Open Access Macedonian Journal of Medical Sciences. 2014 Mar 15; 2(1):83-88. http://dx.doi.org/10.3889/oamjms.2014.015 *Clinical Science*

Effects of Glucose Control on Hematological Indices in Patients with Diabetes Mellitus

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Abstract

Citation: Varım C, Kaya T, Nalbant A, Uçar A, Tamer A. Effects of Glucose Control on Hematological Indices in Patients with Diabetes Mellitus. OA Maced J Med Sci. 2014 Mar 15; 7(1):83-88.

http://dx.doi.org/10.3889/oamjms.2014.015 Key words: Diabetes Mellitus; hemogram;

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Received: 17-Jan-2014; Revised: 30-Jan-2014; Accepted: 04-Feb-2014; Online first: 20-Feb-2014

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Competing Interests: The authors have declared that no competing interests exist.

Aim: We aimed to investigate the effects of diabetes treatment modalities on haematological parameters and leukocyte formula in patients with type 2 diabetes mellitus.

Materials and Methods: The study included 102 patients with type 2 diabetes, out of which 51 receiving insulin treatment and 51 receiving oral antidiabetics (OAD). Hemogram data of insulin and OAD treated groups were compared.

Results HbA1c levels were $11.12 \pm 2.09 \text{ mg/dl}$ in insulin group and $7.94 \pm 2.1 \text{ mg/dl}$ in OAD group p=0.001. Platelet counts were 27866.67 ± 77693 10⁹/L before treatment and 258941.18 ± 69068.2 10⁹/L in OAD group at six months, p: 0.015 whereas; 293011.76 ± 73711.21 10⁹/L before treatment and 289492.86 ± 82631.49 10⁹/L in insulin group at six months p: 0.821. Monocyte counts were 0.47 ± 0.12 10⁹/L before the treatment and 0.57 ± 0.12 10⁹/L in mix insulin therapy subgroup at six months, p:0.004; monocyte percentage was % 6.11 ± 1.74 before the treatment and %7.51 ± 2.57 in mix insulin subgroup at six months p:0.039; Basophiles counts were 0.1 ± 0.02 10⁹/L before treatment and 0.09 ± 0.04 10⁹/L in intensive insulin therapy subgroup at six months, p: 0.005; Lymphocyte and basophils counts were significantly decreased at six months insulin treatment as compared to the pretreatment values.

Conclusion: This study showed that, glucose control effects; blood indices HbA1C, basophiles, eosinophils, platelets and lymphocytes counts.

Introduction

Diabetes Mellitus (DM), is a multisystemic disease characterized by high blood sugar levels causing acute and chronic complications. High blood sugar levels caueses these complications. Diabetes mellitus is known to cause anemia of chronic disease, erythrocyte, leukocyte and platelet dysfunction [1-4]. Oral antidiabetics and insulins are used for the treatment of type 2 DM for many years. These treatment models can be applied separately or in combination. The aim of the treatment is reduction of diabetic complications by providing blood sugar regulation and suppression of chronic inflammatory process.

The effects of the various medical treatment modalities for patients with type 2 DM upon the hematological parameters have been analyzed, showing mainly no significant side effects, the most common side effects being allergic reactions. The increase in the peripheral blood eosinophils, basophils, and leukocyte counts seen in allergic reactions is also seen in chronic inflammatory processes, but the latter can be improved by regulation of blood sugar. This regulation also reduces the number of inflammatory cells in the peripheral blood.

The aim of our study was to evaluate the effects of glucose control on the hematological parameters in the study population.

Materials and Methods

We have performed a retrospective analysis of the data of 102 patients with type 2 diabetes mellitus, which have been followed-up at the Diabetes Polyclinic of our hospital for at least 6 months. The diagnosis of type 2 diabetes was based upon the diagnostic criteria of the American Diabetes

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Association (ADA) from 2010. Venous blood samples were obtained from subjects for hemogram studies. The patients treated with insulin were divided into three subgroups: patients receiving conventional insulin therapy regime (short/intermediate acting insulin mixture 2 times a day), patients on intensive insulin regime (short-acting insulin 3 times a day and basal long-acting insulin) and patients using OAD in combination with basal insulin. The patients using oral antidiabetic therapy were also divided into 3 subgroups: patients using metformin, patients using metformin and sulfonylurea and patients using metformin and DPP4 analogues. Haematologic parameters and HbA1c levels were measured at the initiation of the therapy and after 3 and 6 months

 Table 1: Distributions of the demographic and hematological parameters according to insulin and OAD groups.

| | | insulin (n=51) | OAD (n=51) | ·p | | |
|---------------------|-----------------------|-----------------------|--------------------------|--------|--|--|
| Age (year) | | 55.10 ± 10.86 | 53.51 ± 10.10 | 0.446 | | |
| Gender (male) | | 22 (43.1) | 26 (51.0) | 0.552 | | |
| DM duration (month) | | 117.45 ± 84.94 | 43.31 ± 45.96 | <0.001 | | |
| | Pre-Treatment | 11.12 ± 2.09 | 7.94 ± 2.11 | <0.001 | | |
| | Post-Treat. 3rd Month | 9.17 ± 2.01 | 6.78 ± 1.25 | <0.001 | | |
| Hba1c | Post-Treat. 6th Month | 8.68 ± 1.95 | 7.13 ± 1.10 | <0.001 | | |
| | ² p | <0.001 ^{a,b} | 0.001 ^{a,b} | | | |
| | ³ p | <0.001 | | | | |
| | Pre-Treatment | 8072.55 ± 2050.76 | 8200.00 ± 2068.82 | 0.755 | | |
| | Post-Treat. 3rd Month | 8410.00 ± 2033.62 | 8050.98 ± 2034.44 | 0.377 | | |
| WBC | Post-Treat. 6th Month | 8442.86 ± 2460.14 | 7805.88 ± 1728.86 | 0.147 | | |
| | ² p | 0.192 | 0.320 | | | |
| | ³ p | 0.4 | 44 | | | |
| | Pre-Treatment | 13.02 ± 1.75 | 13.36 ± 1.63 | 0.312 | | |
| | Post-Treat. 3rd Month | 12.98 ± 1.53 | 13.30 ± 1.57 | 0.305 | | |
| Hb | Post-Treat. 6th Month | 12.85 ± 1.56 | 13.21 ± 1.39 | 0.256 | | |
| | ² p | 0.985 | 0.663 | | | |
| | ³ p | 0.166 | | | | |
| | Pre-Treatment | 293011.76 ± 73711.21 | 278666.67 ± 77693.93 | 0.341 | | |
| | Post-Treat. 3rd Month | 285706.00 ± 72484.48 | 276352.94 ± 75820.00 | 0.528 | | |
| Plt | Post-Treat. 6th Month | 289492.86 ± 82631.49 | 258941.18 ± 69068.20 a,b | 0.055 | | |
| | ² p | 0.821 | 0.015 ^{a,c} | | | |
| | ³ p | 0.2 | 87 | | | |
| | Pre-Treatment | 85.12 ± 6.64 | 85.12 ± 6.34 | 1.000 | | |
| | Post-Treat. 3rd Month | 84.88 ± 5.99 | 84.70 ± 5.90 | 0.883 | | |
| MCV | Post-Treat. 6th Month | 84.31 ± 6.32 | 85.37 ± 5.69 | 0.398 | | |
| | ² p | 0.074 | 0.124 | | | |
| | ³ p | 0.8 | 64 | | | |
| | Pre-Treatment | 9.59 ± 11.4 | 8.05 ± 1.27 | 0.341 | | |
| MPV | Post-Treat. 3rd Month | 9.50 ± 11.59 | 7.84 ± 1.12 | 0.311 | | |
| | Post-Treat. 6th Month | 7.87 ± 1.21 | 8.06 ± 1.17 | 0.461 | | |
| | ² p | 0.291 | 0.236 | | | |
| | ³ n | 03 | 12 | | | |

³p 0.312 Data were shown as mean ± standard deviation and n (%). ¹: Results of the comparison between insulin and OAD groups. ²: Results of the comparison among pre-treatment, post-treatment 3rd month and post-treatment 6th month measures. ³: Results of the comparison among two groups according to alterations of hematologic parameters. ⁴: There was statistically significant difference between pre-treatment and post-treatment 3rd month. ⁵: There was statistically significant difference between pre-treatment and post-treatment 6th month. ^c: There was statistically significant difference between post-treatment 3rd month and post-treatment 6th month.

In the study we investigated the effect of the decrease in HbA1c levels to blood count and leukocyte formula in the 6-month treatment period for insulin and OAD using groups and also for the subgroups Patients are started on insulin therapy compared with patients are using oral antidiabetics. We investigate the effect of decrease in HbA1c levels to blood count and leukocyte formula in 6-month treatment period for insulin and OAD using groups and also for subgrups.

All patients were informed for the study and signed an informed consent forms for the study. Ethics committee approval was taken for the study.

Statistical analysis was made using computer software (SPSS version 15.0, SPSS Inc. Chicago, IL, USA). The results of all the parameters of insulin and OAD group are given as mean ± standard deviation. Samples T test was used to compare the data between the two groups. Level of statistical significance of the data interpreted with the 'p' value. p <0.05 was considered statistically significant.

Results

Patients were divided into two groups according to using OAD therapy and using insulin therapy. The average age of the insulin using group is 55.1 years, and 53.5 years of OAD using group. The distribution of gender is, 22 male/33 female patients (n = 55) in insulin using group and 26 male/29 female patients (n = 55) in OAD using group.. Duration of diabetes was found 117.45 ± 84.94 months in insulin using group and 43.31 ± 45.96 months in OAD using group (Table 1).

 Table 2: Distributions of the hematological parameters according to the insulin and OAD groups.

| | | Insulin (n=51) | OAD (n=51) | ¹ p | | | |
|----------|-----------------------|----------------|--------------|----------------|--|--|--|
| | Pre-Treatment | 57.59 ± 9.65 | 58.23 ± 8.65 | 0.724 | | | |
| | Post-Treat. 3rd Month | 57.79 ± 8.76 | 58.68 ± 7.64 | 0.590 | | | |
| Neu % | Post-Treat. 6th Month | 58.79 ± 9.27 | 57.54 ± 7.2 | 0.466 | | | |
| | ² p | 0.465 | 0.548 | _ | | | |
| | ³ p | 8.0 | 195 | | | | |
| | Pre-Treatment | 4.68 ± 1.45 | 4.78 ± 1.46 | 0.730 | | | |
| | Post-Treat. 3rd Month | 4.91 ± 1.63 | 4.76 ± 1.50 | 0.624 | | | |
| Neu | Post-Treat. 6th Month | 4.99 ± 1.77 | 4.52 ± 1.23 | 0.143 | | | |
| | ² p | 0.220 | 0.413 | _ | | | |
| | °p | 0.5 | 518 | | | | |
| | Pre-Treatment | 32.56 ± 9.12 | 32.04 ± 7.26 | 0.753 | | | |
| | Post-Treat. 3rd Month | 32.73 ± 8.41 | 31.64 ± 7.25 | 0.486 | | | |
| Lym % | Post-Treat. 6th Month | 31.35 ± 8.35 | 32.82 ± 7.18 | 0.365 | | | |
| | <u>p</u> | 0.807 | 0.438 | _ | | | |
| | °p | 0.9 | 159 | | | | |
| | Pre-Treatment | 2.72 ± 0.99 | 2.69 ± 1.01 | 0.905 | | | |
| | Post-Treat. 3rd Month | 2.74 ± 0.92 | 2.61 ± 0.98 | 0.486 | | | |
| Lym | Post-Treat. 6th Month | 2.60 ± 0.98 | 2.62 ± 0.88 | 0.935 | | | |
| | <u>²p</u> | 0.597 | 0.631 | _ | | | |
| | °p | 0.8 | 16 | | | | |
| | Pre-Treatment | 6.57 ± 2.35 | 7.01 ± 2.33 | 0.343 | | | |
| | Post-Treat. 3rd Month | 6.55 ± 2.34 | 6.70 ± 2.62 | 0.756 | | | |
| Mono % | Post-Treat. 6th Month | 6.92 ± 2.56 | 6.75 ± 1.87 | 0.710 | | | |
| | <u>p 0.719 0.496</u> | | | | | | |
| | -p | 0.9 | 106 | | | | |
| | Pre-Treatment | 0.50 ± 0.16 | 0.58 ± 0.24 | 0.069 | | | |
| | Post-Treat. 3rd Month | 0.54 ± 0.19 | 0.52 ± 0.2 | 0.601 | | | |
| Mono | Post-Treat. 6th Month | 0.55 ± 0.16 | 0.53 ± 0.19 | 0.507 | | | |
| | р 3- | 0.206 | 0.165 | _ | | | |
| | p Des Transfer est | 0.9 | 0.00 + 0.00 | 0.070 | | | |
| | Pre-Treatment | 2.24 ± 1.6 | 2.38 ± 2.08 | 0.876 | | | |
| E a a 0/ | Post-Treat. 3rd Month | 2.10 ± 1.28 | 1.93 ± 1.27 | 0.508 | | | |
| EOS % | Post-freat. 6th Month | 2.10 ± 1.44 | 2.04 ± 1.35 | 0.631 | | | |
| | <u> </u> | 0.400 | 0.277 | - | | | |
| | Pro-Troatmont | 0.19 ± 0.13 | 0.20 ± 0.22 | 0.970 | | | |
| | Post-Troat 3rd Month | 0.18 ± 0.13 | 0.20 ± 0.22 | 0.070 | | | |
| For | Post-Treat. Std Month | 0.18 ± 0.12 | 0.10 ± 0.12 | 0.479 | | | |
| 205 | ² n | 0.1010.12 | 0.17 ± 0.12 | 0.490 | | | |
| | - <u>P</u> | 0.313 | 68 | - | | | |
| | P Pre-Treatment | 0.90 + 0.38 | 0.88 + 0.39 | 0 776 | | | |
| | Post-Treat 3rd Month | 0.83 + 0.29 | 0.9 + 0.41 | 0.328 | | | |
| Baso% | Post-Treat 6th Month | 0.84 + 0.28 | 0.88 + 0.34 | 0.611 | | | |
| 240070 | ² D | 0 266 | 0.843 | 0.011 | | | |
| | 3 ⁰ | 0.200 | 67 | - | | | |
| | Pre-Treatment | 0.08 ± 0.04 | 0.08 ± 0.06 | 0.551 | | | |
| | Post-Treat, 3rd Month | 0.07 ± 0.05 | 0.08 ± 0.06 | 0.535 | | | |
| Baso | Post-Treat 6th Month | 0.08 + 0.04 | 0.07 + 0.05 | 0.348 | | | |
| Da50 | 2 _D | 0.225 | 0.741 | 5.0-10 | | | |
| | | 0.220 | 700 | - | | | |

Data were shown as mean ±standard deviation. ¹: Results of the comparison between insulin and OAD groups. ²: Results of the comparison among pre-treatment, post-treatment 3rd month and post-treatment 6th month measures. ³: Results of the comparison among two groups according to alterations of hematologic parameters.

Hematological parameters such as platelet and white blood cell count, hemoglobin, hematocrite, mean corpuscular volume (MCV), mean platelet volume (MPV), white blood cell formulas (neutrophils, lymphocytes, eosinophils, basophils, monocytes percentages) and HbA1c levels were examined in both groups. The start of study, 3 and 6 months, these parameters and HbA1c levels were recorded. Changes in these parameters and HbA1c levels were assessed between insulin and OAD using group and as well as its subgroups. HbA1c levels were %11.12 \pm 2.09 mg/dl in insulin group and %7.94 \pm 2.1 in OAD group p = 0.001. Platelet counts were 27866.67 \pm 77693 10⁹/L before treatment and 258941.18 \pm 69068.2 10⁹/L in OAD using group at six months, p: 0.015 whereas; 293011.76 \pm 73711.21 109/L before treatment and 289492.86 \pm 82631.49 10⁹/L, in insulin theraphy group at six months p: 0.821 (Table 1 and 2).

Table 3: Distributions of the demographic and hematological parameters according to the insulin subgroups.

| | | Mix insulin | Intensive insulin | OAD+Basal | |
|---------|--------------------------|--------------------|--------------------|---------------------|-------|
| | | (n=20) | (n=19) | insulin (n=12) | 'p |
| Age (ye | ar) | 55.5 ± 8.38 | 53.37 ± 13.01 | 57.17 ± 11.3 | 0.633 |
| Gender | (male) | 9 (45) | 8 (42.1) | 5 (41.7) | 0.977 |
| DM dur | ation (month) | 126 ± 76.94 | 110 ± 106.26 | 115 ± 62.14 | 0.841 |
| | Pre-Treatment | 11.16 ± 1.86 | 11.68 ± 2.04 | 10.15 ± 2,35 | 0.137 |
| | Post-Treat. 3rd Month | 9.31 ± 2.05 | 9.55 ± 2.27 | 8.34 ± 1.31 | 0.279 |
| Hba1c | Post-Treat. 6th Month | 8.45 ± 1.6 | 9.49 ± 2.28 | 7.88 ± 1.52 | 0.106 |
| | ²p | 0.010 ^b | 0.024 ^b | 0.005 * | |
| | 3р | | 0.100 | | |
| | Pre-Treatment | 7970 ± 1522.84 | 8810.53 ± 2459.43 | 7075 ± 1779.75 | 0.066 |
| | Post-Treat. 3rd Month | 8705 ± 2425.41 | 8678.95 ± 1617.12 | 7409.09 ± 1722.47 | 0.183 |
| WBC | Post-Treat. 6th Month | 8131.25 ± 2448.6 | 9426.67 ± 2449.92 | 7554.55 ± 2223.23 | 0.129 |
| | ² p | 0.161 | 0.298 | 0.631 | |
| | ³ p | | 0.079 | | |
| | Pre-Treatment | 12.86 ± 1.67 | 13.25 ± 1.91 | 12,.3 ± 1.75 | 0.778 |
| | Post-Treat. 3rd Month | 12.91 ± 1.27 | 13.15 ± 1.95 | 12.84 ± 1.22 | 0.836 |
| Hb | Post-Treat. 6th Month | 12.41±1.5 | 13.39 ± 1.8 | 12.72 ± 1.13 | 0.217 |
| | ²p | 0.567 | 0.630 | 0.801 | |
| | 3р | | 0.451 | | |
| | Pre-Treatment | 294000 ± 62839 | 300263.2 ± 72371.6 | 279883.3 ± 95000.8 | 0.760 |
| | Post-Treat. 3rd Month | 284300 ± 65384 | 294789.5 ± 72900.7 | 272572.7 ± 87780.5 | 0.725 |
| Plt | Post-Treat. 6th Month | 290375 ± 54585.6 | 305600 ± 92041.8 | 266245.5 ± 103344.2 | 0.497 |
| | ²p | 0.781 | 0.199 | 0.406 | |
| | 3р | | 0.806 | | |
| | Pre-Treatment | 85.81 ± 7.59 | 85.05 ± 5.52 | 84.07 ± 7.01 | 0.779 |
| | Post-Treat. 3rd Month | 84.97 ± 7.05 | 84.83 ± 5.79 | 84.78 ± 4.57 | 0.996 |
| MCV | Post-Treat. 6th Month | 84.38 ± 6.56 | 84.23 ± 7.45 | 84.32 ± 4.65 | 0.998 |
| | ² p | 0.037 ^a | 0.279 | 0.758 | |
| | 3р | | 0.933 | | |
| | Pre-Treatment | 8.21 ± 1.29 | 11.74 ± 18.58 | 8.49 ± 3.2 | 0.591 |
| MPV | Post-Treat. 3rd Month | 7.88 ± 0.96 | 11.81 ± 18.77 | 8.46 ± 2.5 | 0.550 |
| | Post-Treat. 6th Month | 7.76 ± 0.99 | 7.78 ± 1.26 | 8.16 ± 1.46 | 0.661 |
| | ² p | 0.056 | 0.350 | 0.775 | |
| | 3 | | | | |

Data were shown as mean ±standard deviation and n (%). ¹: Results of the comparison among groups. ²: Results of the comparison among pre-treatment, post-treatment 3rd month and post-treatment 6th month measures. ³: Results of the comparison among three groups according to alterations of hematologic parameters. ³: There was statistically significant difference between pre-treatment and post-treatment 3rd month.

Insulin treatment group was divided into 3 subgroups. These subgroups consisted of patients initiated on mixed insulin therapy, patients started on intensive insulin therapy and patients using OAD therapy in combination with basal insulin. Hematological parameters, white blood cell formulas and HbA1c levels were examined in both subgroups. HbA1c levels were %11.16 ± 1.86 before treatment and % 8.45 ± 1.6 in mix insulin therapy subgroup at six months p: 0.010; HbA1c levels were %11.68 ± 2.04 before treatment and %9.49 ± 2.28 in intensive insulin therapy subgroup at six months p = 0.024; HbA1c levels were %10.15 ± 2.35 before treatment and % 7.88 ± 1.52 in OAD therapy and added basal insulin subgroup at six months p = 0.005.

Table 4: Distributions of the hematological parameters according to the insulin subgroups.

| | | Mix insulin (n=20) | Intensive insulin (n=19) | OAD+Basal insulin (n=12) | ¹ p |
|-----------|------------------------|-----------------------|--------------------------------|--------------------------------|-------------------------|
| Nou | Pre-Treatment | 57.17 ± 10.78 | 56.74 ± 10.03 | 59.65 ± 7.19 | 0.701 |
| | Post-Treat. 3rd Month | 56.98 ± 9.82 | 57.53 ± 7.31 | 59.73 ± 9.54 | 0.703 |
| % | Post-Treat. 6th Month | 55.58 ± 10.32 | 59.27 ± 8.03 | 62.81 ± 8.24 | 0.133 |
| ,0 | ² p | 0.839 | 0.162 | 0.195 | _ |
| | °р | | 0.428 | | |
| | Pre-Treatment | 4.63 ± 1.38 | 5.04 ± 1.66 | 4.21 ± 1.15 | 0.301 |
| | Post-Treat. 3rd Month | 5.07 ± 2.15 | 5.01 ± 1.17 | 4.45 ± 1.19 | 0.568 |
| Neu | Post-Treat. 6th Month | 4.53 ± 1.79 | 5.6 ± 1.68 | 4.82 ± 1.79 | 0.230 |
| | <u></u> | 0.303 | 0.125 | 0.354 | _ |
| | р | | 0.389 | | |
| | Pre-Treatment | 33.99 ± 9.08 | 32.86 ± 9.82 | 29.73 ± 8.15 | 0.443 |
| Lvm | Post-Treat. 3rd Month | 33.74 ± 8.67 | 33.36 ± 6.95 | 29.83 ± 10.26 | 0.436 |
| % | Post-Treat. 6th Month | 33.63 ± 7.72 | 32.05 ± 8.68 | 27.09 ± 7.85 | 0.124 |
| | <u>-p</u> | 0.993 | 0.984 | 0.226 | _ |
| | ⁻ p | 0.00 + 0.00 | 0.209 | 0.40 + 0.70 | 0.000 |
| | Pre-Treatment | 2.68 ± 0.82 | 3.14 ± 1.14 | 2.13 ± 0.73 | 0.020 |
| 1 | Post-Treat. 3rd Month | 2.88 ± 0.94 | 2.91 ± 0.88 | 2.22 ± 0.83 | 0.099 |
| Lym | Post-I reat. 6th Month | 2.59 ± 0.71 | 3.03 ± 1.22 | 2.02 ± 0.66 | 0.029 |
| | | 0.108 | 0.330 | 0.385 | - |
| | P Dro Trootmont | 6 11 + 1 74 | 6.57 + 2.09 | 7 2 ± 1 70 | 0.204 |
| | Pre-Treatment | 6.11±1.74 | 0.57 ± 3.08 | 7.3±1.78 | 0.394 |
| Mono | Post-Treat. 3rd Month | 0.49 ± 1.89 | 0.10 ± 2.0 | 7.30 ± 2.04 | 0.409 |
| % | 2 2 | 7.51±2.57 | 0.470 | 7.30 ± 3.03 | 0.100 |
| | | 0.039 | 0.178 | 0.991 | - |
| | P Pro-Troatmont | 0.47 ± 0.12 | 0.571 | 0.40 ± 0.11 | 0.403 |
| | Post-Troat 3rd Month | 0.47 ± 0.12 | 0.54 ± 0.21 | 0.49 1 0.11 | 0.403 |
| Mono | Post-Treat. Stu Month | 0.50 ± 0.17 | 0.52 ± 0.2 | 0.34 ± 0.2 | 0.810 |
| WIGHO | ² n | 0.004 a.b | 0.00 ± 0.10 | 0.03 1 0.25 | 0.010 |
| | 3n | 0.004 | 0.865 | 0.755 | - |
| | Pre-Treatment | 2 15 + 1 76 | 2 27 + 1 6 | 2 33 + 1 44 | 0 949 |
| | Post-Treat 3rd Month | 1 94 + 0 94 | 2 23 + 1 5 | 2 17 + 1 5 | 0 774 |
| Eos | Post-Treat 6th Month | 2 27 + 1 89 | 1 99 + 1 18 | 1 99 + 0 99 | 0.828 |
| % | ² n | 0.605 | 0.042 ° | 0.687 | |
| | 3 _D | 0.000 | 0.944 | 0.001 | - |
| | Pre-Treatment | 0.16 ± 0.12 | 0.2 ± 0.13 | 0.18 ± 0.14 | 0.635 |
| | Post-Treat, 3rd Month | 0.17 ± 0.08 | 0.18 ± 0.11 | 0.16 ± 0.15 | 0.830 |
| Eos | Post-Treat. 6th Month | 0.19 ± 0.13 | 0.19 ± 0.12 | 0.15 ± 0.09 | 0.550 |
| | ² p | 0.691 | 0.685 | 0.574 | |
| | ³ p | | 0.645 | | - |
| | Pre-Treatment | 0.89 ± 0.38 | 0.85 ± 0.34 | 0.99 ± 0.44 | 0.601 |
| Deee | Post-Treat. 3rd Month | 0.85 ± 0.3 | 0.76 ± 0.3 | 0.92 ± 0.26 | 0.331 |
| Baso | Post-Treat. 6th Month | 0.91 ± 0.24 | 0.83 ± 0.3 | 0.76±0.33 | 0.446 |
| 70 | ² p | 0.737 | 0.239 | 0.143 | _ |
| | ³ p | | 0.759 | | |
| | Pre-Treatment | 0.08 ± 0.04 | 0.1 ± 0.02 | 0.08 ± 0.05 | 0.212 |
| | Post-Treat. 3rd Month | 0.08 ± 0.04 | 0.06 ± 0.05 | 0.08 ± 0.04 | 0.525 |
| Baso - | Post-Treat. 6th Month | 0.09 ± 0.03 | 0.09 ± 0.04 | 0.06 ± 0.05 | 0.029 _{e,f} |
| | ² p | 0.167 | 0.005 ^{a.c} | 0.092 | _ |
| | 3 | | 0 554 | | |

 ^{cp}
 0.554

 Data were shown as mean ±standard deviation and n (%). ¹: Results of the comparison among groups. ²: Results of the comparison among pre-treatment, post-treatment 3rd month and post-treatment 6th month measures. ³: Results of the comparison among three groups according to alterations of hematologic parameters. ⁴: There was statistically significant difference between pre-treatment and post-treatment 3rd month. ⁶: There was statistically significant difference between post-treatment 3rd month. ⁶: There was statistically significant difference between post-treatment 3rd month. ⁶: There was statistically significant difference between post-treatment 3rd month and post-treatment 6th month. ⁶: There was statistically significant difference between group 2 and group 3. ¹: There was statistically significant difference between group 2 and group 3. ¹: There was statistically significant difference between group 2 and group 3. ¹: There was statistically significant difference between group 2 and group 3. ¹: There was statistically significant difference between group 2 and group 3. ¹: There was statistically significant difference between group 2 and group 3. ¹: There was statistically significant difference between group 2 and group 3. ¹: There was statistically significant difference between group 2 and group 3. ¹: There was statistically significant difference between group 2 and group 3. ¹: There was statistically significant difference between group 4 and group 3. ¹: There was statistically significant difference between group 4 and group 3. ¹: There was statistically significant difference between group 4 and group 3. ¹: There was statistically significant difference between group 4 and group 3. ¹: There was statistically significant difference between group 5 and group 3. ¹: There was statistically significant difference between group 5 and group 3. ¹: There was statistically signif

Monocyte counts were $0.47 \pm 0.12 \ 10^{9}/L$ before the treatment and 0.57 ± 0.12 10⁹/L in mix insulin therapy subgroup at six months, p = 0.004; monocyte percentage was % 6.11 ± 1.74 before the treatment and %7.51 ± 2.57 in mix insulin therapy subgroup at six months p = 0.039; Basophils counts were 0.1 \pm 0.02 10⁹/L before treatment and 0.09 \pm $0.04 \ 10^{9}$ /L in intensive insulin therapy subgroup at six months, p: 0.005; Eosinophil percentage was %2.27 ± 1.6 before treatment and %1.99 ± 1.18 in intensive insulin using subgroup at six months, p: 0.042. Lymphocyte and basophils counts were significantly decreased at six months insulin treatment as compared to the pretreatment values. There was no statistically significant difference found in other parameters (Table 3 and 4).

| Table 5: Distributions of the demographic and hematologic | al |
|---|----|
| parameters according to the OAD subgroups. | |

| | | Metformin (n=13) | Metformin + sulfonylurea (n=23) | Metformin + DPP4 analogues (n=15) | ¹ p |
|----------|--------------------------|---------------------|---------------------------------------|---|----------------|
| Age (yea | ar) | 50.23 ± 9.19 | 53.3 ± 9.87 | 56.67 ± 10.88 | 0.245 |
| Gender | (male) | 4 (30.8) | 13 (56.5) | 9 (60) | 0.235 |
| DM dura | ation (month) | 17.92 ± 20.62 | 54.17 ± 54.13 | 47.54 ± 40.26 | 0.077 |
| | Pre-Treatment | 8.17 ± 2.59 | 7.88 ± 1.86 | 7.85 ± 2.15 | 0.910 |
| | Post-Treat. 3rd Month | 5.93 ± 0.8 | 7.42 ± 1.35 | 6.54 ± 0.85 | 0.001 e,f |
| Hba1c | Post-Treat. 6th Month | 6.72 ± 0.92 | 7.32 ± 1.16 | 7.2 ± 1.12 | 0.286 |
| | ²p | 0.025 ° | 0.227 | 0.071 | |
| | 3р | | 0.243 | | |
| | Pre-Treatment | 8130.77 ± 2756.32 | 8386.96 ± 1704.35 | 7973.33 ± 2022.89 | 0.832 |
| | Post-Treat. 3rd Month | 7923.08 ± 3016.39 | 8321.74 ± 1856.43 | 7746.67 ± 1154.41 | 0.681 |
| WBC | Post-Treat. 6th Month | 7430.77 ± 2144.13 | 8052.17 ± 1621.13 | 7753.33 ± 1539.88 | 0.588 |
| | ²p | 0.460 | 0.654 | 0.869 | |
| | ³ p | | 0.654 | | |
| | Pre-Treatment | 13.59 ± 1.76 | 13.36 ± 1.5 | 13.17 ± 1.8 | 0.795 |
| | Post-Treat. 3rd Month | 13.21 ± 1.16 | 13.39 ± 1.57 | 13.24 ± 1.94 | 0.933 |
| Hb | Post-Treat. 6th Month | 13.13 ± 1.18 | 12.95 ± 1.09 | 13.66 ± 1.88 | 0.309 |
| | ²p | 0.075 | 0.075 | 0.493 | |
| | ³ p | | 0.963 | | |
| | Pre-Treatment | 280923.1 ± 79898.3 | 278739.1 ± 76384.9 | 276600 ± 83117.7 | 0.990 |
| | Post-Treat. 3rd Month | 274615.4 ± 68476.9 | 283695.7 ± 86306.2 | 266600 ± 67921.0 | 0.797 |
| Plt | Post-Treat. 6th Month | 265230.8 ± 63989.8 | 262260.9 ± 72357.8 | 248400 ± 71596.9 | 0.782 |
| | ² p | 0.333 | 0.193 | 0.164 | |
| | зр | | 0.883 | | |
| | Pre-Treatment | 84.56 ± 8.41 | 84.84 ± 6.01 | 86.03 ± 4.99 | 0.803 |
| | Post-Treat. 3rd Month | 84.5 ± 7.63 | 84.42 ± 5.02 | 85.31 ± 5.84 | 0.895 |
| MCV | Post-Treat. 6th Month | 84.92 ± 6.71 | 85.03 ± 5.2 | 86.28 ± 5.78 | 0.768 |
| | ² p | 0.766 | 0.448 | 0.322 | |
| | ³ p | | 0.819 | | |
| | Pre-Treatment | 8.19 ± 1.12 | 8.21 ± 1.53 | 7.69 ± 0.89 | 0.421 |
| MPV | Post-Treat. 3rd Month | 7.99 ± 0.97 | 7.91 ± 1.32 | 7.59 ± 0.93 | 0.604 |
| | Post-Treat. 6th Month | 8.39 ± 1.55 | 7.87 ± 0.94 | 8.06 ± 1.12 | 0.439 |
| | ² p | 0.443 | 0.184 | 0.153 | |
| | 4 | | 0.000 | | |

^{*}p 0.579 Data were shown as mean ±standard deviation and n (%). ¹: Results of the comparison among pre-treatment, post-treatment 3rd month and post-treatment 6th month measures. ³: Results of the comparison among three groups according to alterations of hematologic parameters. ⁸: There was statistically significant difference between pre-treatment and post-treatment 3rd month. ⁸: There was statistically significant difference between group 1 and group 3. ¹: There was statistically significant difference between group 1 and group 3. ¹: There was statistically significant difference between group 1 and group 3.

The group receiving oral antidiabetic therapy was divided into 3 subgroups. These subgroups consisted of patients using only metformin, patients using metformin + sulfonylurea and patients using metformin + DPP4 analogues Hematological parameters, white blood cell formulas and HbA1c levels were examined in both subgroups. There was no statistically significant difference in both parameters before treatment and at six months (Table 5 and 6).

Discussion

The results of this study showed that glucose control significantly lowers HbA1c levels together with blood sugar regulation. On the other hand causes a significant decrease in the number of basophils, eosinophils, platelets, lymphocytes. These findings are very important. Significant numerical and proportional decrease in the number of basophils, eosinophils, platelets, lymphocytes suggest that regulation of blood sugar has an anti-inflammatory activity. Diabetic patients are more prone to certain infections than those without DM. Regulation of blood glucose increases the anti-inflammatory activity and reduces predisposition to infections. However, in vitro and in vivo more studies are needed for further investigation. Diabetes Mellitus is a chronic metabolic disorder due to insulin deficiency or defects on the effect of insulin and requiring a continuous medical care. There is disturbance in body metabolism and energy utilization from carbohydrates, fats and proteins insufficiently [5].

| Table | 6: | Distributions | of | the | hematological | parameters |
|--------|------|----------------|-----|------|---------------|------------|
| accord | ling | to the OAD sub | gro | ups. | | |

| | | Metformin (n=13) | Metformin + sulfonylurea (n=23) | Metformin+ DPP4 analogues (n=15) | ¹ p |
|-------------|--|----------------------------|---------------------------------------|---|--------------------|
| | Pre-Treatment | 57.06 ± 7.06 | 57.35 8 .26 | 60.6 ± 10.44 | 0.458 |
| | Post-Treat. 3rd Month | 59.19 ± 10.06 | 58.62 ± 7.71 | 58.32 ± 5.28 | 0.957 |
| Neu % | Post-Treat. 6th Month | 56.81 ± 6.93 | 57.39 ± 8.39 | 58.41 ± 5.64 | 0.840 |
| | ² p ³ p | 0.495 | 0.583 | 0.499 | - |
| | Pre-Treatment | 4.61 ± 1.6 | 4.82 ± 1.2 | 4.87 ± 1.77 | 0.886 |
| | Post-Treat. 3rd Month | 4.75 ± 2.29 | 4.91 ± 1.39 | 4.52 ± 0.66 | 0.740 |
| Neu | Post-Treat. 6th Month | 4.24 ± 1.4 | 4.65 ± 1.3 | 4.57 ± 1 | 0.626 |
| | ² p 3- | 0.576 | 0.638 | 0.668 | - |
| | Pre-Treatment | 32 92 + 6 28 | 0.787 | 31 34 + 6 95 | 0.857 |
| | Post-Treat. 3rd Month | 31.32 ± 9.25 | 31.98 ± 7.6 | 31.39 ± 4.82 | 0.955 |
| Lym % | Post-Treat. 6th Month | 33.29 ± 6.91 | 33.43 ± 8.43 | 31.48 ± 5.36 | 0.698 |
| | ² p | 0.600 | 0.450 | 0.976 | - |
| | °p | 2.04 - 4.50 | 0.892 | 0.5 + 0.00 | 0.500 |
| | Pre-Treatment Post-Treat. 3rd | 2.94 ± 1.52 | 2.67 ± 0.86 | 2.5 ± 0.62 | 0.536 |
| Lvm | Month Post-Treat. 6th | 2.74 ± 1.53 | 2.65 ± 0.78 | 2.45 ± 0.62 | 0.720 |
| , | Month | 2.79 ± 1.42 | 2.65 ± 0.67 | 2.41 ± 0.52 | 0.531 |
| | p | 0.546 | 0.986 | 0.874 | - |
| | Pre-Treatment | 6.95 ± 1.94 | 7.35 ± 2.75 | 6.56 ± 1.95 | 0.600 |
| | Post-Treat. 3rd Month | 6.74 ± 2.18 | 6.51 ± 3.17 | 6.96 ± 2.11 | 0.879 |
| Mono % | Post-Treat. 6th Month | 7.18 ± 1.92 | 6.53 ± 1.97 | 6.7 ± 1.74 | 0.615 |
| | ² p | 0.541 | 0.171 | 0.697 | |
| | p Bro-Troatmont | 0.61 ± 0.3 | 0.959 | 0.51 ± 0.15 | 0.437 |
| | Post-Treat. 3rd Month | 0.49 ± 0.16 | 0.53 ± 0.25 | 0.52 ± 0.16 | 0.866 |
| Mono | Post-Treat. 6th Month | 0.52 ± 0.17 | 0.53 ± 0.21 | 0.52 ± 0.18 | 0.985 |
| | ²p | 0.300 | 0.296 | 0.938 | - |
| | °p Dro Trootmont | 246 + 4.44 | 0.790 | 2 40 + 4 04 | 0.044 |
| | Pre-Treatment Post-Treat. 3rd | 2.16 ± 1.41 1.83 ± 1.44 | 2 ± 1.37 1.64 ± 0.82 | 2.49 ± 1.94 2.44 ± 1.58 | 0.641 |
| Eos % | Post-Treat. 6th | 1.8 ± 1.09 | 1.63 ± 0.74 | 2.84 ± 1.89 | 0.018 ^f |
| | ² p | 0.257 | 0.226 | 0.449 | |
| | ³ p | | 0.118 | | - |
| | Pre-Treatment | 0.17 ± 0.11 | 0.16 ± 0.13 | 0.2 ± 0.18 | 0.731 |
| F ee | Month | 0.14 ± 0.1 | 0.14 ± 0.08 | 0.2 ± 0.16 | 0.219 |
| EOS | Month | 0.14 ± 0.09 | 0.13 ± 0.08 | 0.23 ± 0.16 | 0.032 ^f |
| | <u>р</u> ³ р | 0.241 | 0.290 | 0.529 | - |
| | Pre-Treatment | 0.93 ± 0.31 | 0.91 ± 0.45 | 0.77 ± 0.34 | 0.747 |
| | Post-Treat. 3rd Month | 0.94 ± 0.37 | 0.87 ± 0.47 | 0.91 ± 0.38 | 0.904 |
| Baso% | Post-Treat. 6th Month | 0.93 ± 0.31 | 0.88 ± 0.39 | 0.83 ± 0.3 | 0.729 |
| | <u>²p</u> ³ p | 0.995 | 0.837 | 0.278 | - |
| | Pre-Treatment | 0.09 ± 0.06 | 0.07 ± 0.07 | 0.07 ± 0.05 | 0.850 |
| | Post-Treat. 3rd Month | 0.06 ± 0.07 | 0.09 ± 0.05 | 0.07 ± 0.06 | 0.310 |
| Baso - | Post-Treat. 6th Month | 0.07 ± 0.05 | 0.07 ± 0.04 | 0.07 ± 0.05 | 0.955 |
| | ² p 3 | 0.255 | 0.269 | 1.000 | - |
| | D | | 0.849 | | |

Data were shown as mean ±standard deviation and n (%). ¹: Results of the comparison among groups. ²: Results of the comparison among pre-treatment, post-treatment 3rd month and post-treatment 6th month measures. ⁹: Results of the comparison among three groups according to alterations of hematologic parameters. ¹: There was statistically significant difference between group 2 and group 3.

The incidence of anemia is increased in patients with diabetes. The reason is multifactorial. Chronic hyperglycemia causes abnormal red blood cells and renal sympathetic denervation is associated with oxidative stress and autonomic neuropathy [6]. Hypoxic environment occurs in the renal tubulo interstitial. The amount of erythropoietin produced by the peritubular fibroblasts reduces and becomes inappropriate production. The earliest and the most important reason of anemia in diabetic patients is inappropriately and low erythropoietin levels [1]. Systemic inflammation, functional hematinic deficiencies, erythropoietin resistance and reduced red cell survival also drive anemia in the setting of impaired renal compensation [7].

Mean platelet volume (MPV Mean Platelet Volume), measured by the hematology analyzers, is a marker showing platelet function and activation. Altered platelet morphology and function can be considered as a factor for risk of micro-and macrovascular diseases [2, 8]. Large platelets are younger and more reactive. Therefore, these platelets secrete more serotonin and β -thromboglobulin, contain more intense granules and produce more thromboxane A2 [9-12]. All of these give rise to a procoagulant effect and may cause thrombotic vascular complications. It may talk about the relationship between changes in platelet function especially MPV and diabetic vascular complications [2].

There are many studies showing the relationship between MPV and diabetes mellitus. There are studies showing the relationship between MPV and level of fasting blood glucose, postprandial blood glucose, and impaired fasting glucose and HbA1c levels in diabetic peripheral arterial disease. MPV values was found increased in all of these studies [2, 13-18]. We did not found significant change in MPV values but a significant decrease found in platelet counts in OAD group at six month. Avoid an increase in MPV, falling in platelet counts showed that OAD therapy may be effective in preventing vascular complications of diabetes mellitus.

Diabetes mellitus is a chronic inflammatory disease. In studies, peripheral blood leukocyte count [19], acute phase reactants such as C-reactive protein (CRP) [20], interleukin 6 (IL-6) [21], tumor necrosing factor a (TNF-a) [22], serum-amyloid A (S-AA) [23, 24], was found increased in patients with impaired fasting glucose or insulin resistance. An inverse relationship was shown between HbA1c levels and inflammatory cytokine levels in the blood. This chronic inflammatory process is is supressed by regulation of blood glucose [3]. A prospective study including twenty-study meta-analysis showed that number of peripheral blood leukocytes, basophils, eosinophils and neutrophils increased, no change in the number of monocytes in patients with Type 2 DM [4]. We found significant difference in the number of lymphocytes (p = 0.029) and basophils (p = 0.029) in

insulin using group in 6th month. This result shows that insulin therapy may be effective in suppressing chronic inflammatory process. A statistically significant relationship was showed between blood leukocyte count and glucose intolerance in the study made by Gokulakrishnan K. and et al [25].

References

1. Bosman DR, Winkler AS, Marsden JT, Macdougall IC, Watkins PJ. Anemia with erythropoietin deficiency occurs early in diabetic nephropathy. J Diabetes Care. 2001; 24:495–499.

2. Hekimsoy Z, Payzinb B, Ornek T, Kandoğan G. Mean platelet volume in Type 2 diabetic patients. J Diabetes Complications. 2004; 18:173–176.

3.Mirza S, Hossain M., Mathews C., Martinez P, Pino P, Gay JL, Rentfro A, McCormick JB, Fisher-Hoch SP. Type 2-Diabetes is Associated With Elevated Levels of TNF-alpha, IL-6 and Adiponectin and Low Levels of Leptin in a Population of Mexican American: A Cross-Sectional Study. Cytokine. 2012;57(1):136-142.

4. Gkrania-Klotsas E, Ye Z, Cooper AJ, Sharp SJ, Luben R, Biggs ML, Chen LK, Gokulakrishnan K, Hanefeld M, Ingelsson E, Lai WA, Lin SY, Lind L, Lohsoonthorn V, Mohan V, Muscari A, Nilsson G, Ohrvik J, Chao Qiang J, Jenny NS, Tamakoshi K, Temelkova-Kurktschiev T, Wang YY, Yajnik CS, Zoli M, Khaw KT, Forouhi NG, Wareham NJ, Langenberg C. Differential White Blood Cell Count and Type 2 Diabetes: Systematic Review and Meta-Analysis of Cross-Sectional and Prospective Studies. PLoS One. 2010; 18; 5(10).

5. Satman İ, İmamoğlu Ş, Yılmaz C. TEMD, diagnosis, treatment and follow-up guide of diabetes mellitus and its complications - 2011; 15-15.

6. Waggiallah H, Alzohairy M. The effect of oxidative stress on human red cells glutathione peroxidase, glutathione reductase level, and prevalence of anemia among diabetics. N Am J Med Sci. 2011;3(7):344-347.

7. Thomas MC. Anemia in diabetes: marker or mediator of microvascular disease? Nat Clin Pract Nephrol. 2007;3(1):20-30.

8. Zuberi BF, Akhtar N, Afsar S. Comparison of mean platelet volume in patients with diabetes mellitus, impaired fast. Singapore Med J. 2008;49(2):114-116.

9. Bae SH, Lee J, Roh KH, Kim J. Platelet activation in patients with diabetic retinopathy. Korean J Ophthalmol. 2003; 17:140–144.

10. Colwell JA, Nesto RW. The platelet in diabetes-focus on prevention of ischemic events. Diabetes Care. 2003; 26:2181–2188.

11. Ateş O, Kiki I, Bilen H. Association of Mean Platelet Volume With The Degree of Retinopathy in Patients with Diabetes Mellitus. Eur J Gen Med. 2009; 6:99–102.

12. Chang HA, Hwang HS, Park HK. The Role of Mean Platelet Volume as a Predicting Factor of Asymptomatic Coronary Artery Disease. Korean J Fam Med. 2010; 31:600–606.

13. Szeremeta M, Kemona-Chetnik I, Dymicka-Piekarska V. The relations between platelet count, mean platelet volume and HbA1C in patients with type 2 diabetes. Przegl Lek. 2009; 66(12):1049-1051.

14. Li S, Wang C, Zhong XW, Li HQ, Fu XQ, Ran XW. Variance of mean platelet volume in subjects with normal glucose tolerance, impaired glucose regulation and type 2 diabetic mellitus and its relationship with diabetic peripheral artery disease. Zhonghua Yi Xue Za Zhi. 2012; 31; 92(4):232-235.

15. Demirtunc R, Duman D, Basar M, Bilgi M, Teomete M, Garip T. The relationship between glycemic control and platelet activity in type 2 diabetes mellitus. J Diabetes Complications. 2009;23(2):89-

94.

16. Zuberi BF, Akhtar N, Afsar S. Comparison of mean platelet volume in patients with diabetes mellitus, impaired fasting glucose and non-diabetic subjects. Singapore Med J. 2008;49(2):114-116.

17. Papanas N, Symeonidis G, Maltezos E, Mavridis G, Karavageli E, Vosnakidis T, Lakasas G. Mean platelet volume in patients with type 2 diabetes mellitus. Platelets. 2004; 15(8):475-478.

18. Kodiatte TA, Manikyam UK, Rao SB, Jagadish TM, Reddy M, Lingaiah HK, Lakshmaiah V. Mean Platelet Volume in Type 2 Diabetes Mellitus. J Lab Physicians. 2012;4(1):5-9.

19. Vozarova B, Weyer C, Lindsay RS, Pratley RE, Bogardus C, Tataranni PA. High white blood cell count is associated with a worsening of insülin sensitivity and predicts the development of type 2 diabetes. Diabetes. 2002; 51:455–461.

20. Pradhan AD, Manson JE, Rifai N, Buring JE, Ridker PM. Creactive protein, interleukin 6, and risk of developing type 2 diabetes mellitus. JAMA. 2001; 286:327–334.

21. Pickup JC, Chusney GD, Thomas SM, Burt D. Plasma interleukin-6, tumour necrosis factor alpha and blood cytokine production in type 2 diabetes. Life Sci. 2000; 67:291–300.

22. Rytter E, Vessby B, Asgard R, Johansson C, Sjödin A, Abramsson-Zetterberg L, Möller L, Basu S. Glycaemic status in relation to oxidative stress and inflammation in well-controlled type 2 diabetes subjects. Br J Nutr. 2009; 101:1423–1426.

23. Hu FB, Meigs JB, Li TY, Rifai N, Manson JE. Inflammatory markers and risk of developing type 2 diabetes in women. Diabetes. 2004; 53:693–700.

24. Yang RZ, Lee MJ, Hu H, Pollin TI, Ryan AS, Nicklas BJ, Snitker S, Horenstein RB, Hull K, Goldberg NH, Goldberg AP, Shuldiner AR, Fried SK, Gong DW. Acute-phase serum amyloid A: an inflammatory adipokine and potential link between obesity and its metabolic complications. PLoS Med. 2006; 3:e287

25. Gokulakrishnan K, Deepa R, Sampathkumar R, Balasubramanyam M, Mohan V. Association of leukocyte count with varying degrees of glucose intolerance in Asian Indians: the Chennai Urban Rural Epidemiology Study (CURES-26). Metab Syndr Relat Disord. 2009;7(3):205-210.