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Original Research Article

Study of subcutaneous fat, BMI in diabetic and non-diabetic adolescents**Jagdish Vishnoi¹, B. S. Bomb² and K. Ranjith Babu^{*3}**¹Assistant Professor, Department of General Medicine, Pacific Medical College & Hospital, Udaipur, Rajasthan, India²Professor & Head, Department of General Medicine, Pacific Medical College & Hospital, Udaipur, Rajasthan, India³Assistant Professor, Department of Physiology, Maheshwara Medical College & Hospital, Sangareddy, Telangana, India**QR Code*****Correspondence Info:**Dr. K. Ranjith Babu
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Sangareddy, Telangana, India***Article History:****Received:** 11/08/2017**Revised:** 28/08/2017**Accepted:** 28/08/2017**DOI:** <https://doi.org/10.7439/ijbar.v8i8.4339>**Abstract****Introduction:** Diabetes is a Metabolic Disorder which has got prime focus in the present days. An increase in body fat is generally associated with an increase in risk of metabolic diseases such as type 2 diabetes mellitus, hypertension and dyslipidaemia. Prevalence of childhood and adolescent obesity is further adding the severity. Therefore, any measures that could prevent or delay the development of diabetes are urgently needed. The present study is focused on early detection of the occurrence of Diabetes so that its control can be more effective because of early intervention.**Materials & Methods:** 50 people with increased blood sugar levels of age group of 16-20 years of both the sexes are selected randomly as subjects and 50 people with normal blood sugar levels are randomly selected as control group. Primary data of all the subjects like name, age, sex, occupation, address etc., are obtained as per the proforma. Calculation of Body Fat Percentage (%BF) involves measuring of skin fold thickness at four sites. Body Mass Index (BMI) calculated from Height and Weight of an individual.**Results:** The mean value of BMI in Diabetic group is found to be 25.64Kg/m² (SD ±2.22), while in Non-Diabetic group it is found to be 23.35Kg/m² (SD ±3.95). The mean value of %BF in Diabetic group is 18.20 (SD ± 1.62), while in Non-Diabetic group it is found to be 17.36 (SD ± 1.41). The mean values of RBS in Diabetic group are found to be 157.98mg/dl (SD ± 30.37) and 120.66mg/dl (SD ± 21.44) in Non-Diabetic group.**Discussion:** In the present study, we observed the correlation of increased subcutaneous fat with increased plasma glucose levels. Increased subcutaneous fat is considered to be an aggravating factor for early development of diabetes. Increased subcutaneous fat is also responsible for increasing the severity of diabetes there by, worsening the condition of the person. By knowing the body subcutaneous fat, the chances of occurrence of diabetes can be known.**Keywords:** Diabetes, Subcutaneous Fat, Body Fat, BMI, Blood Sugar.**1. Introduction**

Diabetes is a Metabolic Disorder which has got prime focus in the present days. Focus on Diabetes is not only because of its high prevalence, but also because of the after effects of increased Blood sugar levels in an individual. Even most important aspect is lack of proper management and treatment for Diabetes. An increase in body fat is generally associated with an increase in risk of

metabolic diseases such as type 2 diabetes mellitus, hypertension and dyslipidaemia[1].

There is a great deal of evidence that both genetic and environmental factors are of importance in the pathogenesis of T2DM. Whereas the genetic factors are still poorly understood, numerous studies have shown that obesity (in particular, central obesity), physical inactivity, a

high-fat diet, and a diet rich in saturated fatty acids increase the risk of diabetes [2]. The health risks associated with excessive intraabdominal or visceral adipose tissue deposition have become increasingly well-established [3,4].

Obesity is associated with increased morbidity and mortality and decreased life expectancy. Obesity is associated with increased risk for cardiovascular diseases. These include coronary heart disease, heart failure, and sudden death [5,6]. In addition to cardiovascular diseases, obesity is associated with numerous other medical conditions including type 2 diabetes, dyslipidemia, hypertension, nonalcoholic fatty liver disease, cancers, and sleep apnea [5].

Diabetes is also very common among young people and even children now days. Its contribution is increasing day by day in both developed and developing countries. Prevalence of childhood and adolescent obesity is further adding the severity. Therefore, any measures that could prevent or delay the development of diabetes are urgently needed. The present study is focused on early detection of the occurrence of Diabetes so that its control can be more effective because of early intervention.

The present study is focused mainly on early detection of the chances of occurrence of Diabetes so that development of Diabetes is prevented or delayed.

2. Materials & Methods

The present study is done in a population of the villages surrounding Pacific Medical College & Hospital, Udaipur, Rajasthan during the regular Medical Camps. The study is approved by the Institutional Ethical Committee. 50 people with increased blood sugar levels of age group of 16-20 years of both the sexes are selected randomly as subjects and 50 people with normal blood sugar levels are randomly selected as control group after obtaining informed consent in their mother tongue. Subjects taking any hormonal medications, drugs which interfere with blood sugar levels are excluded from the study.

Primary data of all the subjects like name, age, sex, occupation, address etc., are obtained as per the proforma. Calculation of Body Fat Percentage (%BF) involves measuring of skin fold thickness at four sites. The subjects

were made to stand and skin fold thickness at four sites i.e., Biceps, Triceps, Subscapularis and suprailiac were measured using skin calipers carefully. Biceps fat was measured at the level of nipple line. Triceps skin fat was measured midway between acromion process of scapula and olecranon process. Fat pads at the inferior angle of scapula and superiorly on iliac crest directly in the midaxillary line were measured for subscapular and supra iliac skin fold [7].

The density value was calculated using the equation of Durnin and Womersley [8,9]. The density value can then be converted to Percentage Body Fat using Siri Equation. Body weight of the subjects is measured using KUPPS weighing scale to nearest 0.1Kg with minimal clothing. Height without foot ware was measured using vertical scale (Avery, India) with an accuracy of 0.5cms.

BMI is calculated as:

$$BMI = \frac{\text{weight in Kilograms}}{(\text{height in meters})^2}$$

Waist circumference is measured after all precautions, by placing a measuring tape horizontally around the abdomen at the levels of both iliac crests.

Diabetes mellitus was confirmed based on self-reported responses (i.e. respondent answered yes to 'Has a doctor ever told you that you have diabetes?'). Undiagnosed diabetes mellitus cases were confirmed as per the American Diabetes Association criterion of Fasting Plasma Glucose (FPG) > 125 mg/dl (7.0 mmol/l) [10].

The data was arranged in suitable tables for discussion under different headings. The results were averaged (mean ± standard deviation) for each anthropometrical parameter subgroups separately for Diabetics & Non-Diabetics. One-way analysis of variance was used to test the difference between the groups. Analysis of Variance is a technique by which the total variation is split into two parts, one between groups and other within the groups. Statistical analysis was done using IBM SPSS Statistics 20 package. p-value of <0.05 is considered as statistically significant and p-value of <0.005 is considered as statistically highly significant. Conclusions were drawn based on outcome of this statistical treatment.

3. Observations and Results

Table 1: Mean ± SD of parameters in Diabetic Subjects & Non-Diabetic Control groups.

| Parameter | Diabetics Mean ± SD | Non-Diabetics Mean ± SD | P Value | Significance |
|-----------|---------------------|-------------------------|---------|--------------|
| BMI | 25.64 ± 2.22 | 23.35 ± 3.95 | <0.01 | HS |
| %Body Fat | 18.20 ± 1.62 | 17.36 ± 1.41 | <0.01 | HS |
| RBS | 157.98 ± 30.37 | 120.66 ± 21.44 | <0.01 | HS |

Table 2: Paired Samples Statistics

| | | Mean | N | Std. Deviation | Std. Error Mean |
|-------------|--------|--------|----|----------------|-----------------|
| Blood Sugar | RBS_D | 157.98 | 50 | 30.365 | 4.294 |
| | RBS_ND | 120.66 | 50 | 21.442 | 3.032 |
| %Body Fat | %BF_D | 18.18 | 50 | 1.637 | 0.232 |
| | %BF_ND | 17.32 | 50 | 1.449 | 0.205 |
| BMI | BMI_D | 25.68 | 50 | 2.180 | 0.308 |
| | BMI_ND | 23.34 | 50 | 3.972 | 0.562 |

Table 3: Paired Samples Correlations

| | | N | Correlation | Significance |
|-------------|----------------|----|-------------|--------------|
| Blood Sugar | RBS_D & RBS_ND | 50 | -0.134 | 0.352 |
| %Body Fat | %BF_D&%BF_ND | 50 | 0.199 | 0.166 |
| BMI | BMI_D& BMI_ND | 50 | 0.081 | 0.575 |

Table 4: Paired Samples Test

| | | Paired Differences | | | | | t | df | Significance (2-tailed) |
|----------------|----------------|--------------------|-------------------|--------------------|--|--------|-------|----|----------------------------|
| | | Mean | Std. Deviation | Std. Error Mean | 95% Confidence Interval of the Difference | | | | |
| | | | | | Lower | Upper | | | |
| Blood Sugar | RBS_D - RBS_ND | 37.320 | 39.455 | 5.580 | 26.107 | 48.533 | 6.688 | 49 | 0 |
| % Body Fat | %BF_D - %BF_ND | 0.860 | 1.959 | 0.277 | 0.303 | 1.417 | 3.104 | 49 | 0.003 |
| BMI | BMI_D - BMI_ND | 2.340 | 4.373 | 0.618 | 1.097 | 3.583 | 3.783 | 49 | 0 |

The mean value of BMI in Diabetic group is found to be 25.64Kg/m² (SD ±2.22), while in Non-Diabetic group it is found to be 23.35Kg/m² (SD ±3.95). The mean value of % BF in Diabetic group is 18.20 (SD ± 1.62), while in Non-Diabetic group it is found to be 17.36 (SD ± 1.41). The mean values of RBS in Diabetic group is found to be 157.98mg/dl (SD ± 30.37) and 120.66mg/dl (SD ± 21.44) in Non-Diabetic group.

4. Discussion

In the present study, we observed the correlation of increased subcutaneous fat with increased plasma glucose levels. Normally we expect increased skin thickness and increased body fat, subcutaneous fat in the diabetics. There are relatively limited data on skin thickness in childhood and adolescents. But three other studies have noted an age-related increase in dermis among children [11–13]. Considerably more data exist in adulthood, and several studies have shown a thinning of the dermis with increasing age [13–20]. In particular, our findings are similar to the findings of Tan *et al* showing a linear increase in dermis until the age of 20, with a subsequent decline thereafter [13]. Although Shuster *et al* also found this pattern of decreasing dermal thickness among men, they observed that it was relatively unchanged in women until their 50s after

which it began to decline [21] (probably due to decreasing oestrogen levels after menopause). Increased subcutaneous fat is considered to be an aggravating factor for early development of diabetes. Increased subcutaneous fat is also responsible for increasing the severity of diabetes there by, worsening the condition of the person. By knowing the body subcutaneous fat, the chances of occurrence of diabetes can be known.

With the above observation, it can be clearly confirmed that BMI is more in Diabetics when compared to non-diabetics. It may be the other way; increased BMI resulted in development of diabetes. It can also be noted that increased BMI is one of the favoring factor in development of diabetes. With simple tests like measuring the height and weight of the subject, the increased chances of development of diabetes can be detected early. Once the chances of development of diabetes are detected early, early intervention such as change in life style, food habits and inclusion of healthy habits can be advised. With our current study findings, one can have an idea of chances of developing diabetes based on the increased BMI and Subcutaneous fat. In our further studies, we plan to include more number of subjects from more population and include wide range of age and other parameters.

References

- [1]. World Health Organization. Obesity and Overweight Facts. http://www.who.int/hpr/NPH/docs/gs_obesity.pdf (accessed March 2007).
- [2]. Mokdad AH, Ford ES, Bowman BA, et al. Prevalence of obesity, diabetes, and obesity-related health risk factors, 2001. *JAMA* 2003; 289: 76–9.
- [3]. Ashwell M, Cole TJ, Dixon AK. Obesity: a new insight into the anthropometric classification of fat distribution shown by computed tomography. *Br Med J* 1985; 301: 203-205.
- [4]. DespreÂs J-P, Moorjani S, Lupien PJ, Tremblay A, Nadeau A, Bouchard C. Regional distribution of body fat, plasma lipoproteins, and cardiovascular disease. *Arteriosclerosis* 1990; 10: 495-551.
- [5]. Poirier P., Giles T. D., Bray G. A. et al., “Obesity and cardiovascular disease: pathophysiology, evaluation, and effect of weight loss: an update of the 1997 American Heart Association Scientific Statement on obesity and heart disease from the Obesity Committee of the Council on Nutrition, Physical Activity, and Metabolism,” *Circulation*, 2006; 113 (6): 898–918.
- [6]. Fontaine K. R., Redden D.T., Wang C., Westfall A. O., and Allison D. B. “Years of life lost due to obesity,” *Journal of the American Medical Association*, 2003; 289 (2): 187–193.
- [7]. Willet WC, Dietz WH, Colditz GA. Guidelines for healthy weight. *NEJM* 1999; 341: 427-434.
- [8]. Durnin JVGA, Womersley J. Body fat assessed from total body density and its estimation from skin fold thickness measurements on 481 men and women. *Br J Nutr* 1974; 32: 77-79.
- [9]. Kuriyan R, Petracchi C, Ferro- Luzzi A, Shetty PS, Kurpad AV. Validation of expedient methods for measuring body composition in Indian adults. *Indian J Med Res* 1998; 107: 37-45.
- [10]. American Diabetes Association. Diagnosis and classification of diabetes mellitus. *Diabetes Care* 2006; 29: S43–8.
- [11]. Lo Presti D, Ingegnosi C, Strauss K. Skin and subcutaneous thickness at injecting sites in children with diabetes: ultrasound findings and recommendations for giving injection. *Pediatr Diabetes* 2012; 13: 525–533.
- [12]. Seidenari S, Giusti G, Bertoni L, Magnoni C, Pellacani G. Thickness and echogenicity of the skin in children as assessed by 20-MHz ultrasound. *Dermatology* 2000; 201: 218–222.
- [13]. Tan CY, Statham B, Marks R, Payne PA. Skin thickness measurement by pulsed ultrasound: its reproducibility, validation and variability. *Br J Dermatol* 1982; 106: 657–667.
- [14]. Lasagni C, Seidenari S. Echographic assessment of age-dependent variations of skin thickness. *Skin Res Technol* 1995; 1: 81–85.
- [15]. Bliznak J, Staple TW. Roentgenographic measurement of skin thickness in normal individuals. *Radiology* 1975; 116: 55–60.
- [16]. Laurent A, Mistretta F, Bottiglioli D, Dahel K, Goujon C, et al. Echographic measurement of skin thickness in adults by high frequency ultrasound to assess the appropriate micro needle length for intradermal delivery of vaccines. *Vaccine* 2007; 25: 6423–6430.
- [17]. Branchet MC, Boisnic S, Frances C, Robert AM. Skin thickness changes in normal aging skin. *Gerontology* 1990; 36: 28–35.
- [18]. Levakov A, Vuckovic N, Dolai M, Kacanski MM, Bozanic S. Age-related skin changes. *Med Pregl* 2012; 65: 191–195.
- [19]. Petrofsky JS, McLellan K, Bains GS, Prowse M, Ethiraju G, et al. Skin heat dissipation: the influence of diabetes, skin thickness, and subcutaneous fat thickness. *Diabetes Technol Ther* 2008; 10: 487–493.
- [20]. Petrofsky JS, Prowse M, Lohman E. The influence of ageing and diabetes on skin and subcutaneous fat thickness in different regions of the body. *J Appl Res* 2008; 8: 55.
- [21]. Shuster SAM, Black MM, McVitie EVA. The influence of age and sex on skin thickness, skin collagen and density. *Br J Dermatol* 1975; 93: 639–643.