



UNIVERSITI PUTRA MALAYSIA

***AGREEMENT BETWEEN VARIOUS MEASUREMENTS OF
STANDARDISED UPTAKE VALUE NORMALISED BY LEAN BODY
MASS
IN DETECTING BACKGROUND 18F-FDG ACTIVITY IN PET/CT
ONCOLOGIC IMAGING***

NUR HAFIZAH BINTI MOHAD AZMI

FPSK(M) 2018 10



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By

NUR HAFIZAH BINTI MOHAD AZMI

**Thesis Submitted to the School of Graduate Studies, Universiti Putra
Malaysia, in Fulfilment of the Requirements for the Degree of Master
of Science**

March 2018

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Abstract of thesis presented to the Senate of Universiti Putra Malaysia in
fulfilment of the requirement for the degree of Master of Science

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March 2018

Chair : Subapriya Suppiah, PhD
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PET/CT Scan is a diagnostic imaging tool predominantly used in oncology cases. Standardised uptake value (SUV) is the widely accepted method to quantitatively assess lesions detected on PET/CT. There is a limitation to the utility of this method, however, as this value becomes falsely reduced in overweight patients.

Thus, we propose another quantitative method of using Standard Uptake Lean Body Mass (SUL) which can give a more consistent reading in patients having extremes of body mass index (BMI) values. As the prevalence of obesity is rising in this current decade, the utility of SUL becomes more relevant and necessary.

This study correlated SUV and SUL values using the liver as a baseline reference organ and identified the pattern of distribution across various BMIs. There have been some studies that assessed the variations of SUV and SUL in obese subjects, but there have not been any studies that analysed whether there is a significant difference in SUL in subjects who undergo contrast-enhanced PET/CT.

Interestingly, this study confirmed that SUL reading is consistent even among overweight patients and the utility of contrast media in PET/CT scans does not significantly differ from low dose non-contrast-enhanced scans.

Abstrak tesis yang dikemukakan kepada Senat Universiti Putra Malaysia
sebagai memenuhi keperluan untuk Ijazah Master Sains

**PERSETUJUAN ANTARA BEBERAPA CARA PENGUKURAN
STANDARDISED UPTAKE LEAN BODY MASS DALAM MENGESAN
AKTIVITI BACKGROUND 18F-FDG DALAM KES ONKOLOGI
MELIBATKAN IMBASAN PET/CT**

Oleh

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Imbasan PET/CT adalah alat yang digunakan dalam kes onkologi. Nilai *Standard Uptake Value* (SUV) adalah kaedah kuantitatif yang diterima secara meluas bagi mentafsir kerosakan yang dikesan pada imbasan PET/CT. Terdapat kekurangan ke atas kaedah ini kerana ia berubah pada pesakit yang berlebihan berat badan.

Oleh itu, kami mencadangkan satu kaedah kuantitatif menggunakan *Standard Uptake Lean Body Mass* (SUL) yang memberi bacaan yang lebih konsisten terutama pesakit yang mempunyai peningkatan pada indeks jisim badan (BMI). Malahan, didapati isu obesiti semakin meningkat masa kini maka SUL menjadi lebih relevan dan diperlukan.

Kajian ini mengkaji hubung kait nilai SUV dan SUL menggunakan organ bahagian hati sebagai rujukan asas untuk mengenal pasti nilai kuantitatif bagi pesakit yang berbeza BMI. Terdapat beberapa kajian yang menilai kepelbagaian SUV dan SUL pada pesakit gemuk, tetapi tidak terdapat lagi kajian yang menganalisis sama ada terdapat perbezaan yang ketara bagi nilai SUL pada imbasan PET/CT berkontras.

Menariknya, kajian ini mengesahkan bahawa nilai bacaan SUL adalah konsisten walaupun di kalangan pesakit yang berlebihan berat badan dan obes. Penggunaan media kontras dalam imbasan PET/CT tidak memberi bacaan yang ketara daripada nilai asasnya.

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LIST OF ABBREVIATIONS

BMI	Body Mass Index
CE	Contrast-Enhanced
NCE	Non-Contrast-Enhanced
ROI	Region of Interest
VOI	Volume of Interest
FBS	Fasting blood sugar
ICC	Intra-Class Correlations
LBM	Lean Body Mass
PE	Predictive Equation
PET/CT	Positron Emission Tomography Computed Tomography
SD	Standard Deviation
SUL-CG	SUL computed generated
SUL-James/ SULPE1	SUL derived from the predictive equation for lean body mass by James et al.
SUL-Janma/ SULPE2	SUL derived from the predictive equation for lean body mass by Janmahasatian et al.
SUL	Standardised Uptake Value Normalised for Lean Body Mass
SUV	Standardised Uptake Value
SUV-CG	Standard Uptake Value Computed generated
SUVmax	Maximum Standardised Uptake Value within a tumour / lesion
SUVmean	Averaged value to Standardised Uptake Value within a tumour / lesion
18F-FDG	18 Fluorine – Fluoro-deoxy-glucose
CDNI	Centre for Diagnostic Nuclear Imaging, UPM/ Pusat Pengimejan Diagnostik Nuklear, UPM
PCMC	Prince Court Medical Centre
A	Centre A
B	Centre B
Max	Maximum
Min	Minimum

CHAPTER 1

INTRODUCTION

1.1 Background of Study

Positron Emission Tomography / Computed Tomography (PET/CT) is a hybrid diagnostic imaging tool predominantly used in oncology cases; to diagnose and stage cancers as well as to monitor treatment response. The commonly used radio-isotope in oncology imaging is 2-(18F)-fluoro-2-deoxy-D-glucose (18F-FDG) which is a radioactive isotope tagged to a glucose analogue. Increased utility of glucose by most types of cancer cells forms the basis of PET/CT interpretation of deranged physiological uptake at a cellular level. Therefore, it is important to have a reliable quantification method to assess 18F-FDG uptake in lesions detected by this scan. There are two types of ways to quantify 18F-FDG uptake; namely visual qualitative assessment - comparing uptake in diseased tissue relative to the normal surrounding tissue, and the other is a semi-quantitative assessment using standardised uptake value. Standardised Uptake Value (SUV) is the most widely accepted method to semi-quantitatively assess lesions detected on PET/CT [1],[46],[47],[48] and [49]. This value is dependent on several factors, namely the patient's body habitus and the dose of the injected radioisotope. There is a limitation to the utility of this method, however, as this value becomes falsely increased in overweight and obese patients. This is caused by very little accumulation of FDG in white fat or adipose tissue during fasting state, hence leading to a higher FDG redistribution and uptake in non-fatty tissue [2].

Alterations in 18F-FDG intracellular metabolism is a useful non-invasive biomarker for monitoring treatment response that occurs in cancer cells. A consistent baseline reading is important to enable reliable comparison to be made in serial scans of cancer patients. Accurate measurements of relative tissue uptake of 18F-FDG are necessary. Unfortunately, these measurements are significantly influenced by variations that occur in daily clinical practice, particularly caused by the quantity of injected 18F-FDG dose and patient morphometrics [3]. Thus, we propose another semi-quantitative method of using Standard Uptake Value normalized for Lean Body Mass (SUL) which can give an improved consistency of measurements, particularly in cancer patients who have extremes of body mass index (BMI) values. As the prevalence of obesity is rising in this current decade, the utility of SUL becomes more relevant and necessary.

1.2 Problem Statement and Justification

Semi-quantitative evaluations of most 18F-FDG PET/CT scans in oncology imaging are frequently done using Standardised Uptake Value (SUV) measurements. This can be automatically calculated using a vendor installed computer-generated (CG) software. It indirectly measures the *in vivo* distribution of injected 18F-FDG in cells of body tissue. This reading is reliable in subjects who have a body weight that is within the range of the ideal body mass index (BMI). This reading, however, becomes falsely elevated in overweight and obese subjects [2],[3] and [4]. Thus, there are studies that propose using SUV normalized for lean body mass (SUL) to correct for this, especially in overweight and obese patients [5]. SUL can be acquired through computer-generated automated methods (SUL-CG), via calculations based on estimations of fat weight based on CT data [6] as well as using calculations based on predictive equations for lean body mass (LBM) such as James LBM formula and Janmahasatian LBM formula [5],[7],[38],[39] and [45]. Furthermore, as a large proportion of Malaysians currently fall in the overweight and obese category as evidenced by a recent study [8]. Thus, the issue of improving standardization of PET/CT quantitative calculations becomes even more relevant. Almost all previous studies have been based on low dose non-contrast-enhanced PET/CT scan images. Many centres nowadays perform contrast-enhanced PET/CT scans, thus there is a need to know whether the effect of contrast media would alter the standardization of SUL measurements [52],[55] and [56]. In particular obese patients may have abnormal range of readings. Hence, there is a need to identify whether SUL calculated using computer-generated values are more accurate compared to using estimates based on predictive equations for LBM staging [41].

1.3 Main Aim and Specific Objectives of the Study

The purpose of this study was to answer certain research questions. Research questions included:

1. What is the quantitative range of measurements of SUV and computer generated SUL in the liver area?
2. What is the inter-reader agreement for measurement of
 - a. SUV in non-contrast-enhanced PET/CT studies?
 - b. SUL (Computer generated) in non-contrast-enhanced PET/CT studies?
 - c. SUV in contrast-enhanced PET/CT studies?
 - d. SUL (Computer generated) in contrast-enhanced PET/CT studies?

3. What is the correlation between the measurements of liver SUV and liver SUL (Computer-generated)
 - a. In non-contrast-enhanced PET/CT scans?
 - b. In contrast-enhanced PET/CT scans?
4. What are the independent variables/ factors that affect SUV and SUL measurements in contrast-enhanced PET/CT scans?
5. What is the agreement between two types of methods to measure SUL using predictive equations namely, SUL-James (PE1) and SUL-Janmahasatian (PE2)?
6. What is the correlation between SUL (using contrast-enhanced scans compared to SUL using non-contrast-enhanced scans) with
 - a. BMI groups?
 - b. gender ?

The overall aim of this study was to identify the agreement of SUL (computer generated) and SUL derived by predictive equations at the liver area. This study also assessed the correlation of SUL measurements in contrast-enhanced scans compared to non-contrast-enhanced scans (within different BMI groups) as a semi-quantitative parameter in 18F-FDG PET/CT oncologic imaging for Malaysian population. Subsequent research findings that arise from this study can be a source of reference for improving technique and patient management in oncology imaging in Malaysia.

Specific Objectives of this study were:

1. To identify quantitative measurements of SUV and SUL at the liver area.
2. To measure inter-reader agreement for SUV and SUL measurements.
3. To measure the correlation between the measurements of liver SUV and liver SUL (Computer-generated).
4. To find out the independent variables/ factors that affect SUV and SUL measurements in contrast-enhanced PET/CT scans.
5. To test agreement among three different methods of measuring SUL i.e. computer generated SUL (interchangeably referred to as SUL or SUL-CG), as well as predictive equations for lean body mass based on James (SUL-James) (PE1) and Janmahasatian (SUL-Janma) (PE2).
6. To measure the correlation between SUL (using contrast-enhanced scans compared to SUL using non-contrast-enhanced scans) with BMI groups and gender.

1.4 Organisation of Thesis

The organisation of this thesis is as follows. In the next chapter, we review previous works in studying the various factors that affect SUV and SUL values. We then focus on the effects of body mass index on readings of SUV and SUL. We will explain why the liver was used as a site for baseline measurements. We will then highlight the effects of extremes of BMI upon SUV and discuss the benefits of using SUL. We will also compare the reliability of various methods used for achieving SUL readings i.e. SUL computer generated and SUL derived from predictive equations. We will also discuss the significance of obesity in causing derangement of standardized baseline semi-quantitative PET/CT assessment, which will be highlighted in the last subsection in Chapter 2.

Chapter 3 provides a brief description of how we developed our study protocol. We will explain our materials and methods in conducting this study. We will discuss the need for a multicentre study and explain the statistical methods we used in analysing the study results.

In Chapter 4, we present our results from Centre A i.e. Centre for Diagnostic Nuclear Imaging, Universiti Putra Malaysia and Centre B Prince Court Medical Centre in the form of tables, graphs and PET/CT image figures. Our related work has been published in the Malaysian Journal of Medicine and Health Sciences (2017) and also presented at the International Conference of Translational Molecular Imaging and Aerospace Medicine & Physiology Showcase (2016) at KLIA, Sepang.

In Chapter 5 we highlight the discussion for the very first large scale study conducted to analyse the effects of BMI on semi-quantitative assessment of contrast-enhanced PET/CT scans. We also compare our results based on a multi-centre study involving one government-based centre that performs mainly contrast-enhanced PET/CT scans and the other a private medical centre that mainly performs plain low dose PET/CT scans. In this chapter, we also give a summary of this thesis as well as discuss significant findings from the study and directions for future works. Subsequently, there is a brief description of biographical data of the student.

REFERENCES

- [1] Boellaard R, O'Doherty MJ, Weber WA, Mottaghy FM, Lonsdale MN, Stroobants SG, et al. FDG PET and PET/CT: EANM procedure guidelines for tumour PET imaging: version 1.0. *Eur J Nucl Med Mol Imaging*. Springer-Verlag; 2010;37(1):181–200. Available from: <http://link.springer.com/10.1007/s00259-009-1297-4>
- [2] Malladi A, Viner M, Jackson T, Mercier G, Subramaniam RM. PET/CT mediastinal and liver FDG uptake: Effects of biological and procedural factors. *J Med Imaging Radiat Oncol*. 2013;57(2):169–75. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/23551774>
- [3] Kinahan PE, Fletcher JW. Positron emission tomography-computed tomography standardized uptake values in clinical practice and assessing response to therapy. *Semin Ultrasound CT MR*. NIH Public Access; 2010;31(6):496–505. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/21147377>
- [4] Kim WH, Kim CG, Kim D-W. Comparison of SUVs Normalized by Lean Body Mass Determined by CT with Those Normalized by Lean Body Mass Estimated by Predictive Equations in Normal Tissues. *Nucl Med Mol Imaging* Springer; 2012;46(3):182–8. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/24900058>
- [5] Tahari AK, Chien D, Azadi JR, Wahl RL. Optimum lean body formulation for correction of standardized uptake value in PET imaging. *J Nucl Med*. NIH Public Access; 2014;55(9):1481–4. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/24963129>
- [6] Narita A, Shiomi S, Katayama Y, Yamanaga T, Daisaki H, Hamada K, et al. Usefulness of standardized uptake value normalized by individual CT-based lean body mass in the application of PET response criteria in solid tumours (PERCIST). *Radiol Phys Technol*. 2016;9(2):170–7. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/26873140>
- [7] Erselcan T, Turgut B, Dogan D, Ozdemir S. Lean body mass-based standardized uptake value, derived from a predictive equation, might be misleading in PET studies. *Eur J Nucl Med Mol Imaging* [Internet]. Springer-Verlag; 2002;29(12):1630–8. Available from: <http://link.springer.com/10.1007/s00259-002-0974-3>
- [8] Yang WY, Burrows T, MacDonald-Wicks L, Williams LT, Collins CE, Chee WSS. The Family Diet Study: a cross-sectional study into the associations between diet, food habits and body weight status in Malay families. *J Hum Nutr Diet*. 2016;29(4):441–8. Available from: <http://doi.wiley.com/10.1111/jhn.12356>

- [9] Suppiah S, Chang W, Hassan H, Kaewput C, Asri AA, Saad FA, et al. Systematic review on the accuracy of positron emission tomography/computed tomography and positron emission tomography/magnetic resonance imaging in the management of ovarian cancer: Is functional information really needed? *World J Nucl Med.* 2017;16(3):176. Available from: <http://www.wjnm.org/text.asp?2017/16/3/176/207271>
- [10] Martinez-Outschoorn UE, Lin Z, Trimmer C, Flomenberg N, Wang C, Pavlides S, et al. Cancer cells metabolically "fertilize" the tumour microenvironment with hydrogen peroxide, driving the Warburg effect. *Cell Cycle.* Taylor & Francis; 2011;10(15):2504–20. Available from: <http://www.tandfonline.com/doi/abs/10.4161/cc.10.15.16585>
- [11] Sayre GA, Franc BL, Seo Y. Patient-Specific Method of Generating Parametric Maps of Patlak K(i) without Blood Sampling or Metabolite Correction: A Feasibility Study. *Int J Mol Imaging.* Hindawi Publishing Corporation; 2011 ;2011:185083. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/21912742>
- [12] Wahl RL, Jacene H, Kasamon Y, Lodge MA. From RECIST to PERCIST: Evolving Considerations for PET response criteria in solid tumours. *J Nucl Med [Internet].* NIH Public Access; 2009;50 Suppl 1(Suppl 1):122S–50S. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/19403881>
- [13] Massaro A, Cittadin S, Milan E, Tamiso L, Pavan L, Secchiero C, et al. Reliability of SUVmax vs. SUVmean in FDG PET/CT. *J Nucl Med [Internet].* Society of Nuclear Medicine; 2009 [cited 2017 Apr 2];50(supplement 2):2121–2121. Available from: http://jnm.snmjournals.org/content/50/supplement_2/2121
- [14] Pouya Ziai, Mohammad Reza Hayeri, Aliaksei Salei, Ali Salavati, Sina Houshmand, Abass Alavi, Oleg M. Teytelboym. Role of Optimal Quantification of FDG PET Imaging in the Clinical Practice of Radiology; *RadioGraphics* 2016; 36:481–496.
- [15] Delbeke D, Coleman RE, Guiberteau MJ, Brown ML, Royal HD, Siegel BA, et al. Procedure Guideline for Tumour Imaging with 18 F-FDG PET/CT 1.0*. [cited 2017 Apr 6]; Available from: http://www.zr3.de/tl_files/inhalte/files/download/Artikel_Soc_Nucl_Med_SNM.pdf
- [16] Cheson BD, Fisher RI, Barrington SF, Cavalli F, Schwartz LH, Zucca E, et al. Recommendations for initial evaluation, staging, and response assessment of hodgkin and non-hodgkin lymphoma: The lugano classification. *J Clin Oncol.* 2014;32(27):3059–67.

- [17] Oliver Wong CY, Salem R, Qing F, Wong KT, Barker D, Gates V, Lewandowski R, Elizabeth AH, Dworkin HJ, and Nagle C (2004) Metabolic Response After Intraarterial 90Y-Glass Microsphere Treatment for Colorectal Liver Metastases: Comparison of Quantitative and Visual Analyses by 18F-FDG PET. *The Journal of Nuclear Medicine*. 45(11).
- [18] Lindholm H, Grybäck P, Alej, Sánchez-Crespo ro, Brolin F, Jacobsson H. The FDG-Uptake of Adipose Tissue is Higher in Individuals with Increased Blood Glucose Levels than in Individuals with Normal Levels. *J Nucl Med Radiat Ther* [Internet]. OMICS International; 2014 [cited 2017 Apr 2];5(1). Available from: <http://www.omicsonline.org/open-access/the-fdguptake-of-adipose-tissue-is-higher-in-individuals-with-increased-blood-glucose-levels-than-in-individuals-with-normal-levels-2155-9619.1000167.php?aid=24622>
- [19] Plaxton N, Moncayo V, Barron B, Halkar R. Factors that influence standard uptake values in FDG PET/CT. *J Nucl Med* [Internet]. Society of Nuclear Medicine; 2014 [cited 2017 Apr 2];55(supplement 1):1356–1356. Available from: http://jnm.snmjournals.org/content/55/supplement_1/1356
- [20] Schöder H, Erdi YE, Chao K, Gonen M, Larson SM, Yeung HWD. Clinical implications of different image reconstruction parameters for interpretation of whole-body PET studies in cancer patients. *J Nucl Med* [Internet]. 2004 Apr [cited 2017 Apr 2];45(4):559–66. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/15073250>
- [21] Wiyaporn K, Wiyaporn Msc K, Msc CT, Pusuwan P, Msc TE, Msc SL, et al. Factors Affecting Standardized Uptake Value (SUV) of Positron Emission Tomography (PET) Imaging with 18 F-FDG. *J Med Assoc Thai* [Internet]. 2010 [cited 2017 Apr 2];93(1). Available from: http://www.si.mahidol.ac.th/th/publication/2010/Vol93_No.1_108_5814.pdf
- [22] Jaskowiak CJ, Bianco JA, Perlman SB, Fine JP. Influence of reconstruction iterations on 18F-FDG PET/CT standardized uptake values. *J Nucl Med* [Internet]. Society of Nuclear Medicine; 2005 Mar [cited 2017 Apr 2];46(3):424–8. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/15750154>
- [23] Adams MC, Turkington TG, Wilson JM, Wong TZ. A Systematic Review of the Factors Affecting Accuracy of SUV Measurements. *Am J Roentgenol* [Internet]. American Roentgen Ray Society; 2010 Aug [cited 2017 Apr 2];195(2):310–20. Available from: <http://www.ajronline.org/doi/10.2214/AJR.10.4923>

- [24] Sugawara Y, Zasadny KR, Neuhoff AW, Wahl RL. Reevaluation of the Standardized Uptake Value for FDG: Variations with Body Weight and Methods for Correction. Radiology [Internet]. 1999 Nov [cited 2017 Apr 2];213(2):521–5. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/10551235>
- [25] Tahari AK, Paidpally V, Chirindel A, Wahl RL, Subramaniam RM. Two-Time-Point FDG PET/CT: Liver SUL_{mean} Repeatability. Am J Roentgenol [Internet]. American Roentgen Ray Society ; 2015 Feb [cited 2017 Apr 2];204(2):402–7. Available from: <http://www.ajronline.org/doi/10.2214/AJR.14.12719>
- [26] Fukukita H, Senda M, Terauchi T, Suzuki K, Daisaki H, Matsumoto K, et al. Japanese guideline for the oncology FDG-PET/CT data acquisition protocol: synopsis of Version 1.0. Ann Nucl Med [Internet]. 2010 May 17 [cited 2017 Apr 3];24(4):325–34. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/20401547>
- [27] Paquet N, Albert A, Foidart J, Hustinx R. Within-Patient Variability of 18 F-FDG : Standardized Uptake Values in Normal Tissues. 2004;45(5):784–9.
- [28] Mitumoto T, Taguchi Y, Minamimoto R, Okasaki M, Morooka M, Kubota K, et al. Validation of SUV body weight (SUVbw) vs SUV lean body mass (SUVlbm): The evaluation with each organ of the healthy subjects. J Nucl Med. 2012;53(2626):5–6.
- [29] Kuruva M, Mittal BR, Bhattacharya A, Ydoxhv FDQD, Uhjuhvvlrq O, Vlv D, et al. Multivariate analysis of various factors affecting background liver and mediastinal standardized uptake values Abstract Purpose of the Study : Materials and Methods : Results : Conclusions : Indian J Nucl Med. 2012;3.
- [30] Viner M. Liver SUL mean at FDG PET / CT : Interreader Agreement and Impact of. 2013;267(2).
- [31] Jochimsen TH, Schulz J, Busse H. Lean body mass correction of standardized uptake value in simultaneous whole-body positron emission tomography and magnetic resonance imaging. physic Med Biol. 2013;4651.
- [32] Kono Y, Utsunomiya K, Tanigawa N, Ha-Kawa SK, Ueno Y. Evaluation of lean body mass normalized standard uptake values in PET studies using a predictive equation. J Nucl Med. 2015;56(supplement 3):1787.

- [33] Batallés S, Villavicencio R, Quaranta A, Burgos L, Trezzo S, Staffieri R, et al. Variations of the hepatic SUV in relation to the body mass index in whole body PET-CT studies. *Rev Esp Med Nucl Imagen Mol* [Internet]. 2013 [cited 2017 Apr 2];32(1):26–32. Available from: file:///C:/Users/Dell/Documents/Battales 2012. Linear regression analysis for factors influencing SUV and SUL.pdf
- [34] Abele JT, Fung CI. Effect of hepatic steatosis on liver FDG uptake measured in mean standard uptake values. *Radiology*. 2010;254(3):917–24.
- [35] Han SS, Kim KW, Kim K-I, Na KY, Chae D-W, Kim S, et al. Lean Mass Index: A Better Predictor of Mortality than Body Mass Index in Elderly Asians. *J Am Geriatr Soc* [Internet]. 2010 Feb [cited 2017 Apr 6];58(2):312–7. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/20070416>
- [36] Zasadny KR, Wahl RL. Standardized uptake values of normal tissues at PET with 2-[fluorine-18]-fluoro-2-deoxy-D-glucose: variations with body weight and a method for correction. *Radiology* [Internet]. 1993 Dec [cited 2017 Apr 4];189(3):847–50. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/8234714>
- [37] van Helden EJ, Hoekstra OS, Boellaard R, Roth C, Mulder ER, Verheul HMW, et al. Early 18F-FDG PET/CT Evaluation Shows Heterogeneous Metabolic Responses to Anti-EGFR Therapy in Patients with Metastatic Colorectal Cancer. Rosell R, editor. *PLoS One* [Internet]. Public Library of Science; 2016 May 19 [cited 2017 Apr 2];11(5):e0155178. Available from: <http://dx.plos.org/10.1371/journal.pone.0155178>
- [38] Foster BJ, Platt RW, Zemel BS. Development and Validation of a Predictive Equation for Lean Body Mass in Children and Adolescents. *Ann Hum Biol* [Internet]. Taylor & Francis; 2012 May 23 [cited 2017 Apr 6];39(3):171–82. Available from: <http://www.tandfonline.com/doi/full/10.3109/03014460.2012.681800>
- [39] James W. Research on obesity. London: Her Majesty's Stationary Office; 1976.
- [40] Jackson LB, Henshaw MH, Carter J, Chowdhury SM. Sex-specific lean body mass predictive equations are accurate in the obese paediatric population. *Ann Hum Biol* [Internet]. 2016 Sep 2 [cited 2017 Apr 6];43(5):417–22. Available from: <http://www.tandfonline.com/doi/full/10.3109/03014460.2015.1069893>

- [41] Kim CG, Kim WH, Kim MH, Kim D-W. Direct Determination of Lean Body Mass by CT in F-18 FDG PET/CT Studies: Comparison with Estimates Using Predictive Equations. *Nucl Med Mol Imaging* (2010) [Internet]. 2013 Jun 7 [cited 2017 Apr 6];47(2):98–103. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/24900089>
- [42] Graham MM, Badawi RD, Wahl RL. Variations in PET/CT methodology for oncologic imaging at U.S. academic medical centers: an imaging response assessment team survey. *J Nucl Med* [Internet]. NIH Public Access; 2011 Feb [cited 2017 Apr 2];52(2):311–7. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/21233185>
- [43] WHO International Obesity Task Force. The Asia-Pacific Perspective: Redefining Obesity and its Treatment [Internet]. 1st ed. Inoue S, Zimmet P, editors. Western Pacific Region: Heath Communications Australia Pty Ltd; 2000 [cited 2017 Apr 3]. 1-50 p. Available from: <http://www.wpro.who.int/nutrition/documents/docs/Redefiningobesity.pdf>
- [44] Hamill JJ, Sunderland JJ, LeBlanc AK, Kojima CJ, Wall J, Martin EB. Evaluation of CT-based lean-body SUV. *Med Phys* [Internet]. 2013 Aug 13 [cited 2017 Apr 3];40(9):92504. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/24007180>
- [45] Janmahasatian S, Duffull SB, Chagnac A, Kirkpatrick CMJ, Green B. Lean body mass normalizes the effect of obesity on renal function. *Br J Clin Pharmacol*. 2008;65(6):964–5.
- [46] Groheux D, Espié M, Giacchetti S, Hindié E. Performance of FDG PET/CT in the Clinical Management of Breast Cancer. *Radiology* [Internet]. Radiological Society of North America, Inc.; 2013 Feb [cited 2017 Apr 17];266(2):388–405. Available from: <http://pubs.rsna.org/doi/10.1148/radiol.12110853>
- [47] Vinjamuri S, Ray S. Added value of PET and PET–CT in oesophageal cancer: a review of current practice. *Nucl Med Commun* [Internet]. 2008 Jan [cited 2017 Apr 7];29(1):4–10. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/18049091>
- [48] Suppiah S, Fathinul Fikri AS, Mohad Azmi NH, Nordin AJ. Mapping 18F-Fluorodeoxyglucose metabolism using PET/CT for the assessment of treatment response in Non-Small Cell Lung Cancer patients undergoing Epidermal Growth Factor Receptor inhibitor treatment: A single-centre experience. *Malaysian J Med Heal Sci*. 2017;13(1):23–30.

- [49] Oprea-Lager DE, Vincent AD, van Moorselaar RJA, Gerritsen WR, van den Eertwegh AJM, Eriksson J, et al. Dual-Phase PET-CT to Differentiate [18F]Fluoromethylcholine Uptake in Reactive and Malignant Lymph Nodes in Patients with Prostate Cancer. Rao J, editor. PLoS One [Internet]. Public Library of Science; 2012 Oct 31 [cited 2017 Apr 6];7(10):e48430. Available from: <http://dx.plos.org/10.1371/journal.pone.0048430>
- [50] Delbeke D, P V, J S, C H, M A, ME P. Evaluation of Benign vs Malignant Hepatic Lesions With Positron Emission Tomography. Arch Surg [Internet]. American Medical Association; 1998 May 1 [cited 2017 Apr 17];133(5):510. Available from: <http://archsurg.jamanetwork.com/article.aspx?doi=10.1001/archsurg.133.5.510>
- [51] Hofheinz F, Bütof R, Apostolova I, Zöphel K, Steffen IG, Amthauer H, et al. An investigation of the relation between tumour-to-liver ratio (TLR) and tumour-to-blood standard uptake ratio (SUR) in oncological FDG PET. EJNMMI Res [Internet]. 2011;619(6). Available from: <http://download.springer.com/static/pdf/604/art%253A10.1186%252Fs13550-016-0174-y.pdf?originUrl=http%3A%2F%2Fejnmires.springeropen.com%2Farticle%2F10.1186%2Fs13550-016-0174-y&token2=exp=1492411724~acl=%2Fstatic%2Fpdf%2F604%2Fart%25253A10.1186%25252Fs1355>
- [52] Verburg FA, Kuhl CK, Pietsch H, Palmowski M, Mottaghy FM, Behrendt FF. The influence of different contrast medium concentrations and injection protocols on quantitative and clinical assessment of FDG-PET/CT in lung cancer. Eur J Radiol [Internet]. 2013 Oct [cited 2017 May 3];82(10):e617–22. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/23880426>
- [53] Kim WH, Kim CG, Kim D-W. Comparison of SUVs Normalized by Lean Body Mass Determined by CT with Those Normalized by Lean Body Mass Estimated by Predictive Equations in Normal Tissues. Nucl Med Mol Imaging (2010) [Internet]. 2012 Sep 21 [cited 2017 Apr 3];46(3):182–8. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/24900058>
- [54] Segreto S, Fonti R, Ottaviano M, Pellegrino S, Pace L, Damiano V, et al. Evaluation of metabolic response with 18 F- FDG PET-CT in patients with advanced or recurrent thymic epithelial tumours. Cancer Imaging [Internet]. 2017;17(10):1–8. Available from: <http://download.springer.com/static/pdf/213/art%253A10.1186%252Fs40644-017-0112-x.pdf?originUrl=http%3A%2F%2Fcancerimagingjournal>.

biomedcentral.com%2Farticle%2F10.1186%2Fs40644-017-0112-x&token2=exp=1492412722~acl=%2Fstatic%2Fpdf%2F213%2Fart%25253A10.1186

- [55] Nakamoto Y, Chin BB, Kraitchman DL, Lawler LP, Marshall LT, Wahl RL. Effects of Nonionic Intravenous Contrast Agents at PET/CT Imaging: Phantom and Canine Studies. *Radiology* [Internet]. Radiological Society of North America ; 2003 Jun [cited 2017 Apr 17];227(3):817–24. Available from: <http://pubs.rsna.org/doi/10.1148/radiol.2273020299>
- [56] Cronin CG, Prakash P, Blake MA. Oral and IV Contrast Agents for the CT Portion of PET/CT. *Am J Roentgenol* [Internet]. American Roentgen Ray Society; 2010 Jul [cited 2017 Apr 17];195(1):W5–13. Available from: <http://www.ajronline.org/doi/10.2214/AJR.09.3844>
- [57] Ku CR, Lee N, Hong JW, Kwon IG, Hyung WJ, Noh SH, et al. Intestinal Glycolysis Visualized by FDG PET/CT Correlates with Glucose Decrement After Gastrectomy. *Diabetes* [Internet]. 2016 [cited 2017 Apr 27]; Available from: <http://diabetes.diabetesjournals.org/content/early/2016/11/30/db16-1000>
- [58] Groheux D, Delord M, Rubello D, Colletti PM, Nguyen M-L, Hindié E. Variation of Liver SUV on 18FDG-PET/CT Studies in Women With Breast Cancer. *Clin Nucl Med* [Internet]. 2013 Jun [cited 2017 Apr 27];38(6):422–5. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/23510894>
- [59] Park J, Chang KJ, Seo YS, Byun BH, Choi JH, Moon H, et al. Tumour SUVmax Normalized to Liver Uptake on (18)F-FDG PET/CT Predicts the Pathologic Complete Response After Neoadjuvant Chemoradiotherapy in Locally Advanced Rectal Cancer. *Nucl Med Mol Imaging* (2010) [Internet]. Springer; 2014 Dec [cited 2017 Apr 27];48(4):295–302. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/26396634>
- [60] Devriese J, Beels L, Maes A, Van de Wiele C, Pottel H. Evaluation of CT-based SUV normalization. *Phys Med Biol* [Internet]. IOP Publishing; 2016 Sep 7 [cited 2017 Apr 27];61(17):6369–83. Available from: <http://stacks.iop.org/0031-9155/61/i=17/a=6369?key=crossref.761feda4ca4b9626f090070c2a2f8148>
- [61] Park JK, Kim SK, Cho IH, Kong EJ. Measurement of SUVs-maximum for normal region using VOI in PET/MRI and PET/CT. *ScientificWorldJournal* [Internet]. Hindawi Publishing Corporation; 2014 [cited 2017 Apr 27];2014:194925. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/24672297>

- [62] Chirindel A, Alluri KC, Tahari AK, Chaudhry M, Wahl RL, Lodge MA, et al. Liver standardized uptake value corrected for lean body mass at FDG PET/CT: Effect of FDG uptake time. Clin Nucl Med [Internet]. 2015;40(1):e17–22. Available from: <http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L600782541>
- [63] Munro BH (2000) Statistical Methods for Health Care Research (4th Edition). Lippincott Williams & Wilkins: New York.
- [64] Bullmer, MG (1979), Principles of Statistics. NY: Dover Books on Mathematics.
- [65] Kevin P. Balanda and HL MacGillivray, " Kurtosis: A Critical Review". The American Statistician 42:2[May 1988]. Pp111-119.
- [66] Toothaker and Larry E (1993) Multiple Comparison Procedures(Quantitative Applications in the Social Sciences) (2nd ed). Newbury Park, CA: Chapman and Hall/CRC.pp.27-45.ISBN0-803-94177-3.