers of healthcare</li> </ul>		
Understand the principles of recruitment and appointment procedures	CbD, PGD	1,2,3,4
Understand the need to be non-discriminatory in the recruitment process	CbD, PGD	1,2,3,4
<b>Skills</b>		
Participate in managerial meetings	CbD, MSF	1,2,3
Take an active role in promoting the best use of healthcare resources	CbD, MSF	1,2,3
Work with stakeholders to create and sustain a patient-centred service	CbD, MSF	1,2,3
Employ new technologies appropriately, including information technology	CbD, MSF	1,2,3
Be involved in business case development for new equipment	CbD, MSF	1,2,3
<b>Behaviours</b>		
Recognise the importance of just allocation of healthcare resources	CbD, MSF	2,3,4
Recognise the role of doctors as active participants in healthcare systems	CbD, MSF	2,3,4
Respond appropriately to health service targets and take part in the development of services	CbD, MSF	2,3,4
Recognise the role of patients and carers as active participants in healthcare systems and service planning	CbD, MSF	2,3,4
Show willingness to improve managerial skills (e.g. management courses) and engage in management of the service	CbD, MSF	2,3,4

## Basic Science and Regulations

### Level 1 Competencies

<b>The trainee will learn the basic physics and mathematics as related to the delivery of safe nuclear medicine</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Structure and modes of decay of radioactive atoms	PGD	1,2
Interaction of emissions from radioactive atoms with matter	PGD	1,2
Biological implications of and radiation hazards from ionising radiation	PGD	1,2
Molecular biology	PGD	1,2
Probability theory	PGD	1,2
Parametric and non-parametric statistics	PGD	1,2
Understand how radioisotopes are manufactured and problems of supply and transportation	PGD	1,2
Appropriate mathematics and physics applied to radionuclide tracer theory, modelling of tracer kinetics and quantitative imaging	PGD	1,2
<b>Skills</b>		
Understand how these are to be applied on a daily basis	PGD	1,2
<b>Behaviours</b>		
Be willing to learn	PGD, MSF	1,2,3,4
Understand their limitations and ask for assistance from non-medical colleagues	PGD, MSF	1,2,3,4

<b>The trainee will learn the methods by which radiation is detected in nuclear medicine and the principles of the computer based image analysis and display including artefacts</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Theory of systems used to detect and analyse emissions from radioactive atoms	CbD, PGD	1,2
Knowledge of how detection systems are used, calibrated and tested in Nuclear Medicine	CbD, PGD	1,2
Principles of collimation and practical experience with the use of collimators	CbD, PGD	1,2
Understanding of the tracer principle in particular the use of time activity analysis	CbD, PGD	1,2
Principles of single-photon emission tomography and co-incidence counting	CbD, PGD	1,2
Principles of image reconstruction	CbD, PGD	1,2
Understand how artefacts occur and how their effect can be reduced if possible and if not their effect on image quality and clinical report	CbD, PGD	1,2
Understand the principles of positron imaging	CbD, PGD	1,2

Understand how cross sectional imaging is performed, The parameters used to reduce radiation dose to the patients. How images are registered, and how images can be mis-registered and how this may effect the resultant fused image	CbD, PGD	1,2
Kinetics of radioactive tracers used in Nuclear Medicine	CbD, PGD	1,2
Use of principles of kinetics and modelling techniques to calculate parameters such as glomerular filtration rate etc	CbD, PGD	1,2
Physiological principles of tracer techniques	CbD, PGD	1,2
Errors associated with quantitative measurements	CbD, PGD	1,2
<b>Skills</b>		
Understand how these are to be applied on a daily basis	PGD, mini-IPX	1,2
Able to apply these techniques	PGD, mini-IPX	1,2
Able to understand how problems with these techniques can lead to deterioration of image quality and how this may be affect clinical reporting	PGD, mini-IPX	1,2
Recognise and if possible correct mis-registration in hybrid imaging and if such correction is not possible know the possible effect on image interpretation	PGD, mini-IPX	1,2
Be able to check the validity of non imaging tests such as a GFR estimation	PGD, mini-IPX	1,2
<b>Behaviours</b>		
Be willing to learn	PGD, MSF	1,2,3,4
Understand their limitations and ask for assistance from non-medical colleagues	PGD, MSF	1,2,3,4

**The trainee will learn the principles of radiation biology and radiation protection and know the legal framework in which nuclear medicine is practised**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Theory of biological effects of high and low-level radiation from unsealed sources	PGD	1,2
Calculation of radiation dose from radiopharmaceuticals (Effective dose [ED])	PGD	1,2
Know the importance of the upper limits of activities of radioactivity as defined by ARSAC for each investigations	PGD	1,2
Know the circumstances in which these upper limits may be breached	PGD	1,2
Know the recommended adjustments made in the administered activity for children	PGD	1,2
Know advice must be given to lactating women receiving radioisotopes	PGD	1,2
Know when radioisotopes can be given to women who are pregnant	PGD	1,2
Basic principles of radionuclide therapy	PGD	1,2
Nature of the cancerous process and the radiobiological basis of cancer radionuclide therapy	PGD	1,2

Management of radiation accidents such as spills relating to Nuclear Medicine	PGD	1,2
Know those laws affecting the practice of nuclear medicine	PGD	1,2
Understand the role of the regulatory authorities such as ARSAC, The Environment Agency, The Health and Safety Executive, The Health Commission and their effect on practice and understand that infringements of these regulations represent a criminal offence. National and international regulatory requirement on the practice of nuclear medicine including: IRR99; MARS legislation; ARSAC; RSA 93; IR(ME)R 2000; Medical guidance notes; Product licenses and other appropriate legislation	PGD	1,2
Regulatory requirements which apply to the design and operation of radiopharmacies GMP 1997 (Orange Guide)	PGD	1,2
Regulations controlling transport of radioactive materials in the UK RM(Road Transport) (GB) R 1996	PGD	1,2
Mechanism by which the regulations are applied and policed within the UK	PGD	1,2
Health and safety regulations governing safe practice e.g. COSHH Regulations 1999	PGD	1,2
ALARA (as low as reasonably achievable) and ALARP (as low as reasonably practical)		
Know the role of the RPS in Nuclear Medicine	PGD	1,2
<b>Skills</b>		
Understand how these are to be applied on a daily basis	PGD, MSF	1,2,3,4
Learn to work in a way which is safe for self and colleagues	PGD, MSF	1,2,3,4
Understand the importance of complying with the legal framework of nuclear medicine	PGD, MSF	1,2,3,4
<b>Behaviours</b>		
Be willing to learn	PGD, MSF	1,2,3,4
Understand their limitations and ask for assistance from non-medical colleagues	PGD, MSF	1,2,3,4
Be willing to self report incidents such as spills and mal administration to the RPS	PGD, MSF	1,2,3,4
Be willing to report any witnessed infringements of the regulations to the proper authorities	PGD, MSF	1,2,3,4

## Level 2 Competencies

The trainee will learn the methods by which radiopharmaceuticals are prepared		
Knowledge	Assessment Methods	GMP
Production of radionuclides using reactors, cyclotrons and generators	PGD	1,2
Physical properties of radionuclides, clinical applications	PGD	1,2
Physicochemical and biological properties of different radiopharmaceuticals in routine clinical practice, clinical trials and under development	PGD	1,2
Different formulations used in Nuclear Medicine	PGD	1,2
Cell Labelling techniques	PGD	1,2
Principles of Quality Assurance (QA) in the radiopharmacy	PGD	1,2
Quality control parameters which determine the quality of radiopharmaceuticals including radionuclide & radiochemical purity	PGD	1,2
Principles of aseptic preparation	PGD	1,2
Skills		
Be able to measure the activity of radiopharmaceutical given correctly	PGD, DOPS	1,2
Be able to draw up a patient dose using aseptic technique	PGD, DOPS	1,2
Be able to perform and understand a simple quality assurance test	PGD, DOPS	1,2
Behaviours		
Be willing to learn	PGD, MSF	1,2,3,4
Be willing to use techniques that reduce the risk of cross infection between patients and ensure those around also comply	PGD, MSF	1,2,3,4
Understand their limitations and ask for assistance from non-medical colleagues	PGD, MSF	1,2,3,4

## Diagnostic Nuclear Medicine

### Level 1 Competencies

<b>The trainee will learn how to report correctly a ventilation/perfusion lung scan</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of thrombo-embolic disease	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand the methodology of Ventilation perfusion imaging including the different methods of ventilation and their advantages and disadvantages	CbD, PGD, mini-IPX	1,2
Understand the special circumstances of imaging women during pregnancy and lactation	CbD, PGD, mini-IPX	1,2
Understand how the images are displayed for reading	CbD, PGD, mini-IPX	1,2
Understand the criteria for positivity and negativity in the diagnosis of pulmonary embolism	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Give an effective and safe injection of Tc-99 MAA	CbD, PGD, mini-IPX	1,2
Recognise the patterns of positivity and negativity for Pulmonary Embolism	CbD, PGD, mini-IPX	1,2
Recognise the patterns of abnormality that suggest an alternate diagnosis such as COPD or lung cancer	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Be willing to transmit important urgent results to the patient and their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the fears and concerns of patients concerning radiation especially if pregnant	CbD, MSF	1,2,3,4
Be able to impart to the patient relevant public health information for example smoking cessation	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly a bone scan in a patient with metastases</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of malignant disease and how it affects the bones	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand those conditions that can affect the result of the bone scintigraphy	CbD, PGD, mini-IPX	1,2
Understand the methodology of bone imaging	CbD, PGD, mini-IPX	1,2
Understand how the images are displayed for reading	CbD, PGD, mini-IPX	1,2
Understand the criteria for positivity and negativity in the diagnosis of bone metastases	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
Know when SPECT may be helpful	CbD, PGD, mini-IPX	1,2
If applicable have sufficient knowledge of non-contrast CT to interpret SPECT-CT image of the spine and pelvis	CbD, PGD, mini-IPX	1,2
Know when to suggest treatment with bone palliation radionuclide methods	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Recognise the patterns of positivity and negativity for bone metastases	CbD, PGD, mini-IPX	1,2
Recognise the patterns of abnormality that suggest an alternate diagnosis such as degenerative disease. Know how to use alternative imaging such as CT or SPECT-CT to determine this	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Be willing to transmit important urgent results to the patient and their referring clinician especially if the patient was not known to have a history of cancer	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the fears and concerns of patients with known or possible cancer	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly a bone densitometry (optional)</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of osteoporosis and the factors which influence bone density	CbD, PGD, mini-IPX	1,2
Know the methodology of dual photon densitometry	CbD, PGD, mini-IPX	1,2
Understand the criteria for osteoporosis and low bone density	CbD, PGD, mini-IPX	1,2
Know current guidelines for use of bone densitometry	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Ensure current regions of interest have been drawn	CbD, PGD, mini-IPX	1,2
Know the relevance of T and Z scores	CbD, PGD, mini-IPX	1,2
Know what recommendations for treatment should be given	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing bone densitometry	CbD, MSF	1,2,3,4
Be able to discuss with patients those life style factors which may reduce the risk of osteoporosis	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly a static renal DMSA scan</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of reflux nephropathy	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Know the latest guidelines on if and when Tc-99m DMSA should be performed in children	CbD, PGD, mini-IPX	1,2
Know the time interval required between last urinary tract infection and scanning	CbD, PGD, mini-IPX	1,2
Understand the methodology of DMSA imaging including the different views taken, the meaning of quantification and in adults the possible use of SPECT	CbD, PGD, mini-IPX	1,2
Understand how the images are displayed for reading	CbD, PGD, mini-IPX	1,2
Understand the criteria for positivity and negativity in the diagnosis of renal scars and other space occupying lesions	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2

<b>Skills</b>		
Recognise abnormal patterns of relative renal function, and obstruction	CbD, PGD, mini-IPX	1,2
Recognise the patterns of abnormality that suggest an alternate diagnosis such as cyst or tumour	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the parents for children undergoing the test including why there may a considerable delay until imaging can be performed	CbD, MSF	1,2,3,4
Understanding the Ghillick rules on consent in children	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly dynamic renography</b>		
	<b>Assessment Methods</b>	<b>GMP</b>
<b>Knowledge</b>		
Understand the pathophysiology of renal clearance	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Know the latest guidelines on if and when Dynamic renography should be performed in children	CbD, PGD, mini-IPX	1,2
Understand the methodology of dynamic renography imaging including whether or not Frusemide should be given and the timing of that injection	CbD, PGD, mini-IPX	1,2
Understand how the images are displayed for reading	CbD, PGD, mini-IPX	1,2
Understand the criteria for interpreting dynamic renography, for example the criteria for diagnosing/excluding obstruction	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Give an effective and safe injection of Tc-99m DTPA/MAG3 via a central line if required	CbD, PGD, mini-IPX	1,2
Recognise abnormal patterns of relative renal function or obstruction	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patient and parents of children undergoing the test	CbD, MSF	1,2,3,4
Understanding the Ghillick rules on consent in children	CbD, MSF	1,2,3,4

Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly a simple GI transit study</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of the problems that can occur in the GI concerning transit including scleroderma, diabetes and severe constipation	CbD, PGD, mini-IPX	1,2
Understand the preparation requirement for each study	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient and examine the patient as required	CbD, PGD, mini-IPX	1,2
Determine the correct study for the indication and correct type of meal to be administered	CbD, PGD, mini-IPX	1,2
Understand the methodology of imaging different types of GI transit including that of the oesophagus, stomach and colon	CbD, PGD, mini-IPX	1,2
Understand how the images and any computer analysis are displayed for reading	CbD, PGD, mini-IPX	1,2
Understand the criteria for positivity and negativity in the diagnosis of GI dysmotility	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Recognise the patterns of abnormality suggestive of GI dysmotility in the oesophagus, stomach and colon	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing the test undergoing radionuclide tests	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn how to report correctly a thyroid scan**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of thyroid disease	CbD, PGD, mini-CEX, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-CEX, mini-IPX	1,2
Understand those factors than can effect thyroid scinigtigraphy and be able to decide if the study can proceed	CbD, PGD, mini-CEX, mini-IPX	1,2
Understand the methodology of thyroid imaging including any additional views taken, including the possible use of SPECT and SPECT-CT	CbD, PGD, mini-CEX, mini-IPX	1,2
Understand how the images are displayed for reading	CbD, PGD, mini-CEX, mini-IPX	1,2
Understand what may influence the result of a thyroid scan and how abnormality can be diagnosed	CbD, PGD, mini-CEX, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-CEX, mini-IPX	1,2
<b>Skills</b>		
Recognise the patterns of abnormality suggestive of Grave's disease, multi-nodular goitre, toxic nodule, "cold" nodule and thyroiditis	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients	CbD, mini-IPX, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician especially if a cold nodule is found	CbD, mini-IPX, MSF	1,2,3,4
Be able to recommend I-131 if relevant	CbD, mini-IPX, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, mini-IPX, MSF	1,2,3,4
Be willing to learn	CbD, mini-IPX, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, mini-IPX, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, mini-IPX, MSF	1,2,3,4
Understand the need for audit and research	CbD, mini-IPX, MSF	1,2,3,4

**The trainee will be able to perform myocardial stressing as required for myocardial perfusion scintigraphy-uncomplicated patient**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of coronary artery disease	CbD, PGD, DOPS	1,2
Be able to take an appropriate history from the patient and examine the patient as required	CbD, PGD, DOPS	1,2
Understand the methodology of adenosine/dobutamine and physical stress.	CbD, PGD, DOPS	1,2

Understand correct patient preparation for each type of test and relevant drug interactions	CbD, PGD, DOPS	1,2
Understand the contra-indications for each type of stress and alternates that can be offered in such circumstances	CbD, PGD, DOPS	1,2
Understand the time parameters such as how long should stress be administered, when during that stress should the radiopharmaceutical be given, when should the stress be terminated, what time should the patient be scanned and any instructions to the patient between stressing and imaging	CbD, PGD, DOPS	1,2
<b>Skills</b>		
Have their ALS certificate	CbD, PGD, DOPS	1,2
Recognise abnormalities on a resting ecg that mean the stress can proceed safely or should not be done	CbD, PGD, DOPS	1,2
Understand what constitutes an appropriate and sufficient stress for a diagnostic test	CbD, PGD, DOPS	1,2
Recognise those symptoms and signs that should result in termination of the stress test including changes in vital signs and arrhythmias	CbD, PGD, DOPS	1,2
Be able to communicate to the patient what is happening and what to expect to happen and the duration of any side effects	CbD, PGD, DOPS	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing radionuclide tests and stress tests	DOPS, MSF	1,2,3,4
Be willing to explain why the stress test is required for the management of their condition and obtain oral/written consent as required	DOPS, MSF	1,2,3,4
Be open in discussion with the patient about those measures that can be taken to reduce the risk of ischaemic heart disease including dietary advice and smoking cessation	DOPS, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	DOPS, MSF	1,2,3,4
Be willing to learn	DOPS, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	DOPS, MSF	1,2,3,4
Understand the need for self reflection of skills	DOPS, MSF	1,2,3,4

## Level 2 Competencies

The trainee will learn how to report correctly a myocardial perfusion SPECT		
Knowledge	Assessment Methods	GMP
Understand the pathophysiology of coronary artery disease	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand when attenuation correction should be applied and the consequences of that action	CbD, PGD, mini-IPX	1,2
Understand when gating should be performed and the usefulness of a gated study including normal ranges for parameters such as LVEF, EDV and ESV	CbD, PGD, mini-IPX	1,2
Understand how the images analysed and are displayed for reading	CbD, PGD, mini-IPX	1,2
Understand those factors which may influence image quality, including attenuation and left bundle branch block	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
Understand the role of SPECT-CT and the relevance of calcium scoring	CbD, PGD, mini-IPX	1,2
Skills		
Understand the strengths and weaknesses and the biodistribution of the main agents used in MPS –Tl-201 chloride, Tc-99m MIBI and Tc-99m tetrofosmin	CbD, PGD, mini-IPX	1,2
Recognise the patterns of abnormality that define a positive and negative test	CbD, PGD, mini-IPX	1,2
Be able to determine if there are artefacts such as attenuation affecting the scan and interpret how these may change the result	CbD, PGD, mini-IPX	1,2
Be able to use a gated study to improve accurate reading of the MPS	CbD, PGD, mini-IPX	1,2
In a gated study be able to identify dyskinesia and the presence of apical aneurysm	CbD, PGD, mini-IPX	1,2
Be able to identify abnormal activity outside of the heart on the MPS or abnormality on the CT (if performed) that could represent other significant thoracic abnormality	CbD, PGD, mini-IPX	1,2
Behaviours		
Understand the concerns of the patients undergoing radionuclide tests	CbD, MSF	1,2,3,4
Be willing to repeat imaging if this will improve diagnostic accuracy and be able to communicate why this must be done to the patient	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4

Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly bone scintigraphy in a range of benign diseases</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of bone disease including congenital disease, trauma infection, inflammatory disease, hamartomas, degenerative and metabolic diseases and benign bone tumours	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand the methodology of 3 phase bone scanning, SPECT and SPECT-CT	CbD, PGD, mini-IPX	1,2
Have sufficient knowledge of cross-sectional imaging to be able to read a SPECT-CT study	CbD, PGD, mini-IPX	1,2
Understand how the images are analysed and displayed for reading	CbD, PGD, mini-IPX	1,2
Understand the criteria for normal and abnormal and the clinical relevance of these results	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
Understand the legal consequences of diagnosing a non-accidental injury in both a child and a vulnerable adult	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Recognise the patterns of abnormality which can be seen within bones	CbD, PGD, mini-IPX	1,2
Understand the non-specific nature of bone scintigraphy and the importance of pattern recognition in reading a scan	CbD, PGD, mini-IPX	1,2
Recognise potentially life threatening conditions such as discitis, septic arthritis	CbD, PGD, mini-IPX	1,2
Recognise the appearances of non-accidental injury in both children and adults	CbD, PGD, mini-IPX	1,2
Be able to use CT (and other available imaging) to improve accuracy of reporting	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing radionuclide tests	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
If non accidental injury is suspected be willing and able to contact the proper authorities	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4

Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn how to report correctly and understand dynamic renography with interventions**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of vesico-ureteric reflux	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand the methodology of indirect and direct (optional) radionuclide micturating cystography	CbD, PGD, mini-IPX	1,2
Understand how images are analysed and are displayed for reading	CbD, PGD, mini-IPX	1,2
Understand the criteria for reflux and the clinical relevance of grade III/IV reflux	CbD, PGD, mini-IPX	1,2
Understand the pathophysiology of renovascular hypertension	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient and examine the patient as required	CbD, PGD, mini-IPX	1,2
Know the drugs that will interfere with the test and the times they need to be stopped	CbD, PGD, mini-IPX	1,2
Understand the physiological differences that occur when giving ACE inhibitors	CbD, PGD, mini-IPX	1,2
Understand the criteria for renovascular hypertension as seen on a pre and post ACE inhibitor dynamic renography	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Recognise how the images may need to be manipulated to see reflux	CbD, PGD, mini-IPX	1,2
Be able to apply any additional mathematic analysis such as de-convolution which will aid in the diagnosis of renovascular hypertension	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing radionuclide tests	CbD, MSF	1,2,3,4
To be able to communicate with patients the importance of complying with anti-hypertensive medication if this has been prescribed	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn how to report correctly and understand non-imaging tests done in nuclear medicine**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the historical uses of red cell survival studies and schilling tests	CbD, PGD	1,2
Understand the present role of the following non-imaging studies: Glomerular filtration rate, red cell mass, plasma volume and bile salt absorption	CbD, PGD	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD	1,2
Understand the methodology of Glomerular filtration rate, red cell mass and bile salt absorption	CbD, PGD	1,2
Understand how data is analysed and possible causes of error, and how if possible these can be minimised and corrected	CbD, PGD	1,2
<b>Skills</b>		
Recognise the significance of Glomerular filtration rate, red cell mass and bile salt absorption tests	CbD, PGD	1,2
<b>Behaviours</b>		
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn how to report correctly and understand hepatobiliary scintigraphy**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of biliary disease in particular the causes of neonatal jaundice and cholecystitis. Also in post liver transplant(optional)	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand the methodology of HIDA and preparation for imaging in biliary atresia and in suspected gall bladder disease	CbD, PGD, mini-IPX	1,2
Understand how images are analysed and displayed for reading	CbD, PGD, mini-IPX	1,2
Understand the criteria for a positive study in biliary atresia and for a positive study in gall bladder disease	CbD, PGD, mini-IPX	1,2
Know when it is correct to give a fatty meal or CCK stimulus	CbD, PGD, mini-IPX	1,2

Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Be able to make a definitive diagnosis of biliary atresia or other causes of neonatal jaundice	CbD, PGD, mini-IPX	1,2
Be able to give a fatty meal or CCK safely	CbD, PGD, mini-IPX	1,2
Understand the results of hepatobiliary scintigraphy	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing radionuclide tests	CbD, MSF	1,2,3,4
Be able to deal with the concerns of parents of very sick infants	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn understand how single photon emission tomography (SPET) can be used in the management of CNS disease**

	<b>Assessment Methods</b>	<b>GMP</b>
<b>Knowledge</b>		
Understand the pathophysiology of a wide range of disease affecting the brain including tumour, cerebrovascular disease, degenerative disease and Parkinson's syndromes	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Have a basic understanding of neuroanatomy with knowledge of the sites of major intra-cerebral features as seen on cross sectional imaging and how this correlates with SPET	CbD, PGD, mini-IPX	1,2
Understand which nuclear medicine test should be applied for a particular situation	CbD, PGD, mini-IPX	1,2
Know which medications may need to be stopped or reduced for I-123 iopflupane imaging	CbD, PGD, mini-IPX	1,2
Know the normal distribution of Tc-99m MIBI/TI-201, Tc-99m HMPAO and I-123 iopflupane	CbD, PGD, mini-IPX	1,2
Recognise and understand the changes in distribution in disease	CbD, PGD, mini-IPX	1,2
Identify when and what quantification may help diagnosis	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
As appropriate be able to make a diagnosis of intracerebral tumour with TI-201/Tc-99m MIBI	CbD, PGD, mini-IPX	1,2

Be able to identify the difference between Alzheimer's and multi-infarct dementia on Tc-99m HMPAO imaging	CbD, PGD, mini-IPX	1,2
Be able to identify Parkinson's syndromes with I-123 iopflupane and determine the severity of disease	CbD, PGD, mini-IPX	1,2
Understand those factors that can result in artefacts that can reduce the accuracy of the test and how these can be corrected or their effect reduced	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing radionuclide tests	CbD, MSF	1,2,3,4
Be able to work with vulnerable adults and understand the legal requirements of consent in the mentally frail	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn how to report correctly and understand radionuclide imaging of infection and inflammation**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of infection and inflammation; in particular, be aware of the patterns of infection that can occur in osteomyelitis, infected joint prosthesis and in patients who are immunocompromised. Know the criteria for diagnosis of fever of unknown origin	CbD, PGD, mini-IPX	1,2
In addition be aware of the causes and clinical presentation of inflammatory bowel disease	CbD, PGD, mini-IPX	1,2
Know when to apply a specific infection/inflammation study or when a simpler less specific study may be sufficient	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand the strengths and weaknesses of Ga-67 citrate, In-111 or Tc-99m HMPAO labelled leucocytes and Tc-99m Selusomab. Know when each should be used	CbD, PGD, mini-IPX	1,2
Be aware of the imaging times for each radiopharmaceutical and their normal distribution	CbD, PGD, mini-IPX	1,2
Be aware of any medications which may interfere with radionuclide infection /inflammation imaging and decide if their use should be stopped or modified	CbD, PGD, mini-IPX	1,2
Understand how images are analysed and displayed for reading	CbD, PGD, mini-IPX	1,2

Understand the criteria for a positive study with each radiopharmaceutical in each clinical scenario	CbD, PGD, mini-IPX	1,2
Be able to decide whether alternate or additional imaging is required	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Be able to identify the difference between normal distribution and pathology in Ga-67 imaging	CbD, PGD, mini-IPX	1,2
Be able to identify the variable results of Ga-67 in sarcoid and their significance	CbD, PGD, mini-IPX	1,2
Be able to determine the presence of inflammatory bowel disease with labelled leucocytes	CbD, PGD, mini-IPX	1,2
Know when SPET or SPET/CT may be helpful and be able to interpret those results	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing radionuclide tests	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly and understand parathyroid localisation</b>			
	<b>Assessment Methods</b>	<b>GMP</b>	
<b>Knowledge</b>			
Understand the pathophysiology of parathyroid disease and its clinical importance. Take note of the possibility of ectopic sites of adenomas	CbD, PGD, mini-IPX	1,2	
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2	
Understand the different methodologies for imaging parathyroid adenomas and know one of these in detail	CbD, PGD, mini-IPX	1,2	
Know when SPET or SPET/CT may be of use	CbD, PGD, mini-IPX	1,2	
Understand how images are analysed and displayed for reading	CbD, PGD, mini-IPX	1,2	
Be able to decide the relationship between radionuclide imaging and ultrasound in localisation of a parathyroid adenoma	CbD, PGD, mini-IPX	1,2	
<b>Skills</b>			
Be able to decide if a study is positive for a parathyroid adenoma and be able to identify its site (possibly with the aid of SPET-SPET/CT)	CbD, PGD, mini-IPX	1,2	
<b>Behaviours</b>			
Understand the concerns of the patients undergoing radionuclide	CbD, MSF	1,2,3,4	

tests		
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn how to report correctly and understand radionuclide imaging of adrenal adenoma**

	<b>Assessment Methods</b>	<b>GMP</b>
<b>Knowledge</b>		
Understand the pathophysiology of medullary and cortical adrenal tumours. Know the probability of bilateral disease or malignant spread	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand the methodologies for imaging the adrenal gland with I-123/I-131 nor cholesterol and I-123/I-131 mIBG	CbD, PGD, mini-IPX	1,2
Know which medications may interfere with imaging and be able to determine if they can be stopped or modified before imaging	CbD, PGD, mini-IPX	1,2
Know when the use of renal imaging and/or SPET or SPET/CT may be of use for localisation	CbD, PGD, mini-IPX	1,2
Understand how the images analysed and are displayed for reading	CbD, PGD, mini-IPX	1,2
Be able to decide the relationship between radionuclide imaging in adrenal tumours and other imaging modalities including CT and MRI	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Be able to decide if a study is positive and be able to identify its site (possibly with the aid of SPET-SPET/CT)	CbD, PGD, mini-IPX	1,2
Be able to differentiate between physiological and pathological activity of tracer	CbD, PGD, mini-IPX	1,2
Be able to use any available quantification to determine positivity	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing radionuclide tests	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4

Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly and understand sentinel node localisation</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of malignant disease, know how it spreads and in which cancers sentinel node localisation is both possible and useful	CbD, PGD, DOPS, mini-IPX	1,2
Be aware of national and international guidelines concerning sentinel node localisation	CbD, PGD, DOPS, mini-IPX	1,2
Be able to take an appropriate history from the patient and examine the patient as required	CbD, PGD, DOPS, mini-IPX	1,2
Understand the different methodologies for injection and imaging sentinel nodes depending on the primary cancer and learn at least one of these in detail.	CbD, PGD, DOPS, mini-IPX	1,2
Know when and if SPET/CT may be of use		
Understand how the images analysed and are displayed for reading, including the use of a "shadowgram"	CbD, PGD, DOPS, mini-IPX	1,2
<b>Skills</b>		
Be able to explain to the patient that the test is a form of staging and not diagnostic	CbD, PGD, DOPS, mini-IPX	1,2
Be able to give a sentinel node injection correctly to ensure good lymph flow in a timely manner	CbD, PGD, DOPS, mini-IPX	1,2
Be able to identify the site of a sentinel node and mark it for surgery and communicate the results of marking effectively with the referring surgeon	CbD, PGD, DOPS, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients undergoing radionuclide tests	CbD, MSF	1,2,3,4
Understand the fears of a patient with a potentially fatal cancer	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn how to report correctly and understand tumour specific imaging with single photon agents**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of cancers that can be imaged with radionuclide techniques and understand the specific nature of those techniques in tumour sites and how nuclear medicine techniques are used to compliment CT and MRI	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand the different methodologies for imaging Cancers with single photon including the following; <ul style="list-style-type: none"> <li>• I-123/I-131 mIBG in neuroblastoma</li> <li>• I-123/I-131 mIBG in malignant pheochromocytoma</li> <li>• I-123/I-131 mIBG in neuroendocrine tumours</li> <li>• In-111 pentetretotide in neuroendocrine tumours</li> <li>• I-123/I-131 in thyroid cancer</li> <li>• Tc-99m DMSA (V) in medullary cell thyroid cancer</li> <li>• Tc-99m MIBI in breast cancer (optional)</li> <li>• Tc-99m depreotide in lung cancer (optional)</li> <li>• Tl-201 in Kaposi's sarcoma (optional)</li> <li>• Tc-99m MIBI in sarcomas (optional)</li> </ul>	CbD, PGD, mini-IPX	1,2
Know those that are not optional in detail	CbD, PGD, mini-IPX	1,2
Know when SPET or SPET/CT may be of use	CbD, PGD, mini-IPX	1,2
Know the clinical situation in which test may be applied and understand the clinical imaging schedule for each referring to national and EANM guidelines as required	CbD, PGD, mini-IPX	1,2
Know what medication must be stopped/modified before administration of the radiopharmaceuticals This includes the drugs that interfere with uptake of I-123/I-131 mIBG, injected somatostatins, In-111 pentetretotide imaging and thyroid preparations when imaging with I-123/I-131. Know how to give rTSH and when it should be used for imaging with I-123/I-131	CbD, PGD, mini-IPX	1,2
Know when specialist imaging techniques such as scintimammography should be used (optional)	CbD, PGD, mini-IPX	1,2
Understand how the images analysed and are displayed for reading	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Be able to decide if a study is positive and be able to identify its site (possibly with the aid of SPET-SPET/CT )	CbD, PGD, mini-IPX	1,2
Know what can produce false positive results and how to communicate that possibility to the referring clinician	CbD, PGD, mini-IPX	1,2
Understand the consequences of a test being negative in a patient with known disease and how this is determined by tumour biology. Know if this affects prognosis	CbD, PGD, mini-IPX	1,2
Be able to present results at MDT	CbD, PGD, mini-IPX	1,2

<b>Behaviours</b>		
Understand the concerns of the patients (and for neuroblastoma the parents/guardians of children) undergoing radionuclide tests	CbD, MSF	1,2,3,4
Deal sensitively with patients with cancer (or their parents in children with neuroblastoma)	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly F-18 FDG PET-CT in diagnosis and staging of lung cancer</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of lung cancer	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand how F-18 FDG may be used to diagnose lung cancer in a patient with a single pulmonary nodule or stage a patient which CT suggests is operable	CbD, PGD, mini-IPX	1,2
Know the health economic arguments concerning the use of PET-CT in diagnosing and staging lung cancer	CbD, PGD, mini-IPX	1,2
Know the causes of a false negative or false positive result	CbD, PGD, mini-IPX	1,2
Be able to identify other unsuspected pathology on the PET or CTstudy	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Be able to run a glucose clamp in a diabetic patient if required	CbD, PGD, mini-IPX	1,2
Be able to decide if a study is positive for lung cancer and also to be able to determine if a patient with known lung cancer is operable	CbD, PGD, mini-IPX	1,2
Recognise issues related to mis-registration of fusion images and able to determine how the effect of this may be reduced	CbD, PGD, mini-IPX	1,2
Be able to recognise the causes of false positive uptake of F-18 FDG in the chest and if possible how to differentiate this uptake from cancer	CbD, PGD, mini-IPX	1,2
Know when additional tests may be required	CbD, PGD, mini-IPX	1,2
Be able to confidently present the results in an MDT	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients with suspected or known cancer undergoing radionuclide tests	CbD, MSF	1,2,3,4

Understand the causes of lung cancer and if appropriate impart advise to the patient on smoking cessation	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will be able to perform myocardial stressing as required for myocardial perfusion scintigraphy-complicated patient**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Be able to take an appropriate history from the patient and examine the patient as required	CbD, PGD, DOPS	1,2
Understand the methodology of adenosine/dobutamine and physical stress	CbD, PGD, DOPS	1,2
Understand the problems of stressing patients with the following <ul style="list-style-type: none"> <li>• within 6 weeks of myocardial infarction</li> <li>• patients with left bundle branch block</li> <li>• patients with untreated arrhythmias</li> <li>• patients with 1<sup>st</sup> and 2<sup>nd</sup> degree heart block</li> <li>• patients with other severe co-morbidities</li> </ul>	CbD, PGD, DOPS	1,2
Understand correct patient preparation for each type of test and clinical situation and relevant drug interactions	CbD, PGD, DOPS	1,2
Understand the contra-indications for each type of stress and alternates that can be offered in such circumstances	CbD, PGD, DOPS	1,2
Decide if it is safe to stress patient	CbD, PGD, DOPS	1,2
Understand the time parameters such as how long should stress be administered, when during that stress should the radiopharmaceutical be given, when should the stress be terminated, what time should the patient be scanned and any instructions to the patient between stressing and imaging	CbD, PGD, DOPS	1,2
<b>Skills</b>		
Recognise abnormalities on a resting and stress ecg that mean the stress can proceed safely or should not be done	CbD, PGD, DOPS	1,2
Understand what constitutes an appropriate and sufficient stress for a diagnostic test	CbD, PGD, DOPS	1,2
Recognise those symptoms and signs that should result in termination of the stress test including changes in vital signs and arrhythmias	CbD, PGD, DOPS	1,2
Be able to communicate to the patient what is happening and what to	CbD, PGD, DOPS	1,2

expect to happen and the duration of any side effects

### Behaviours

Understand the concerns of the patients undergoing radionuclide tests and stress tests	DOPS, MSF	1,2,3,4
Be willing to explain why the stress test is required for the management of their condition and obtain oral/written consent as required in particular be able to discuss the risks associated with having the test or not having the test	DOPS, MSF	1,2,3,4
Be open in discussion with the patient about measures that can be taken to reduce the risk of ischaemic heart disease including dietary advice and smoking cessation	DOPS, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	DOPS, MSF	1,2,3,4
Be willing to learn	DOPS, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	DOPS, MSF	1,2,3,4
Understand the need for self reflection of skills	DOPS, MSF	1,2,3,4

## Level 3 Competencies

The trainee will learn how to report correctly F-18 FDG PET-CT in diagnosis and staging of lymphoma		
Knowledge	Assessment Methods	GMP
Understand the pathophysiology of different forms of lymphoma including how lymphoma is staged using F-18 FDG PET	CbD, PGD, mini-IPX	1,2
Understand the mechanism of uptake of F-18 FDG and what may lead to a false negative or false positive study	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand how F-18 FDG may be used to stage and re-stage a patient with lymphoma	CbD, PGD, mini-IPX	1,2
Know the health economic arguments concerning the use of PET-CT in staging and re-staging lymphoma	CbD, PGD, mini-IPX	1,2
Be aware of the timing issues concerning when scanning should occur both during chemotherapy treatment and after completion of chemotherapy/radiotherapy	CbD, PGD, mini-IPX	1,2
Know the EORTC or similar criteria for response measurement using PET	CbD, PGD, mini-IPX	1,2
Skills		
Be able to run a glucose clamp in a diabetic patient if required.	CbD, PGD, mini-IPX	1,2
Be able to decide if a study is positive for lymphoma and be able to describe the site of abnormality using cross sectional imaging	CbD, PGD, mini-IPX	1,2
Be able to recognise common causes of physiological activity of F-18 FDG including brown fat, thymic and bone marrow uptake. Know how it is possible to reduce the effect of these on the scan	CbD, PGD, mini-IPX	1,2
Recognise issues related to mis-registration of fusion images and able to determine how the effect of this may be reduced	CbD, PGD, mini-IPX	1,2
Be aware of other pathologies that can have uptake of F-18 FDG especially in the immunocompromised	CbD, PGD, mini-IPX	1,2
Be able to identify other important unsuspected pathologies on the PET study or the CT study provided	CbD, PGD, mini-IPX	1,2
Be able to confidently present the results in an MDT	CbD, PGD, mini-IPX	1,2
Know when any additional test may be required	CbD, PGD, mini-IPX	1,2
Behaviours		
Understand the concerns of the patients with suspected or known lymphoma undergoing radionuclide tests	CbD, MSF	1,2,3,4
Keep up to date with the latest research finding and recommendations on the use of F-18 FDG PET in lymphoma	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4

Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn how to report correctly F-18 FDG PET-CT in diagnosis and staging of other cancers**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of different forms cancer and why they may or may not have uptake of F-18 FDG	CbD, PGD, mini-IPX	1,2
Understand the mechanism of uptake of F-18 FDG and what may lead to a false negative or false positive study	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand how F-18 FDG may be used to diagnose, stage and re-stage a patient with a variety of cancers	CbD, PGD, mini-IPX	1,2
Know when additional imaging such as a full leg view is required in some cancers such as melanoma	CbD, PGD, mini-IPX	1,2
Know the health economic arguments concerning the use of PET-CT in diagnosing staging and re-staging cancer	CbD, PGD, mini-IPX	1,2
Be aware of the timing issues concerning when scanning should occur both during chemotherapy treatment and after completion of chemotherapy/radiotherapy or surgery	CbD, PGD, mini-IPX	1,2
Know the EORTC or similar criteria for response measurement using PET	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Be able to run a glucose clamp in a diabetic patient if required	CbD, PGD, mini-IPX	1,2
Be able to decide if a study is positive for cancer and be able to describe the site of abnormality using cross sectional imaging	CbD, PGD, mini-IPX	1,2
Be able to recognise common causes of physiological activity of F-18 FDG including brown fat, thymic and bone marrow uptake. Know how it is possible to reduce the effect of these on the scan	CbD, PGD, mini-IPX	1,2
Know and be able to report common cause of uptake of F-18 FDG in unexpected sites including the thyroid and colon and recommend further actions	CbD, PGD, mini-IPX	1,2
Recognise issues related to mis-registration of fusion images and able to determine how the effect of this may be reduced	CbD, PGD, mini-IPX	1,2
Be aware of other pathologies that can have uptake of F-18 FDG especially in the immunocompromised	CbD, PGD, mini-IPX	1,2
Be able to identify other important unsuspected pathologies on the PET study or the CT study provided	CbD, PGD, mini-IPX	1,2
Be able to confidently present the results in an MDT	CbD, PGD, mini-IPX	1,2
Know when any additional test may be required	CbD, PGD, mini-IPX	1,2

<b>Behaviours</b>		
Understand the concerns of the patients with suspected or known cancer undergoing radionuclide tests	CbD, MSF	1,2,3,4
Keep up to date with the latest research finding and recommendations on the use of F-18 FDG PET in different cancer groups	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

<b>The trainee will learn how to report correctly cardiac PET</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand the pathophysiology of coronary artery disease	CbD, PGD, mini-IPX	1,2
Understand the mechanism of uptake of F-18 FDG and Rb-82 and what may lead to a false negative or false positive study	CbD, PGD, mini-IPX	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD, mini-IPX	1,2
Understand what is being measured in the heart by uptake of these two agents and how they can be used to establish a diagnosis in heart disease	CbD, PGD, mini-IPX	1,2
Know when additional imaging such calcium scoring and CT angiography is available during cardiac PET-CT	CbD, PGD, mini-IPX	1,2
Know the health economic arguments concerning the use of PET-CT in diagnosing staging heart disease	CbD, PGD, mini-IPX	1,2
Be aware of the protocols for stress and rest imaging	CbD, PGD, mini-IPX	1,2
<b>Skills</b>		
Be able to run a glucose clamp in a diabetic patient if required	CbD, PGD, mini-IPX	1,2
Be able to read the studies and use any quantification to improve the accuracy of reporting cardiac PET-CT	CbD, PGD, mini-IPX	1,2
Recognise issues related to mis-registration of fusion images and able to determine how the effect of this may be reduced	CbD, PGD, mini-IPX	1,2
Be able to identify other important unsuspected pathologies on the PET study or the CT study provided	CbD, PGD, mini-IPX	1,2
Know when any additional test may be required	CbD, PGD, mini-IPX	1,2
<b>Behaviours</b>		
Understand the concerns of the patients with suspected or known cancer undergoing radionuclide tests	CbD, MSF	1,2,3,4

Keep up to date with the latest research finding and recommendations on the use of PET-CT in cardiology	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

**The trainee will learn how to report correctly at least ONE example of non-oncological F-18 FDG PET-CT**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand how F-18 FDG PET-CT can be used in a variety of diseases including: <ul style="list-style-type: none"> <li>Brain metabolism studies</li> <li>Vasculitis</li> <li>Infection or inflammation</li> </ul>	CbD, PGD	1,2
Understand the mechanism of uptake of F-18 FDG and what may lead to a false negative or false positive study	CbD, PGD	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD	1,2
<b>Skills</b>		
Be able to run a glucose clamp in a diabetic patient if required	CbD, PGD	1,2
Be able to read the studies	CbD, PGD	1,2
Recognise issues related to mis-registration of fusion images and able to determine how the effect of this may be reduced	CbD, PGD	1,2
Be able to identify other important unsuspected pathologies on the PET study or the CT study provided	CbD, PGD	1,2
Know when any additional test may be required	CbD, PGD	1,2
<b>Behaviours</b>		
Keep up to date with the latest research finding and recommendations on the use of PET-CT in non-oncological disease	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4
Understand the need for self reflection of skills	CbD, MSF	1,2,3,4

Understand the need for audit and research	CbD, MSF	1,2,3,4
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**The trainee will learn how to report correctly at least ONE example of non- F-18 FDG PET-CT (this is a fast moving field and examples will be given but other examples can be used).**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Understand PET can be used in a variety of different diseases using a variety of PET pharmaceuticals to look at a different types of disease including: <ul style="list-style-type: none"> <li>• F-18 skeletal disease</li> <li>• F-18 FLT cancer cell turnover</li> <li>• F-18 choline Renal cell cancer Prostate cancer</li> <li>• F-18 DOPA pancreatic neuroendocrine tumours Parkinson's syndrome</li> <li>• F-18 FMISO Hypoxia</li> <li>• C-11 methionine Brain primary tumours</li> <li>• Ga-68 somatostatins Neuroendocrine tumours</li> </ul>	CbD, PGD	1,2
Understand the mechanism of uptake of each agent and what may lead to a false negative or false positive study	CbD, PGD	1,2
Understand the imaging protocol for each agent and for each indication	CbD, PGD	1,2
Be aware of the legal framework in place when using novel tracers	CbD, PGD	1,2
Know when such an agent must be used as part of an approved research project	CbD, PGD	1,2
Be able to take an appropriate history from the patient (or their parents) and examine the patient as required	CbD, PGD	1,2
<b>Skills</b>		
Be able to read the studies and how this information may be used	CbD, PGD	1,2
Recognise issues related to mis-registration of fusion images and able to determine how the affect of this may be reduced	CbD, PGD	1,2
Be able to identify other important unsuspected pathologies on the PET study or the CT study provided	CbD, PGD	1,2
Know when any additional test may be required	CbD, PGD	1,2
<b>Behaviours</b>		
Keep up to date with the latest research finding and recommendations on the use of PET-CT with different tracers	CbD, MSF	1,2,3,4
Be willing to transmit important urgent results about the patient to their referring clinician	CbD, MSF	1,2,3,4
Show a knowledge and willingness to comply with rules of IR(ME)R and other regulations governing use of radioisotopes	CbD, MSF	1,2,3,4
Be willing to learn	CbD, MSF	1,2,3,4
Understand their limitations and ask for assistance from medical and non-medical colleagues	CbD, MSF	1,2,3,4

Understand the need for self reflection of skills	CbD, MSF	1,2,3,4
Understand the need for audit and research	CbD, MSF	1,2,3,4

## Therapy with Radio-Isotopes

To provide the trainee with the knowledge, skills and attitudes to prescribe, administer and monitor the use of radiopharmaceuticals for therapy.

### Level 2 Competencies

The trainee will be able to deliver I-131 therapy for hyperthyroidism		
Knowledge	Assessment Methods	GMP
Pathophysiology of different causes of hyperthyroidism	PGD, mini-CEX, mini-IPX, MSF	1
Different treatment options for patients with hyperthyroidism	PGD, mini-CEX, mini-IPX, MSF	1
Appropriate selection of patients with hyperthyroidism for I-131	PGD, mini-CEX, mini-IPX, MSF	1
Understand appropriate follow-up required for patients having been treated with I-131	PGD, mini-CEX, mini-IPX, MSF	1
Reading pre-therapy radioisotope studies to determine if treatment is appropriate with I-131	PGD, mini-CEX, mini-IPX, MSF	1
Understand both the dosimetric and empirical methods used in treating hyperthyroidism with I-131	PGD, mini-CEX, mini-IPX, MSF	1
Understand the legislation concerning the safe delivery of I-131 including radiation protection for self, other staff and the patient's carers	PGD, mini-CEX, mini-IPX, MSF	1
Understand special requirements for treatment of patients under the age of 18	PGD, mini-CEX, mini-IPX, MSF	1
Skills		
Be able to take relevant history and perform relevant clinical examination within thyroid clinic	CbD, PGD, mini-CEX	2,3
Recognise those complications that would be a contra-indication to treatment with I-131	CbD, PGD, mini-CEX	2,3
Be able to explain the treatment and obtain consent for treatment with special reference to female patient's concerns about fertility and contraception	CbD, PGD, mini-CEX	2,3
Be able to advise on management of thyroid eye disease	CbD, PGD, mini-CEX	2,3
Give advice on termination and re-commencement of anti-thyroid medication	CbD, PGD, mini-CEX	2,3
Arrange appropriate follow-up and further management of the patient	CbD, PGD, mini-CEX	2,3
Behaviours		
Be responsive to the concerns of the patient and their carers concerning treatment	CbD, DOPS, mini-CEX, MSF	1,3,4
Show a professional attitude in interactions with patient and colleagues	CbD, DOPS, mini-CEX, MSF	1,3
Communicate essential information in an appropriate and timely way	CbD, DOPS, mini-CEX, MSF	1,3

Be aware of issues concerning fertility and contraception in different ethnic cultures and how that impacts on patient care	CbD, DOPS, mini-CEX, MSF	1,3
Work well with other team members, be willing to take advice from the RPS and RPA	CbD, DOPS, mini-CEX, MSF	1,3,4
Show awareness of the importance of research and audit	CbD, DOPS, mini-CEX, MSF	1,2

## Level 3 Competencies

<b>The trainee will be able to deliver I-131 therapy for treatment of patients with thyroid cancer</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Pathophysiology of thyroid cancer	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Different treatment options for patients with thyroid cancer	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Appropriate selection of patients with hyperthyroidism for I-131. Understand the need for ablation of thyroid remnant	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Understand the long term prognosis of the disease in patients treated or not with I-131	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Understand appropriate follow-up required for patients having been treated with I-131 for thyroid cancer	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Reading pre-therapy radioisotope studies to determine if treatment is appropriate with I-131 including I-123 and F-18 FDG PET imaging	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Understand both the dosimetric and empirical methods used in treating thyroid cancer with I-131	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Understand the advantages and disadvantages and methodology of use of withdrawal of thyroid hormone supplementation and/or TSH stimulation in preparation for therapy	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Understand the role of thyroglobulin in the long term follow-up of patients with thyroid cancer	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Understand the legislation concerning the safe delivery of I-131 including radiation protection for self, other staff and the patient's carers	CbD, PGD, mini-CEX, mini-IPX, MSF	1
Understand special requirements for treatment of patients under the age of 18	CbD, PGD, mini-CEX, mini-IPX, MSF	1
<b>Skills</b>		
Be able to take relevant history and perform relevant clinical examination within thyroid cancer clinic	CbD, PGD, mini-CEX	1,3
Work with the thyroid cancer MDT to determine best management of the patient	CbD, PGD, mini-CEX	1,3,4
Recognise those complications that would be a contra-indication to treatment with I-131	CbD, PGD, mini-CEX	1

Be able to prepare the patient for therapy with I-131 including use of low-iodine diets and side effects of thyroid hormone supplement withdrawal	CbD, PGD, mini-CEX	1,3,4
Be able to explain the treatment and obtain consent for treatment with special reference to female patient's concerns about fertility and contraception	CbD, PGD, mini-CEX	1,3
Give advice on termination and re-commencement of thyroxine replacement therapy	CbD, PGD, mini-CEX	1
Arrange appropriate follow-up and further management of the patient	CbD, PGD, mini-CEX	1,3
<b>Behaviours</b>		
Be responsive to the concerns of the patient and their carers concerning treatment in particular reference to concerns the patient may have about cancer	CbD, DOPS, mini-CEX, MSF	1,3
Show a professional attitude in interactions with patient and colleagues	CbD, DOPS, mini-CEX, MSF	1,3
Communicate essential information in an appropriate and timely way	CbD, DOPS, mini-CEX, MSF	1,3
Be aware of issues concerning fertility and contraception in different ethnic cultures and how that impacts on patient care	CbD, DOPS, mini-CEX, MSF	1,3
Work well with other team members, be willing to take advice from the RPS and RPA	CbD, DOPS, mini-CEX, MSF	1,3
Show awareness of the importance of research and audit	CbD, DOPS, mini-CEX, MSF	1,3,4

<b>The trainee will be able to deliver radionuclide synovectomy</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Pathophysiology of different causes of inflammatory joint disease	PGD, mini-CEX, mini-IPX, MSF	1
Different treatment options with inflammatory joint disease	PGD, mini-CEX, mini-IPX, MSF	1
Appropriate selection of patients for treatment with radionuclides	PGD, mini-CEX, mini-IPX, MSF	1
Understand appropriate follow-up required for patients having been treated with radiation synovectomy including awareness of complications including infection and radionecrosis	PGD, mini-CEX, mini-IPX, MSF	1
Know the European Association of Nuclear Medicine guidelines on appropriate radio-isotope and activity to be given depending on joint and the number of joints that can be treated at any given time	PGD, mini-CEX, mini-IPX, MSF	1
Understand the need for immobilisation of the joint for 24 hours after treatment	PGD, mini-CEX, mini-IPX, MSF	1
Understand the legislation concerning the safe delivery of Y-90, Re-186 and Eu-169 including radiation protection for self, other staff and the patient's carers	PGD, mini-CEX, mini-IPX, MSF	1
Understand special requirements for treatment of patients under the age of 18	PGD, mini-CEX, mini-IPX, MSF	1

<b>Skills</b>		
Be able to take relevant history and perform relevant clinical examination of patient with joint disease	CbD, PGD, DOPS, mini-CEX	2,3
Be able to explain procedure to patient and obtain consent	CbD, PGD, DOPS, mini-CEX	2,3
Be able to ensure correct activity of radiopharmaceutical has been drawn up	CbD, PGD, DOPS, mini-CEX	2,3
Be able to have skills to inject joints using a sterile technique or use other clinicians such as radiologists and rheumatologist to obtain access to the joint	CbD, PGD, DOPS, mini-CEX	2,3
Be able to withdraw an appropriate amount of fluid from the joint and give corticosteroids if indicated	CbD, PGD, DOPS, mini-CEX	2,3
Able to give radioisotopes without contamination of patient, self or colleagues	CbD, PGD, DOPS, mini-CEX	2,3
Give advice post therapy complication and suggest appropriate actions	CbD, PGD, DOPS, mini-CEX	2,3
Ensure patient's joint is appropriately immobilised for at least 24 hours	CbD, PGD, DOPS, mini-CEX	2,3
Arrange appropriate follow-up and further management of the patient	CbD, PGD, DOPS, mini-CEX	2,3
<b>Behaviours</b>		
Be responsive to the concerns of the patient and their carers concerning treatment	CbD, DOPS, mini-CEX, MSF	1,3
Be able to explain to patient criteria for treatment success and failure including expected time scale for response	CbD, DOPS, mini-CEX, MSF	1,3
Show a professional attitude in interactions with patient and colleagues	CbD, DOPS, mini-CEX, MSF	1,3
Communicate essential information in an appropriate and timely way	CbD, DOPS, mini-CEX, MSF	1,3
Be aware of issues concerning fertility and contraception in different ethnic cultures and how that impacts on patient care	CbD, DOPS, mini-CEX, MSF	1,3
Work well with other team members, be willing to take advice from the RPS and RPA	CbD, DOPS, mini-CEX, MSF	1,3
Show awareness of the importance of research and audit	CbD, DOPS, mini-CEX, MSF	1,3

<b>The trainee will be able to deliver P-32 in haematological malignancy</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Pathophysiology of polycythaemia and essential thrombocythaemia	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of probable success of treatment compared to alternative therapies. In addition possible side effects compared to alternate treatments and long term prognosis including risk of myelofibrosis and acute leukaemia	CbD, PGD, DOPS, mini-CEX, MSF	1
Appropriate selection of patients for P-32 treatment	CbD, PGD, DOPS,	1

Understand appropriate follow-up required for patients having been treated with P-32 working closely with haematologist	mini-CEX, MSF CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of guidelines of the European Association Nuclear Medicine guidelines for treatment with P-32	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the legislation concerning the safe delivery of P-32 including radiation protection for self, other staff and the patient's carers	CbD, PGD, DOPS, mini-CEX, MSF	1
<b>Skills</b>		
Be able to discuss appropriate use of P-32 with haematological colleagues including within an MDT	CbD, PGD, DOPS, mini-CEX	1,3
Recognise those complications that would be a contra-indication to treatment with P-32	CbD, PGD, DOPS, mini-CEX	1
Be able to explain the treatment and obtain consent for treatment with special reference to female patient's concerns about fertility and contraception	CbD, PGD, DOPS, mini-CEX	1,3
Communicate to the patient a realistic view of outcomes in this palliative treatment	CbD, PGD, DOPS, mini-CEX	1,3
Arrange appropriate follow-up and further management of the patient	CbD, PGD, DOPS, mini-CEX	1,3,4
<b>Behaviours</b>		
Be responsive to the concerns of the patient and their carers concerning treatment	CbD, DOPS, mini- CEX, MSF	1,3
Show a professional attitude in interactions with patient and colleagues	CbD, DOPS, mini- CEX, MSF	1,3
Communicate essential information in an appropriate and timely way	CbD, DOPS, mini- CEX, MSF	1,3
Be aware of issues concerning fertility and contraception in different ethnic cultures and how that impacts on patient care	CbD, DOPS, mini- CEX, MSF	1,3
Work well with other team members, be willing to take advice from the RPS and RPA	CbD, DOPS, mini- CEX, MSF	1,3
Show awareness of the importance of research and audit	CbD, DOPS, mini- CEX, MSF	1,3

<b>The trainee will be able to deliver Radiolabelled antibodies in haematological malignancy</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Pathophysiology of lymphoma, leukaemia and myeloma, and where radiolabelled antibodies can be used to treat these diseases	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of probable success of treatment compared to alternative therapies. In addition possible side effects compared to alternate treatments and long term prognosis including risk of myelofibrosis and acute leukaemia	CbD, PGD, DOPS, mini-CEX, MSF	1
Appropriate selection of patients with haematological malignancy for treatment with these agents	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of the use of immunohistochemistry in identifying patients	CbD, PGD, DOPS,	1

appropriate for treatment	mini-CEX, MSF	
Know in which clinical situations pre-scanning with a tracer dose is needed for dosimetric assessment or to determine suitability for treatment	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of the indications for use of Y-90 tiuxetan ibritumomab and other available agents	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of the dosing regimes for use of Y-90 tiuxetan ibritumomab (or other agents)	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of the need for conditioning with un-radiolabelled antibodies such as Rituximab and the required timings for these treatments	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware if the treatment will be performed in isolation or in combination with other anti-cancer drugs or bone marrow transplant	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the legislation concerning the safe delivery of Y-90 and I-131 products including radiation protection for self, other staff and the patient's carers	CbD, PGD, DOPS, mini-CEX, MSF	1
<b>Skills</b>		
Be able to discuss appropriate use of Y-90 tiuxetan ibritumomab or alternate agents with haematological colleagues including within an MDT	CbD, PGD, DOPS, mini-CEX	1,3
Recognise those complications that would be a contra-indication to treatment with these agents	CbD, PGD, DOPS, mini-CEX	1,3
Be happy to administer these drugs via a central line catheter using an aseptic technique	CbD, PGD, DOPS, mini-CEX	1,2
Be able to explain the treatment and obtain consent for treatment with special reference to female patient's concerns about fertility and contraception	CbD, PGD, DOPS, mini-CEX	1,3
Communicate to the patient a realistic view of outcomes in these treatments	CbD, PGD, DOPS, mini-CEX	1,3
Understand the experimental nature of some of these treatments	CbD, PGD, DOPS, mini-CEX	1,3
Be prepared to treat acute anaphylaxis or other less acute immune reactions	CbD, PGD, DOPS, mini-CEX	1,3
Arrange appropriate follow-up and further management of the patient	CbD, PGD, DOPS, mini-CEX	1,3
<b>Behaviours</b>		
Be responsive to the concerns of the patient and their carers concerning treatment	CbD, DOPS, mini-CEX, MSF	1,3
Show a professional attitude in interactions with patient and colleagues	CbD, DOPS, mini-CEX, MSF	1,3
Communicate essential information in an appropriate and timely way	CbD, DOPS, mini-CEX, MSF	1,3
Be aware of issues concerning fertility and contraception in different ethnic cultures and how that impacts on patient care being sensitive to those patients who may have lost fertility as part of their treatment	CbD, DOPS, mini-CEX, MSF	1,3
Work well with other team members, be willing to take advice from the RPS and RPA	CbD, DOPS, mini-CEX, MSF	1,3
Show awareness of the importance of research and audit	CbD, DOPS, mini-CEX, MSF	1,3

<b>The trainee will be able to deliver radionuclide treatment for painful bone metastases</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Pathophysiology of bone metastases and the methods used to treat bone pain	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the relevance and usefulness of diagnostic imaging with Tc-99m MDP/HDP in selecting patients for therapy	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of probable success of treatment compared to alternative therapies. In addition possible side effects compared to alternate treatments and long term prognosis including risk of bone marrow suppression	CbD, PGD, DOPS, mini-CEX, MSF	1
Appropriate selection of patients for treatment via site specific MDT	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand appropriate preparation of patient for treatment of painful bone metastases including whether or not it will be given in combination with chemotherapy drugs and/or bisphosphonates	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of recommendations for activities to be given for both the beta emitters Sr-89, Sm-153 EDTMP, Re-186/Re-188 HEDP and the alpha emitter Ra-223	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the appropriate dosing regimes including standard dose and weight related dosing including minimum time intervals for repeat treatments	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the legislation concerning the safe delivery of these products and the different requirements for radiation protection for self, other staff and the patient's carers with each agent	CbD, PGD, DOPS, mini-CEX, MSF	1
<b>Skills</b>		
Be able to discuss appropriate use agents used to treat painful bone metastases with colleagues including within an MDT	CbD, PGD, DOPS, mini-CEX	1,3
Recognise those complications that would be a contra-indication to treatment with each agent with particular reference to possible haematological toxicity	CbD, PGD, DOPS, mini-CEX	1
Understand that some contra-indications such as risk of long bone and vertebral fracture may be treated and then the patient presented for therapy	CbD, PGD, DOPS, mini-CEX	1
Be able to explain the treatment and obtain consent for treatment with special reference to female patient's concerns about fertility and contraception (where relevant)	CbD, PGD, DOPS, mini-CEX	1,3
Communicate to the patient a realistic view of outcomes in this palliative treatment	CbD, PGD, DOPS, mini-CEX	1,3
Be able to explain the possibility of a flare reaction, the best methods to treat and expected duration	CbD, PGD, DOPS, mini-CEX	1,3
Explain how success in treatment is determined including the use of pain diaries, the expected duration of treatment and the time when a repeat treatment may be given	CbD, PGD, DOPS, mini-CEX	1,3
Arrange appropriate follow-up and further management of the patient	CbD, PGD, DOPS, mini-CEX	1,3,4
<b>Behaviours</b>		

Be responsive to the concerns of the patient and their carers concerning treatment	CbD, DOPS, mini-CEX, MSF	1,3
Show a professional attitude in interactions with patient and colleagues	CbD, DOPS, mini-CEX, MSF	1,3
Communicate essential information in an appropriate and timely way	CbD, DOPS, mini-CEX, MSF	1,3
Be aware of issues concerning fertility and contraception in different ethnic cultures and how that impacts on patient care	CbD, DOPS, mini-CEX, MSF	1,3
Work well with other team members, be willing to take advice from the RPS and RPA in particular when dealing with alpha emitters	CbD, DOPS, mini-CEX, MSF	1,3
Show awareness of the importance of research and audit	CbD, DOPS, mini-CEX, MSF	1,3

<b>The trainee will be able to deliver I-131 mIBG therapy</b>		
	<b>Assessment Methods</b>	<b>GMP</b>
<b>Knowledge</b>		
Pathophysiology of those tumours including neuroblastoma, phaeochromocytoma, paraganglioma and neuroendocrine tumours in which I-131 mIBG may be useful	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the relevance and usefulness of diagnostic imaging with I-123/I-131 mIBG in selecting patients for therapy	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of probable success of treatment compared to alternative therapies. In addition possible side effects compared to alternate treatments and long term prognosis including risk of bone marrow suppression and effects on the thyroid	CbD, PGD, DOPS, mini-CEX, MSF	1
Appropriate selection of patients for treatment with I-131 mIBG	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand appropriate follow-up required for patients having been treated with I-131 mIBG with appropriate referring clinician	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of guidelines of the European Association Nuclear Medicine guidelines for treatment with I-131 mIBG	CbD, PGD, DOPS, mini-CEX, MSF	1
In particular be aware of the dosimetric and empirical approach to treatment	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the legislation concerning the safe delivery of I-131 mIBG including radiation protection for self, other staff and the patient's carers	CbD, PGD, DOPS, mini-CEX, MSF	1
<b>Skills</b>		
Be able to discuss appropriate use of I-131 mIBG with oncological colleagues including within an MDT	CbD, PGD, DOPS, mini-CEX	1
Know how patients should be prepared for therapy for example the stopping or reduction of drugs which interfere with uptake and the need to give appropriate cover with potassium iodide	CbD, PGD, DOPS, mini-CEX	1
Recognise those complications that would be a contra-indication to treatment with I-131 mIBG for example when and where cardiovascular monitoring is required	CbD, PGD, DOPS, mini-CEX	1
Be able to deal with any resultant cardiovascular side effect	CbD, PGD, DOPS, mini-CEX	1

Be able to explain the treatment and obtain consent for treatment with special reference to female patient's concerns about fertility and contraception	CbD, PGD, DOPS, mini-CEX	1,3
Communicate to the patient a realistic view of outcomes in this palliative treatment	CbD, PGD, DOPS, mini-CEX	1,3
Arrange appropriate follow-up and further management of the patient	CbD, PGD, DOPS, mini-CEX	1,3,4
Be able to deal with the special concerns in treating children including the fears and hopes of the patient's family/guardians	CbD, PGD, DOPS, mini-CEX	1,3
<b>Behaviours</b>		
Be responsive to the concerns of the patient (and parent/guardian) and their carers concerning treatment	CbD, DOPS, mini-CEX, MSF	1,3
When treating children be able to communicate in a manner appropriate for the child's age and development	CbD, DOPS, mini-CEX, MSF	1,3
Show a professional attitude in interactions with patient and colleagues	CbD, DOPS, mini-CEX, MSF	1,3
Communicate essential information in an appropriate and timely way	CbD, DOPS, mini-CEX, MSF	1,3
Be aware of issues concerning fertility and contraception in different ethnic cultures and how that impacts on patient care	CbD, DOPS, mini-CEX, MSF	1,3
Work well with other team members, be willing to take advice from the RPS and RPA	CbD, DOPS, mini-CEX, MSF	1,3
Show awareness of the importance of research and audit	CbD, DOPS, mini-CEX, MSF	1,3

<b>The trainee will be able to deliver radiolabelled somatostatin therapy</b>		
<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Pathophysiology of those tumours including phaeochromocytoma, paraganglioma and neuroendocrine tumours in which radiolabelled somatostatins may be useful	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the relevance and useful of diagnostic imaging with In-111 penteretide/Ga-68 DOTATATE/NOC/TOC PET in selecting patients for therapy and how this helps patient selection	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the relationship between the diagnostic and therapeutic peptides used	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of the legislation required to perform radiolabelled somatostatin therapy	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of the different peptides available and the characteristics of Y-90 and Lu-177 and how selections are made on the combination used for therapy	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of probable success of treatment compared to alternative therapies. In addition possible side effects compared to alternate treatments and long term prognosis including risk of bone marrow suppression and renal failure and the need for co-administration of anionic amino acids	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand appropriate follow-up required for patients having been	CbD, PGD, DOPS,	1

treated with radiolabelled somatostatins with the appropriate referring clinician	mini-CEX, MSF	
In particular be aware of published dosing regimes	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the legislation concerning the safe delivery of both Y-90 and Lu-177 labelled somatostatins including radiation protection for self, other staff and the patient's carers	CbD, PGD, DOPS, mini-CEX, MSF	1
<b>Skills</b>		
Be able to discuss appropriate use of radiolabelled somatostatins with oncological colleagues including within an MDT	CbD, PGD, DOPS, mini-CEX	1,3
Know how patients should be prepared for therapy for example the stopping or reduction of short acting or long acting somatostatins and starting amino acid infusions at least 1 hour prior to therapy and providing anti-emetics	CbD, PGD, DOPS, mini-CEX	1
Be able to explain the treatment and obtain consent for treatment with special reference to female patient's concerns about fertility and contraception. Also explain the dosing regime (normally 3-4 cycles every 6-12 weeks)	CbD, PGD, DOPS, mini-CEX	1,3
Communicate to the patient a realistic view of outcomes in this palliative treatment	CbD, PGD, DOPS, mini-CEX	1,3
Arrange appropriate follow-up and further management of the patient	CbD, PGD, DOPS, mini-CEX	1
Be able to deal with the special concerns in treating children including the fears and hopes of the patient's family/guardians	CbD, PGD, DOPS, mini-CEX	1
<b>Behaviours</b>		
Be responsive to the concerns of the patient and their carers concerning treatment	CbD, DOPS, mini-CEX, MSF	1,3
When treating children be able to communicate in a manner appropriate for the child's age and development	CbD, DOPS, mini-CEX, MSF	1,3
Show a professional attitude in interactions with patient and colleagues	CbD, DOPS, mini-CEX, MSF	1,3
Communicate essential information in an appropriate and timely way	CbD, DOPS, mini-CEX, MSF	1,3
Be aware of issues concerning fertility and contraception in different ethnic cultures and how that impacts on patient care	CbD, DOPS, mini-CEX, MSF	1,3
Work well with other team members, be willing to take advice from the RPS and RPA	CbD, DOPS, mini-CEX, MSF	1,3
Show awareness of the importance of research and audit	CbD, DOPS, mini-CEX, MSF	1,3

**The trainee will be able to deliver intra-arterial therapy of liver primary cancer/metastatic disease**

<b>Knowledge</b>	<b>Assessment Methods</b>	<b>GMP</b>
Pathophysiology of primary and secondary cancers within the liver	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the relevance and use of diagnostic imaging with CT/MRI and PET in selecting patients for therapy	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of probable success of treatment compared to alternative therapies. In addition possible side effects compared to alternate treatments and long term prognosis including risk of bone marrow suppression and effects on the thyroid	CbD, PGD, DOPS, mini-CEX, MSF	1
Appropriate selection of patients for treatment including size and site of tumour(s) and the presence or absence of portal vein thrombosis	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand appropriate follow-up required for patients having been treated with these agents with appropriate referring clinician	CbD, PGD, DOPS, mini-CEX, MSF	1
Be aware of guidelines of the European Association Nuclear Medicine guidelines for treatment with I-131 Lipiodol and Y-90 particulates	CbD, PGD, DOPS, mini-CEX, MSF	1
In particular be aware of the dosimetric and empirical approach to treatment	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand that with Y-90 labelled particulates a pre-dosing intra-arterial Tc-99m MAA scan should be performed to determine both the possibility of shunting to the lungs (must be less than 20%) and effect on the administered activity	CbD, PGD, DOPS, mini-CEX, MSF	1
Understand the legislation concerning the safe delivery of both I-131 and Y-90 labelled products including radiation protection for self, other staff and the patient's carers	CbD, PGD, DOPS, mini-CEX, MSF	1
<b>Skills</b>		
Be able to discuss appropriate use of Lipiodol and Y-90 particulates with colleagues including within an MDT	CbD, PGD, DOPS, mini-CEX	1,3
Know how patients should be prepared for therapy for example the requirements for intra-arterial cannulation including clotting screen and platelet count	CbD, PGD, DOPS, mini-CEX	1,3
Be able to give product safely within the sterile facilities of X-ray special suite	CbD, PGD, DOPS, mini-CEX	1,2
Be able to explain the treatment and obtain consent for treatment with special reference to female patient's concerns about fertility and contraception	CbD, PGD, DOPS, mini-CEX	1,3
Communicate to the patient a realistic view of outcomes in this palliative treatment	CbD, PGD, DOPS, mini-CEX	1,3
Be able to deal with the special concerns in treating children including the fears and hopes of the patient's family/guardians	CbD, PGD, DOPS, mini-CEX	1,3
<b>Behaviours</b>		
Be responsive to the concerns of the patient (and parent/guardian) and their carers concerning treatment	CbD, DOPS, mini-CEX, MSF	1,3
When treating children be able to communicate in a manner appropriate for the child's age and development	CbD, DOPS, mini-CEX, MSF	1,3

Show a professional attitude in interactions with patient and colleagues	CbD, DOPS, mini-CEX, MSF	1,3
Communicate essential information in an appropriate and timely way	CbD, DOPS, mini-CEX, MSF	1,3
Be aware of issues concerning fertility and contraception in different ethnic cultures and how that impacts on patient care	CbD, DOPS, mini-CEX, MSF	1,3
Work well with other team members, be willing to take advice from the RPS and RPA	CbD, DOPS, mini-CEX, MSF	1,3
Show awareness of the importance of research and audit	CbD, DOPS, mini-CEX, MSF	1,3