Clinical practice guideline for body composition assessment based on upper abdominal magnetic resonance images annotated using artificial intelligence

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Introduction

Upper abdominal magnetic resonance (MR) imaging is appropriate for body composition analysis.1 Especially for individuals with obesity, it is of great value to quantify the hepatic proton density fat fraction (PDFF) and the amount of abdominal adipose tissue during clinical evaluation and for research on obesity-related risks. Analytical results may be used to determine the optimal choice of surgical procedure and evaluate treatment outcomes. Multiple artificial intelligence (AI) algorithms and systems have been developed for the automated measurement of body composition. The basis of AI development and application is to have uniform standards for clinical data acquisition and management. The uneven quality of MR images is one of the major obstacles to AI system development and analytical results. A standardized process of MR scanning and clinical data management is urgently needed.

Purpose and Target Audience

This guideline aims to standardize data acquisition, utilization, and storage for AI systems that target the automatic quantification of body composition. This guide is recommended for surgeons, clinical researchers, and radiologists who focus on body composition analysis and obesity-related topics, for example, type 2 diabetes, metabolic syndrome, and bariatric surgery.

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**Data Acquisition and Evaluation**

The standardized data acquisition process guarantees high-quality MR images for AI analysis. An image used for AI labeling and clinical diagnosis must follow the basic process mentioned in the following sections.

**Subject preparation**

Patients should have an empty stomach before MR scanning. Metal articles should be removed. For overweight subjects with a large waist circumference, wide-bore MR equipment is preferred. According to clinical practice experience, if the patient is >125 kg, doctors or technicians should carefully evaluate the feasibility of upper abdominal MR examination. One challenge is that the bore may not be sufficiently large to accommodate the subject’s abdomen. Furthermore, it may be difficult for the patient to hold their breath during MR examination, leading to significant motion artifacts.

**MR parameter setting**

A 3.0-T or 1.5-T MR device is preferred for data acquisition. The standard parameters for MR examination for AI analysis are listed in Supplementary Table 1, http://links.lww.com/CM9/A951. PDFF is a reliable measure that can be used to accurately evaluate hepatic steatosis. The Dixon image is 3-dimensional with high resolution. The fat image at the axial level of lumbar 1 to lumbar 2 (L1–L2) intervertebral disc on the Dixon image is considered the best choice for quantifying adipose tissue. If a 1.5-T MR device cannot perform 3-dimensional Dixon imaging, dual-echo scanning is also available.

**Availability of the MR image for AI quantification**

All acquired images should be saved in the Digital Imaging and Communications in Medicine (DICOM) format, as the slice thickness and other important information can be stored. The image quality required by the AI analysis is similar to that required for clinical diagnosis. Overall, images with significant artifacts are judged not acceptable. The other requirements for different applications are listed in the following sections.

The image quality required for PDFF quantification is usually high. For AI analysis of hepatic PDFF, the first step is to recognize the margin of the liver parenchyma. For patients with hepatic steatosis over grade 2 (PDFF >17.4%), the signal intensity of the liver parenchyma is significantly higher than that of the vessels and adjacent organs. As a result, the margin of the liver is relatively easy for the AI system to recognize. However, for patients with hepatic steatosis grade 1 (6.4% < PDFF ≤ 17.4%), the margin of the liver parenchyma is difficult to recognize. Therefore, AI annotation may not be precise. For subjects without hepatic steatosis (PDFF ≤ 6.4%), AI annotation may fail if only based on PDFF images. The principles and examples of different degrees of hepatic steatosis are listed in Supplementary Table 2, http://links.lww.com/CM9/A951.

According to the recently developed appropriateness criteria, a Dixon sequence image of fat at the axial level of the L1–L2 intervertebral disc can be used for abdominal adipose tissue quantification. It is essential to cover the skin of the abdomen; otherwise, subcutaneous adipose tissue (SAT) cannot be measured. Owing to an insufficient field of view, parallel acquisition, or an uneven magnetic field, the quality of the image may vary. The principles and examples of different image quality degrees are shown in Supplementary Table 3, http://links.lww.com/CM9/A951.

**Comprehensive evaluation of MR image quality**

Quality evaluation of both the PDFF and fat images should be considered. These principles are listed in Supplementary Table 4, http://links.lww.com/CM9/A951.

**Annotation standards**

**Image for PDFF quantification**

The whole liver parenchyma should be included. Large vessels, local lesions, regions beyond the margin of the liver, and imaging artifacts should be avoided (Figure 1A).

The average PDFF value can be calculated by averaging the values of all voxels included in the region of interest. Since the PDFF value and its change after bariatric surgery vary in different parts of the liver, AI systems are being developed to record the values of different liver segments (Figure 1B).

**Image for abdominal adipose tissue quantification**

Visceral adipose tissue (VAT) and SAT can be recognized and labeled. Different regions of interest, for example, muscle, can also be defined for analysis (Figure 1C and 1D).
The images can be labeled automatically using AI or manually using ITK-SNAP 3.8.0 software (http://www.itksnap.org/). Since a single slice of an MR image is volumetric, the value acquired after labeling is influenced by slice thickness. The volume is calculated as the Supplementary formulas, http://links.lww.com/CM9/A951.

Database management
For clinical follow-up and research, it is recommended to set up a database to manage clinical data and MR images.

Data registry
The clinical and radiological data should be registered in a standardized database. For example, there is a prospective national registry database named the “Greater China Metabolic and Bariatric Surgery Database” (GC-MBD) (Clinicaltrial.gov: NCT03800160) where data of >10,000 cases have been recorded.

Data quality control
The database needs a committee to hold regular meetings to discuss issues concerning quality control. It is highly recommended that a multidisciplinary team achieve a consensus on the variables in the database. For example, according to the consensus of surgeons, radiologists, clinical researchers, and statisticians, variables that should be documented in the GC-MBD include, but are not limited to, structured demographic information, laboratory tests, PDFF values, VAT and SAT values, biological sample information, and adverse event records. Upper abdominal MR images in the DICOM format should also be uploaded. Before data entry, it is essential for the team’s main participants to undergo training. The manager of the database should check and verify the authenticity, accuracy, and integrity of all information according to the source data. Data modification traces should be recorded in the system. After verification, the data should be locked.

Discussion
There are three key points in AI analysis of body composition using upper abdominal MR images: uniform data acquisition standards, imaging annotation, and database management. This guideline will promote the development and application of AI systems for the automatic quantification of PDFF and abdominal adipose tissue.

The PDFF value can significantly influence the availability of PDFF images. For patients without hepatic steatosis, the grayscale contrast between the hepatic parenchyma and vessels is insufficient to train the neural network of the AI system. New strategies may solve this problem. For example, the liver may need to be registered with other higher contrast structures (eg, axial T1-weighted imaging or portal venous phase of contrast-enhanced imaging) to achieve margin recognition for the AI system. As such, additional MR sequences and related parameter standards are required.

Whole-body MR imaging can precisely quantify the volume of adipose tissue. For timesaving, scanning and analyzing a single-slice abdominal MR is preferred. Since imaging annotation for AI analysis requires only a single slice image of fat, we can acquire a single slice at the axial level of the L1–L2 intervertebral disc during the Dixon sequence acquisition to achieve further reduction of scanning time.

Clinical practice guideline registration and ethical approval
This guideline was registered on the International Practice Guideline Registry (IPGRP-2021CN177).

This work was approved by the Ethics Committees of Beijing Friendship Hospital, Capital Medical University (No. 2018-P-022-01).

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Conflicts of interest
None.

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