

## Special Article – Breast Cancer- Risk Factors, Diagnosis and Treatment

The Benefit of  $^{18}\text{F}$ -FDG PET/CT in Breast Cancer Patient Staging and Treatment Planning: A Case Report

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## Abstract

A 50 year old female patient was treated with left mastectomy plus axillary dissection for ductal infiltrating carcinoma mpT1c pN1(1/15) M0 in October 2009. The patient was staged post-operatively with  $^{18}\text{F}$ -FDG PET/CT and unexpected internal mammary chain lymph node metastases were detected. The patient received adjuvant chemo/hormonal therapy and radiotherapy on regional lymph nodes up to a dose of 50 Gy with a volume modulated arc (VMAT) technique, reaching a complete response. At 5 year follow-up the patient is still alive in complete response without radiotherapy late toxicity.

**Keywords:**  $^{18}\text{F}$ -FDG PET/CT; Breast cancer; Internal mammary chain; VMAT

## Introduction

The role of  $^{18}\text{F}$ -FDG PET/CT in the staging of breast cancer is still controversial. Its use in early stage cases is not recommended, whereas in locally advanced cases it can lead to a change in the staging phase from 6.7% to 52% and consequently to a change in the treatment approach from 5.6% to 56% of the cases [1,2].

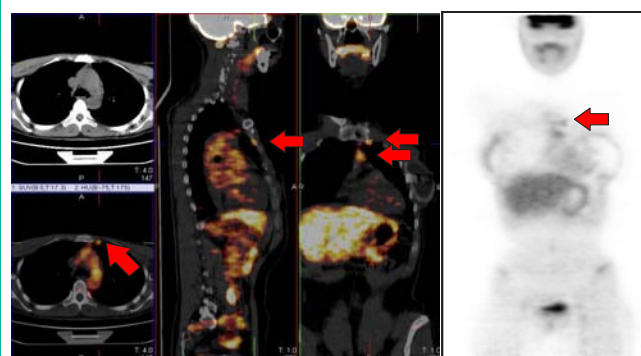
$^{18}\text{F}$ -FDG PET/CT improves the diagnosis of undetected lymph node metastases outside the axilla (i.e. infraclavicular, supraclavicular and internal mammary nodes) or occult and asymptomatic distant metastases [3].

## Case Presentation

A 50 year old female patient detected a lump in the inner-superior quadrant of left breast. Subsequent mammography, ultrasound and fine needle aspiration confirmed a ductal infiltrating carcinoma and she was treated with mastectomy, simultaneous reconstruction with prosthesis and contralateral mastopasty. The final diagnosis confirmed a ductal infiltrating carcinoma ER95%, PR80%, Ki67 20%, c-erbB2 score 2 (FISH neg), lymphovascular invasion, mpT1c pN1(1/15) M0 associated to peritumoral DCIS G3 (5-10%). A post-operative  $^{18}\text{F}$ -FDG PET/CT was performed and the final stage was modified in pN3b due to the detection of metastases in the internal mammary chain lymph nodes (IMC LN). The diagnose was confirmed by a fine needle aspiration on the largest lymph node. Therefore the patient received 4 adjuvant chemotherapy cycles (AC schedule), LH-RH analogues for 2 years, and aromatase inhibitors for 5 years and radiotherapy at the end of chemotherapy.

The radiotherapy treatment was performed with VMAT Rapid Arc (Varian Medical System, Palo Alto, California) from April 2010 until May 2010. The prescribed total dose was of 50 Gy, 5 fractions per week, 2 Gy each. The post-operative  $^{18}\text{F}$ -FDG PET/CT was merged with the planning CT. The contoured Clinical target Volume (CTV) included the supraclavicular/infraclavicular lymph nodes and the IMC. The chest wall was excluded due to fact that the tumor size was <2 cm and due to the presence of the prosthesis. The  $^{18}\text{F}$ -FDG PET/CT allowed a correct detection of the IMC LN, not visible on

the planning CT (Figure 1). The Planning Target Volume (PTV) was obtained extending the CTV up to 0.8 cm and cropped 0.2 cm from the body surface. As Organs at Risk (OARs), the contralateral breast, the ipsilateral prosthesis, the lungs, the spine, the heart, the thyroid and the left anterior descending coronary artery (LAD), were considered. A dedicated structure was contoured for minimizing the dose spillage outside the PTV, named Healthy Tissue (HT). Such structure was drawn at 0.2 cm distance from the PTV and extended up to 2.0 cm. The treatment plan was designed on the Varian Eclipse Treatment Planning System (platform 8.6) for a Varian iX linear accelerator, 6 MV energy beam. Two partial arcs (CW and CCW) were applied in order to avoid beam entrances from the contralateral lung and contralateral breast. Inverse planning was performed by means of the Progressive Resolution Optimizer III, version 10.0.28, employing the intermediate dose options in order to increase the accuracy of optimization and the agreement between optimization and final dose calculation. Dose-volume constraints during the optimization were defined to match the planning goals of having more than 95% of the



**Figure 1:**  $^{18}\text{F}$ -FDG PET/CT pre-irradiation.

On the upper corner of the left hand-side a coronal view of the planning CT where no lymph nodes are visible; the same CT plane is shown in the lower corner blended with the  $^{18}\text{F}$ -FDG PET which makes the IMC lymph nodes clearly localizable. The median part of the image displays the sagittal and frontal views of the whole planning CT blended with the  $^{18}\text{F}$ -FDG PET, with red arrows indicating the IMC lymph nodes. On the right hand-side the  $^{18}\text{F}$ -FDG PET alone, with red arrows indicating the same IMC lymph node.

**Table 1:** Dosimetric parameters.

Dosimetric parameters resulting from the treatment planning of the patient for the PTV and OARs.  $V_x$  represents the percentage of volume receiving an x% (or x Gy) level of dose;  $D_y$  represents the dose received by a percentage y of the volume.

Organ	Dosimetric Parameters
PTV	$V_{95\%} = 95\%$ $V_{107\%} = 0\%$ $D_{2\%} = 104\%$
Ipsilateral lung	Mean Dose = 8.4 Gy $V_{20Gy} = 10\%$
Ipsilateral breast prosthesis	$V_{30\%} = 12\%$
Controlateral lung	Mean Dose = 7.4 Gy
Controlateral breast	Mean Dose = 2.7 Gy
Heart	Mean Dose = 8.0 Gy
LAD	Mean Dose = 4.8 Gy $D_{2\%} = 12.6$ Gy

PTV volume receiving >95% of the prescribed dose ( $V_{95\%} > 95\%$ ) and of minimizing the dose to the ipsilateral lung ( $V_{20Gy} < 20\%$ ), to the heart and the LAD. The obtained dosimetric parameters are listed in Table 1. The patient concluded the scheduled treatment. Orthogonal X-Ray imaging was performed every day for a correct patient set-up.

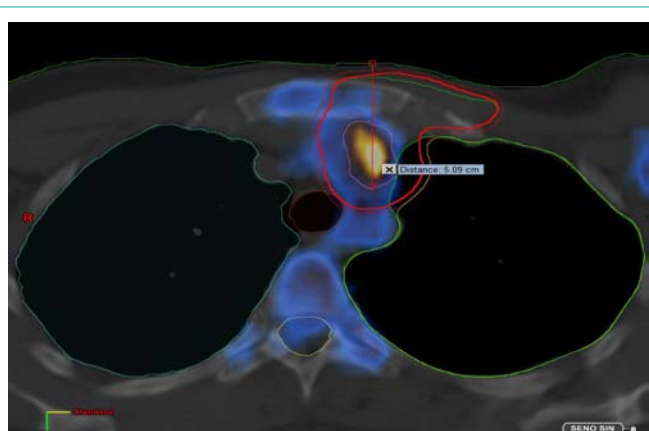
Only grade I cutaneous toxicity (EORTC/RTOG scale) was detected during treatment.

In June 2010 a  $^{18}\text{F}$ -FDG PET/CT confirmed the complete response on macroscopic IMC but a mammography detected a doubtful area in the upper external quadrant of the right breast. Since a fine needle aspiration was suspicious for DCIS the patient underwent right mastectomy and simultaneous reconstruction with prosthesis, in July 2010. No cancer cells were found in the mastectomy specimen, while a papilloma associated with ductal hyperplasia was detected.

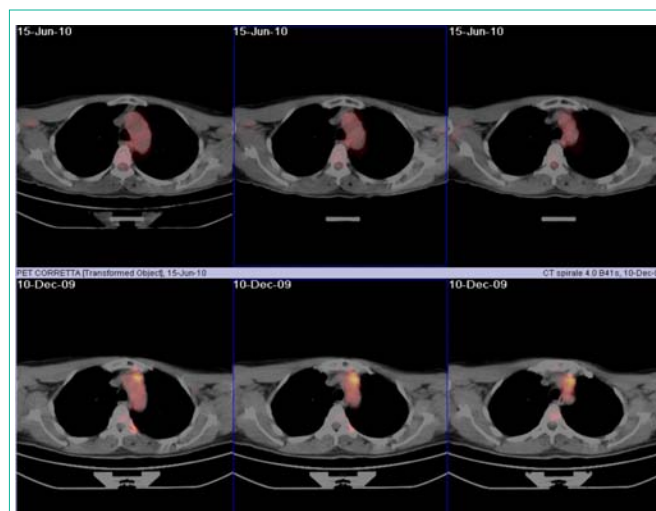
Up to date the patient has been regularly followed up. No late toxicity was registered. Noteworthy the left prosthesis profile was not altered and the controlateral mastectomy was performed without any complications, despite their inclusion in the irradiated volume.

## Discussion

$^{18}\text{F}$ -FDG PET/CT was requested, after a multidisciplinary



**Figure 2:** Location of one of the IMC lymph nodes. Coronal view of the planning CT blended with the post-operative  $^{18}\text{F}$ -FDG PET. In red the PTV volume contoured on the basis of the PET image information. The red line measures the depth of the detected lymph node from the body surface (around 5 cm).



**Figure 3:**  $^{18}\text{F}$ -FDG PET/CT pre- and post-irradiation. In the first row, three coronal views of the  $^{18}\text{F}$ -FDG PET/CT acquired in June 2010 after irradiation; in the second row, the same slices are shown from the  $^{18}\text{F}$ -FDG PET/CT pre-irradiation (December 2009). In the latter images, an up-take of  $^{18}\text{F}$ -FDG is clearly visible in correspondence of the IMC lymph nodes, whereas the former images reveal a complete remission of the patient.

discussion, because of the breast cancer site (inner quadrant). It is known from literature that the risk of IMC metastases is around 3% in upper outer quadrant breast cancer stage I, and >65% in internal inner quadrant in case of positive axillary lymph nodes [4].

This case report strengthened the role of high sensitive and sensible diagnostic tool, such as  $^{18}\text{F}$ -FDG PET/CT, in the localization of IMC LN not adequately detectable by other diagnostic approaches. This patient was treated according to the real stage both through systemic and loco-regional therapy.

The IMC localization has always been problematic and controversial due to the lack of a non-invasive accurate diagnostic tool: lymphoscintigraphy and CT (indirect detection) can reach an accuracy of 5 mm and 6 mm in depth, and of 6 mm and 7 mm in lateral direction, respectively; ultrasonography is not accurate enough [5,6]. Consequently in multicentric randomized trials, investigating the role of IMC radiotherapy, no specific recommendations regarding the IMC site has been given. As an example, according to the EORTC 22922/10925 trial [7], the IMC is located in the parasternal region, and approximately 85% of the lymph nodes lie within 4 cm lateral to the midline and within 4 cm depth in the first 3 intercostal spaces. The top of the IMC lies behind the insertion of the sternocleidomuscle in the medial supraclavicular area. According to the French trial [8] the IMC is located in the first five intercostal spaces. In the current case the  $^{18}\text{F}$ -FDG PET/CT identified the presence of IMC LN even at 5 cm depth and 4.5 cm laterally from the midsternum line (Figure 2), allowing to draw the target volume for irradiation with definitively high confidence. To remark is also the fact that the same imaging conducted in June 2010 revealed a complete response (Figure 3) in contrast with the suspicious mammography. The report of the  $^{18}\text{F}$ -FDG PET/CT was then confirmed by pathology specimen. In conclusion, the here reported case is a strong example of the clinical efficacy of  $^{18}\text{F}$ -FDG PET/CT for diagnose and staging of IMC LN.

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