



A DESCRIPTIVE ANALYSIS FOR TLC AND RBC FOR TYPE 2 DIABETIC

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ABSTRACT

Red cell distribution (RDW) width has been established as being affected in patients with diabetes, especially with poor glycemic control. Further, studies have shown that there is a certain degree of anemia, even in young males with type 2 diabetes, as associated to healthy volunteers. High RDW designates impairment of erythropoiesis, reflecting chronic inflammation and increased levels of oxidative stress. In this study, the mean HB was 12.81 mg/dl with 95% CI of 12.56 - 13.07, which reflects mild anemia. A mean Hb of 11.17 ± 4.42 g/dl in diabetics as compared to 14.11 g/dl in the non-diabetic Indian population. The mean MCV in this study population was 84.52 fl while the mean MCH was 27.59 pg, which are within normal limits, indicating that the anemia observed is probably a normocytic normochromic anemia due to chronic disease. In the present study, the mean RDW was 13.76 (95% CI 13.45-13.95) in the study population. This study also observed that Red cell distribution width (RDW) was meaningfully higher in diabetic patients than in control subjects.

Key Words:- RDW, CI, Hb, MCV, Erythropoiesis, Glycemic control.

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INTRODUCTION

In this section presents introduction of this research work. A major headache with diabetes is that it is an Iceberg disease. Inapparent, the undiagnosed proportion of subjects with diabetes is on the increase in emerging economies like India due to the less developed healthcare systems. (Alberti et al, 1998) The recent IDF atlas estimates that in India alone, around 36.0 million people are living undiagnosed with diabetes, which also

has led to the increasing burden of complications which are irreversible in the long term like the vascular complications especially affecting the eyes, the heart, the kidneys and the nerves. (Thomas CC, Philipson LH, 2015) The study on subjects aged between 20 and 85 years assessed the occurrence of noncommunicable diseases in South India in a classical undeveloped village. (Garcia SD et al, 2013). They observed that the proportion of population classified, based on the HbA1c criteria as Diabetes and Prediabetes was more than 50%. They also observed that in known hypertensive subjects, the proportion with suboptimal control were 40%. (Shaw JE et al, 2010) About one-third of the subjects had elevated cystatin C levels, Dyslipidemia. (Danaei G et al., 2011) In comparison with ICMR-INDIAB study in rural Tamil Nadu, the burden observed in their study was higher. They concluded that 1/3rd to 1/2 of this study population was at risk of cardiovascular disease, with poor control of preexisting cardiovascular risk factors. (Anjana RM et al., 2011) Platelet distribution width was wider in T2DM (0.93, 0.09-1.76; N = 471). They concluded that subjects with Type 2 DM tends to have higher mean platelet volume, higher platelet distribution width values, but a similar platelet count on comparison with subjects without T2DM and the use of these indices

as biomarkers of CVD biomarker needs additional investigation. (Huxley R et al., 2006)

In these articles represents sector 2 of these articles explains the feature on the related works. In section 3 presents the materials and methods adopted and section 4 presents the particulars of the experimentations and discussions. Finally segment 5 accomplishes the articles by allocation our implications and upcoming strategies.

RELATED WORKS

In this segment represents focuses the related works of this research work. The Population with diabetes in India arose from about 32.7 million in 2003 to about 35.5 million in 2006, 40.9 million in 2007, 50.8 million in 2010, 61.3 million in 2011, 65.1 million in 2013 and recently about 69.2 million in 2015. The estimates for 2040 is around 123.5 million. In a multicenter survey conducted from 1972 to 1975 by ICMR across six cities and the adjoining rural areas, the overall national prevalence was projected to be around 2.1%. The chief complication in type 2 diabetes mellitus and in pre-diabetics. (Niskanen L et al., 1998) They made a web search including PubMed, EMBASE, and Web of Science published through 2014 in English and picked up selected case-control and cross-sectional studies providing data on mean platelet volume, width of the platelet distribution, or platelet count in T2DM subjects, IFG, IGT, or metabolic syndrome and compared them with healthy controls. (Unnikrishnan RI et al, 2007) (Pradeepa R et al., 2008) Then, in the year 2001, the National Urban Diabetes Survey which was conducted across six large cities of India, reported an age-standardized prevalence of around 12.1%. The WHO-ICMR study on NCD Risk Factor Surveillance conducted between 2003 to 2005, in 44,523 subjects who were between 15–64 years of age in urban and rural areas of 6 states, estimated the proportion of subjects with self-reported diabetes was 4.5%. (Mohan V et al., 2001)

The large ICMR-INDIA DIABETES is also known as ICMR-INDIAB study. It is an epidemiological study which describes the prevalence at national levels. (Rema M et al., 2005) It estimated the proportion of subjects with diabetes and prediabetes in India by Facing capillary and 2-hour post glucose load. (Demirtas L, et al., 2015) The results of Phase I study reported the proportion of subjects with diabetes among 14000 people in 2011 as 10.4%, 8.4%, 5.3% and 13.6% in states of Tamil Nadu, Maharashtra, Jharkhand, and Chandigarh, separately. They observed that in these states

the prevalence of diabetes in urban areas was greater than in rural areas. Centered on their report in 2011, the total proportion of people living with diabetes was projected to be around 62.4 million in India. Besides this, this study also revealed that India had a large proportion of subjects with prediabetes, about 77 million people, with the impending potential to progress to type 2 DM. In Phase I of the ICMR-INDIAB study, it was imminent that the age for commencement of diabetes was set around 25–34 years. (Panzer S, et al., 1983) Supporting this, the Chennai Urban Rural Epidemiology Study known as CURES study, which was showed in Chennai, south India, about 5% were suffering from diabetes in the age group of 25–34 years. The CURES study, which is a larger and also a more representative study, estimated that the incidence rates per 1000 person-years were around 22.2 for diabetes, about 29.5 for prediabetes and it was about 51.7 for any dysglycemia. And also in the same study population, about 19.4% of the subjects who were normal glucose tolerant and 58.9% of the subjects who were prediabetic advanced to diabetes. (Lippi G, et al., 2014)

MATERIALS AND METHODS

In this part presents the materials and strategies for this exploration work. Spellbinding investigation was done by mean and standard deviation for quantitative factors, recurrence, and extent for absolute factors. Information was likewise spoken to utilizing proper graphs like bar chart, pie outline, and box plots. The affiliation among glycemic control boundaries and different quantitative result factors was surveyed by plotting the information on dissipate plots and figuring individual relationship coefficient. P esteem < 0.05 was considered factually huge. IBM SPSS variant 22 was utilized for factual examination.

STUDY AREA:

Present study was designed to conduct in the clinical settings of Sri Ramakrishna Multi-Specialty Hospitals, Coimbatore, Tamil Nadu, India.

RESULTS AND DISCUSSIONS

In this section focuses the results and discussions of this research work. Various RBC, WBC, and platelet-related hematological indices were considered as primary outcome variables. Blood glucose related parameters were considered as primary explanatory parameters.

Table 1. Descriptive analysis for Total & differential leucocyte count in study population (N=160)

Parameter	Mean ±STD	Median	Min	Max	95% C.I. for EXP(B)	
					Lower	Upper
TLC	8403.13 ± 1401.19	8400	5200	11100	8184.35	8621.90
NEUTROPHILS	59.08 ± 9.37	59.40	32.00	81.00	57.62	60.55
LYMPHOCYTES	28.75 ± 7.38	28.70	8.90	49.70	27.60	29.90
MONOCYTES	6.26 ± 1.62	6.20	2.00	10.70	6.01	6.51
EOSINOPHILS	3.74 ± 2.71	3.10	0.00	17.00	3.31	4.16
BASOPHILES	0.47 ± 0.29	0.40	0.00	1.80	0.42	0.51

Table 2. Descriptive analysis for Red blood cell counts & indices in study population (N=160)

Parameter	Mean ±STD	Median	Min	Max	95% C.I. for EXP(B)	
					Lower	Upper
RBC	4.65 ± 0.53	4.61	3.45	6.38	4.57	4.73
HB	12.81 ± 1.62	12.80	8.90	16.90	12.56	13.07
PCV	39.22 ± 5.42	38.75	20.50	51.90	38.37	40.06
MCV	84.52 ± 6.56	85.20	65.80	99.50	83.50	85.55
MCH	27.59 ± 2.7	27.60	19.40	35.50	27.17	28.01
MCHC	32.38 ± 1.14	32.35	25.60	38.00	32.20	32.56
RDW	13.7 ± 1.6	13.40	9.00	19.60	13.45	13.95

The mean TLC was 8403 ± 1401.19 cells with minimum 5200 cells and the maximum 11100 cells (95% CI 8184.35 to 8621.90). The mean Neutrophils was 59.08 ± 9.37 cells with minimum 32.00 and maximum 81 (95% CI 57.62 to 60.55). The mean Lymphocytes was 28.75 ± 7.38 cells with minimum 8.90 and maximum 49.7 (95% CI 27.60 to 29.90). The mean Monocytes was 6.26 ± 1.62 cells with minimum 2.0 and maximum 10.7 (95% CI 6 to 7.44). The mean Eosinophils was 3.74 ± 2.71 cells with minimum 0 and maximum 17 (95% CI 3.31 to 4.16). The mean Basophils was 0.47 ± 0.29 cells with minimum 0.40 and maximum 1.80 (95% CI 0.42 to 0.51). in the study population.

The mean RBC was 4.65 ± 0.53 cells with minimum 3.45 and the maximum 6.38 (95% CI 4.57 to 4.73). The mean HB was 12.81 ± 1.62 g/dL with minimum 8.90 and maximum 16.90 (95% CI 12.56 to 13.07). The mean PCV was 39.22 ± 5.42 % with minimum 20.50 and maximum 51.90 (95% CI 38.37 to 40.06). The mean MCV was 84.52 ± 6.56 fL with minimum 65.80 and maximum 99.50 (95% CI 83.50 to 85.55). The mean MCH was 27.59 ± 2.7 pg with minimum 25.60 and maximum 35.5 (95% CI 27.17 to 28.01). The mean MCHC was 32.28 ± 1.14 % with minimum 25.60 and maximum 38 (95% CI 32.20 to 32.56). The mean RDW was 13.76 ± 1.6% with minimum 9.00 and maximum 19.60 (95% CI 13.45 to 13.95) in the study population.

In order to establish a relationship between inflammation and control of diabetes, the present study evaluated changes in neutrophil-lymphocyte ratio (NLR) with changes in FBS, PPBS and HbA1c. It was found that there was a statistically significant association among and all three parameters namely, FBS, PPBS and

HbA1c and Neutrophil lymphocyte ratio and this correlation was found to be statistically significant (P value < 0.05). Similar to this, Shiny et al. (2014)⁽¹⁹⁾ also observed a significant positive correlation. Buyukkaya E et al. (2014)⁽⁵⁸⁾ also observed a significant correlation between the Metabolic syndrome criteria and inflammation as described by increase in NLR. In a current study from india by khandare et al (2017)⁽⁶⁶⁾, NLR was found to be higher in patient with diabetic nephropathy. hence it can be inferred that NLR can predict poor glycemic control and diabetic patients with high NLR should be carefully monitored to prevent diabetic complications. The RDW tended to linearly increase with increase in HbA1c levels. Nada A M (2015)⁽²⁰⁾ in their study also observed that Red cell distribution width (RDW) was higher in patients with uncontrolled glycemia (HbA1c >7%) than those with good control (HbA1c ≤7%; P=0.035). Lippi G et al (2014)⁽¹⁵⁾ also demonstrated that HbA1c was significantly positively related with RDW values even after alteration for age and gender.

CONCLUSION

Finally this work concludes that the Platelet lymphocyte ratio (PLR) is being increasingly recognized as a prophet of outcome in patient hospitalized with diabetes. Mertoglu et al⁽⁶⁵⁾ reported that NLR meaningfully upsurges in prediabetic and diabetic patients, while PLR meaningfully decreases in prediabetes and early stages of diabetes but upsurges in later stages. The present study, however, failed to demonstrate a statistically significant correlation between PLR and glycemic parameters (FBS, PPBS and HbA1c).

This may be because the study is not adequately powered to detect the association.

In the present study, there was no statistically significant association among platelet count and glycemic

control. The relationship between HbA1c and platelet indices such as Mean platelet volume and Platelet distribution width were not evaluated in this study.

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