

A Study on Clinical Profile of Metabolic Syndrome of Adults

Dr.N.Kirubanand M.D., Dr.Sakthi Srinivas M.D.,

Assistant Professor, Department of General Medicine, Government Thiruvannamalai Medical College and Hospital

Assistant Professor, Department of General Medicine, Government Thiruvannamalai Medical College and Hospital

Abstract

INTRODUCTION: Defining metabolic syndrome has always produced problems and controversies. Many organisations such as the WHO, IDF and the NCEP-ATPIII Has proposed different definitions, and many studies have been done comparing these definitions. And there is no agreement about which criteria for the diagnosis of metabolic syndrome are the best to use. Despite various definitions and criteria's the metabolic syndrome is an important predictor of future catastrophic events such as stroke and cardiovascular disease.

AIMS AND OBJECTIVES:

1, To assess the in hospital prevalence of metabolic syndrome using the IDF criteria among adults who are admitted in GTVMCH.TIRUVANNAMALAI.

2, To assess the clinical profile of metabolic syndrome in patients admitted in medical wards.

MATERIALS AND METHODS

Cross sectional observational study
Patients who are getting admitted on Sunday, Tuesday, Wednesday in medical wards.

INCLUSION CRITERIA:

Consenting patients., Patients getting admitted from February 2018– January 2019. Patients who are between the age group of 30– 65 yrs of age.

EXCLUSION CRITERIA:

Non consenting patients. Patients with hypothyroidism, Cushing's syndrome and congenital obesity syndromes, Critically ill patients, Patients with end organ damage.

DATA ANALYSIS: Data collected from each individual was entered after coding of variables & appropriate analysis was done with help of EPI- INFO version 7. Qualitative data was analysed & depicted as percentages. Quantitative data was analysed and depicted as means and median. Odds ratio was calculated using chi-square test and p value was calculated

SUMMARY: In this study the prevalence of metabolic syndrome was 27 percent with the prevalence being more common in the female population which coincided with the earlier studies which were done in India. With the rise in metabolic syndrome proper education and life style modications along with its risk factors should be told to the patients.

KEYWORDS: Metabolic syndrome , Syndrome X, insulin resistance syndrome.

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I. Introduction

An increase in the Urbanisation and Westernisation has altered our food habits as well as our lifestyle .This has led to an increase in the non-communicable diseases like diabetes mellitus, hypertension, chronic lung diseases and coronary artery diseases. The magnitude of the problem is evident when a study shows that 64% of the total deaths in 23 low income countries are due to non-communicable diseases¹.

India has also fallen into this category, with an eccentric increase in urbanisation and westernisation, there has been an abrupt increase in NCDs (Non communicable diseases) like diabetes, and coronary artery diseases. Metabolic syndrome is a cluster of risk factors and diseases (which will be explained in the review of literature) which has predisposed the patients to increased risk of CVD and stroke. If identified early these patients with metabolic syndrome can avoid life threatening disease and circumstances. The importance of this syndrome is that all the components which are present are modifiable risk factors. So early identification of the patient and proper education and treatment is needed for these patients .

Vague in 1947 first observed an association between central adiposity and some metabolic abnormalities present in patients with diabetes mellitus and cardiovascular diseases². Later Reaven outlined the clinical importance of this syndrome .He didn't consider central obesity as a risk factor for the development of metabolic syndrome, but he described the syndrome as a group of metabolic abnormalities in which the insulin resistance had a causative role and he called it as Syndrome X .

Metabolic syndrome has been called with different names such as “Syndrome X”, the “Deadly quartet” and the “insulin resistance syndrome” over the past few decades. But

Defining this syndrome has always produced problems and controversies. Many organisations such as the WHO, IDF and the NCEP-ATPIII Has proposed different definitions, and many studies have been done comparing these definitions. And there is no agreement about which criteria for the diagnosis of metabolic syndrome are the best to use. Despite various definitions and criteria’s the metabolic syndrome is an important predictor of future catastrophic events such as stroke and cardiovascular disease.

II. Aims And Objectives:

- 1) To assess the in hospital prevalence of metabolic syndrome using the IDF criteria among adults who are admitted in GTVMCH.TIRUVANNAMALAI.
- 2) To assess the clinical profile of metabolic syndrome in patients admitted in medical wards.

III. Materials And Methods

STUDY DESIGN:

Cross sectional observational study. **STUDY SETTING:**

Patients who are getting admitted on Sunday, Tuesday, Wednesday in medical wards.

INCLUSION CRITERIA:

- 1) Consenting patients.
- 2) Patients getting admitted from February 2018– January 2019.
- 3) Patients who are between the age group of 30 – 65 yrs of age.

EXCLUSION CRITERIA:

- 1) Non consenting patients.
- 2) Patients with hypothyroidism, Cushing’s syndrome and congenital obesity syndromes.
- 3) Critically ill patients.
- 4) Patients with end organdamage.

SAMPLE SIZE AND METHODS:

With the prevalence of 25.8%, with confidence interval of 95%, absolute precision of 7% the sample size calculated was 165. First patient getting admitted on Sunday, Tuesday and Wednesday were selected for the study. The most common age group getting admitted in our hospital is between 30 – 65 yrs of age and hence this age group was selected for the study. For all the patients who were included in the study anthropometric measurements including waist circumference, height, bmi and weight were measured using standard techniques.

Waist circumference was measured using the non-stretchable inch tape at the smallest horizontal girth between the costal margins and the iliac crest during minimal respiration.

Height and weight were measured with the participant wearing light clothing and no shoes, and body mass index was calculated using the formula weight (kg)/height (m²).

Blood pressure was measured using a standard sphygmomanometer in the right upper limb. Pulse rate was counted for one whole minute.

Relevant blood investigations like haemoglobin, total count, fasting blood glucose and fasting lipid profile was done using standard measurements.

Patients with coronary artery disease were identified with the history or previous treatment details or presence of pathological q waves in the ECG.

History for risk factors such as hypertension, diabetes, smoking and alcohol intake was asked using a structured questionnaire to obtain information.

Patients who had consumed > than 14 units/week of alcohol in woman and > than 21 units/week of alcohol in men (AASLD guidelines) were 1 unit is equal to 8 grams of alcohol, were considered to be taking alcohol in excess and was included in the history.

Patients who were not meeting the recommended levels of physical activity (WHO) of at least 150 minutes of moderate-intensity aerobic physical activity throughout the week or do at least 75 minutes of vigorous-intensity aerobic physical activity throughout the week or an equivalent combination of moderate- and vigorous-intensity activity were considered sedentary.

DATA ANALYSIS:

Data collected from each individual was entered after coding of variables & appropriate analysis was done with help of EPI- INFO version 7. Qualitative data was analyse & depicted as percentages. Quantitative data was analysed and depicted as means and median. Odds ratio was calculated using chi-square test and p value was calculated.

IV. Results And Analysis

PREVALENCE OF METABOLIC SYNDROME:

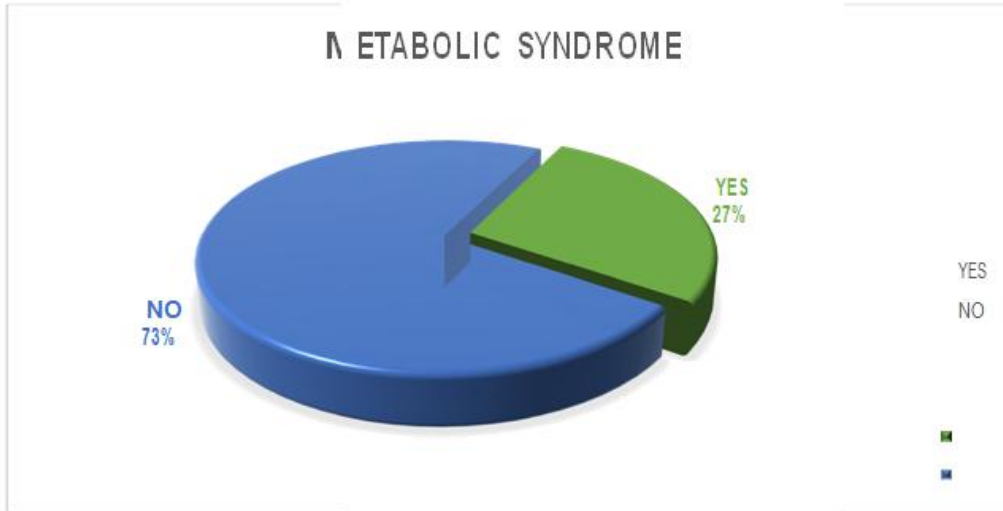


FIGURE NO: 1

TABLE NO: 2

METABOLIC SYNDROME	Frequency	Percent
NO	120	72.73 %
YES	45	27.27 %
TOTAL	165	100.00 %

SEX DISTRIBUTION OF THE STUDY POPULATION:

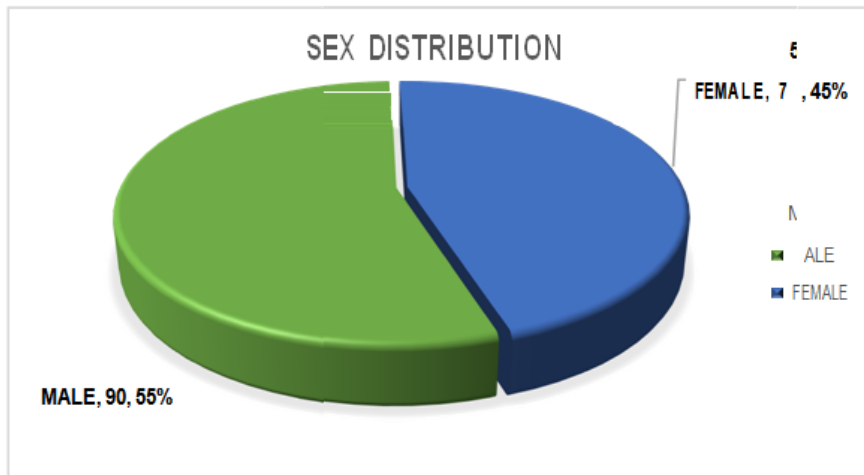


FIGURE NO: 2

TABLE NO: 3

sex	Frequency	Percent
female	75	45.45 %
male	90	54.55 %
TOTAL	165	100.00 %

AGE DISTRIBUTION OF THE STUDY POPULATION:

TABLE NO: 4

	OBSERVATIONS	MEAN	SD	MEDIAN 25 – 75 IQR
AGE	165	49.87	9.38	50(45 – 58)

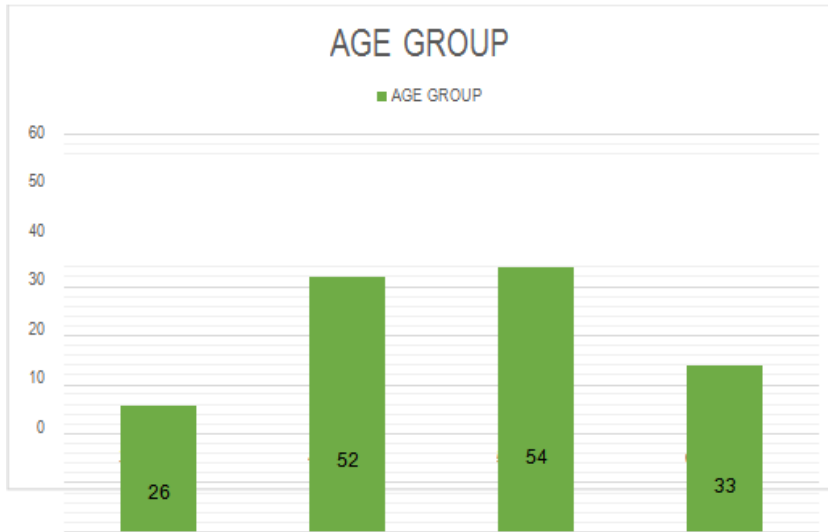


FIGURE NO: 3

TABLE NO: 5

age group	Frequency	PERCENT
30 – 39 yrs	26	15.76 %
40 – 49 yrs	52	31.52%
50 – 59 yrs	54	32.73%
60 – 65 yrs	33	20.00%
TOTAL	165	100%

METABOLIC SYNDROME DISTRIBUTION ACCORDING TO SEX:

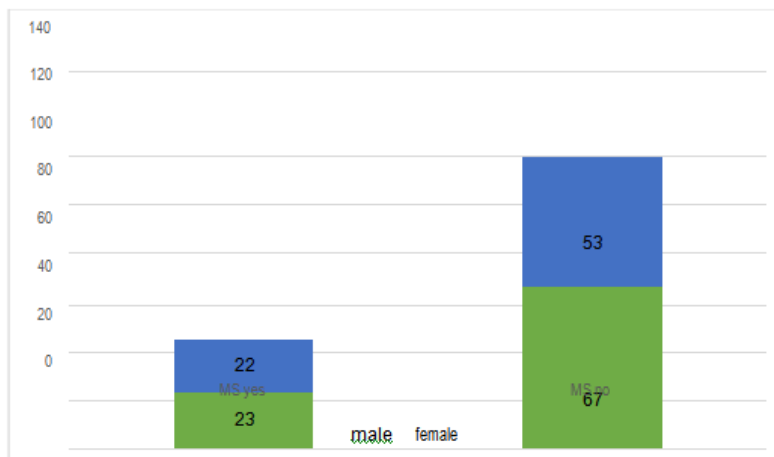


FIGURE NO: 4

TABLE NO: 6

	MALE	FEMALE
METABOLIC SYN YES	23	22
METABOLIC SYN NO	67	53

PREVALENCE OF METABOLIC SYNDROME ACCORDING TO SEX:

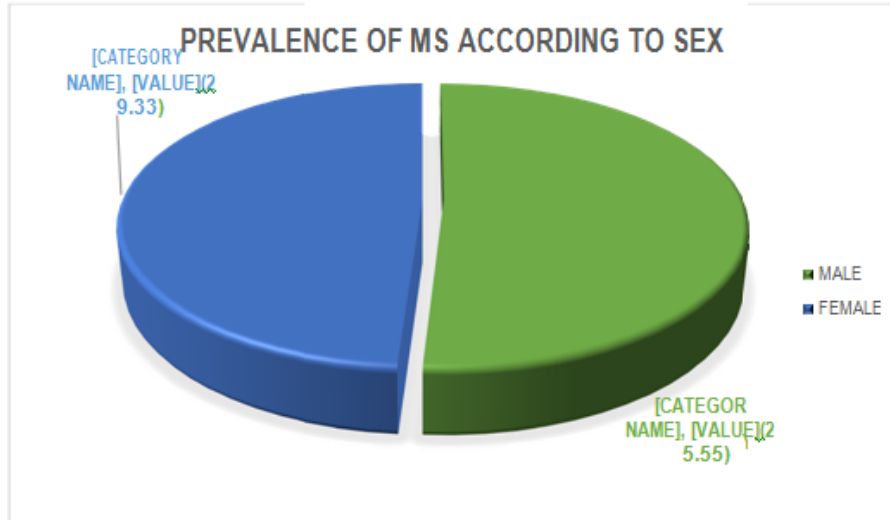


FIGURE NO: 5

TABLE NO: 7

METABOLIC SYNDROME	
MALE	23 (25.55%)
FEMALE	22 (29.33%)
TOTAL	45 (27.27%)

AGE GROUP FREQUENCY OF METABOLIC SYNDROME:

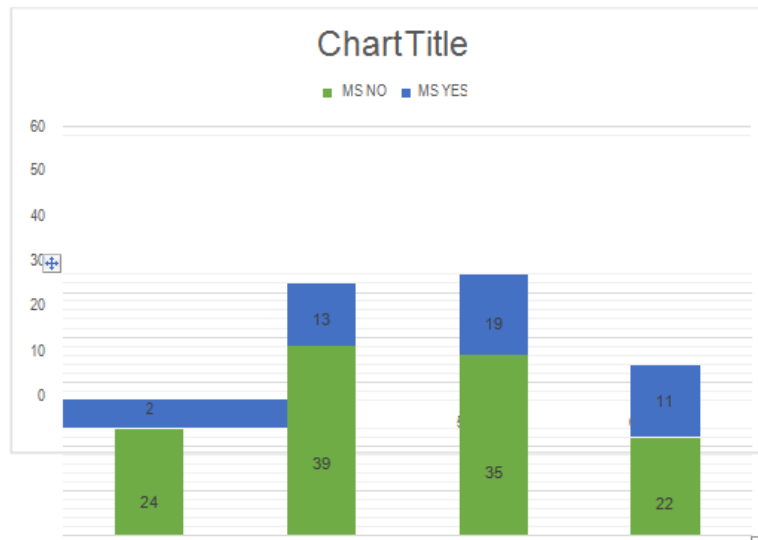


FIGURE NO: 6

TABLE NO: 8

METABOLIC SYNDROME			
Age	NO	YES	TOTAL
30-39 YRS	24	2(7.69%)	26(15.76%)
40-49 YRS	39	13(25.00%)	52(31.52%)
50-59 YRS	35	19(35.19%)	54(32.73%)
60-65 YRS	22	11(33.33%)	33(20.00%)
TOTAL	120	45	165

RELATIONSHIP OF TRIGLYCERIDES TO METABOLIC SYNDROME:

TABLE NO: 9

PARAMETER	MEAN	SD	MEDIAN 25 – 75 IQR
TRIGLYCERIDE	142.30	25.27	141(129 – 156)

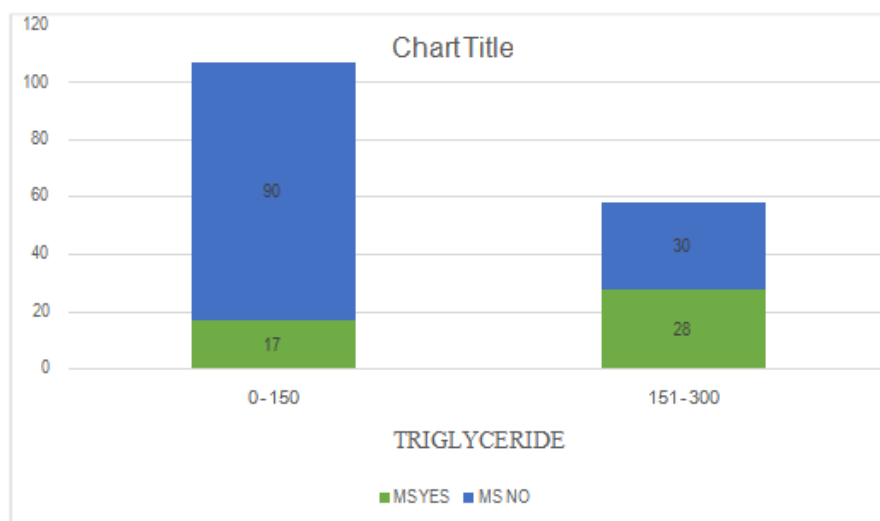


FIGURE NO: 7

TABLE NO: 10

METABOLIC SYNDROME			
TRIGLYCERIDE	YES	NO	Total
151-300	28	30	58
Row %	48.28 %	51.72 %	100.00 %
Col %	62.22 %	25.00 %	35.15 %
0 - 150	17	90	107
Row %	15.89 %	84.11 %	100.00 %
Col %	37.78 %	75.00 %	64.85 %
Total	45	120	165
Row %	27.27 %	72.73 %	100.00 %
Col %	100.00 %	100.00 %	100.00 %

Odds ratio was 4.94

TRIGLYCERIDE AGE WISE DISTRIBUTION:

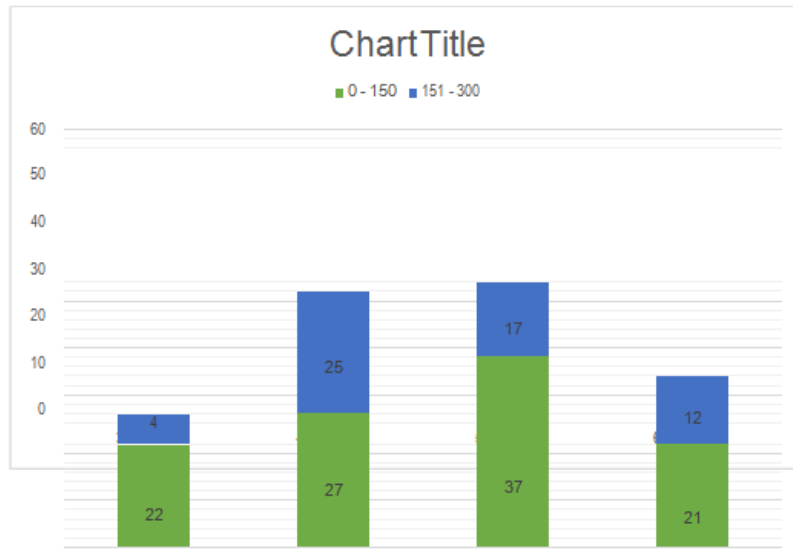


FIGURE NO: 8

TABLE NO: 11

	0 - 150	151 - 300	Total
30 - 39 yrs	22	4	26
40 - 49 yrs	27	25	52
50 - 59 yrs	37	17	54
60 - 65 yrs	21	12	33
	107	58	165

FASTING BLOOD SUGAR TO METABOLIC SYNDROME:

TABLE NO: 12

PARAMETER	MEAN	SD	MEDIAN 25-75 IQR
FBS	105.62	23.38	97 (94 - 105)

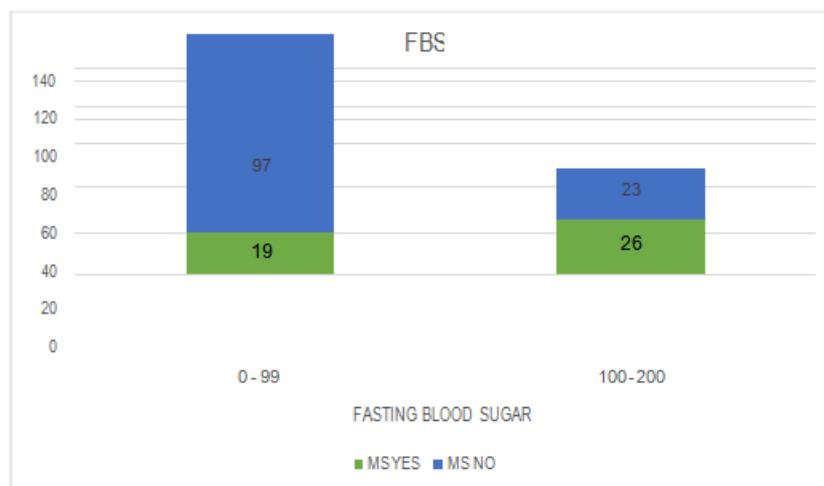


FIGURE NO: 9

TABLE NO: 13

METABOLIC SYNDROME			
FBS	YES	NO	Total
100-200	26	23	49
Row %	53.06 %	46.94 %	100.00 %
Col %	57.78 %	19.17 %	29.70 %
0 - 99	19	97	116
Row %	16.38 %	83.62 %	100.00 %
Col %	42.22 %	80.83 %	70.30 %
Total	45	120	165
Row %	27.27 %	72.73 %	100.00 %
Col %	100.00 %	100.00 %	100.00 %

Odds ratio was 5.77

AGE GROUP FREQUENCY OF HIGH FASTING BLOOD SUGAR:

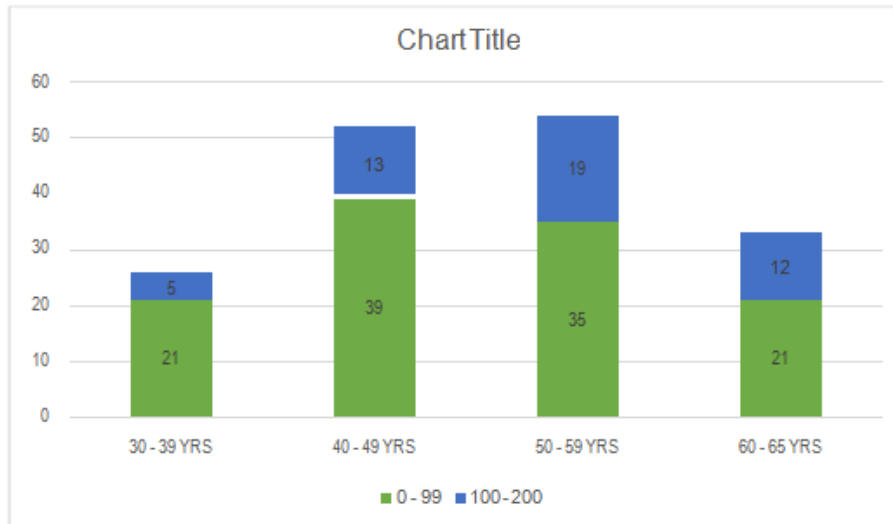


FIGURE NO: 10

TABLE NO: 14

FASTING BLOOD GLUCOSE			
Age	0 - 99	100 - 200	TOTAL
30 - 39 yrs	21	5	26
40 - 49 yrs	39	13	52
50 - 59 yrs	35	19	54
60 - 65 yrs	21	12	33
TOTAL	116	49	165

BMI AND METABOLIC SYNDROME

TABLE NO: 15

PARAMETER	MEAN	SD	MEDIAN 25 – 75 IQR
BMI	24.32	2.58	24.06 (22.66 – 25.76)

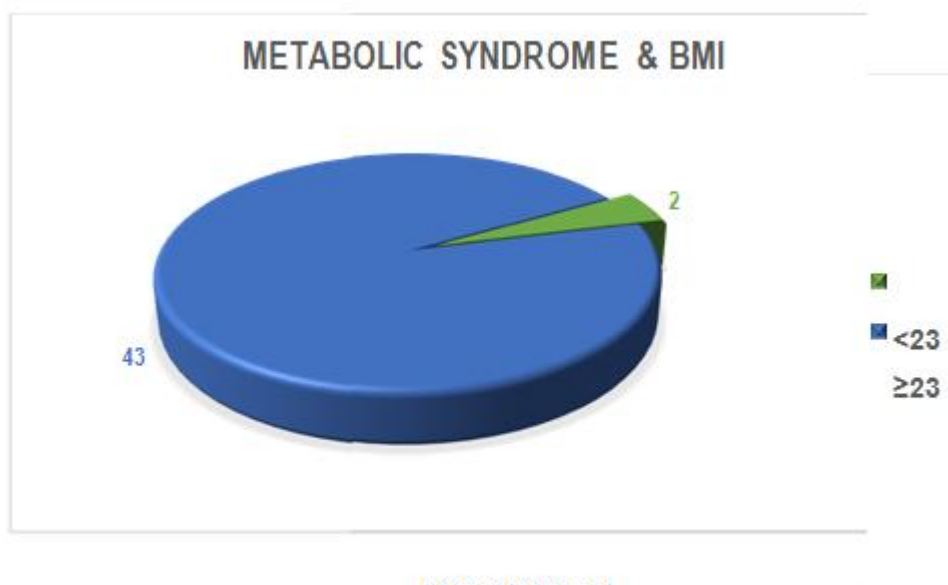


FIGURE NO: 11

TABLE NO: 16

BMI	METABOLIC SYNDROME
< 23	2
≥ 23	43

Odds ratio was 12.9

RELATION OF HDL FOR MEN TO METABOLIC SYNDROME:

TABLE NO: 17

PARAMETER	MEAN	SD	MEDIAN 25 – 75 IQR
HDL	39.78	3.80	40 (38 – 42)

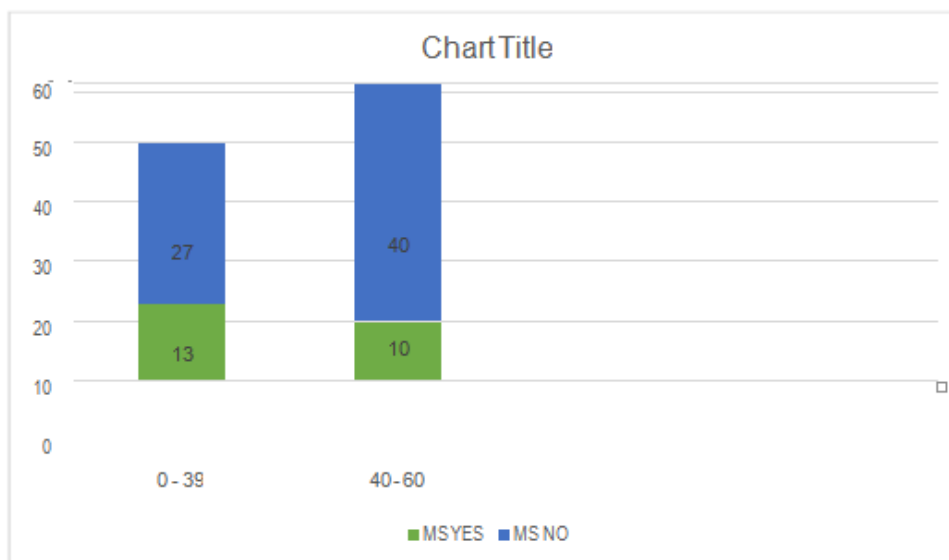


FIGURE NO: 12

TABLE NO: 18

METABOLIC SYNDROME

HDL	YES	NO	Total
0 - 39	13	27	40
Row %	32.50 %	67.50 %	100.00 %
Col %	56.52 %	40.30 %	44.44 %
40 - 60	10	40	50
Row %	20.00 %	80.00 %	100.00 %
Col %	43.48 %	59.70 %	55.56 %
Total	23	67	90
Row %	25.56 %	74.44 %	100.00 %
Col %	100.00 %	100.00 %	100.00 %

Odds Ratio 1.9259

AGE GROUP FREQUENCY OF HDL IN MEN:

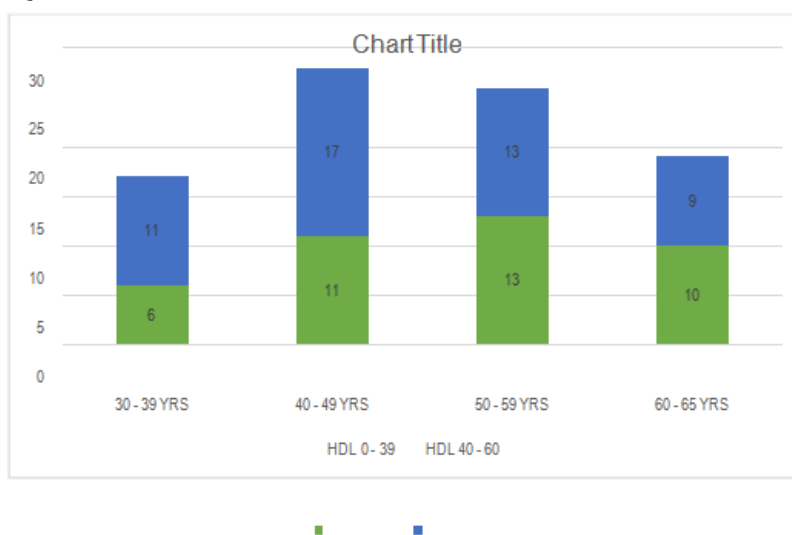


FIGURE NO: 13

TABLE NO: 19

AGE	HDL 0 – 39	HDL 40 – 60	Total
30 – 39 yrs	6	11	17
40 – 49 yrs	11	17	28
50 – 59 yrs	13	13	26
60 – 65 yrs	10	9	19
Total	40	50	90

RELATIONSHIP OF HDL FOR WOMAN TO METABOLIC SYNDROME:

TABLE NO: 20

PARAMETER	MEAN	SD	MEDIAN 25 – 75 IQR
HDL	43.64	5.81	43 (41 – 48)

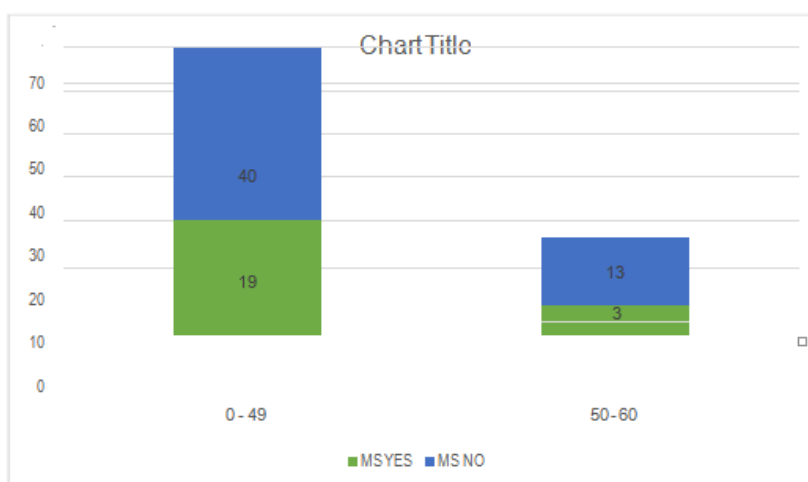


FIGURE NO: 14

TABLE NO: 21

Metabolic Syndrome			
HDL	YES	NO	Total
0 - 49	19	40	59
Row %	32.20 %	67.80 %	100.00 %
Col %	86.36 %	75.47 %	78.67 %
50 - 60	3	13	16
Row %	18.75 %	81.25 %	100.00 %
Col %	13.64 %	24.53 %	21.33 %
Total	22	53	75
Row %	29.33 %	70.67 %	100.00 %
Col %	100.00 %	100.00 %	100.00 %

Odds ratio was 2.05

AGE GROUP FREQUENCY OF HDL IN WOMEN:

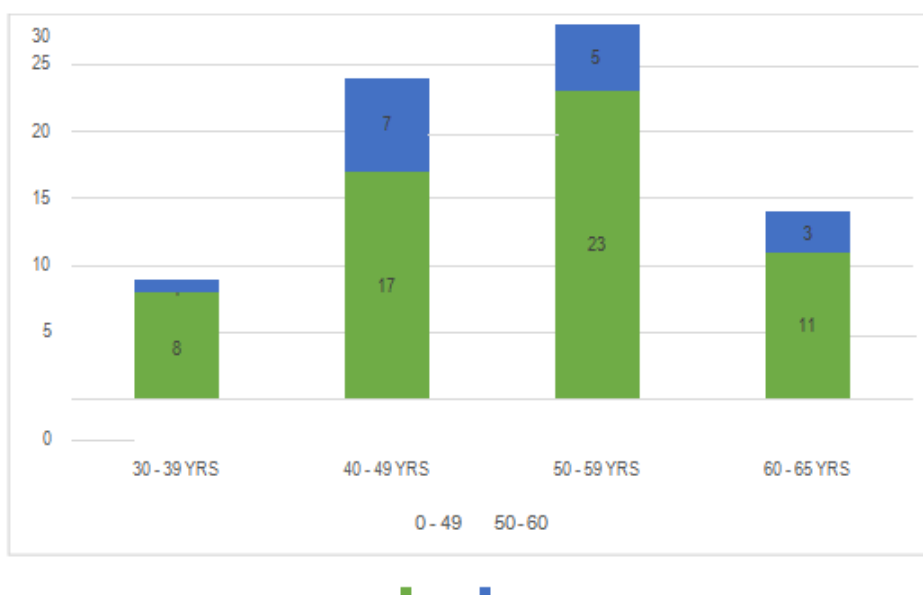


FIGURE NO: 15

TABLE NO: 22

AGE	HDL 0 - 49	HDL 50 - 60	TOTAL
30 - 39 yrs	8	1	9
40 - 49 yrs	17	7	24
50 - 59 yrs	23	5	28
60 - 65 yrs	11	3	14
TOTAL	59	16	75

RELATIONSHIP OF SYSTOLIC BLOOD PRESSURE TO METABOLIC SYNDROME:

TABLE NO: 23

PARAMETER	MEAN	SD	MEDIAN 25 – 75 IQR
SBP	123.43	15.72	120 (110 – 132)

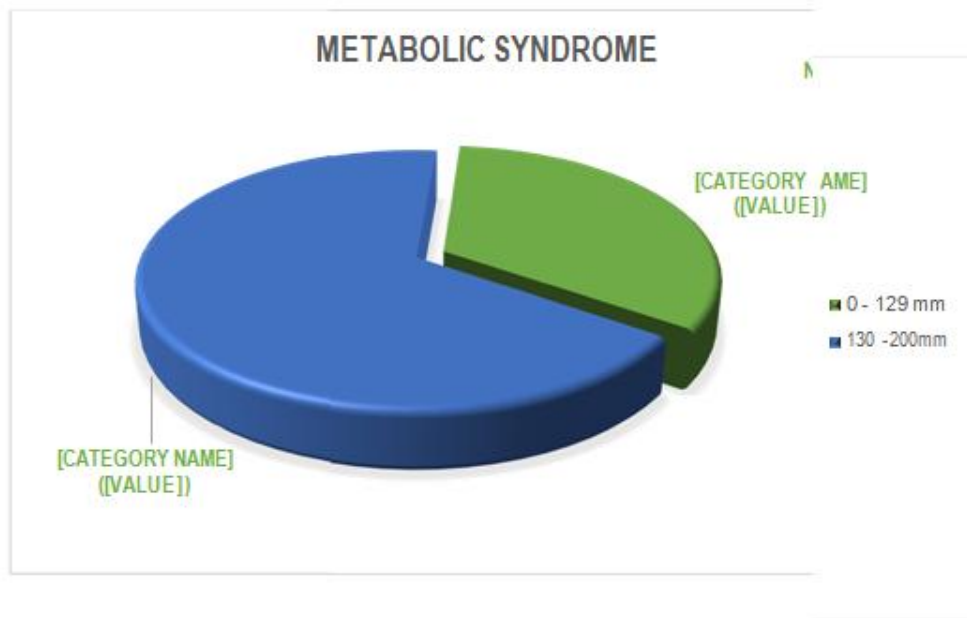


FIGURE NO: 16

TABLE NO: 24

METABOLIC SYNDROME	
BLOOD PRESSURE	YES
130 - 200	30
0 - 129	15
Total	45

AGE GROUP FREQUENCY OF SYSTOLIC BLOOD PRESSURE:

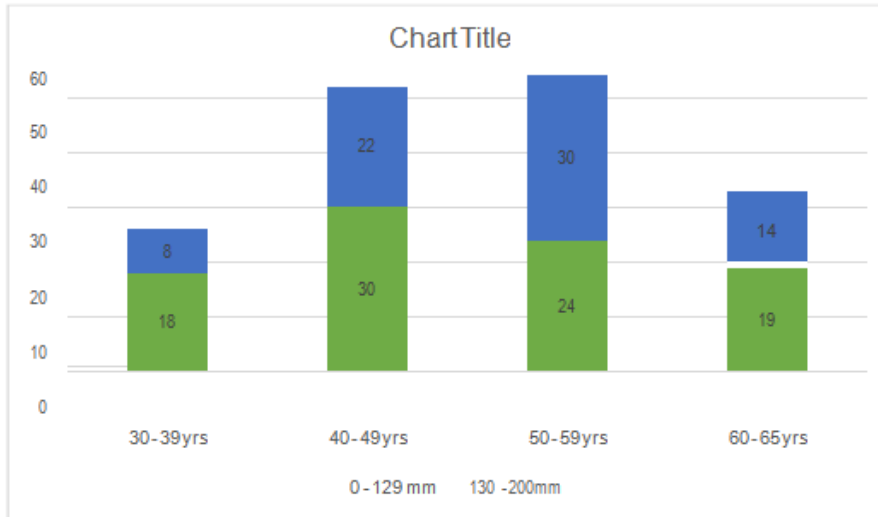


FIGURE NO: 17

TABLE NO: 24

Age	SBP 0 – 129 mm hg	SBP 130 – 200 mm hg	TOTAL
30 – 39 yrs	18	8	26
40 – 49 yrs	30	22	52
50 – 59 yrs	24	30	54
60 – 65 yrs	19	14	33
TOTAL	91	74	165

RELATIONSHIP OF DIASTOLIC BLOOD PRESSURE TO METABOLIC SYNDROME:

TABLE NO: 26

PARAMETER	MEAN	SD	MEDIAN 25 – 75 IQR
DBP	77.64	9.70	80 (70 – 80)

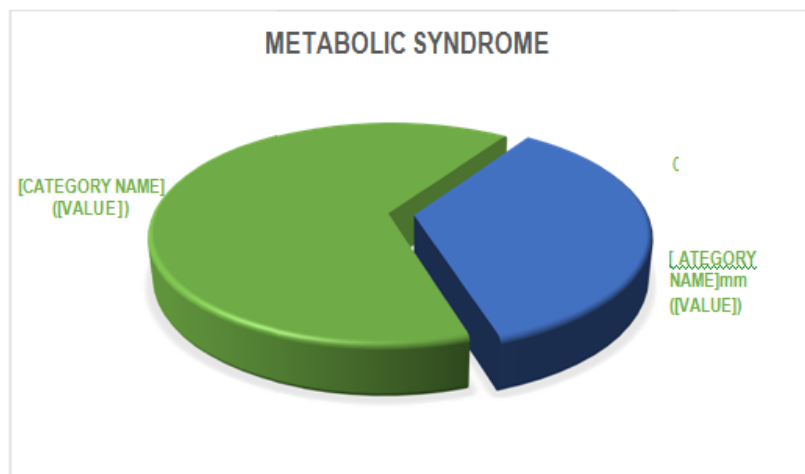


FIGURE NO: 18

TABLE NO: 27

METABOLIC SYNDROME	
BLOOD PRESSURE	YES
0 – 84	29
85 – 120	16
Total	45

AGE GROUP FREQUENCY OF DIASTOLIC BLOOD PRESSURE:

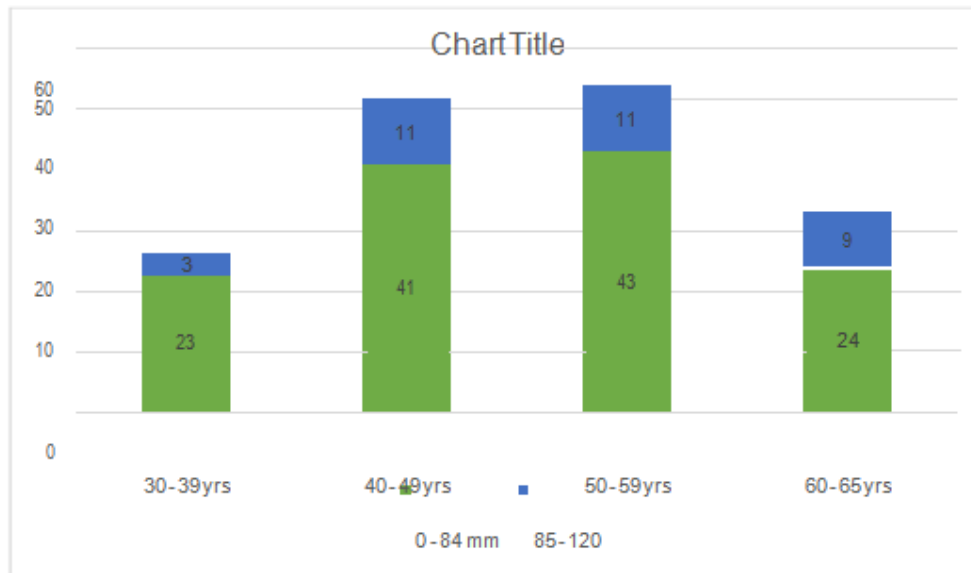


FIGURE NO: 19

TABLE NO: 28

Age	DBP 0 – 84 mm hg	DBP 85 – 120 mm hg	TOTAL
30 – 39 yrs	23	3	26
40 – 49 yrs	41	11	52
50 – 59 yrs	43	11	54
60 – 65 yrs	24	9	33
TOTAL	131	34	165

RELATION BETWEEN CAD AND METABOLIC SYNDROME:

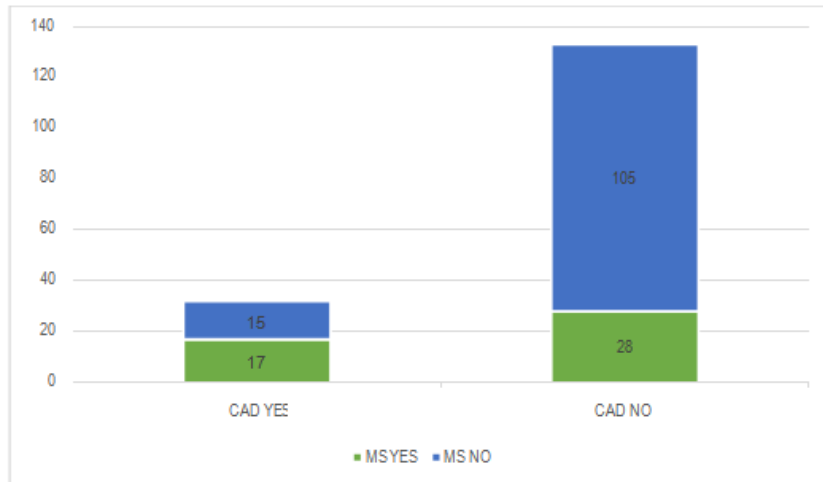


FIGURE NO: 20

TABLE NO: 29

cad	METABOLIC SYNDROME		Total
	YES	NO	
YES	17	15	32
Row %	53.13 %	46.88 %	100.00 %
Col %	37.78 %	12.50 %	19.39 %
NO	28	105	133
Row %	21.05 %	78.95 %	100.00 %
Col %	62.22 %	87.50 %	80.61 %
Total	45	120	165
Row %	27.27 %	72.73 %	100.00 %
Col %	100.00 %	100.00 %	100.00 %

Odds ratio was 4.25

RELATIONSHIP OF SEDENTARY LIFE STYLE AND METABOLIC SYNDROME:

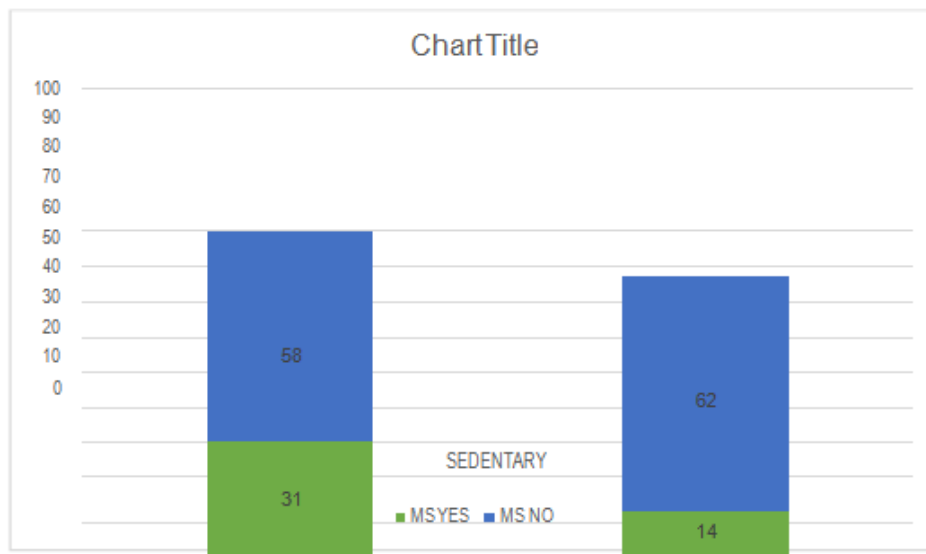


FIGURE NO: 21

TABLE NO: 30

METABOLIC SYNDROME			
Sedentary	YES	NO	Total
YES	31	58	89
Row %	34.83 %	65.17 %	100.00 %
Col %	68.89 %	48.33 %	53.94 %
NO	14	62	76
Row %	18.42 %	81.58 %	100.00 %
Col %	31.11 %	51.67 %	46.06 %
Total	45	120	165
Row %	27.27 %	72.73 %	100.00 %
Col %	100.00 %	100.00 %	100.00 %

Odds ratio was 2.36

RELATIONSHIP OF SMOKING TO METABOLIC SYNDROME:

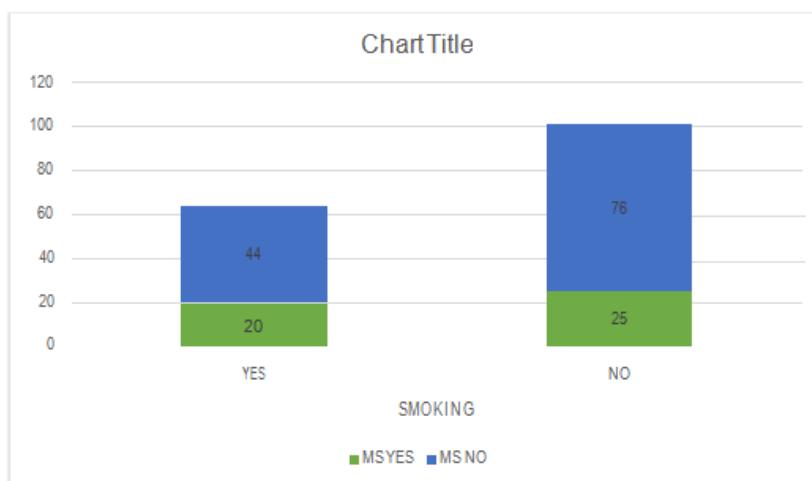


FIGURE NO: 22

TABLE NO: 31

METABOLIC SYNDROME			
smoking	YES	NO	Total
YES	20	44	64
Row %	31.25 %	68.75 %	100.00 %
Col %	44.44 %	36.67 %	38.79 %
NO	25	76	101
Row %	24.75 %	75.25 %	100.00 %
Col %	55.56 %	63.33 %	61.21 %
Total	45	120	165
Row %	27.27 %	72.73 %	100.00 %
Col %	100.00 %	100.00 %	100.00 %

Odds ratio was 1.38

RELATIONSHIP BETWEEN ALCOHOL CONSUMPTION AND METABOLIC SYNDROME:

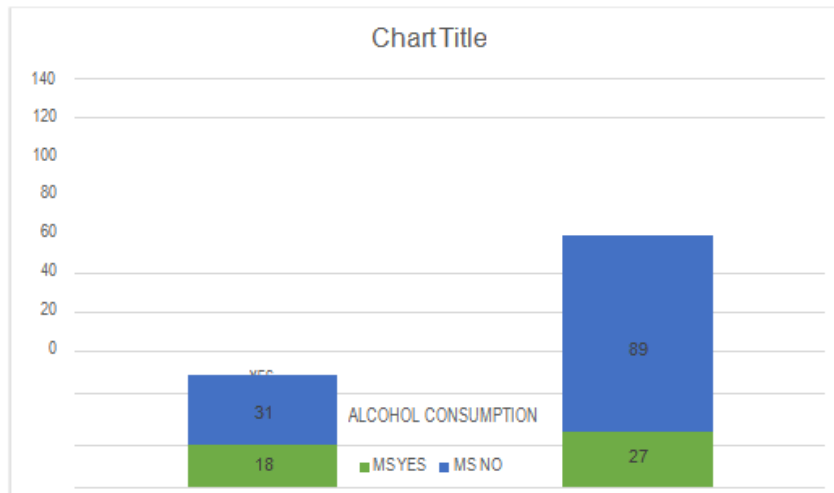


FIGURE NO: 23

TABLE NO: 32

METABOLIC SYNDROME

alcohol consumption	YES	NO	Total
YES	18	31	49
Row %	36.73 %	63.27 %	100.00 %
Col %	40.00 %	25.83 %	29.70 %
NO	27	89	116
Row %	23.28 %	76.72 %	100.00 %
Col %	60.00 %	74.17 %	70.30 %
Total	45	120	165
Row %	27.27 %	72.73 %	100.00 %
Col %	100.00 %	100.00 %	100.00 %

Odds ratio was 1.91

PREVALENCE OF OBESITY (WAIST CIRCUMFERENCE) IN MALES:

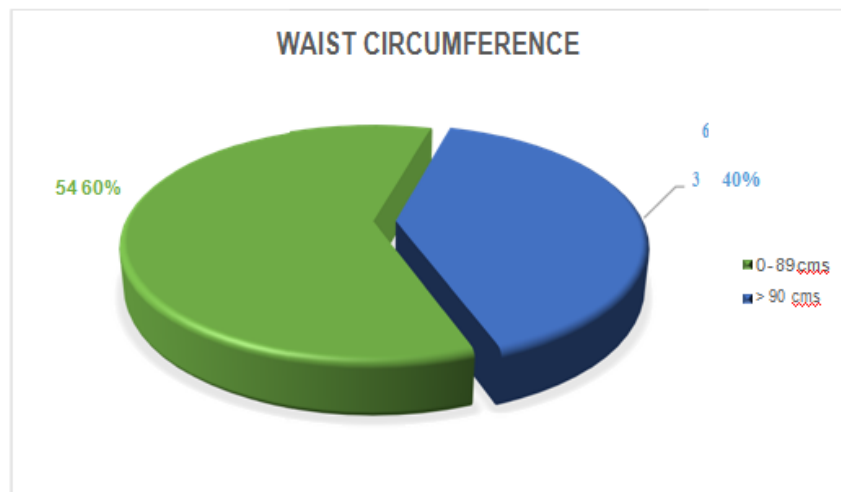


FIGURE NO: 24

TABLE NO: 33

MEASUREMENT	WAIST CIRCUMFERENCE
0 – 89 cms	54 (60%)
≥ 90 cms	36 (40%)

PREVALENCE OF OBESITY (WAIST CIRCUMFERENCE) IN FEMALES:

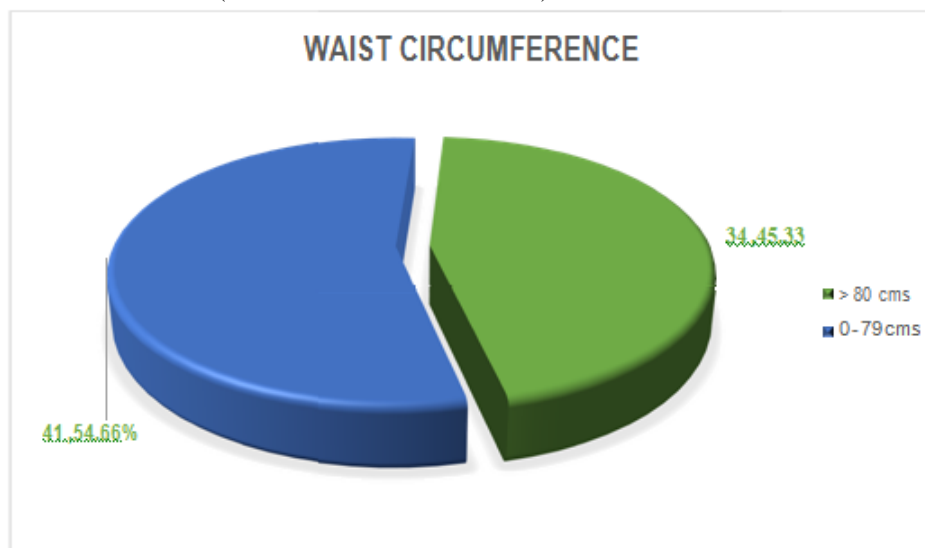


FIGURE NO: 25

TABLE NO: 34

MEASUREMENT	WAIST CIRCUMFERENCE
0 – 79 cms	41 (54.66%)
≥ 80 cms	34 (45.33%)

MAJOR HEAMATOLOGICAL PARAMETERS:

TABLE NO: 35

PARAMETERS	MEAN	SD	MEDIAN 25 – 75 IQR
HAEMOGLOBIN	12.06	1.68	12 (11 – 13.1)
TOTAL COUNT	8312.72	2217.97	7900 (6700 – 9700)

ANTHROPOMETRIC PARAMETERS:

TABLE NO: 36

PARAMETERS	MEAN	SD	MEDIAN 25 – 75 IQR
WAIST CIRCUMFERENCE	85.13	7.26	85 (79 – 90)
WC (MALE)	87.63	6.77	88 (84 – 93)
WC (FEMALE)	82.13	6.71	79 (78 – 86)
HEIGHT	161.03	9.63	159 (154 – 169)

OTHER LIPID PARAMETERS:

TABLE NO: 37

PARAMETER	MEAN	SD	MEDIAN 25 – 75 IQR
LDL	135.47	30.32	130 (118 – 146)

V. Discussion:

Among the 165 sample size study population 54.55% were males and 45.45% were females and the median age group was 50 with an interquartile range of 45 and 58. Among the age distribution 31.52% was present in the 40 to 49 age group and 32.73 % was present in the 50 to 59 age group. The prevalence of metabolic syndrome in this study was 27.27 %,and among the females the prevalence of metabolic syndrome was 29.33 % (22) and among the male population the prevalence of metabolic syndrome was 25.55% (23). According to a study done in south India by Deepa et al the prevalence of metabolic syndrome was 25.8% using the IDF criteria and the prevalence among the males and the females were 23.1% and 28.2%¹⁸.According to a study done in India the prevalence of metabolic syndrome was 33.3 in males and 40.4 in females⁵⁵. Anationwide population based survey in Taiwan observed 20.4% of men and 15.3 % of women with metabolic syndrome⁵⁶.So as told earlier different studies show different prevalence rate according to the ethnicity age andpopulation.

Among the age group distribution maximum prevalence of MS was present in the 50 to 59 (35.19 %) age group and the 60 to 65(33.33%) age group, according to Ervin et al, males and females aged 40 to 59 yrs were three times more likely to develop metabolic syndrome than the 20 to 39 yrs age group⁵⁷, and according to a study done by Yousefzadeh et al and Sheikhvatan et al the prevalence of metabolic syndrome peaked in the age range of 51 to 60 years among both males and females⁵⁸which was coinciding with our study.

Among the components of metabolic syndrome the median waist circumference is 85 with an interquartile range of 79 and 90. Whereas in the male population the median waist circumference is 88 with an interquartile range of 84 to 93, and in the females the median waist circumference is 79 with an interquartile range of 78 and 86.Overall prevalence of obesity in males was 40% and the prevalence of obesity in females was 45% and in the study done by Deepa et al the prevalence of obesity in males and females was 38.5% and 58.3%¹⁸and in a study done in India high waist circumference was present in 35.7% of males and 57.5% of females⁵⁹. Body mass index more than 23 was present in 71 percent of the study population, and 36% of the population was associated with metabolic syndrome (ICMR guidelines normal for Indian population 18.5 – 22.9), and according to a study done by Sawant et al the prevalence of high body mass index was 79.01% (>23kg/m²)⁶⁰. The overall prevalence of hyper triglyceridemia in the study population was 35.15% and was high in the 40 to 49 age group (48.08%) and the 60 to 65 age group (36.36%). ODD's ratio was 4.94.The median value was 141 with an interquartile range of 129 and 156.

According to deepa et al the prevalence of hypertriglyceridemia in the study population was 25.2%, according to deedwania et al the prevalence of hypertriglyceridemia was 41.2% in males and 31.5% in females

⁵⁷ Impaired fasting glucose was prevalent in 29.7% of the study population and the odd's ratio was 5.77, with the prevalence of impaired fasting blood glucose level increasing in the 50 to 59 age group (35%) and in the 60 to 65 age group (36%). Median value of IFG was 97 with an interquartile range of 94 and 105, and an Indian study showed a prevalence 20.9% ¹⁸ and according to Thakur et al the prevalence of fasting blood glucose was 32.6% ⁶¹. HDL levels (in men) less than < 40 was present in 44.44% of the study (male) population, the median value of HDL was 40 with an inter quartile range of 38 and 42 and odds ratio was 1.9 for those patients who had HDL < 40 for developing metabolic syndrome. Here HDL less than 40 was present in high percentage in the 50 to 59 age group (50%) and in the 60 to 65 age group (52%).

For women those who were having HDL levels less than 50, the median value was 43 with an interquartile range of 41 and 48, 78% of the study (female) population had HDL values less than 50 with an odds ratio of 2.05. HDL levels less than 50 were present in high percentage in the 40 to 49 age group (70.83%), 50 to 59 age group (82.14%) and 60 to 65 age group (78.57%) and there is an Indian study which shows a prevalence of 70.4% of decreased HDL ⁵⁹. Systolic blood pressure ≥ 130 elevated in 44.85% of the study population with an odds ratio of 3.45, the median value of SBP was 120 with an interquartile range of 110 and 132, high prevalence of systolic blood pressure ≥ 130 was found in the 50 to 59 age group (55.56%) and the 60 to 65 age group (42.42%).

Diastolic blood pressure ≥ 85 was present in 20 % of the study population with an odd's ratio of 3.12 and high prevalence of diastolic blood pressure ≥ 85 was found in the 50 – 59 age group (20.37%) and the 60 to 65 age group (27.27%). Median value for DBP was 80 with an interquartile range 70 and 80. And according to an Indian study the prevalence of metabolic syndrome in hypertensive patients was 63.6% (according to the IDF criteria).

Around 19.39% of the study population had coronary artery disease and 53% of patients with CAD were associated with metabolic syndrome, with an odds ratio of 4.25, according to an Indian study 61.06% of CAD patients were associated with metabolic syndrome ⁶².

In the study population 53.94% of patient's lead sedentary life style and among them 34.83 were associated with metabolic syndrome with an odds ratio of 2.36.

Among smokers (predominantly men) 31.25% of them were associated with metabolic syndrome and odds ratio was 1.38. Among people who were consuming excess amount of alcohol 36.73% were having metabolic syndrome with an odds ratio of 1.9. According to a study done in Puerto Rico, 42.7% of smokers and 48.4% of ex-smokers were associated with metabolic syndrome ⁶³ and a study done in India shows a prevalence of 24.6% of metabolic syndrome among those who consume alcohol in excess ⁶⁴.

Median value of haemoglobin was 12 with an interquartile range of 11 and 13.1, and the median value of total count was 7900 with an interquartile range of 6700 and 9700.

The median value of LDL was 130 with an interquartile range of 118 and 146.

VI. Conclusion

- 1) Here in this study the prevalence of metabolic syndrome was present in about 27% of the study population, with the prevalence of metabolic syndrome more in the female population.
- 2) In the age group distribution of metabolic syndrome the highest percentage was present in the sixth decade of life.
- 3) Among the individual components of metabolic syndrome decreased HDL was the commonest followed by high blood pressure, elevated triglycerides and IFG, and increased waist circumference was of course a definitive criteria for the diagnosis of metabolic syndrome according to the IDF criteria.
- 4) People with metabolic syndrome were four times more prone to develop coronary artery disease.
- 5) People who are smoking and consuming excess amount of alcohol are more prone to develop metabolic syndrome.
- 6) People who are leading a sedentary life without physical activity are two times more prone to develop metabolic syndrome.

So in this study the prevalence of metabolic syndrome was 27 percent with the prevalence being more common in the female population which coincided with the earlier studies which were done in India. With the rise in metabolic syndrome proper education and life style modications along with its risk factors should be told to the patients.

LIMITATIONS OF THE STUDY:

Limitations of the study are the small sample size which may not represent the general population. Systematic random sampling may cause the sampling error in this study. A larger sample may be needed to represent the general population. All the risk factors could not be studied properly. The other limitation is that this is a hospital study where many confounding factors are present which may alter the study.

References

- [1]. Alwan A, Maclean DR, Riley LM, d'Espaignet ET, Mathers CD, Stevens GA, et al. Monitoring and surveillance of chronic non-communicable diseases: progress and capacity in high-burden countries. *Lancet*. 2010;376(9755):1861-8.
- [2]. Unwin N. The metabolic syndrome. *Journal of the Royal Society of Medicine*. 2006;99(9):457-62.
- [3]. Reaven GM. Role of insulin resistance in human disease (syndrome X). An expanded definition. *Ann Rev Med*. 1993;44:121-3.
- [4]. Somaa B, Almgren P, Tuomi T, et al. Cardiovascular morbidity and mortality associated with the metabolic syndrome. *Diabetes care* 2001; **24**: 683–689.
- [5]. Chetna Mangat, NK goel and Dinesh K Walia. Metabolic syndrome: a challenging health issue in highly urbanized union territory of north India. *Diabetology and Metabolic Syndrome*. 2010;2:19.
- [6]. Kassi E, Pervanidou P, Kaltsas G, Chrousos G (2011) Metabolic syndrome: definitions and controversies. *BMC Med* 9: 48.
- [7]. Yamaoka K, Tango T (2012) Effects of lifestyle modification on metabolic syndrome: a systematic review and meta-analysis. *BMC Med* 10: 138.
- [8]. Alberti KG, Zimmet PZ: Definition, diagnosis and classification of diabetes Mellitus and its complications. *Diagnosis and classification of diabetes mellitus provisional report of a WHO consultation*. *Diabet Med* 1998,15:539-553.
- [9]. Executive Panel on Detection Evaluation and Treatment of High Blood Cholesterol in Adults: Executive Summary of the Third Report of the National Cholesterol Education
- [10]. Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III). *JAMA* 2001, 285:2486-2497.
- [11]. Alberti KGMM, Zimmet PZ, Shaw JE, IDF Epidemiology Task Force Consensus Group: The metabolic syndrome: a new worldwide definition from the International Diabetes Federation consensus. *Lancet* 2005,366:1059-1062.
- [12]. Hildrum B, Mykletun A, Hole T, Midthjell K, Dahl AA: Age-specific prevalence of the metabolic syndrome defined by the International Diabetes Federation and the National Cholesterol Education Program: the Norwegian HUNT 2 study. *BMC Public Health* 2007, 7:220.
- [13]. Athyros VG, Ganotakis ES, Bathianaki M, Monedias I, Goudevenos IA, Papageorgiou AA: Awareness, treatment and control of the metabolic syndrome and its components: A Multicentre Greek Study. *Hellenic J Cardiol* 2005, 46:380-386.
- [14]. Galassi A, Reynolds K, He J. Metabolic syndrome and risk of cardiovascular disease: a metaanalysis. *Am J Med* 2006;119:812-819.
- [15]. Park YW, Zhu S, Palaniappan L, Heshka S, Carnethon MR, Heymsfield SB. The metabolic syndrome: prevalence and associated risk factor findings in the U.S. population from the Third National Health and Nutrition Examination Survey, 1988-1994. *Arch Intern Med* 2003;163:427-436.
- [16]. Alkerwi A, Donneau AF, Sauvageot N, Lair ML, Scheen A, et al. (2011) Prevalence of the metabolic syndrome in Luxembourg according to the Joint Interim Statement definition estimated from the ORISCAV-LUX study. *BMC Public Health* 11: 4.
- [17]. Ford ES, Giles WH, Mokdad AH (2004) Increasing prevalence of the metabolic syndrome among U.S. Adults. *Diabetes Care* 27: 2444–2449.
- [18]. Mohan V, Gokulakrishnan K, Sandeep S, Srivastava BK, Ravikumar R, et al. (2006) Intimal media thickness, glucose intolerance and metabolic syndrome in Asian Indians—the Chennai Urban Rural Epidemiology Study (CURES -22). *Diabet Med* 23: 845–850.
- [19]. Wen-Harn Pan, Wen-Ting Yeh, and Lu-Chen Weng, Epidemiology of metabolic syndrome in Asia, *Asia Pac J Clin Nutr* 2008;17(S1):37-42.
- [20]. Gupta R, Deedwania PC, Gupta A, Rastogi S, Panwar RB, Kothari K. Prevalence of metabolic syndrome in an Indian urban population. *Int J Cardiol*. 2004;97:257-61.
- [21]. Deepa M, Farooq S, Datta M, Deepa R, Mohan V (2007) Prevalence of metabolic syndrome using WHO, ATP III and IDF definitions in Asian Indians: the Chennai Urban Rural Epidemiology Study (CURES-34). *Diabetes Metab Res Rev*. 23: 127-134.
- [22]. Nakagawa T, Hu H, Zharikov S, Tuttle KR, Short RA, Glushakova O, Ouyang X, Feig DI, Block ER, Herrera-Acosta J, Patel JM, Johnson RJ (2006). "A causal role for uric acid in fructose-induced metabolic syndrome". *Am J Phys Renal Phys* 2006;290: F625–F631.
- [23]. Motala A, Mbanya JC, Ramaiya K, Metabolic Syndrome in Sub-Saharan Africa. *Ethn Dis*. 2009;Suppl 2:S2-8-S2-10.
- [24]. Chen, J. Tian, Z. Q. Zhang, W. G. Chen, J. H. Yan, Z. C. Ni, Y. X. Zhong, J. Jin, J. Zhao, Z. G. Mu, H. Zhu, Z. M. Relationship between visceral adipose tissue and prevalence of metabolic syndrome MS in patients with MS, and hypertension and/or diabetes, English Abstract Research Support, Non-U.S. Gov't China 2006/10/27 09:00 *Zhonghua Yi Xue Za Zhi*. 2006 Aug 15;86(30):2110-3.
- [25]. Lear, S. A. Chockalingam, A. Kohli, S. Richardson, C. G. Humphries, K. H. Elevation in cardiovascular disease risk in South Asians is mediated by differences in visceral adipose tissue. *Obesity (Silver Spring)*. 2012 Jun;20(6):1293-300. doi: 10.1038/oby.2011.395.
- [26]. Taylor H, Liu J, Wilson G, Golden SH, Crook E, Brunson CD, et al: Distinct component profiles and high risk among African Americans with metabolic syndrome: the Jackson Heart Study. *Diabetes Care* 2008, 31(6):1248-53.
- [27]. Scuteri A, Najjar SS, Morrell CH, et al. The metabolic syndrome in older individuals: prevalence and prediction of cardiovascular events. *Diabetes Care* 2005; **28**: 882–887.
- [28]. Reaven GM, Chen YD, Jeppesen J, Maheux P, Krauss RM. Insulin resistance and hyperinsulinemia in individuals with small, dense low density lipoprotein particles. *J Clin Invest* 1993;92:141-146.
- [29]. Goldstein BJ. Insulin resistance as the core defect in type 2 diabetes mellitus. *Am J Cardiol* 2002;90:3G-10G.
- [30]. Ginsberg HN, Zhang YL, Hernandez-Ono A. Regulation of plasma triglycerides in insulin resistance and diabetes. *Arch Med Res* 2005;36:232-240.
- [31]. Grundy SM. Atherogenic dyslipidemia associated with metabolic syndrome and insulin resistance. *Clin Cornerstone* 2006;8Suppl 1:S21-7.
- [32]. Kahn R, Buse J, Ferrannini E, et al. The metabolic syndrome: Time for a critical appraisal: Joint statement from the American Diabetes Association and the European Association for the Study of Diabetes. *Diabetes Care* 2005;28:2289–2304.
- [33]. Eckel RH, Grundy SM, Zimmet PZ. The Metabolic Syndrome. *The Lancet* 2005; 365(9468):1415-1428.
- [34]. Miranda PJ, DeFronzo RA, Califf RM, Guyton JR. Metabolic Syndrome: definition, pathophysiology, and mechanisms. *Am Heart J* 2005; 149(1):33-45.
- [35]. Park HS, Oh SW, Cho SI, Choi WH, Kim YS. The Metabolic Syndrome and associated lifestyle factors among South Korean adults. *Int J Epidemiol* 2004; 33(2):328-336.
- [36]. Eliasson B, Attvall S, Taskinen M, Smith U. The insulin resistance syndrome in smokers is related to smoking habits. *Arterioscler Thromb Vasc Biol* 1994; 14(12):1946-1950.
- [37]. Cryer PE, Haymond MW, Santiago JV, Shah SD. Norepinephrine and epinephrine release and adrenergic mediation of smoking-
- [38].

- associated hemodynamic and metabolic events. *N Engl J Med* 1976; 295: 573–7.
- [39]. Kirschbaum C, Wust S, Strasburger CJ (1992) 'Normal' cigarette smoking increases free cortisol in habitual smokers. *Life Sci* 1992; 50: 435–42.
- [40]. Bergman RN, Ader M. Free fatty acids and pathogenesis of type 2 diabetes mellitus. *Trends EndocrinolMetab* 2000; 11: 351–6.
- [41]. Fan AZ, Russell M, Dorn J, Freudenheim JL, Nochajski T, Hovey K, et al. Lifetime Alcohol Drinking Pattern is Related to the Prevalence of Metabolic Syndrome. The Western New York Health Study (WNYHS). *Eur J Epidemiol* 2006; 21(2):129-138.
- [42]. Alkerwi A, Boutsen M, Vaillant M, Barre J, Lair ML, Albert A, et al. Alcohol consumption and the prevalence of Metabolic Syndrome : A meta-analysis of observational studies. *Atherosclerosis* 2009; 204(2):624-635.
- [44]. Kahn SE, Hull RL, Utzschneider KM. Mechanisms linking obesity to insulin resistance and type 2 diabetes. *Nature* 2006;444:840-846.
- [45]. Fujimoto WY. The importance of insulin resistance in the pathogenesis of type 2 diabetes mellitus. *Am J Med* 2000;108Suppl 6a:9S-14S.
- [46]. Grundy SM, Brewer HB, Jr, Cleeman JI, Smith SC, Jr, Lenfant C, American Heart Association., National Heart, Lung, and Blood Institute. Definition of metabolic syndrome: Report of the National Heart, Lung, and Blood Institute/American Heart Association conference on scientific issues related to definition. *Circulation* 2004;109:433-438.
- [47]. Ferrannini E. Is insulin resistance the cause of the metabolic syndrome? *Annals of medicine*. 2006;38(1):42-51.
- [48]. Drager L, F. Queiroz, E. L. Lopes, H. F. Genta, P. R. Krieger, E. M. Lorenzi-Filho, G. Obstructive sleep apnea is highly prevalent and correlates with impaired glycemic control in consecutive patients with the metabolic syndrome. *J CardiometabSyndr*. 2009 Spring;4(2):89-95.
- [49]. Tortosa A, Bes-Rastrollo M, Sanchez-Villegas A, Basterra- Gortari FJ, Nunez-Cordoba JM, Martinez-Gonzalez MA. Mediterranean Diet Inversely Associated with the Incidence of Metabolic Syndrome: the Sun Prospective Cohort. *Diabetes care*. 2007;30(11):2957-2959.
- [50]. Meydani M. A Mediterranean-style diet and metabolic syndrome. *Nutrition reviews* 2005;63:312-4.
- [51]. Prasad, D. S. Kabir, Z. Dash, A. K. Das, B. C. Prevalence and risk factors for metabolic syndrome in Asian Indians: A community study from urban Eastern India. *J Cardiovasc Dis Res* vol (3) issue 3 2012 ; 204-11
- [52]. Hwang LC, Bai CH, Chen CJ, Chien KL. Gender difference on the development of metabolic syndrome: a population based study in Taiwan. *Eur J Epidemiol* 2007; 22:899- 906.
- [53]. Ervin, R. B. Prevalence of metabolic syndrome among adults 20 years of age and over, by sex, age, race and ethnicity, and body mass index: United States, 2003-2006. *Natl Health Stat Report* 2009; Issue 13 1-7.
- [54]. Yousefzadeh G, Sheikhatan M. Age and gender differences in the clustering of metabolic syndrome combinations: A prospective cohort research from the Kerman Coronary Artery Disease Risk Study (KERCADRS). *Diabetes & metabolic syndrome*. 2015;9(4):337-42.
- [55]. McFarlane SI, Banerji M, Sowers JR. Insulin resistance and cardiovascular disease. *J Clin EndocrinolMetab* 2001;86:713-718.
- [56]. Anderson EA, Hoffman RP, Balon TW, Sinkey CA, Mark AL. Hyperinsulinemia produces both sympathetic neural activation and vasodilation in normal humans. *J Clin Invest* 1991;87:2246-2252.
- [57]. Facchini, F. S. Stoohs, R. A. Reaven, G. M. Enhanced sympathetic nervous system activity. The linchpin between insulin resistance, hyperinsulinemia, and heart rate. *Am J Hypertens*;9 Issue 10 Pt 1 : 1013-7
- [58]. Mancia G, Bombelli M, Corrao G, Facchetti R, Madotto F, Giannattasio C, Treveno FQ, Grassi G, Zanchetti A, Sega R. Metabolic syndrome in the Pressioni Arteriose Monitorate E Loro Associazioni (PAMELA) study: daily life blood pressure, cardiac damage, and prognosis. *Hypertension* 2007;49:40-47.
- [59]. Lohsoonthorn V, Dhanamun B, Williams M A. Prevalence of Metabolic syndrome & its relationship to white blood cell count in population of Thai men & women receiving routine health examinations. *Am J Hypertens* 2006; 19: 339-45.
- [60]. Rabin, K. R. Kamari, Y. Avni, I. Grossman, E. Sharabi, Y. Adiponectin: linking the metabolic syndrome to its cardiovascular consequences. *Expert Rev Cardiovasc Ther*. Vol 3, Issue 3. 2005 May;3(3):465-71.
- [61]. Lizardi-Cervera, J. Laparra, D. I. Chavez-Tapia, N. C. Ostos, M. E. Esquivel, M. U. Prevalence of NAFLD and metabolic syndrome in asymptomatic subjects. *Rev Gastroenterol Mex* .2006 Oct-Dec;71(4):453-9.
- [62]. Deedwania PC, Gupta R, Sharma KK, Achari V, Gupta B, Maheshwari A, et al. High prevalence of metabolic syndrome among urban subjects in India: a multisite study. *Diabetes & metabolic syndrome*. 2014;8(3):156-61.
- [63]. Sawant A, Mankeshwar R, Shah S, Raghavan R, Dhongde G, Rajee H, et al. Prevalence of metabolic syndrome in urban India. *Cholesterol*. 2011;2011:920983.
- [64]. Thakur S, Raina S, Thakur S, Negi PC, Verma BS. Prevalence of metabolic syndrome among newly diagnosed hypertensive patients in the hills of Himachal Pradesh, India. *Indian journal of endocrinology and metabolism*. 2013;17(4):723-6.
- [65]. Garg PR, Kabita S, Sinha E, Kalla L, Kaur L, Saraswathy KN. The association of non-HDL cholesterol with the presence of metabolic syndrome in North Indian subjects with and without CAD. *Annals of human biology*. 2013;40(1):111-5.
- [66]. Calo WA, Ortiz AP, Suarez E, Guzman M, Perez CM, Perez CM. Association of cigarette smoking and metabolic syndrome in a Puerto Rican adult population. *Journal of immigrant and minority health / Centre for Minority Public Health*. 2013; 15(4):810-6.
- [67]. Mattoo SK, Chakraborty K, Basu D, Ghosh A, Vijaya Kumar KG, Kulhara P. Prevalence & correlates of metabolic syndrome in alcohol & opioid dependent inpatients. *The Indian journal of medical research*. 2011; 134:341-8.
- [68]. *Harrisons principles of internal medicine* 19 th edition. Volume 2 page no 2450-2452.

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