

Ultrasonographic Measurement of Skin and Subcutaneous Thickness at Insulin Injection Sites among Adult Filipinos with Diabetes

Irene Catambing and Michael Villa

Section of Endocrinology and Metabolism, Department of Internal Medicine, St. Luke's Medical Center, Quezon City, Philippines

Abstract

Objectives. To determine the skin and subcutaneous thickness of adult diabetic Filipinos using ultrasonography.

Methodology. We studied 293 Filipino diabetic adults who had ultrasonographic measurements of their skin thickness (ST) and subcutaneous thickness (SCT) at common insulin injection sites.

Results. The mean ST ranges from 1.76 mm to 2.75mm. The mean SCT ranges from 6.91 mm to 19.1 mm. The anterior thigh area has the thinnest mean ST and SCT. On the other hand, the buttocks have the thickest mean ST and area of abdomen has the thickest mean SCT.

On multiple regression analysis, the predictors for skin thickness are injection site, age, BMI, gender and insulin use, however the overall influence of all these factors on skin thickness is variable at best.

Conclusion. Skin thickness among Filipinos varies marginally depending on injection site being thickest at the buttocks and thinnest at the thighs. On the other hand, there is greater variability in SCT depending on the injection site, being thickest at the abdominal area and thinnest at the anterior thigh.

Key words: *diabetes mellitus, subcutaneous injection, skin thickness, subcutaneous thickness*

INTRODUCTION

The proper insulin injection technique is an established factor in optimizing blood sugar control among diabetic patients. Several studies suggest that injection technique is of similar importance to the type and dose of insulin delivered to attain good glycemic control.¹ Other factors that affect the pharmacokinetics/pharmacodynamics of insulin include: patient's age, gender, body mass index (BMI), dose volume, insulin formulation, insulin antibodies and individual patient response differences, as well as variations in injection technique. Injection technique includes factors intended to facilitate the most consistent, least painful delivery of insulin into subcutaneous tissues (SC) and includes choice of injection site and needle length, angle of needle insertion and the use of lifted skin fold. Errors in insulin injection technique curtails the ability of many patients to attain glycemic goals.²

The pharmacokinetic behavior of injected insulin varies with the nature of the tissue into which it is deposited, whether intradermal, or through fat, muscle or the peritoneal cavity, and is reflected in the different rates of appearance of insulin in the blood and ultimately, in blood glucose control.³

Studies indicate that the absorption of insulin is altered in intramuscular injection compared to true subcutaneous placement. These provide evidence that if insulin is injected accidentally and intermittently into the muscle, this will contribute to day to day variability in the control of blood glucose.³

Commonly used anatomical sites for SC insulin injections include the upper arm (triceps areas), antero-lateral upper thigh, abdomen (with the exception of a circle with a 2 inch radius around the navel) and the buttocks, typically the upper outer quadrant. Site selection should take into consideration the variable absorption between sites.

The difference in the rate of absorption at various anatomical sites is attributed to the number of blood capillaries and/or their permeability, as well as differing blood flow.^{4,5}

Ultrasound examination of the skin for experimental and clinical use has been recognized since the early 1990's. In principle, the high frequency ultrasound wave (over 10 MHz) is emitted off from a transducer, becomes reflected in the skin from the interfaces between media of different acoustic properties, and returns back to the probe. The time lag between the emitted and the reflected signal enables the calculation of the distance between reflecting

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*Corresponding author: Irene C. Catambing, MD
Diabetes, Thyroid and Endocrine Center
12th floor, Cathedral Heights Building
St. Luke's Medical Center
#279 E. Rodriguez Blvd., Quezon City, Philippines
Tel/Fax No.: +632-7230101 local 5210
E-mail: catambing.md@gmail.com*

objects, whereas the amount of reflected energy characterizes the echogenicity of a given object.⁶

Two dimensional scanning, called B-mode scanning, is predominantly used, and in this mode, a two dimensional, cross sectional skin image is obtained. The first reflectant layer in the skin is the epidermis, the echogenicity of which depends mainly on the thickness of stratum corneum and the amount of air contained between the keratotic scales. Dermal echoes, however, are many and variable. Dermal echogenicity depends on the architecture of skin fiber network and on the amount of contained water.⁶

The reproducibility of ultrasound technique for the determination of skin thickness was already investigated by Tan et al, as early as 1980.⁷ The study involved 2 independent observers who measured the skin thickness on the flexor aspect of both mid-forearms of 20 subjects. Each observer took five readings from each site. The study demonstrated that skin thickness measurement determined by ultrasound was highly reproducible with only small variability between observers.⁷

The thickness of SC tissues in the adult patient varies widely by gender, body site and body mass index, whereas the thickness of the skin (epidermis and dermis) is quite constant, averaging approximately 1.9-2.4 mm across injection sites, ages, races, BMI and gender; it is rarely >3.0 mm at injection sites for insulin.⁶ This was the same conclusion by Gibney et al., who conducted a study on the skin and subcutaneous layer thickness on 388 diabetic subjects, 16% of which were Asians living in the United States.² To date, there is no published normative data on diabetic Filipinos living in the Philippines.

GENERAL OBJECTIVE

To determine the skin and subcutaneous thickness of adult Filipinos with diabetes mellitus using ultrasonography, and ascertain its implications for needle length recommendation.

SPECIFIC OBJECTIVES

1. To determine the average skin and subcutaneous thickness of adult Filipinos with diabetes mellitus using ultrasonography
2. To determine the effect on skin and subcutaneous tissue of factors such as body mass index, gender, age, injection site and insulin use

METHODOLOGY

This is a cross-sectional analytic study. The study enrolled adult patients with diabetes mellitus from the outpatient department of the Section of Endocrinology, Diabetes and Metabolism. The measurements were conducted at the

Diabetes, Thyroid and Endocrine Center from July 2012 to November 2012.

Participants of the study included: Adult patients (>19 years old), diagnosed with type 1 or type 2 diabetes mellitus. The following participants were excluded: Pregnant patients or those with any pathologic condition which might affect the distribution of or thickness of SC tissue such as: Cushing's syndrome, generalized edema, soft tissue inflammation (at any site of measurement), and presence of ascites.

The following data were included in the study documentations: the subjects' general information (e.g., name, age, sex) brief medical history: past illnesses, present illnesses, duration of diabetes and presence of complications as self reported by the patients as well as their current diabetes regimen). The subjects' weight and height were measured on the day of sonographic measurement.

Ultrasound was selected to measure the skin thickness and subcutaneous thickness due to its utility, safety and reproducibility. The SonoSite M-Turbo portable ultrasound unit with a 13-6 MHz, 38 mm broadband linear array transducer probe (SonoSite Ltd.) was utilized for the study. A single ultrasound technician trained and skilled with the features and operation of the machine performed all the sonographic procedures to minimize operational bias. Ultrasound gel was applied, the probe was placed perpendicularly to the predetermined sites (see below). As necessary, the probe may be moved within areas of interest to obtain clear images. Skin thickness and subcutaneous thickness were measured. Three frames and measurement per site were made. The average of the three measurements were taken and used.

A radiosonologist independently reviewed the images for appraisal and verification of the images and values.

Skin and subcutaneous thickness at the common sites for insulin injection were measured. Specific locations for measurements based on bony and other superficial body landmarks were used to reduce inter-subject measurement variability. The skin and subcutaneous thickness was measured bilaterally with the exception of the abdominal area at the following sites:

- a) Posterior upper arm: deltoid area, mid-distance between the acromion and olecranon processes
- b) Anterior upper thigh: quadriceps area, mid-distance between the anterior superior iliac spine and the top edge of the patella
- c) Anterior abdomen: mid-distance between the umbilicus and the anterior superior iliac spine
- d) Upper buttocks: mid-distance between the inter-trochanteric spine and the tip of the coccyx

The ultrasound technician measured the sites following a specific sequence to reduce positional and technical

variability: right upper arm, left upper arm, abdomen, right thigh, left thigh, left buttocks then finally right buttocks.

STATISTICAL METHODS

Sample size was estimated by the following: an estimated effect size (f2) of 0.15, with desired power of 80%, 6 predictors and probability level of 0.05. A minimum sample size of 97 was calculated. (123 if 90% power)

Descriptive statistics for skin thickness (ST) and subcutaneous adipose layer thickness (SCT) include graphical display, mean, median, standard deviation, minimum and maximum, and are provided for the entire population, and for each BMI subgroup and by gender. One way ANOVA with Bonferroni post hoc analysis was done to determine statistical difference in the mean ST and SCT by injection site. Multivariate analyses were done to better weigh the statistical and clinical impact of the various inputs such as injection site, gender, age, BMI and insulin use for both ST and SCT. The threshold for statistical significance is $\alpha = 0.05$.

RESULTS

A total of 293 subjects participated in the study. Table 1 shows the demographic and clinical data of the subjects. Majority of the participants (67%) were females, with a mean age of 58.57 ± 10.09 years. Half of the participants were elderly, belonging to the 60 to 85 year old age group. Based on the WHO BMI category, almost half of the study population will fall within the normal BMI range (<25), 38% were overweight (BMI 25- 29.9), while 13% were obese. However, if we apply the Asia Pacific BMI classification for overweight and obesity, the majority of the patients will fall within the overweight and obese class I category (24.5% and 38.6% respectively). On average, study participants had been diagnosed with diabetes for 11 years.

In this study, ultrasound measurements for ST and/or for SCT were obtained for a total of 293 subjects. However, there were some participants with no ST and/or SCT measurement in some injection sites. These included patients who opted not to have their buttocks and thighs measured due to modesty and/or restrictive clothing. One subject was afflicted with polio and the investigators opted not to have his buttocks and thighs measured.

Table 3 shows the mean (SD) skin thickness by injection site. The results show that the thighs have the lowest mean skin thickness measurement at 1.76 mm while the buttocks have the highest mean skin thickness measurement at 2.75 mm. One way analysis of variance shows that there is a statistically significant difference in the mean skin thickness measurement between the different sites of injection. Bonferroni post hoc analysis showed that mean

skin thickness measurement of the buttocks significantly differs from the mean skin thickness of the abdomen, arm, and thigh, while the mean skin thickness of the abdomen and the arm are not significantly different.

Table 1. Demographic characteristics of the entire study population (n = 293)

Number with measurements	Skin thickness	
	• Upper arm (R/L)	291/291
	• Abdomen	291
	• Thigh (R/L)	284/283
	• Buttocks (R/L)	282/281
	Subcutaneous adipose layer thickness	
	• Upper arm (R/L)	291/291
	• Abdomen	291
	• Thigh (R/L)	283/282
	• Buttocks (R/L)	279/277
Gender	Males (%)	98 (33.4)
	Females (%)	195 (66.6)
Age (years)	Mean (SD)	58.57 (10.09)
	Min/Max	29/84
Age category (years)	18 – 39 (%)	13 (4.5)
	40 – 59 (%)	133 (45.9)
	60 – 85 (%)	144 (49.7)
Insulin	Yes (%)	130 (44.8)
	No (%)	160 (55.2)
BMI (kg/m2)	Mean (SD)	25.79 (4.29)
	Min/Max	18.00/41.70
BMI category (WHO)	< 25 (%)	138 (47.6)
	25 – 29.9 (%)	114(38.9)
	≥ 30 (%)	38 (13.1)
(Asia Pacific Category)	Underweight < 18.5	1 (0.3)
	Normal 18.5 to 22.9	66 (22.8)
	Overweight/at risk 23.0 to 24.9	71 (24.5)
	Obese I 25.0 to 29.9	112 (38.6)
	Obese II >30.0	40 (13.8)
Number of years diagnosed with diabetes	Mean (SD)	10.74 (7.63)
	Min/Max	0/45
Proportion of participants with complications	Retinopathy, n (%)	162 (71.1)
	Nephropathy, n (%)	117 (51.3)
	Coronary Artery disease, n (%)	27 (11.8)
	Cerebrovascular disease, n (%)	34 (14.9)

Table 2. Demographic profile of the study participants by gender

Characteristic	Male, n = 98	Female, n = 195
Age (years)		
• Mean (SD)	56.39 (10.61)	59.66 (9.66)
• Min/Max	33/83	29/84
Age category		
• 18 – 39 (%)	7 (7.2)	6 (3.1)
• 40 – 59 (%)	54 (55.7)	79 (40.5)
• 60 – 85 (%)	36 (37.1)	108 (55.4)
Insulin use, n (%)	46 (46.9)	84 (43.8)
BMI		
• Mean (SD)	25.54 (4.03)	25.92 (4.42)
• Min/Max	18.09/37.40	18.00/41.70
BMI category		
• < 25 (%)	42 (42.9)	96 (50.0)
• 25 – 29.9 (%)	48 (49.0)	66 (34.4)
• ≥ 30 (%)	8 (8.2)	30 (15.6)
Underweight < 18.5	0 (0)	1 (0.5)
Normal 18.5 to 22.9	25 (25.5)	41 (21.4)
Overweight/at risk 23.0 to 24.9	17 (17.3)	54 (28.1)
Obese I 25.0 to 29.9	48 (49.0)	64 (33.3)
Obese II >30.0	8 (8.2)	32 (16.7)
Number of years diagnosed with DM		
• Mean (SD)	10.20 (7.83)	11.02 (7.53)
• Min/Max	1/35	0/45
Proportion of participants with complications		
• Retinopathy, n (%)	38 (64.4)	124 (73.4)
• Nephropathy, n (%)	40 (67.8)	77 (45.6)
• Coronary Artery disease, n (%)	7 (11.9)	20 (11.8)
• Cerebrovascular disease, n (%)	13 (22.0)	21 (12.4)

Table 3. Skin thickness (mm) by injection site

Site	N	Mean (SD)	95% CI	One-way ANOVA p value
Arm	291	2.12 (0.45)	2.07 – 2.18	0.000
Abdomen	291	2.25 (0.46)	2.20 – 2.31	
Thigh	284	1.76 (0.33)	1.72 – 1.80	
Buttock	281	2.75 (0.52)	2.69 – 2.82	

Figure 1 demonstrates that there is little variation in the arm skin thickness among patients of different BMI categories. Similarly, Figure 2 also demonstrates little variation in the skin thickness of the abdomen among these groups of patients.

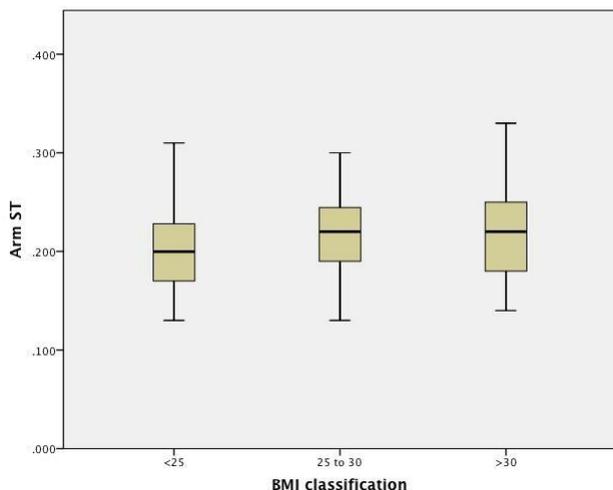


Figure 1. Arm skin thickness using WHO BMI category

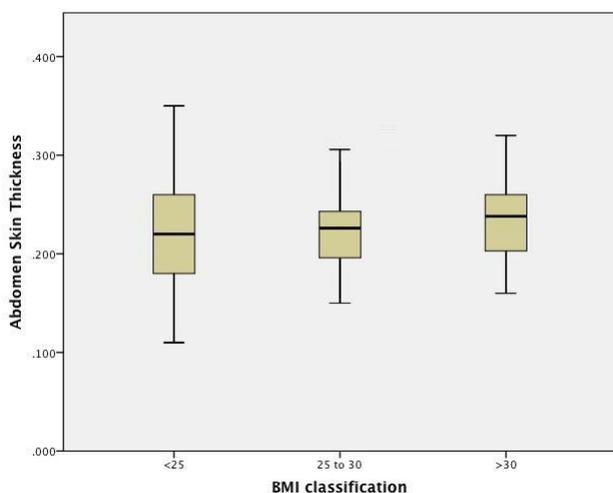


Figure 2. Abdomen skin thickness using WHO BMI category

Figure 3 demonstrates an apparent trend of increasing thigh skin thickness as the BMI increases. On the other hand, Figure 4 demonstrates that there is little variation in the buttock skin thickness among patients of different BMI categories

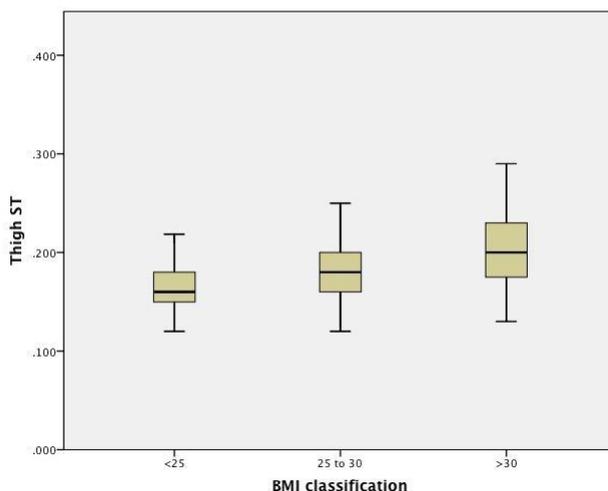


Figure 3. Thigh skin thickness using WHO BMI category

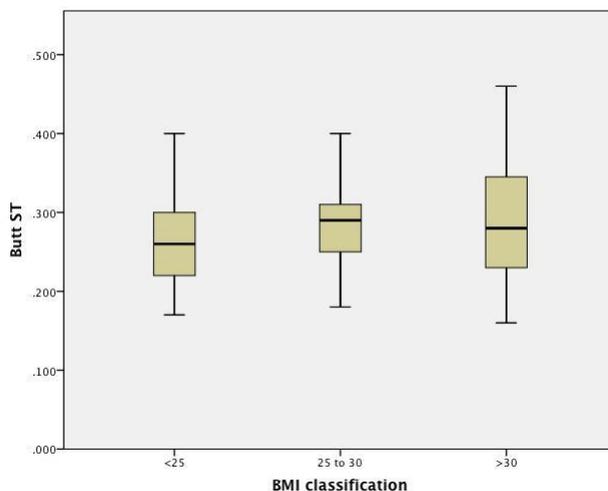


Figure 4. Buttock skin thickness using WHO BMI category

Table 4 shows the mean (SD) subcutaneous adipose layer thickness by injection site. It can be seen that the thighs have the lowest mean subcutaneous adipose layer measurement (6.91 ± 3.48mm), while the buttocks have the highest mean subcutaneous adipose layer thickness measurement at 12.91mm (5.42).

Table 4. Subcutaneous adipose layer thickness (mm) by injection site

Site	N	Mean (SD)	95% CI	One-way ANOVA p value
Arm	291	8.04 (3.82)	7.60 – 8.49	0.000
Abdomen	291	19.1 (1.264)	1.76 – 2.056	
Thigh	283	6.91 (3.48)	6.50 – 7.32	
Buttock	276	12.91 (5.42)	12.27 – 13.55	

One way analysis of variance shows that there is a statistically significant difference in the mean SCT measurement according to site of injection. Bonferroni post hoc analysis showed that mean SCT measurement of the buttocks is significantly different from the mean skin thickness of the abdomen, arm, and thigh, while the mean

SCT of the thigh and the arm are not significantly different.

Figure 5 demonstrates that there is little variation in the mean arm subcutaneous adipose tissue thickness among patients of different BMI categories, however there is a wide confidence interval seen among obese patients . Similarly, Figure 6 demonstrates that there is little variation in the mean abdominal subcutaneous adipose tissue thickness among patients of different BMI categories, however a wide confidence interval is likewise seen among obese patients.

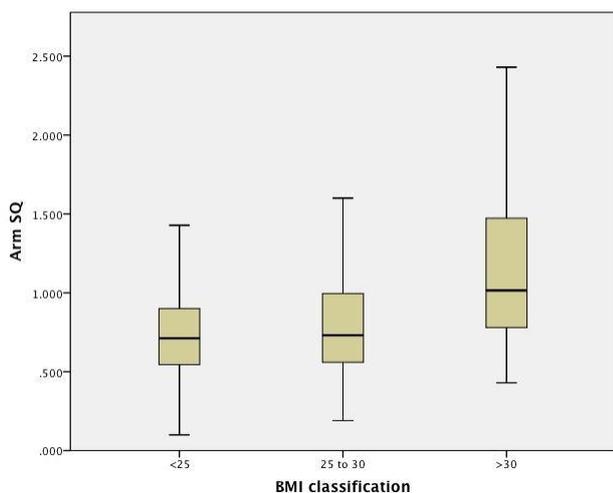


Figure 5 . Arm SCT according to WHO BMI category

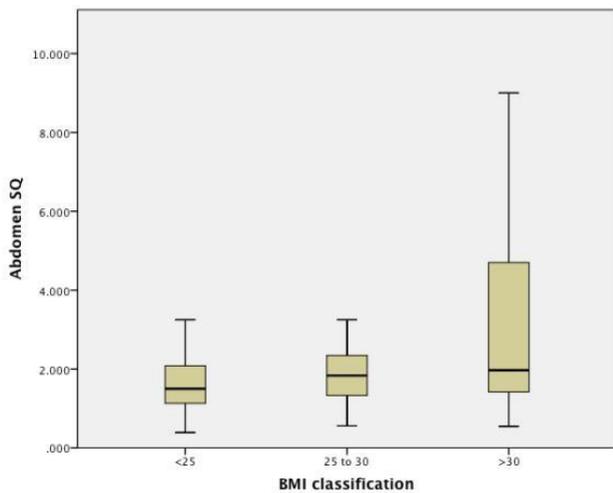


Figure 6. Abdomen SCT according to WHO BMI category

Figure 7 demonstrates that there is little variation in the mean thigh subcutaneous adipose tissue thickness among patients of different BMI categories, however a wider confidence interval is again seen among obese patients.

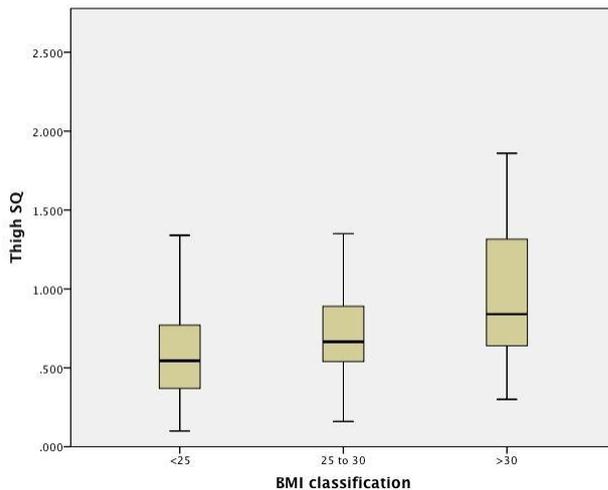


Figure 7. Thigh SCT according to WHO BMI category

Figure 8 demonstrates that there is little variation in the mean buttocks subcutaneous adipose tissue thickness among patients of different BMI categories.

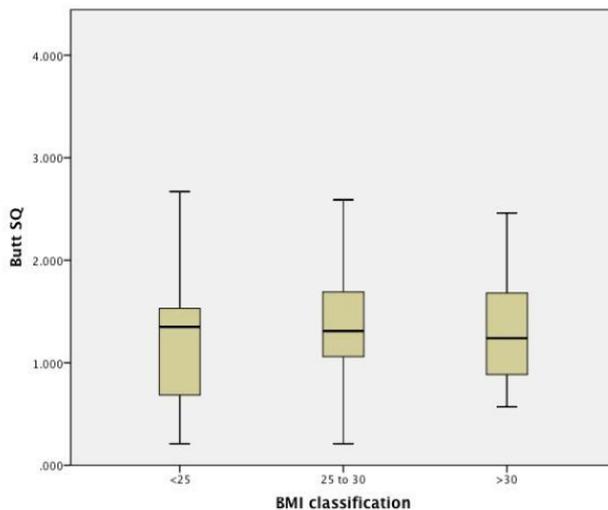


Figure 8. Buttock SCT according to WHO BMI category

Utilizing multiple regression analysis to predict skin thickness, we included injection site as the main predictor variable, and injection site, age, gender, years diabetic, BMI, years diabetic and insulin use were considered as confounders. The final model (see Table 5) shows that all these variables are significant predictors of skin thickness. The adjusted R² of 0.487 indicates that this model predicts 48.7% of the variance in skin thickness is due to the above model.

Table 5. Multiple regression results for predicting skin thickness (final model)

Predictor variable	β coefficient	Standardized β coefficient	p value	R ²	Adjusted R ²
<u>Skin thickness</u>					
(Constant)	0.316		0.000	0.490	0.487
Injection site	-0.031	-0.612	0.000		
Age in years	-0.001	-0.186	0.000		
Gender	0.027	0.226	0.000		
BMI	0.001	0.109	0.000		
Years diabetic	0.000	0.055	0.025		
Insulin use	-0.008	-0.067	0.005		

The remaining 51.3% is due to individual variation and might be explained by other factors that were not taken into account in the analysis or that the sample size is inadequate.

To determine the factors affecting subcutaneous thickness, we utilized multiple regression analysis and again included injection site as the main predictor variable, and injection site, age, gender, years diabetic, BMI, years diabetic and insulin use were considered as confounders. The final model (see Table 6) shows that the significant predictors of subcutaneous adipose layer thickness are injection site, age in years, BMI, and gender. Years diabetic and insulin use are not significant predictors of subcutaneous adipose layer thickness.

Table 6. Multiple regression results for predicting subcutaneous adipose layer thickness (final model)

Predictor variable	β coefficient	Standardized β coefficient	p value	R ²	Adjusted R ²
<u>Subcutaneous adipose layer thickness</u>					
(Constant)	1.000		0.000	0.469	0.215
Injection site	-0.297	-0.370	0.000		
Age in years	-0.006	-0.071	0.013		
BMI	0.053	0.256	0.000		
Gender	-0.171	-0.091	0.001		
Years diabetic	0.000	0.001	0.968		
Insulin use	-0.038	-0.021	0.469		

The adjusted R² of 0.215 indicates that this model predicts 21.5% of the variance in subcutaneous adipose layer thickness. The injection site, age, gender, and years diabetic are the factors that account for this variance while remaining 78.5% is due to individual variation and might be explained by other factors that were not taken into account in the analysis. These include the effect of minute differences in hydration status, and use of lotion or emollients on the skin. This effect is expected to be small, especially if there is no overt sign of dehydration.

Most of the study participants live in urban areas with possibly more limited sun exposure compared to those living in the rural areas. The amount of sun exposure is a possible factor affecting skin thickness.⁷

Although use of insulin was taken into consideration, the duration of insulin use and the injection site preferred by the patient were not evaluated in this study.

Using the Asia-Pacific BMI Classification (See Table 7A), an additional 114 patients will be classified as obese. These patients were classified as overweight under the WHO classification. In addition, 71 participants who were classified to have normal weight under WHO category, are overweight under the Asia Pacific Classification. The implications of this re-classification of participants are analyzed below.

Post hoc analysis using the Asia-Pacific BMI Classification

Table 7A. Patient Division per BMI Category (n = 293)

BMI category (Asia-Pacific Classification)	Underweight < 18.5	1 (0.3)
	Normal 18.5 to 22.9	66 (22.8)
	Overweight/at risk 23.0 to 24.9	71 (24.5)
Obese I 25.0 to 29.9		112 (38.6)
	Obese II >30.0	40 (13.8)
BMI category (WHO Classification)	< 25 (%)	138 (47.6)
	25 – 29.9 (%)	114(86.9)
	≥ 30 (%)	38 (13.1)

Table 7B. Study participants by gender per BMI Category

	Male, n = 98	Female, n = 195
BMI category (WHO Classification)		
• < 25 (%)	42 (42.9)	96 (50.0)
• 25 – 29.9 (%)	48 (49.0)	66 (34.4)
• ≥ 30 (%)	8 (8.2)	30 (15.6)
BMI category (Asia-Pacific Classification)		
• Underweight < 18.5	0 (0)	1 (0.5)
• Normal 18.5 to 22.9	25 (25.5)	41 (21.4)
• Overweight/at risk 23.0 to 24.9	17 (17.3)	54 (28.1)
• Obese I 25.0 to 29.9	48 (49.0)	64 (33.3)
• Obese II >30.0	8 (8.2)	32 (16.7)

Table 7C. Study participants by insulin use per BMI Category n = 290

	Insulin requiring, n = 130	Not insulin requiring, n = 160
BMI category (Who Classification)		
• < 25 (%)	51 (39.5)	86 (54.4)
• 25 – 29.9 (%)	59 (45.7)	53 (33.5)
• ≥ 30 (%)	19 (14.7)	19 (12.0)
BMI category (Asia-Pacific Classification)		
• Underweight < 18.5	0 (0)	1 (0.6)
• Normal 18.5 to 22.9	23 (17.8)	42 (26.6)
• Overweight/at risk 23.0 to 24.9	28 (21.7)	43 (27.2)
• Obese I 25.0 to 29.9	59 (45.7)	51 (32.3)
• Obese II >30.0	19 (14.7)	21 (13.3)

Figures 1-4 show the mean skin thickness of participants per BMI category based on the WHO Category. While Figures 9-12 show the mean skin thickness of participants per BMI category based on the Asia Pacific classification. When these figures were compared side-by-side, no significant difference can be clearly demonstrated. In both analysis, little variation in skin thickness is seen among the participants of the different BMI category, whether they were groups based on the WHO or Asia-Pacific Classification.

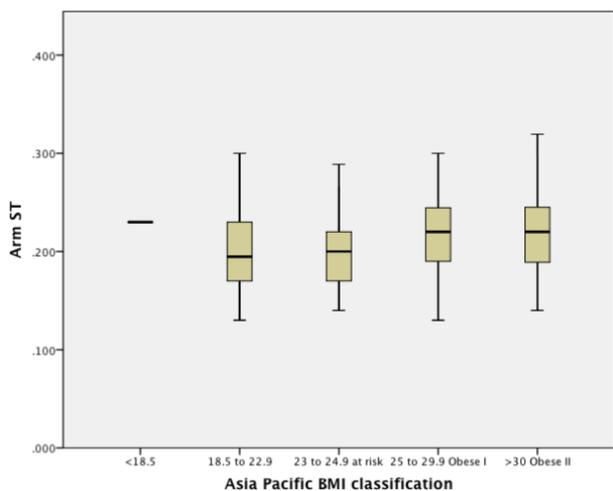


Figure 9. Arm skin thickness using Asia Pacific BMI category

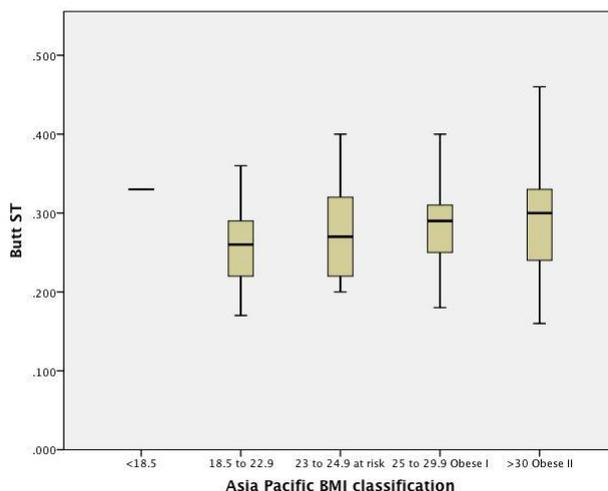


Figure 12. Buttock skin thickness using Asia Pacific BMI category

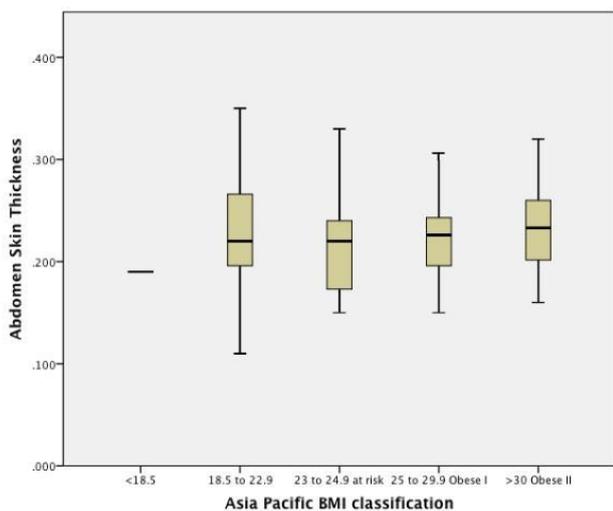


Figure 10. Abdomen skin thickness using Asia Pacific BMI category

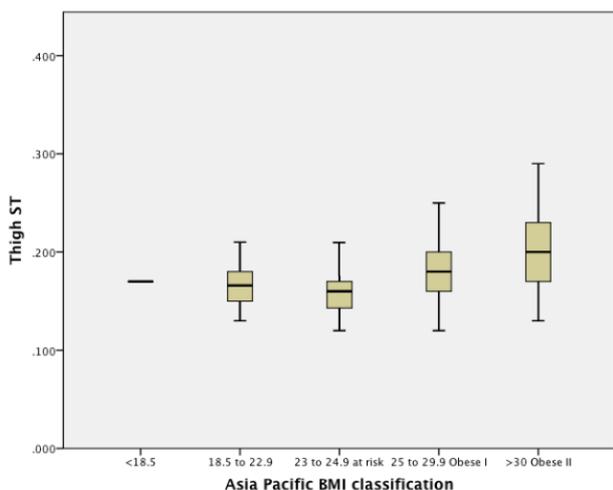


Figure 11. Thigh skin thickness using Asia Pacific BMI category

Figures 5-8 show the mean subcutaneous thickness of participants per BMI category based on the WHO Category while Figures 13-16 show the mean subcutaneous thickness of participants per BMI category based on the Asia Pacific guidelines. When these figures were compared side-by-side, no significant difference can be clearly demonstrated. In both analysis, little variation in subcutaneous thickness is seen among the participants of different BMI category, whether they were groups based on the WHO or Asia-Pacific Classification.

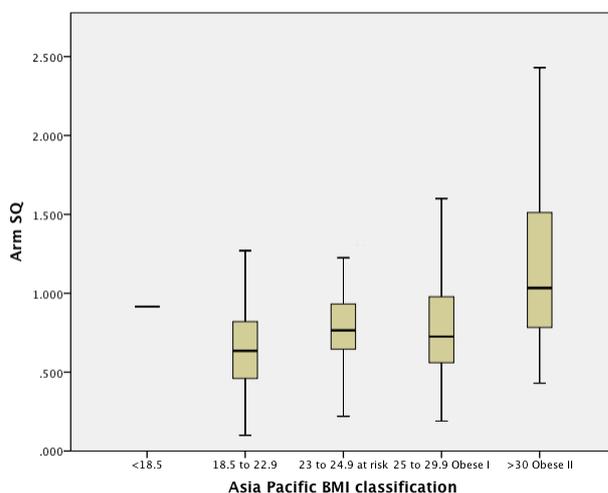


Figure 13. Arm SCT according to Asia Pacific BMI category

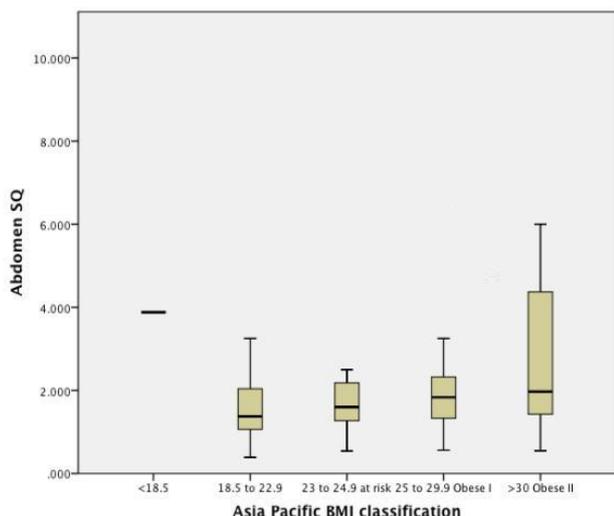


Figure 14. Abdomen SCT according to Asia Pacific BMI category

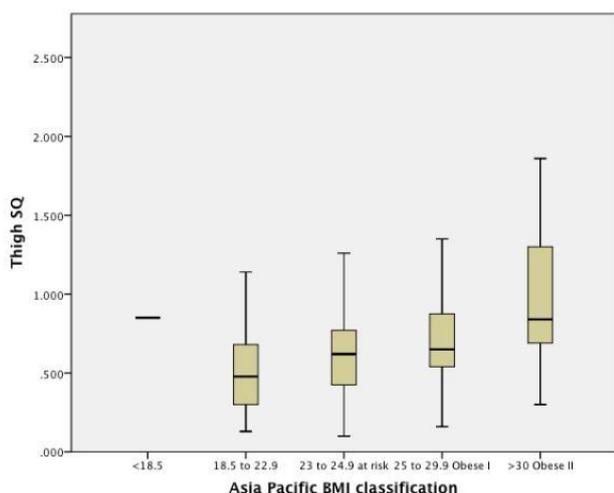


Figure 15. Thigh SCT according to Asia Pacific BMI category

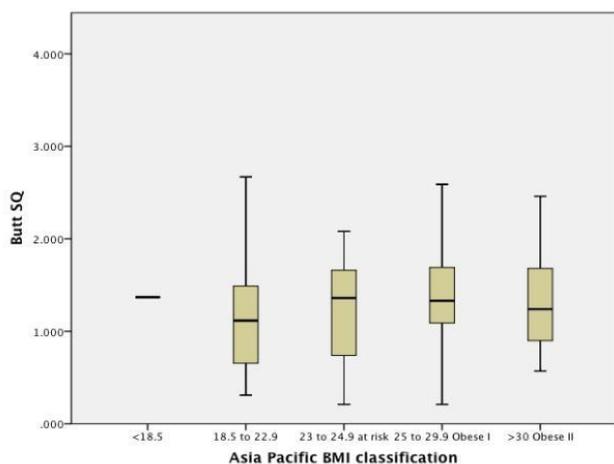


Figure 16. Buttock SCT according to Asia Pacific BMI category

DISCUSSION

Insulin can ameliorate many of the metabolic abnormalities in diabetes. Insulin therapy is shrouded by firmly held fallacies and myths, especially among less informed patients. The primary resistance to insulin therapy is the stigma of an injectable maintenance medication. A visual demonstration of short insulin needles will allay most of the patient’s fears, especially of the pain and discomfort accompanying the injection.

The American Diabetes Association recognizes the influence of needle length on glycemic control and has recommended that physicians and patients note variability in blood sugars when changing from one needle length to another.

Several studies have undertaken to measure skin and subcutaneous adipose layer thickness by ultrasound to establish the appropriate needle length for insulin injections while minimizing the risk of intramuscular injection.^{2,3} It is established that the skin thickness varies considerably within and between different anatomical areas. Variations in skin thickness according to age, ethnic origin, gender and body site were factors investigated that influence effective insulin delivery. Most of these studies indicate that skin thickness is very consistent across subjects with diverse demographic features.

Similarly, subcutaneous adipose tissue varies considerably and is affected by particular characteristics such as body site, body mass index and gender. The skin thickness in our study across body sites and BMI categories ranged from 1.7 to 2.699 mm. These values are similar to the findings of other races involving Caucasians.^{2,3}

Similar to other findings, the thigh has the thinnest mean lower bound 95% CI at 1.7 mm and the thickest mean upper bound 95% CI was 2.69 mm in the buttocks.^{2,3}

Gibney demonstrated in his study of 388 diabetic adults in the United States that the mean skin thickness for the common insulin injection sites ranged from 1.9 mm to 2.44 mm, with the thinnest thickness at the thigh and the buttocks being the thickest among the measurements. Mean subcutaneous measurement for this series ranged from 10.4 mm to 15.4 mm at the thighs.⁷

Gibney’s findings were comparable with Laurent, who conducted a normative study of the skin thickness among non-diabetic subjects among three ethnic groups, including Asians. Measurements that would correspond to insulin injection sites were skin thickness at the level of the waist was 1.91 mm and 1.55 mm in the thigh.^{2,10}

Lee and Hwang in 2001 studied 452 Korean patients and measured 28 different sites including: abdomen (site # 10), buttocks (site #13), back of the arm (site # 16), and front of

the thigh (site # 21). In this study, the authors noted that among females the buttocks measured 1.728 mm, the abdomen 1.225 mm, the posterior arm was 1.351 mm and the front of the thigh was 1.077 mm. Among males, the skin thickness in the abdomen was 1.43mm, the buttocks measured 1.678 mm, the back of the arm and the front of the thigh measured 1.0 mm and 1.162 mm respectively.¹¹

This study demonstrated relatively similar skin thickness measurements across increasing BMI categories. Subcutaneous measurement differences across BMI categories were more marked. The greatest skin thickness measurement of 2.82 mm in the buttock area supports the recommendation that short needles are appropriate and safe for diabetic Filipinos on insulin therapy. The average measurements at the injections sites are as follows: posterior arm 2.12 mm, anterior thigh 1.76 mm, abdomen 2.25 mm and buttocks 2.75 mm also indicate that short needles (4 mm, 5 mm) are sufficient to adequately penetrate the skin and deliver insulin to the subcutaneous layer. This would also minimize inadvertent intramuscular injection.

CONCLUSION AND RECOMMENDATIONS

Skin thickness among Filipinos varies minimally depending on injection site, being thickest at the buttocks and thinnest at the thigh area. On the other hand, there is greater variability in subcutaneous tissue depending on the injection site, being thickest at the abdominal area and thinnest at the anterior thigh.

LIMITATIONS OF THE STUDY

The study was limited to diabetic patients seen at the OPD during the duration of the study period and certain subgroups of patients (e.g., <40 year old, type 1 DM, those with low or normal BMI) were not well represented in this analysis

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