

# Magnetic Resonance Imaging

The purpose of continuing qualifications requirements (CQR) is to assist registered technologists in documenting their continued qualifications in the disciplines of certification and registration held. To accomplish this purpose the continuing qualifications requirements are presented in three parts: the professional profile, the structured self assessment (SSA) and continuing education (CE).

The purpose of the CQR SSA is to assist registered technologists identify gaps in the knowledge and cognitive skills underlying the intelligent performance of the tasks typically required for practice within the disciplines of certification and registration held and help direct their professional development efforts.

The Structured Self Assessment Content Specifications for Magnetic Resonance Imaging is provided to assist magnetic resonance imaging (MRI) technologists during their CQR compliance period. Its purpose is to prepare MRI technologists for the SSA and to help education providers develop coursework for the MRI technologists who need to address specified areas with targeted continuing education. Targeted CE is assigned only if a standard is not met in a category on the SSA.

The SSA is composed of sets of questions that are designed to evaluate an individual's knowledge in topics related to current practice. Participants have a maximum of 80 minutes to complete the SSA. Please allow an additional eight minutes for the tutorial, two minutes for the non-disclosure agreement (NDA), and 10 minutes for a follow-up survey.

The table below presents the major categories and subcategories covered on the SSA. The number of questions in each category are listed in bold and number of questions in each subcategory in parentheses. The potential number of targeted CE credits that would be prescribed if the standard is not met, are across from each subcategory, with the maximum amount listed at the bottom. Specific topics within each category are addressed in the content outline, which makes up the remaining pages of this document.

Content Category	<b>Number of Questions</b>	Potential CE Credits
Patient Care	10	
Patient Interactions and Management (10)		3
Safety	10	
MRI Screening and Safety (10)		4
Image Production	30	
Physical Principles of Image Formation (10)		7
Sequence Parameters and Options (10)		6
Data Acquisition, Processing, and Storage (10)		5
Procedures	30	
Neurological (10)		5
Body (10)		3
Musculoskeletal (10)		3
	Total 80	Maximum CE 36



## **Patient Care**

## 1. Patient Interactions and Management

- A. Ethical and Legal Aspects
  - 1. patients' rights
    - a. informed consent (\*e.g., written, oral, implied)
    - b. confidentiality (HIPAA)
    - c. American Hospital Association (AHA) Patient Care Partnership (Patients' Bill of Rights)
      - 1. privacy
      - 2. extent of care (e.g., DNR)
      - 3. access to information
      - 4. living will, health care proxy, advance directive
      - 5. research participation
  - 2. legal issues
    - a. verification (e.g., patient identification, compare order to clinical indication)
    - b. common terminology (e.g., battery, negligence, malpractice, beneficence)
    - c. legal doctrines (e.g., respondeat superior, res ipsa loquitur)
    - d. restraints versus immobilization
  - 3. ARRT Standards of Ethics
- B. Interpersonal Communication
  - 1. modes of communication
    - a. verbal/written
    - b. nonverbal (e.g., eye contact, touching)
  - 2. challenges in communication
    - a. interactions with others
      - 1. language barriers
      - 2. cultural and social factors
      - 3. physical or sensory impairments
      - 4. age
      - 5. emotional status, acceptance of condition
    - b. explanation of medical terms
    - c. strategies to improve understanding

- 3. patient education
  - a. explanation of current procedure (e.g., purpose, exam length)
  - b. pre- and post-procedure instructions (e.g., preparations, diet, medications, discharge instructions)
  - c. respond to inquiries about other imaging modalities
  - d. communication with patient during procedure
- C. Physical Assistance and Monitoring
  - 1. patient transfer and movement
    - a. body mechanics (e.g., balance, alignment, movement)
    - b. patient transfer techniques
  - 2. assisting patients with medical equipment
    - a. infusion catheters and pumps
    - b. oxygen delivery systems
    - c. other (e.g., nasogastric tubes, urinary catheters, tracheostomy tubes)
  - 3. routine monitoring
    - a. vital signs
    - b. physical signs and symptoms
    - c. fall prevention
    - d. documentation
    - e. sedated patients
    - f. claustrophobic patients
- D. Medical Emergencies
  - allergic reactions (e.g., contrast media, latex)
  - 2. cardiac/respiratory arrest (e.g., CPR)
  - 3. physical injury, trauma, or RF burn
  - 4. other medical disorders (e.g., seizures, diabetic reactions)
- \* The abbreviation "e.g.," is used to indicate that examples are listed in parenthesis, but that it is not a complete list of all possibilities.

(Patient Care continues on the following page.)



## **Patient Care (continued)**

- E. Infection Control
  - 1. chain of infection (cycle of infection)
    - a. pathogen
    - b. reservoir
    - c. portal of exit
    - d. mode of transmission
      - 1. direct
        - a. droplet
        - b. direct contact
      - 2. indirect
        - a. airborne
        - b. vehicle-borne (fomite)
        - c. vector-borne (mechanical or biological)
    - e. portal of entry
    - f. susceptible host
  - 2. asepsis
    - a. equipment disinfection
    - b. equipment sterilization
    - c. medical aseptic technique
    - d. sterile technique
  - 3. CDC Standard Precautions
    - a. hand hygiene
    - b. use of personal protective equipment (e.g., gloves, gowns, masks)
    - c. safe handling of contaminated equipment/surfaces
    - d. disposal of contaminated materials
      - 1. linens
      - 2. needles
      - 3. patient supplies
      - 4. blood and body fluids
    - e. safe injection practices
  - 4. transmission-based precautions
    - a. contact
    - b. droplet
    - c. airborne
  - 5. additional precautions
    - a. neutropenic precautions (reverse isolation)
    - b. healthcare associated (nosocomial) infections

- F. Handling and Disposal of Toxic or Hazardous Material
  - 1. types of materials
    - a. chemicals
    - b. chemotherapy
  - 2. safety data sheet (e.g., material safety data sheets)
- G. Pharmacology
  - 1. patient history
    - a. medication reconciliation (current medications)
    - b. premedications
    - c. contraindications
    - d. scheduling and sequencing examinations
  - 2. administration
    - a. routes (e.g., IV, oral)
    - b. supplies (e.g., needles)
    - c. procedural technique (e.g., venipuncture)
    - d. dose calculation
    - e. power injector
      - 1. fluoro-triggering
      - 2. timing bolus
      - 3. automatic bolus tracking
  - 3. contrast media types and properties (e.g., gadolinium, linear versus macrocyclic, ionic versus non-ionic)
  - 4. appropriateness of contrast media to examination
    - a. patient condition
    - b. patient age and weight
    - c. laboratory values (e.g., BUN, creatinine, eGFR)
  - 5. complications/reactions
    - a. local effects
      - (e.g., extravasation/infiltration, phlebitis)
    - b. systemic effects
      - 1. mild
      - 2. moderate
      - 3. severe
    - c. emergency medications
    - d. technologist's response and documentation



## **Safety**

## 1. MRI Screening and Safety

- A. Screening and Education (patients, personnel, non-personnel)
  - 1. biomedical implants
    - a. identify and document device, year, make, model
    - research and verify device labeling (MR safe, MR conditional, MR unsafe)
    - c. identify device specific parameters
    - d. scanning conditional implants
  - 2. ferrous foreign bodies
  - 3. medical conditions (e.g., pregnancy)
  - 4. prior diagnostic or surgical procedures
  - topical or externally applied items (e.g., tattoos, medication patches, body piercing jewelry, monitoring devices, clothing)
  - 6. level 1 and level 2 MR personnel
- B. Electromagnetic Fields
  - 1. static field
    - a. translational and rotational forces
    - b. magnetohydrodynamic effect
    - c. magnetohemodynamic effect
    - d. magnetic shielding
    - e. spatial gradient of the static magnetic field
    - f. FDA guidelines
  - 2. radiofrequency (RF) field
    - a. thermal heating (specific absorption rate [SAR])
    - b. conductive loops
    - c. proximity burns
    - d. RF shielding
    - e. FDA guidelines
  - 3. gradient field
    - a. current induction
    - b. acoustic noise
    - c. peripheral neurostimulation
    - d. magnetophosphenes
    - e. FDA guidelines

### C. Equipment

- placement of conductors
   (e.g., ECG leads, coils, cables)
- 2. cryogen safety
- 3. ancillary equipment (MR safe, MR conditional, MR unsafe)

#### D. Environment

- climate control (temperature, humidity)
- 2. designated MR safety zones
- 3. gauss lines
- 4. emergency procedures (e.g., quench, fire)



## **Image Production**

## 1. Physical Principles of Image Formation

- A. Instrumentation
  - electromagnetism

     (e.g., Faraday's law)
  - 2. static magnet
    - a. types (superconductive, resistive, permanent)
    - b. magnetic field strength
    - c. shim coils
  - 3. RF system
    - a. coil configuration
    - b. surface coils
    - c. phased array coils
    - d. transmit and receive coils
    - e. transmit and receive bandwidth
    - f. pulse profile
  - 4. gradient system
    - a. gradient coil configuration
    - b. slew rate
    - c. rise time
    - d. duty cycle
- B. Fundamentals
  - 1. nuclear magnetism
    - a. Larmor equation
    - b. precession
    - c. gyromagnetic ratio
    - d. resonance
    - e. RF pulse
    - f. equilibrium magnetization
    - g. energy state transitions
    - h. phase coherence
    - i. free induction decay (FID)
    - j. magnetic susceptibility (e.g., diamagnetism, paramagnetism, superparamagnetism, ferromagnetism)
  - 2. tissue characteristics
    - a. T1 recovery
    - b. T2 decay (relaxation)
    - c. T2\* (susceptibility)
    - d. proton (spin) density (PD)
    - e. flow
    - f. diffusion
    - g. perfusion

- 3. spatial localization
  - a. vectors
  - b. X, Y, Z coordinate system
  - c. physical gradient
  - d. slice select gradient
  - e. phase-encoding gradient
  - f. frequency (readout) gradient
  - g. sampling frequency/rate
  - h. k-space (raw data)
- C. Artifacts (Cause, Appearance, and Compensation)\*\*
  - 1. aliasing
  - 2. Gibbs, truncation
  - 3. chemical shift
  - 4. chemical misregistration
  - 5. magnetic susceptibility
  - 6. radiofrequency (e.g., zipper)
  - 7. motion and flow (e.g., patient motion, ghosting)
  - 8. partial volume averaging
  - 9. crosstalk
  - 10. cross excitation
  - 11. moiré pattern
  - 12. parallel imaging artifacts
  - 13. eddy currents
  - 14. dielectric effect
- D. Quality Control
  - 1. slice thickness
  - 2. spatial resolution
  - 3. contrast resolution
  - 4. signal to noise
  - 5. center frequency
  - 6. transmit gain
  - 7. geometric accuracy
  - 8. equipment inspection (e.g., coils, cables, door seals)

(Image Production continues on the following page.)

<sup>\*\*</sup> The subsequent list of artifacts is not a complete list of all possibilities.



## **Image Production (continued)**

## 2. Sequence Parameters and Options

- A. Imaging Parameters
  - 1. repetition time (TR)
  - 2. echo time (TE)
  - 3. inversion time (TI)
  - 4. number of signal averages (NSA, NEX)
  - 5. flip angle (e.g., Ernst angle)
  - 6. field of view (FOV)
  - 7. matrix
  - 8. pixel
  - 9. voxel
  - 10. number of slices
  - 11. slice thickness and gap
  - 12. phase and frequency
  - 13. echo train length (ETL)
  - 14. effective TE
  - 15. bandwidth (transmit, receive)
  - concatenations (number of acquisitions per TR)
  - 17. b-value
  - 18. velocity encoding (VENC)
- B. Image Contrast
  - 1. T1 weighted
  - 2. T2 weighted
  - 3. PD weighted
  - 4. T2\* weighted
  - 5. diffusion weighted imaging (DWI)
  - 6. susceptibility weighted imaging (SWI)
- C. Imaging Options
  - 1. 2D/3D
  - 2. slice order

(sequential, interleaving)

- 3. spatial saturation pulse/band
- 4. gradient moment nulling
- 5. suppression techniques (e.g., fat, water, Dixon method)
- 6. physiologic gating and triggering
- 7. in-phase/out-of-phase
- 8. rectangular FOV
- 9. anti-aliasing
- 10. parallel imaging
- 11. filtering

#### **FOCUS OF QUESTIONS:**

Questions will address the interdependence of the imaging parameters, weightings, and options listed on the left, and how they affect image quality.

## Image Quality

- contrast to noise ratio (CNR, C/N)
- signal to noise ratio (SNR, S/N)
- · spatial resolution
- · acquisition time

(Image Production continues on the following page.)



## **Image Production (continued)**

## 3. Data Acquisition, Processing, and Storage

- A. Pulse Sequences
  - 1. spin echo (SE)
    - a. conventional spin echo
    - b. fast spin echo (FSE)
  - 2. inversion recovery (IR) (e.g., STIR, FLAIR)
  - 3. gradient echo (GRE)
    - a. conventional gradient echo
    - b. spoiled gradient echo
    - c. coherent gradient echo
    - d. steady state free precession (SSFP)
    - e. fast gradient echo
    - f. MRA/MRV
      - 1. flow dynamics
      - 2. time-of-flight (TOF)
      - 3. phase contrast
      - 4. contrast enhanced
  - 4. echo planar imaging (EPI)
  - 5. diffusion weighted imaging (DWI)
  - 6. susceptibility weighted imaging (SWI)
  - 7. perfusion
  - 8. spectroscopy

## B. Data Manipulation

- k-space mapping and filling (e.g., centric, spiral, keyhole)
- 2. fast Fourier transformation (FFT)
- 3. post-processing
  - a. maximum intensity projection (MIP) reformation
  - b. multiplanar reformation (MPR)
  - c. subtraction
  - d. apparent diffusion coefficient (ADC) mapping
  - e. CINE

#### C. Informatics

- hard/electronic copy (e.g., DICOM file format)
- 2. archive
- PACS and electronic medical record (EMR)
- 4. security and confidentiality
- 5. networking



## **Procedures**

## 1. Neurological

- A. Head and Neck
  - 1. brain
  - 2. brain for MS
  - 3. brain for seizure
  - 4. infant brain (less than one year old)
  - 5. brain for CSF Flow
  - 6. IACs
  - 7. pituitary
  - 8. orbits
  - 9. cranial nerves (non IACs)
  - 10. vascular head (MRA)
  - 11. vascular head (MRV)
  - 12. brain perfusion
  - 13. brain spectroscopy
  - 14. sinuses
  - 15. soft tissue neck (e.g., parotids, thyroid)
  - 16. vascular neck

### B. Spine

- 1. cervical
- 2. thoracic
- 3. lumbar
- 4. sacrum-coccyx
- 5. sacroiliac (SI) joints
- 6. whole spine
- 7. brachial plexus
- 8. lumbar plexus

#### **FOCUS OF QUESTIONS**

Questions about each of the studies listed on the left may focus on any of the following factors:

## Anatomy and Physiology

- · imaging planes
- · pathological considerations
- protocol considerations
- patient considerations (e.g., pediatric, geriatric, bariatric, trauma)

### Patient Set-Up

- · patient data input
- · coil selection and position
- · patient orientation
- landmarking
- · physiologic gating and triggering

#### Contrast Media

· effect on images

## **Additional Procedures**

- CINE (e.g., CSF flow study, TMJs)
- · surgical planning

(Procedures continue on the following page.)



## **Procedures (continued)**

## 2. Body

- A. Thorax
  - 1. chest (non cardiac)
  - 2. breast
  - 3. vascular thorax

#### B. Abdomen

- 1. liver
- 2. pancreas
- 3. MRCP
- 4. adrenals
- 5. kidneys
- 6. enterography
- 7. vascular abdomen

#### C. Pelvis

- 1. soft tissue pelvis (e.g., bladder, rectum)
- 2. female soft tissue pelvis (e.g., uterus)
- 3. male soft tissue pelvis (e.g., prostate)
- 4. vascular pelvis (e.g., femoral, iliac)

## 3. Musculoskeletal

- A. Temporomandibular Joints (TMJs)
- B. Sternum
- C. Sternoclavicular (SC) Joints
- D. Shoulder
- E. Long Bones (upper extremity)
- F. Elbow
- G. Wrist
- H. Hand
- I. Fingers (non thumb)
- J. Thumb
- K. Bony Pelvis
- L. Hip
- M. Long Bones (lower extremity)
- N. Knee
- O. Ankle
- P. Foot
- Q. Arthrogram
- R. Vascular Extremities

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