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Dr. Mukesh Singla
Professor, SPGOI, Rohtak,
Haryana, India

Imaging techniques: A review

Dr. Mukesh Singla

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Abstract

Computed Tomography (CT) and Positron Emission Tomography (PET) are nowadays within the most employed imaging techniques for the screening and the identification of multiple pathologies, including lung cancer. Medical imaging is a field in continuous development to increase the employment of non-invasive techniques in diagnosis. For the screenings or detection of tumor masses, the use of these techniques can reduce the stress for the patient avoiding a surgical intervention and its risks. Moreover, imaging can be used as a preclinical tool for the decision making process for the most suitable treatment without the necessity of directly intervene in an invasive way.

Keywords: Computed tomography, positron emission tomography, Fluoro-Deoxy-glucose

1. Introduction

1.1 Nodules

A solitary pulmonary nodule ^[12] (parenchymal, non-pleural nodule) is a small, round or egg-shaped lesion in the lungs. Juxtapleural pulmonary nodule is a small, worm-shaped lesion connected to pleura. (Figure 1)

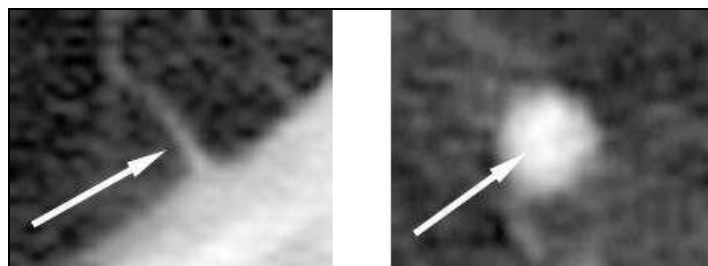


Fig 1: Nodule: Juxta-pleural nodule (left), parenchymal nodule (right).

Nodules are typically asymptomatic, and they are usually noticed by chance on a chest X-ray that has been done for another reason. They are usually smaller than 3-4 cm in diameter (no larger than 6 cm) and are always surrounded by normal, functioning lung tissue. Their intensity in CT scans is from -300 to 0 HU. Nodules are fairly common abnormalities on chest X-ray images: nearly one of every 500 chest X-rays shows a newly diagnosed nodules ^[12]. In the United States, physicians are challenged each year by more than 150,000 new cases. Sixty percent of all nodules are benign. In certain geographical areas where the infectious agents (especially fungi) that cause nodules occur, the percentage of benign nodules increases remarkably (in some areas as high as 90% to 95%). Malignant nodules may be primary lung cancer tumors or metastases from other parts of the body. If the lesion is suspected to be benign, serial chest X-rays or CT scans may be taken on a regular basis for observation of the lesion. If the affected person is at high risk for lung cancer or if the CT scan appearance of the lesion suggests it is malignant, surgical removal of the lesion is recommended.

1.2 Tomography

Tomography is a method to obtain a cross-sectional images (transversal slices) of given object. In Computed Tomography images of objects (patients) are obtained by X-ray projection ^[11]. The mathematical basis for tomographic imaging was laid down by Johann

Correspondence
Dr. Mukesh Singla
Professor, SPGOI, Rohtak,
Haryana, India

Radon (December 16, 1887 (Litoměřice)–May 25, 1956). By applying his theorem slices of human body at various angles can be reconstructed.

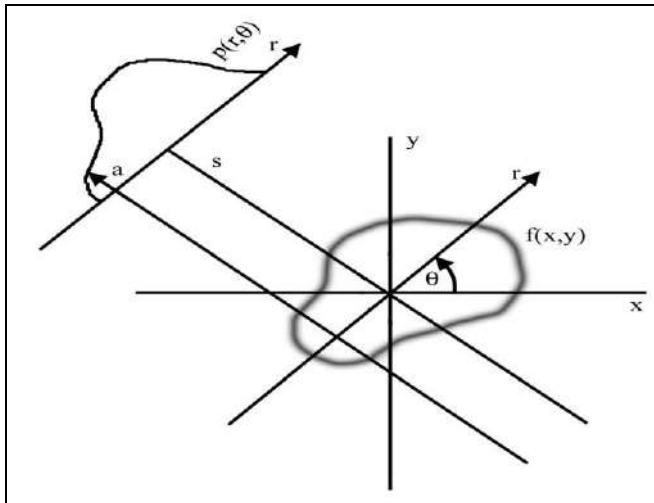


Fig 2: Parallel beam geometry

In X-ray CT, the line integral represents a logarithm of the total attenuation of the beam of X-rays as it travels in a straight line through the object.

2. Imaging techniques

In the medical field the identification of pathologies in the early stages is a crucial open problem that is at the base of a correct diagnosis and a subsequent decision of the most appropriate therapy. An invasive method for lung cancer detection is the employment of biopsy, which is an extraction of a tissue sample from the living patient [1]. It is clear that this approach is painful for the subject since it involves surgery. In order to avoid or reduce the employment of this dangerous technique some alternatives have been proposed, such as imaging based assessment. For example, in the past, one of the most used technologies was the X-rays imaging [2], while in recent years it has been used the F-fluoro-deoxy-glucose Positron Emission Tomography/Computed Tomography imaging method, known also as FDG PET/CT. This examination allows the fusion of information from multiple sources giving, in this way, a more precise and accurate reproduction of the inside of the human body. Next Sections describe more in detail both the CT and the PET.

2.1 Computed tomography

The Computed Tomography (CT) is a clinical imaging technique that exploits a X-rays beam in order to acquire signals, that after being elaborated by a computer, generate images generally called slices. The images are spatial representation of the attenuation coefficient of the rays in a section of the scanned object and can be considered as tomographic reconstructions of the body that contain more information compared to the traditional X-rays. The obtained images can also be stacked in order to form a 3-D reconstruction of the body district investigated, facilitating the visualization and the identification of organs or abnormal structures by a specialist [3]. The main difference between the conventional X-rays and the CT is that the first one uses a fixed source, while the CT has multiple sources that rotate around the patient inside a circular structure

called gantry. A CT scan is a clinical examination that consists of a collection of multiple contiguous images of a specific part of the body in accordance to the area of interest of the clinician, as shown in Figure 3. During the acquisition, the patient lies supine on a table, that can be moved in or out of the gantry while the X-rays sources rotate around the subject [4]. The sources constantly release beams of X-rays that, passing through the human body and reaching digital detectors, which are positioned opposite to the sources, are able to create the images. The creation of a single 2-D slice is possible only after a complete rotation of the sources. In fact, in order to reconstruct the image, it is necessary to apply to the signals collected a mathematical procedure called back-projection reconstruction [5]. This step is done through the Radon transform, which, considering the angle of acquisition of each signal, is capable of mixing them in the most suitable way for reducing the creation of artifacts. This procedure is repeated for each movement of the table so that the entire region of interest is covered. The thickness of each slice can be decided by the physician, but it usually varies between 1 to 10 millimeters and the dimensions of each pixel are around few millimeters, allowing a very good spatial resolution [6]. Due to these intrinsic geometric characteristics, the CT is often preferred to the X-rays and has become a common tool for the identification of lesions and tumors in the abdomen, lungs, head and also for inspection of the heart [7]. Consisting in the same technology of the X-ray, the CT can easily discriminate between hard and soft tissues due to the fact that structures such as bones stop the rays creating light spot in the image, while soft organs result in darker shape in the reconstructed image. For the aforementioned reasons, the CT is a good standard for structural investigation; however, Positron Emission Tomography, which gives details about the metabolic active structures, allows a more precise identification of tumors.

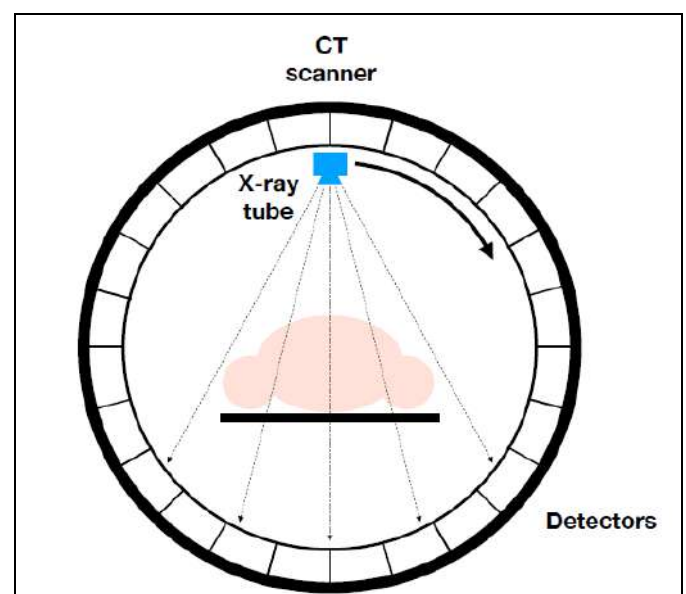


Fig 3: Basic scheme of Computed Tomography

2.2 The Positron Emission Tomography (PET)

PET is an imaging technology used in nuclear medicine that allows the assessment of structures functionalities employing radioactive substances. The PET creates 3-D images using radio-pharmaceuticals as tracers, which decay

releasing positrons, which are then used for the creation of the images [8]. Positrons are particles that have similar mass to the one of electrons but with opposite charge, so passing through the human body they combine with the electrons and annihilate one another. This chemical reaction releases energy and two photons which produce two rays that, as shown in Figure 4, are shot in opposite direction and by colliding with the detectors are employed for the creation of the acquired images [9]. The main limitation of the PET is that it has an intrinsic low spatial resolution mainly due to the physical width of the detectors in combination with the decoding of the signal and the penetration. In fact, as highlighted in [3, 4], if the rays are not perfectly perpendicular incident onto the detectors, they can interact with more than one detector so the signal can be associated to the wrong one degrading the resolution of the final image. The tracers used for a PET scan are formed by carrier molecules bonded to radioactive atoms and usually are administrated to the patient by injection, inhalation or ingestion [11]. The total amount of the tracer is so low that it does not influence the normal function of the system: for these reasons the PET is defined as physiologic tomography. This methodology is

based on the emission of positrons e^+ from the decay of the isotopes, which have a very short life, so, after a short path, (for example in lungs where there is the lowest tissue density), these positrons can reach maximum few millimeters. In the tissues, positrons annihilate with a negative electron generating in this way two rays of 511 KeV (Kilo electron Volt) emitted in opposite directions (180°) [9]. Positioning a couple of detectors is possible to exactly identify the line along which positrons have been emitted, shown in Figure 4. In addition, if two logic impulses are super imposed in time, the system recognizes annihilation and records also the integral value of the line joining the two activated crystals, which is then coded in the image. The aim of PET imaging technique is to identify cancer, monitor it and verify the efficiency of treatments, in addition to the detection of metastases. As tracer is usually used glucose because cells or tissues that have a high metabolic activity, such as dividing cancer cells, request a huge amount of energy, which means an increase in the glucose consumption. In fact, the more the cancer is aggressive, the more rapidly it will utilize glucose. For these motivation radiolabeled glucose is employed as tracer for

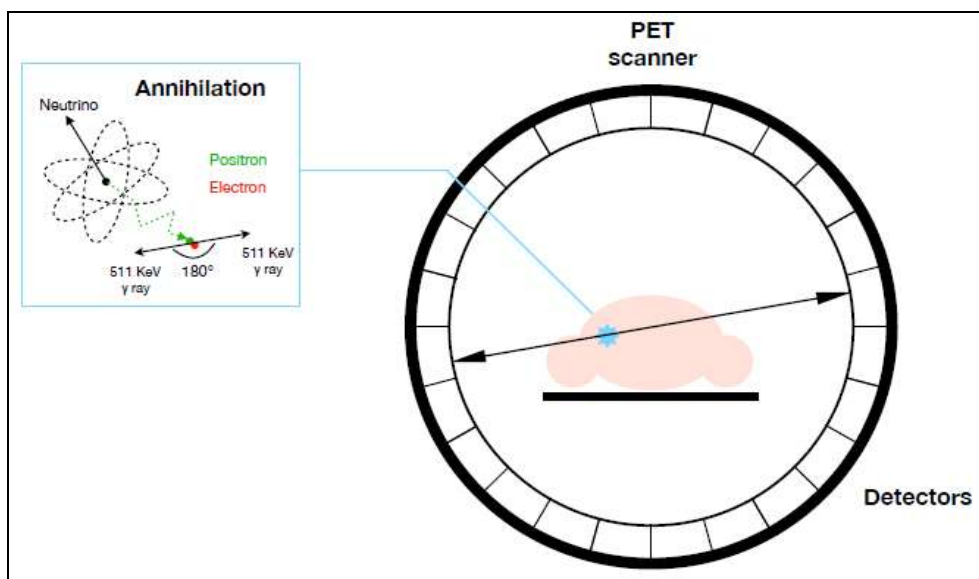


Fig 4: Basic scheme of Positron Emission Tomography

the detection of cancer and metastases spread in the body [12]. In the proposed work the tracer used for the PET acquisition is F-fluoro-deoxy-glucose (FDG). Summarizing what has been said above it is possible to conclude that the CT provides images with a good spatial resolution giving interesting details about geometry and mechanical properties of the internal structures. On the other hand, the PET, which lacks in spatial resolution, is able to highlight the metabolic active areas allowing the identification of the structures that are requesting a considerable amount of blood such as the heart or tumors. As confirmed by [13] scanning the same body district with both in one examination (PET/CT imaging) has become a highly employed tool for the identification and the staging of cancer all around the world thanks to the combination of anatomical (CT) and functional (PET) information. For the aforementioned reason, this thesis employs data acquired by the FDGPET/CT, in this way, the proposed pipeline is able not only to analyze the selected dataset, but also other dataset acquired with this widespread technology.

3. Conclusion

CT and PET can be used separately but more and more frequently are jointed to combine information both from a structural and a functional point of view. In fact, the CT has a good spatial resolution giving the possibility to highlight the details of the inside of the body district investigated. The good resolution allows discrimination between the bones, the air inside the body and soft tissues; this classification is done exploiting the density of each structure, where a dense one appears lighter on the resulting image, while a hollow results as black. Considering the aforementioned characteristics the CT can be used for obtaining anatomical details that are useful for the identification of multiple diseases but cannot be enough when detailed functional information are needed. PET, on the other hand, thanks to the used radio-tracers, provides information about the metabolic activity of tissues. In fact, more active is the structure, more metabolic request it has and more blood containing the tracer it beckons resulting in a bright spot on the image.

4. References

1. Biopsy. <https://www.cancer.gov/publications/dictionaries/cancer-terms/def/biopsy>. Accessed: 2018-04-03.
2. X-rays. <https://www.cancer.gov/publications/dictionaries/cancer-terms/def/x-ray>. Accessed: 2018-03-22.
3. Computed tomography. <https://www.nibib.nih.gov/science-education/science-topics/computed-tomography-ct>. Accessed: 2018-03-21.
4. Positron emitting products. <https://www.fda.gov/Radiation-EmittingProducts/Radiation-Emitting-Products-and-Procedures/Medical-Imaging/Medical-X-Rays/ucm115317.htm>. Accessed: 2018-03-21.
5. Miqueles E, Koshev N, Helou ES. A backprojection slice theorem for tomographic reconstruction. *IEEE Transactions on Image Processing*, 2018; 27(2):894-906.
6. Ct resolution. <http://www.ctlab.geo.utexas.edu/about-ct/resolution-and-size-limitations/>. Accessed: 2018-03-21.
7. Liguori C, Frauenfelder G, Massaroni C, Saccomandi P, Giurazza F, Pitocco F *et al.* Emerging clinical applications of computed tomography. *Medical Devices (Auckland, NZ)*. 2015; 8:265.
8. Nuclear medicine. <https://www.nibib.nih.gov/science-education/science-topics/nuclear-medicine>. Accessed: 2018-03-21.
9. Shibuya K, Yoshida E, Nishikido F, Suzuki T, Tsuda T, Inadama N *et al.* Annihilation photon acollinearity in pet: volunteer and phantom FDG studies. *Physics in Medicine & Biology*. 2007; 52(17):5249.
10. Moses WW. Fundamental limits of spatial resolution in pet. *Nuclear Instruments and Methods in Physics Research Section A: Accelerators, Spectrometers, Detectors and Associated Equipment*. 2011; 648:S236-S240.
11. Wernick MN, Aarsvold JN. *Emission tomography: the fundamentals of PET and SPECT*. Elsevier, 2004.
12. Kaira K, Oriuchi N, Sunaga N, Ishizuka T, Shimizu K, Yamamoto N. A systemic review of pet and biology in lung cancer. *American journal of translational research*, 2011; 3(4):383.
13. Szyszko TA, Yip C, Szlosarek P, Goh V, Cook GJ. The role of new pet tracers for lung cancer. *Lung Cancer*, 2016; 94:7-14.