
Contents

SECTION I

INTRODUCTION AND FUNDAMENTALS

- 1 Introduction to FDG PET/CT, 1
- 2 FDG PET/CT Performance and Reporting, 5

SECTION II

MUSCULOSKELETAL SYSTEM AND BODY WALL

- 3 Skeleton on FDG PET/CT, 9
- 4 Muscle and Nerve on FDG PET/CT, 33
- 5 Skin and Breast on FDG PET/CT, 43

SECTION III

HEAD AND NECK

- 6 Brain on FDG PET/CT, 51
- 7 Head and Neck on FDG PET/CT, 63

SECTION IV

THORAX

- 8 Lung on FDG PET/CT, 87
- 9 Pleura on FDG PET/CT, 99
- 10 Heart on FDG PET/CT, 103
- 11 Thymus Masses on FDG PET/CT, 109

SECTION V

ABDOMINAL SOLID ORGANS

- 12 Hepatobiliary FDG PET/CT, 113
- 13 Spleen on FDG PET/CT, 127

- 14 Pancreas on FDG PET/CT, 133

- 15 Adrenal Glands on FDG PET/CT, 143

SECTION VI

GASTROINTESTINAL TRACT AND PERITONEUM

- 16 Gastrointestinal Tract on FDG PET/CT, 151

- 17 Peritoneum on FDG PET/CT, 171

SECTION VII

GENITOURINARY

- 18 Urinary Tract on FDG PET/CT (Kidneys, Ureters, Bladder), 179

- 19 Female Pelvis on FDG PET/CT, 191

- 20 Male Pelvis on FDG PET/CT, 205

SECTION VIII

ADDITIONAL ISSUES IN FDG PET/CT

- 21 Lymph Nodes on FDG PET/CT, 211

- 22 Measuring Treatment Response on FDG PET/CT, 225

- 23 Artifacts on FDG PET/CT, 231

SECTION IX

RADIOTRACERS OTHER THAN FDG FOR ONCOLOGIC PET/CT

- 24 Radiotracers Other Than FDG for Oncologic PET/CT, 235

CHAPTER 1

Introduction to FDG PET/CT

¹⁸F-fluorodeoxyglucose positron emission tomography/computed tomography (FDG PET/CT) provides noninvasive metabolic and anatomic imaging. The radioisotope, fluorine-18, has a short half-life allowing for imaging with limited patient dose. Fluorine-18 is used to chemically replace a hydroxyl group on glucose, and the resultant FDG is taken up into cells analogous to glucose. As tumor cells often uptake more glucose than normal tissues, FDG allows effective imaging of tumors.

The majority of this textbook focuses on hybrid FDG PET/CT for oncology. More than 1.7 million PET examinations are performed each year in the United States. There are several interesting facts about these PET examinations.

1. Greater than 95% of PET scans are performed with FDG as the radiotracer.
2. 95% of PET scans are performed for oncology (3% for cardiology, 2% for neurology).
3. Greater than 95% of PET scans are performed as hybrid PET/CT studies. (Data from IMV 2015 PET Imaging Market Summary Report, <http://www.imvinfo.com/index.aspx?sec=p&sub=dis&siteid=200083>.)

Let's look at each point.

1. Greater than 95% of PET scans are performed with FDG as the radiotracer.
FDG is by far the most commonly used radiotracer for PET imaging. There are other PET radiotracers that are U.S. Food and Drug Administration (FDA) approved for imaging, and many more which are not FDA approved, but the vast majority of PET scans are currently performed with FDG as the tracer. Thus the majority of this book focuses on the interpretation of FDG PET. The final chapter introduces other radiotracers with strong potential for increased utilization in the future.

2. 95% of PET scans are performed for oncology.

There are important applications of PET for cardiology and neurology; however, the vast majority of PET examinations are performed to evaluate patients with malignancy. Thus this book focuses on oncologic FDG PET.

3. Greater than 95% of PET scans are performed as hybrid PET/CT studies.

There are several reasons for this. First, PET scans are easier to interpret when they are corrected for attenuation. A PET camera counts 511 keV photons that are received by its detectors, but this is not the number of photons that are actually emitted. Many photons are attenuated while passing through the body before they reach the camera. In general, the deeper the photons originate within the body, the more tissue they need to pass through, and the more attenuation occurs. Thus the camera sees a greater percentage of the photons that originate near the body surface, less from those that originate deeper within the body. We can create an image from the number of photons actually detected by the PET camera. This is called the nonattenuation corrected image (Fig. 1.1).

Notice how the surface of the body appears to have more counts than areas deeper in the body. For example, the superficial portion of the liver (*arrow*) appears to have more counts than the deeper portions of the liver in the nonattenuation corrected image (see Fig. 1.1). However, we know that in a normal liver, the number of photons emitted from the liver cells should be about equal. This limits visualization of the deeper structures in the body.

When we "correct" for attenuation, the FDG images appear the way we are used to seeing them (Fig. 1.2).

Now the liver appears homogeneous in the attenuation corrected image (see Fig. 1.2). How is this attenuation correction done? The PET/CT camera uses the data from CT photon attenuation as a map to "correct" for attenuation of the photons created by

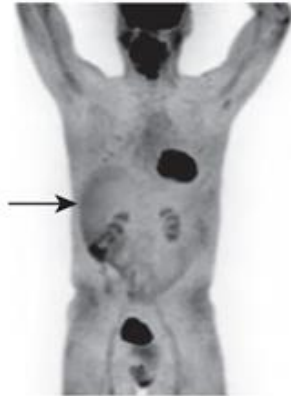


FIG. 1.1 Nonattenuated corrected maximum intensity projection image from an FDG PET scan. Arrow points at the liver surface, which appears to have more photon counts than the deeper portions of the liver.

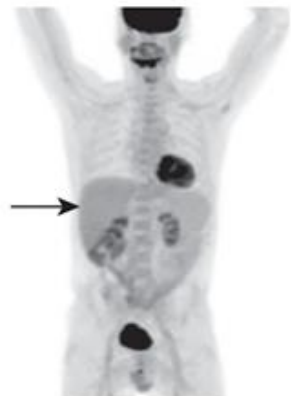


FIG. 1.2 Attenuated corrected maximum intensity projection image from the same FDG PET scan as Fig. 1.1. Arrow points at the liver surface, which after attenuation correction appears to have the same photon counts as the deeper portions of the liver.

positron annihilation. In general, the attenuation corrected image is much easier to interpret. How does a PET-only camera correct for attenuation? A PET-only camera uses a transmission positron source. Thus both PET-only and PET/CT cameras can effectively correct for attenuation. However, the CT scan takes only seconds to acquire on a PET/CT, but the transmission data may take 30 minutes to acquire

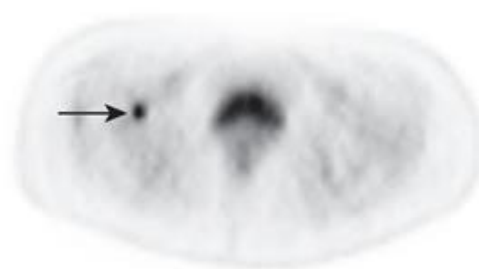


FIG. 1.3 Axial FDG PET through the pelvis in a patient with breast cancer. There is an FDG focus in the right pelvis (arrow). The midline FDG avidity is urine in the bladder.

on a PET-only camera. Thus the use of hybrid PET/CT allows much faster patient throughput than a PET-only camera.

Second, the CT component of a PET/CT allows for lesion localization. Sometimes it is difficult to determine where a PET focus is in the body (Fig. 1.3). Fig. 1.3 depicts a patient with breast cancer. There is an FDG focus in the right pelvis. What this FDG focus represents depends on where this focus is. If it is within a bone, then it is suspicious for an osseous metastasis. However, if it is outside the bone in a muscle, then it is probably physiologic muscle and is benign. The ability to fuse the FDG PET and CT images (Fig. 1.4) allows for localization of the FDG focus to the bone, and thus the FDG focus is suspicious for osseous metastasis. This biopsy was proven to be an osseous metastasis.

Third, the CT component of a PET/CT helps with lesion characterization. The corresponding findings on the CT of a PET/CT can help determine what an FDG focus represents (Fig. 1.5). In Fig. 1.5, a patient with lymphoma has an FDG-avid focus in the right inguinal region, which at first glance is suspicious for an FDG-avid lymph node. However, look at the corresponding CT image. The lesion is lower in attenuation than expected for a lymphoma node. The Hounsfield units of this lesion are 0, equal to water. This CT finding would be unusual for a lymphoma node and thus prompts further investigation. The same patient in sagittal projection is shown in Fig. 1.6. On the sagittal PET image, there is again an FDG focus in the right inguinal region; however, on the sagittal CT image, the bladder can be seen to be herniating through an inguinal hernia. The FDG focus in the right inguinal region is due to is a benign bladder hernia, rather than nodal lymphoma. This

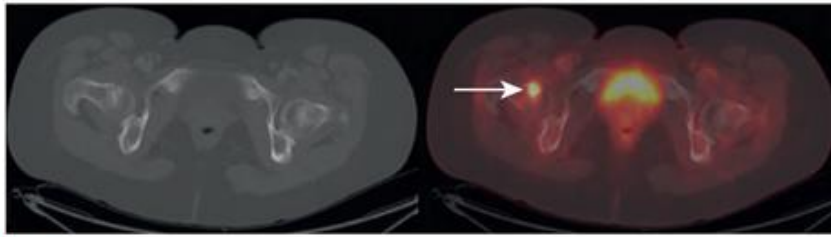


FIG. 1.4 Axial CT and fused FDG PET/CT that correspond with Fig. 1.3. The FDG focus in the right pelvis localizes to the right femur (*arrow*) and is suspicious for an osseous metastasis. Biopsy proved this was an osseous metastasis.

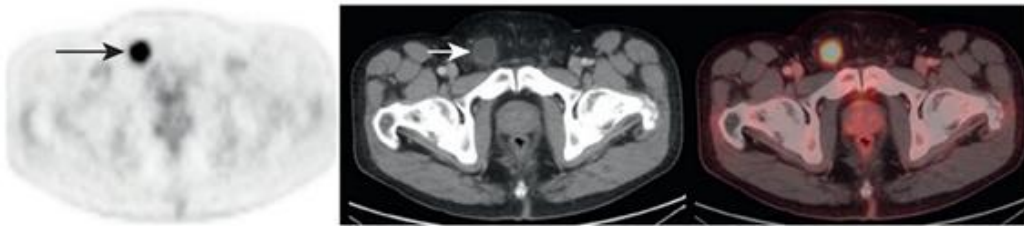


FIG. 1.5 Axial FDG PET, CT, and fused FDG PET/CT through the pelvis of a patient with lymphoma. The FDG-avid focus on the PET-only image (*arrow*) is at first glance suspicious for a malignant node. However, on the corresponding CT image, the lesion has attenuation equal to water (*short arrow*). This is unusual for a lymphoma node and prompts further evaluation.

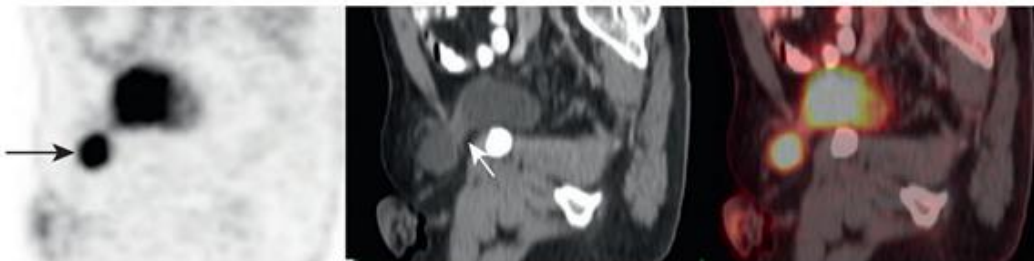


FIG. 1.6 Sagittal FDG PET, CT, and fused FDG PET/CT through the pelvis of the same patient as Fig. 1.5. The FDG focus in the right inguinal region is again seen on the PET-only images (*arrow*). The sagittal CT image demonstrates the herniation of the bladder through an inguinal hernia (*short arrow*). The CT helps characterize the FDG focus in the right inguinal region as a benign bladder hernia, rather than malignancy.

explains the Hounsfield units of 0 on CT. The FDG "lesion" is actually only urine within a herniated bladder. The CT component of the PET/CT was instrumental in characterizing the PET finding and preventing misdiagnosis.

Given the advantages for (1) rapid attenuation correction, (2) lesion localization, and (3) lesion characterization, almost all PET scans are performed as hybrid PET/CT scans. Thus this book focuses on hybrid PET/CT.

In summary, the vast majority of PET scans currently performed are hybrid FDG PET/CT for oncology, and thus the majority of this book focuses on these examinations. This content will cover organ system by organ system throughout the body, emphasizing how to integrate FDG PET and CT findings to arrive at the best interpretation of FDG PET/CT studies. My goal is to provide you with an organized, systematic approach to reading oncologic FDG PET/CT.