Incorporation of Imaging-Based Functional Assessment Procedures into the DICOM Standard
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I. Purpose

Drawing from the profile development of the QIBA-fMRI Technical Committee, the purpose of the QIBA-fMRI DICOM Subcommittee is to develop DICOM extensions supporting imaging-based functional assessment, specifically functional MRI (fMRI).

This working paper is intended to capture the concepts surrounding task-based fMRI, generalizing from the shared clinical and research experience of the QIBA-fMRI. Secondarily it should support other functional imaging studies (e.g. connectivity analysis) and other modalities (e.g. MEG). The level of detail should be sufficient to permit creation of DICOM object and relationship definitions as well as procedure steps describing the workflow, inputs, and outputs of functional assessment with imaging.

II. Background

As previously described [1], the fMRI workflow consist of multiple steps surrounding the actual imaging component, involving roles of patient, trainer, tester, processor and clinical user. Ultimately, we would like to capture both the steps of the workflow and the data items themselves in the DICOM framework.

1. Imaging-based functional assessment refers to the measurement of cortical activation resulting from a) extrinsic, functional tasks or b) intrinsic functional connectivity.

2. Although fMRI based on blood oxygen level dependence (BOLD) is the focus of the QIBA-fMRI committee, assessment of cortical activation may be performed with other imaging sequences and modalities (e.g., MEG). Therefore, in striving to define the most generally useful DICOM extensions, we will strive to avoid terminology specific to a given imaging method.

3. Some of the familiar terms of fMRI will appear to be missing, e.g., ‘block paradigm,’ ‘event-related.’ The proposed presentation model scheme is intended to support fixed or randomized timing and stimulus selection, hence a superset of current methods. Phases can represent classic stimulus/control or periods in which stimulus events of a given class are scheduled to happen.

DICOM coding is not a ‘presentation language’ but a means of promoting interoperability among imaging, processing and PACS systems. Expression of

functional assessment explicitly in DICOM terms is intended to validate the proposal, not suggest that paradigms will be ‘written’ in DICOM.

III. Framework

The following is an outline of the major proposed records and data objects. These would translate to DICOM entities. See Dictionary below for more details. This framework will be supplemented by entity relationship graphs in the style of the DICOM standard.

The framework is divided into three high-level records: Specification for a paradigm, Execution of the paradigm Specification, and Analysis of a paradigm Execution.

Most stimulus ‘files’ (e.g. JPG) may be mapped to existing DICOM objects; alternatively, file system paths to external files can be employed. At this time many data formats are left unspecified. In some cases they may map to existing DICOM records and tags created for other purposes. Before getting too far into those details, would be best to

1. Paradigm Specification
   a. Identification
      i. Title: Text description
      ii. Class (one of): Motor, Hearing, Vision, Language, Cognitive, Memory, etc.
      iii. Difficulty (one of): Nominal, Fast/Hard, Slow/Easy, etc.
      iv. Natural Language: English, etc.
      v. Author
      vi. Creation date
      vii. Revision
   b. Imaging Model
      i. Modality & scan type
      ii. Scan length
      iii. Scan parameters
   c. Statistical Model
      i. Relates epoch phases to expected cortical activation time course
   d. Stimulus Set(s), each a set of Stimulus Objects, as follows:
      i. Stimulus file UID
      ii. Type (one of): Image (JPG, PNG, etc.), Movie (MP4, etc.), Sound (WAV), text (TXT), etc. Note that this may be defined by the UID above.
iii. Inherent length (msec); either presentation time (stimulus file length if applicable) or zero to represent indefinite, continuous performance

iv. Response(s) expected (multiple allowed):
   1. Response window [msec post-start, msec post-end]
   2. Response period length, msec
   3. Response Type (one of): key-press, eye tracking, physiological change, etc.
   4. Expected Response Value

e. Presentation model
   i. Instructions to Tester
   ii. Instructions to Subject
   iii. Timer definition(s)
   iv. Selector definition(s)
   v. Timeline, consisting of multiple Epochs, each defined as:
      1. Phase (one of): Stimulus/Control, A/B/C…
      2. Epoch length, msec
      3. Stimulus Pattern(s), one or more, each containing:
         a. List of one or more fixed-timing Stimulus Object(s)

or

b. Variable presentation: a Stimulus Set, chosen from using Selector, with timing determined by a Timer

2. Paradigm Execution
   a. Patient
   b. Ordering clinician
   c. Performing clinician (radiologist, neuropsychologist, etc.)
   d. Training/Testing Staff (technologist or clinician)
   e. Paradigm Specification instance UID
   f. Use (Training, Test, Re-test)
   g. Paradigm Execution for Training, Instance UID
      (if Test or Re-test, the record of the corresponding Training)
h. Patient Record Attachments
   (other test results, e.g. handedness survey, neuro evaluation, etc.)

i. System QA (equipment checklist, scanner QA, etc.)

j. Staff comments & instructions

k. Assessment of Patient performance by Staff

l. Self-assessment of performance by Patient

m. Assessment of paradigm execution
   i. Probably embeds limitations of the methodology (e.g. BOLD signal
      response versus MEG) and the physical implementation (e.g. visual
      frame rate, audio frequency range, etc.)

n. Epochs performed, series of
   i. Timestamp
   ii. Phase

o. Stimuli presented, series of
   i. Timestamp
   ii. Stimulus Object UID
   iii. Stimulus presentation length, msec

p. Responses received, series of
   i. Timestamp
   ii. Type
   iii. Value

q. Performance Metric (multiple allowed)
   i. Title
   ii. Type (e.g. attention probe, response accuracy, post-test memory, etc.)
   iii. Number of trials
   iv. Number of correct trials
   v. Accuracy

3. Paradigm Analysis
   a. Paradigm Execution for testing, Instance UID
   b. Paradigm Execution for training, Instance UID
      (might be optional)
c. Processing Staff (technologist or clinician)

d. Epoch Evaluation (time-series editing)
   i. Epoch timestamp
   ii. Phase
   iii. Disposition (one of)
      1. Analyzed
      2. Rejected (reason)

e. Imaging distortion correction

f. Motion correction
   i. Algorithm
   ii. Results
      E.g. statistics: time course of deviation removed in multiple translations & rotations

g. Statistical model applied
   i. Ideal time course, this test instance
      This may include reference waveforms

h. Activation time course
   i. Sampling volume method (e.g. strongest cluster, atlas segmentation, hand-drawn VOI)
   ii. Sampling volume description (3D mask)
   iii. Activation curve

i. Map (multiple); each is an image series
   i. Type (one of)
      1. Functional activation, statistical parameter (e.g. t, r, F)
      2. Functional activation, AMPL
      3. Cardiovascular reactivity
      4. Functional connectivity (a/k/a resting state)
      5. other.
   ii. Parametric Threshold
   iii. Spatial Filtering applied
   iv. Clustering applied
v. Color palette (applied or suggested)
vi. Other features
j. Contrast-Noise map
k. Sample image volume from pre-processed time series
l. Performance Metric (multiple allowed)
i. Title
ii. Type (e.g. attention probe, response accuracy, post-test memory, etc.)
iii. Number of trials
iv. Number of correct trials
v. Accuracy
m. Other analyses
i. Laterality

IV. Dictionary

This informal description would be supplemented by a dictionary of DICOM tags and records. Presently this is offered in order of appearance in the above framework.

Elements not listed here such as Patient, Ordering Clinician, Timestamp, image data, etc. are assumed to align with DICOM objects already available in the specification.

1. Paradigm: An assessment task, in which stimuli and tasks are related to cortical activation.

2. Paradigm Specification: A model for a functional paradigm, composed of a statistical model relating the paradigm task to cortical activation, stimuli employed, and a presentation model scheduling stimuli and expected responses through time.

3. Paradigm Execution: A record of the execution of a paradigm, including a timeline of the actual stimuli presented, responses elicited, and other observations about the run. This may be captured for training and testing (scanning).

4. Paradigm Analysis: A record of the analysis of Paradigm Execution results, including processing steps and results (e.g. motion correction), QA measures (e.g. epoch editing), activation maps, select activation time course(s), etc.

5. Stimulus: Digital representation of audio, visual, tactile etc. information delivered to the subject during the course of a paradigm.
6. **Stimulus Object**: A particular stimulus (either in DICOM format or as a file system path to an external file), along with some properties. Properties include type (image (JPG, PNG, etc.), audio (WAV), tone, movie (MPG), text, etc.); presentation length (inherent length of audio and movie files, or the specified time for text, images, tones, etc.); and expected response(s) and window for response.

7. **Stimulus Set**: An ordered collection of stimulus objects of the same type sharing one or more characteristics (e.g., a set of ‘Famous Faces’).

8. **Timer**: Specification for determining stimulus presentation timing. Timers can be reused multiple times in a paradigm execution, either restarting to reuse the same timing, or continuing a timing sequence.
   a. **Mode**:
      FIXED: specified msec of presentation.
      RANDOM: [min max] msec, seed:
      A pseudo-random uniform distribution (random characteristics, but guaranteed to produce the same sequence from a given seed). Timers are abstracted from the presentation model itself so they may be used multiple times in a paradigm execution (e.g., to match the timing of faces in a stimulus epoch to the matched non-faces in a control epoch).

9. **Selector**: Specification for choosing a sequence of selection indices from a range (1…n) corresponding to entries in a stimulus set. The selector is used to choose a particular subset of stimuli from a Stimulus Set, and defines their order of presentation. Selectors can be reused multiple times in a paradigm execution, either restarting to reuse the same selection, or continuing a selection.
   a. **Mode**:
      LINEAR: start, end, increment (+/-)
      RANDOM-REPLACEMENT: [min max] indices, seed, or
      RANDOM-NO-REPLACEMENT: [min max] indices, seed:
      A pseudo-random uniform distribution (random characteristics, but guaranteed to produce the same sequence of indices from a given seed). Selectors are abstracted from the presentation model itself so the may be used multiple times in a paradigm (e.g., to select stimuli paired in multiple classes, such as ‘Face normal versus ‘Face scrambled’). Selection without replacement prevents duplicate use of a given stimulus until the Selector is reset.

10. **Timeline**: The part of the presentation model defining the series of epochs making up the assessment. It is assumed that epochs are of predefined (though not necessarily equal) length, since they usually must be tied to the scan sequence of the imager.
11. **Epoch**: A period of time in the presentation corresponding to a particular phase of activation (e.g. movement versus no movement, or visual language versus aural language). During the epoch, one or more stimulus patterns are followed to elicit the desired activation. Multiple patterns might be executed in parallel during the epoch (e.g. supporting multi-media stimulation).

12. **Stimulus pattern**: A script for stimulus presentation composed of a Timer defining the start of each stimulus event, a Selector choosing the Stimulus Object to present in the event, and a Stimulus Set from which the Stimulus Object is selected.

13. **Event**: Logging of something that happened during the actual execution of a paradigm, consisting of a timestamp, event type, and the event details.

14. **Epoch Event**: Start of an epoch, logged in the Execution of the paradigm.

15. **Stimulus Event**: Event corresponding to presentation of a stimulus, including a timestamp, the UID of the stimulus object, and the actual presentation length.

16. **Response Event**: Event corresponding to a patient response during stimulus presentation, including a timestamp, response type, and response value. If the paradigm execution system supports it, the stimulus object whose window the response falls in may be recorded as well.

17. **Performance Metric**: Real-time measurement of patient performance during paradigm execution. Some presentation systems may analyze patient performance as the paradigm is executed (e.g. real-time accuracy in answering questions), but since all results are captured this could also be analyzed retrospectively. Hence it appears under both the Execution and Analysis records.

V. **Implementation issues**

There are many details concerning paradigm execution and analysis that generally elude the standards writers. In general, validation of paradigm design will be beyond the scope of DICOM. However, this is a list for discussion.

1. **Stimulus specification**: This is a slightly unusual DICOM ‘instance’ in that it exists prior to association with a particular patient. This may be merely a terminology issue; Color Palettes are presumably in a similar state, existing independently of a particular image instance.

2. **Epoch length versus stimulus presentation**: Utilizing pseudo-random timing introduces implementation issues beyond the DICOM standard. E.g., it may be necessary to define that if a stimulus will extend beyond the end of an epoch, it is not started, versus being cut off.
3. **Presentation length:** If the inherent length of a stimulus (e.g. WAV of a story, movie) is longer than the presentation time defined by a timer, it is presumably cut off. If the inherent length of the stimulus is shorter, should the stimulus end, or repeat?

4. **Presentation scaling:** The image and video representations may or may not have inherent physical scaling. Some paradigms may require particular scaling (e.g. text readability, stimulation of a particular visual field angle). This needs to be expressed somehow and then translated in the presentation system at the imaging system. Similarly for audio volume.

5. **Presentation precision:** temporal, spatial (image), and audio (frequency) precision are all subject to the limitations of the presentation hardware. Expectations regarding the presentation system capabilities should probably be part of the statistical model.

6. **Presentation method:** Some paradigm schemes which can be expressed within the standard would be beyond the capabilities of current presentation software (e.g. refresh rate and resolution of displays).

7. **Imaging method:** Different methods of imaging for functional assessment will have different inherent limitations (e.g. temporal resolution of BOLD versus MEG) which will affect paradigm execution.

8. **Oddball experiments:** These could be implemented by defining a Stimulus Set per epoch, with the oddball(s) included in the set. If the set size and timing are defined to ensure the entire set will be used in an epoch, then RANDOM-NO-REPLACEMENT should guarantee that the oddball will appear once at a random time in the epoch. The disadvantage of this approach is that a stimulus set must be defined for each unique epoch (trial). This could be circumvented with another layer of flexibility in the paradigm specification, e.g. a probabilistic selector between stimulus sets, but further discussion would be advisable first.

VI. **Real-World Examples**

To be supplied...

*Prepared for the QIBA-fMRI DICOM Subcommittee*

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