A Comparative Study between the Effect of Ropinirole and Metformin on Metabolic Syndrome in Women
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Abstract

Background: Metabolic syndrome is a multiplex risk factor that arises from hyperglycemia /insulin resistance accompanied with abnormal fat function &central obesity& possibility to develop diabetes mellitus type-2, atherosclerosis & various diseases. Aim of study: To determine the effect of ropinirole & metformin on glycemic Parameters, insulin sensitivity and lipid profile associated with metabolic syndrome. Patients and methods: The present study was performed in diabetic and endocrine center from May 2016 to march 2018. Ninety female patients with metabolic syndrome participate in this study .They were allocated to three groups (each group contain 30 patients) first group put on diet restriction physical exercise only. Second group received metformin 750mg extended release orally once daily at night with diet restriction & physical exercise. Third group received ropinirole 0.25 mg orally after meal in morning with diet restriction & physical exercise. The duration of the study extended to 12 weeks. Blood samples were taken from each patient at zero time to determine base line level & after 12 weeks of treatment. The measured parameters were WBC, BMI, FBS, FI, HOMA-IR, Hb A1c, TC, TG, HDL, LDL. Results: Showed highly significant lowering effect of both ropinirole & metformin on the levels of the all parameters except HDL with high significant elevation, at the mean time all the parameters of the control were significantly changed. Conclusion: Both drugs improve glycemic parameters, insulin sensitivity and lipid profile in metabolic syndrome but metformin is slightly more powerful on these parameters.

Keywords: Ropinirole, metformin, metabolic syndrome, insulin resistance, lipid profile.

INTRODUCTION

Metabolic syndrome (M.S) is a clustering of clinical components including hyperglycemia/insulin resistance obesity and dyslipidemia [1]. The international diabetes foundation (IDF) published new criteria for definition & diagnosis of metabolic syndrome. It include increase the waist circumference (WTC) and FBS &TG levels and decrease of HDL level and rise of blood pressure [2]. The present study was performed to explore the possible effect of Ropinirole & metformin on M. S. Ropinirole is a selective non ergoline pure dopamine D2 receptor agonist, it has negligible affinity for dopamine D1 receptor it used in treatment of Parkinsonism [3]. While metformin is an antidiabetic drug related to biguanides group, its primary effect is to activate the enzyme AMP that activated protein kinase (AMDK) and to reduce hepatic glucose production used in polycystic ovary disease [4].

PATIENTS & MATERIALS & METHODS

The present study conducted at endocrine & diabetes center from may 2016 to march 2018 after approval by ethical committee of Al-nahrain College of medicine.

Chemicals
All the chemicals used in the present study were of analytic grade. Ropinirole 0.5 mg tab. Supplied by IDIR Ranbaxy Company. Metformin 750 mg XR supplied by Merck Germany. Kits for TC,HDL,LDL,TG purchased by accent 2000 poland. HBA1c supplied by Bio-Rad lab USA. Insulin Elisa. Tohso.

Patients
Ninety female patients participated in this study. They were selected according to the definition of IDF for M.S. They were informed about the aim of the study. The range of the patient’s age was between 26-45
years old. All the patients with other diseases were excluded from the study. Patients were allocated to three groups (each group contain 30 patients). The first group (control) received no medication but put on diet restriction & physical exercise only. Second group: received metformin 750 mg extended release orally once daily at night in addition to diet restriction & physical exercise. Third group: received Ropinirole 0.25 mg orally after meal in the morning with diet restriction and physical exercise. The following parameters were measured in all the participants starting from zero time (base line) & continued for 12 weeks for each parameter: WTC (waist circumference) BMI (body mass index) FBS (fasting blood sugar) FI (fasting insulin) lipid profile total cholesterol, TG, HDL, LDL, VLDL. Hemoglobin A1c (HBA1c). HOMA-IR (hemostatic mode assessment for insulin resistance) blood pressure was measured every 4 hours. After fasting the patient for 12 hours, 10ml of venous blood collected at each visit. 2 ml of the whole blood anticoagulant tube used for HBA1c measured at zero time & after 12 weeks. The other part was obtained in plasma tube & centrifuged. 10 min to separate the serum in order to be used in spectrophotometer method [5].

**Statistical Analysis**

The results of the study were analyzed by using (SPSS) methods.

The paired T- Dependent test used for comparing the results of same group & independent T-test used for comparing between different groups& the statistical signification was p <0.05.

**RESULTS**

The obtained results revealed that both metformin & Ropinirole groups produce highly significant reduction P<0.001 in the levels of WTC and BMI & FBS and FI also TC, TG, LDL, VLDL, except HDL which is highly elevated significantly p<0.001 when the results of the base line compared with the results after 12 weeks of treatment. In general metformin has more potent effect than Ropinirole on these parameters. At the meantime the control group produce significant lowering effect of FBS, FI, HBA1c, HOMA-IR levels p<0.05 with highly significant effect of WTC, BMI & LIPID profile p<0.001 see Table 1, 2, 3, 4.

Table-1: The effect of control, metformin and ropinirole groups on the WTC and BM index after 12 weeks of treatment

| Parameters                  | Groups       | Period       | Control       | Metaformin    | Ropinirole    |
|-----------------------------|--------------|--------------|---------------|---------------|---------------|
| Waist circumference (WTC) (cm) | Base line    | 122.73± 1.20 | 125.26± 1.68  | 125.90± 1.78  |
|                             | 12 weeks     | 104.32±1.01  | 93.95± 1.26   | 96.94±1.37    |
|                             | ∆ WTC        | -18.41± 0.17 | -31.13± 0.42  | -30.21± 0.42  |
|                             | baseline     | 35.59±0.41   | 36.98± 1.17   | 38.53±1.45    |
|                             | 12 weeks     | 30.61±0.35   | 27.73± 0.88   | 32.54±1.27    |
|                             | ∆ BMI        | -4.98± 0.31  | -9.24±0.29    | -5.99± 0.35   |

*Significant at p< 0.05 **highly significant at p<0.001 comparing with the base line level by using a paired t-test a significant at P< 0.05 ab highly significant at P< 0.001 comparing changes of drugs and control by using unpaired t-test. ∆ changing in the base line.

Table-2: The effect of control, metformin and ropinirole groups on the FBS and FI levels after 12 weeks of treatment

| Parameters                  | Groups       | Period       | Control       | Metaformin    | Ropinirole    |
|-----------------------------|--------------|--------------|---------------|---------------|---------------|
| Fasting blood sugar FBS mg/dl | Base line    | 115.2 ± 5.74 | 117.1 ± 5.74  | 119.06 ± 5.74 |
|                             | 12 weeks     | 110.62 ± 0.66 | 93.68 ± 0.72  | 108.59 ± 0.68 |
|                             | ∆ FBS        | - 4.57 ± 1.29 | - 23.42 ± 0.18 | - 18.47 ± 0.10 |
| Fasting insulin FI IU/ml    | baseline     | 48.36 ± 1.08 | 49.26 ± 1.51  | 48.40 ± 1.33  |
|                             | 12 weeks     | 38.77 ± 0.92 | 27.63 ± 0.75  | 29.04 ± 0.79  |
|                             | ∆ FI         | - 9.59 ± 1.66 | - 24.63 ± 0.75 | - 19.36 ± 0.53 |

*Significant at p< 0.05 **highly significant at p<0.001 comparing with the base line level by using a paired t-test a significant at P< 0.05 ab highly significant at P< 0.001 comparing changes of drugs and control by using unpaired t-test. ∆ changing in the base line.
In the present study all the patients with M.S were put on diet restriction & physical exercise but patients of group1(control) received no medication, the diet and exercise therapy are change in group1 to produce significant lowering effect in body weight, WTC & BMI considerably after 12 weeks of the treatment when compared with zero time (base line). The exercise therapy activates AMDK enzyme to induce favorable metabolic changes in muscle& adipose tissue using fat as energy source instead of storing it [6] meanwhile FBS, Fl, HBA1c, HOMA-IR were significantly reduced when compared with base line after 12 weeks of diet changes & exercise therapy due to increase fuel utilization & enhance glycolysis in skeletal muscles [7] our results were compatible with the results of others [8-10].

At the same time the results of lipid profile revealed highly significant reduction in TC, TG, and LDL levels with highly significant increase in HDL level these results were compatible with results of others for the same reasons [11].

Metformin is exerted highly significant lowering effect on body weight, BMI & WTC (group 2) due to its multiple effects in suppressing glucose hepatic production, inhibition of glucose absorption from the gut & improvement of glycolysis with anorexic effect [12]. Metformin extended release make the body accelerates fatty acid oxidation in the cell especially at night to compensate the reduction in glucose which is important substrate of energy [13]. Metformin promoted weight loss & BMI&WTC in patients with obesity in DM-type 2 after 12 weeks of treatment. Our results were similar to the results of others [14]. Metformin showed highly significant reduction in FBS, Fl, HOMA-IR & HBA1c after 12 weeks of treatment due to inhibition of hepatic glucose production & decline in insulin level these results were similar to results of others [15].

Metformin improve insulin sensitivity by several mechanisms include increasing insulin receptor tyrosine kinase activity, enhancing glycogen synthesis.
& increasing expression of GLUT & glucose transporter [16].

Metformin showed remarkable reduction in serum cholesterol, TG, VLDL, LDL with highly significant elevation in HDL level in the base line level. These effects attributed to insulin resistance that lead to increase amount of free fatty acid infiltrate to the liver. Therefor increase TG as well as VLDL production. These results were compatible with the results of other researchers [17].

Metformin therapy may increase the lipoprotein, lipase production. Therefore increase HDL that reduce the risk of atherosclerosis [18].

Administration of Ropinirole to patients with insulin resistance produced high significant lowering effect of body weight, BMI, WTC in group-3 this reduction attributed to the ability of D2-receptor agonist in shifting the energy utilization toward fatty acid oxidation as a result more stored fat will consumed that lead to weight loss & change of BMI ,WTC. These results were similar to results of others when they used cabergoline as D2-receptor agonist [19]. Ropinirole has an effect on appetite producing a state of anorexia & nausea that lead to weight loss [20].

In the present study ropinirole caused highly significant decline in FBS & F1 levels compared with base line during 12 weeks of treatment, this lead to decrease HOMA-IR which is index of insulin resistance and also HBA1c decreased. These results compatible with the results of others [21] when they used cabergoline in study that cause significant decrease in FBS & HBA1c.

Ropinirole showed highly significant changes in lipoproteins after 12 weeks of treatment decreasing TC, TG, LDL, VLDL and increasing HDL levels, these changes related to improving insulin sensitivity, decreasing body weight, shifting the energy utilization, lowering of FFA oxidation & inhibition of lipolysis, these results were compatible with results of [22, 23] who noticed that cabergoline reducing TG and increasing HDL.

CONCLUSION
Both metformin & Ropinirole improve glycemic parameters, insulin sensitivity & lipid profile in metabolic syndrome but metformin has more powerful effects on these parameters.

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