

SERUM CORTISOL AND INSULIN HORMONE LEVELS AND THEIR ROLE IN NORTH INDIAN MEN AND WOMEN

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ABSTRACT

Objective

To study cortisol and insulin hormone serum levels and their role in north Indian men and women having Type 2 Diabetes.

Research Design and Methods

For the analyses, (n=200) subjects including (n=94) males and (n=106) females, out of which 100 diagnosed cases and 100 age and sex matched healthy controls were studied. Only diagnosed cases of diabetes type 2 (50 men and 50 women) aged 45–75 years undergoing glucose profile testing in outdoor clinics in the hospital PGIMS, Rohtak (2011-2013) were included following a detailed protocol. Patients with acute complications like coma and acidosis, pregnant women, postmenopausal women on hormone replacement therapy, use of steroids since past six months, type 1 diabetes were excluded. Early morning fasting samples were collected and serum analysed for cortisol, insulin, fasting blood glucose and HbA1c. Serum Insulin^{1,2} levels (normal healthy adults <25 μ IU/ml) were done using solid phase sandwich Enzyme linked immunosorbent assay, serum Cortisol levels^{3,4} (Reference range: 50-230 ng/ml) were done using DRG Cortisol ELISA kit -a solid phase enzyme linked immunosorbent assay, and HbA1c levels (normal=4-5.6% in normal people, <6.5% -target for control in diabetics) were measured on Auto analyser via Immunoassay Kits. The results were analysed and compared.

Results

Overall analysis showed that diabetic men had low testosterone values (287.50 \pm 61.09) ng/dL as compared to controls (409.38 \pm 113.23) ng/dL (p<0.001) and raised HbA1c, whereas diabetic women had raised testosterone (52.35 \pm 41.09) ng/dL values (p<0.001) and raised HbA1c as compared to controls (25.00 \pm 16.99) ng/dL (p<0.001). Diabetic Women had mean estradiol levels (47.00 \pm 53.36) pg/ml lower as compared to control females (69.31 \pm 57.51) pg/ml, (p <0.05), also they negatively correlated with HbA1c. Men showed no significant difference in estradiol levels in diseased and controls and showed no correlation between estradiol and HbA1c levels.

Conclusions

In North India - Diabetes type 2 is associated with Cortisol and Insulin levels significantly higher in cases as compared to controls, also fasting blood glucose and HbA1c levels were higher in diabetics irrespective of sex. Such

associations suggest possible clinical applications of hormone biomarkers in potentially adding prospective risk information. More prospective studies are needed to better define risk levels.

KEYWORDS: Serum Cortisol and Insulin Hormone Levels

INTRODUCTION

Cortisol directly plays role in glucose metabolism and affects insulin's actions as well. Cortisol stimulates gluconeogenesis (formation, in the liver, of glucose from certain amino acids, glycerol, lactate and/or propionate).

Cortisol counteracts insulin, contributes to hyperglycemia-causing hepatic gluconeogenesis and inhibits the peripheral utilization of glucose (insulin resistance) by decreasing the translocation of glucose transporters (especially GLUT4) to the cell membrane.⁵ However, cortisol increases glycogen synthesis (glycogenesis) in the liver.

Chiodini found that in type 2 diabetic subjects, hypothalamic-pituitary-adrenal activity is enhanced in patients with diabetic complications and the degree of cortisol secretion is related to the presence of diabetes complications. They evaluated cortisol secretion in hundred and seventy, type 2 diabetic subjects. They evaluated the presence of chronic complications (incipient nephropathy, asymptomatic neuropathy, background retinopathy, and silent macroangiopathy) and found that cortisol levels were higher in diabetics with complications than without them and controls.⁶

Reynolds analysed serum cortisol levels in type 2 DM cases and found higher fasting cortisol levels to be associated with greater estimated cognitive decline. Therefore, strategies targeted at lowering cortisol action may be useful in ameliorating cognitive decline in individuals with type 2 diabetes.⁷

From a clinical perspective, the consistent findings among both men and women of significant associations for cortisol and insulin suggest possible clinical applications of hormone biomarkers in potentially adding predictive risk information. More prospective hormonal investigations are needed to better define risk levels. More prospective hormonal investigations are needed to better define risk levels.

For inconsistency observed in the studies so far, the present study has been planned to study cortisol, insulin and HbA1c levels in the patients of Type 2 DM and find the correlation with glycemic control and complications.

RESEARCH DESIGN AND METHODS

The study was conducted in the hospital PGIMS, Rohtak (2011-2013) in department of biochemistry in collaboration with department of medicine. Only diagnosed cases of diabetes type 2 undergoing glucose profile testing in outdoor clinics were included following a detailed protocol. 100 patients with Type 2 DM and 100 age and sex matched healthy controls were taken. Out of 100 cases, 55 were males and 45 were females whereas in 100 controls 51 were males and 49 females. Patients of age group 45-75 years were included in the study. Patients with acute complications like hyperglycaemic hyperosmolar coma, comorbid conditions like testicular tumor, prostate or breast cancer, lipidemias, PCOS (polycystic ovarian syndrome) and CAH (congenital adrenal hyperplasia), Insulin therapy,

Intake of drugs (known to interfere with HPA axis or with autonomic nervous system) like β -blockers, α -blockers, and cholinergic agonists and antagonists; hormone-modulating therapies or topical/systemic glucocorticoids within 3 months, chronic debilitating disease such as severe depression or psychiatric illness, head trauma, renal failure, haemochromatosis, cirrhosis, hepatitis C, HIV, congenital hypogonadotropic hypogonadism or panhypopituitarism,

pregnant and lactating women were excluded.

History was taken from all diabetic patients and control subjects and complete general and systemic physical examination was performed. All patients and controls were subjected to anthropometric measurements, routine and special investigations. Anthropometry included measurement of weight, height, waist circumference, hip circumference, BMI and waist hip ratio. Informed consent was taken from all subjects and all hazards were explained. The study was approved by ethical committee of the University of Health Sciences, Rohtak where the study was carried out. Routine investigations included haemoglobin, total leukocyte count, blood urea, serum creatinine and fasting blood glucose levels. Special investigation performed were glycosylated haemoglobin, serum cortisol and serum insulin.

5ml overnight fasting blood sample was collected from the antecubital vein aseptically without anticoagulant and allowed to clot. Serum was separated by centrifugation of the sample and was used for the assays (sample were stored at 2-8°C for 1day, and at -20°C if storage was required for more than 1 day).1 ml blood sample was collected in EDTA vial separately irrespective of time and meal for estimation of glycosylated haemoglobin. All the patients with diabetes mellitus type-2 as well as control were subjected to serum investigations.

Glycosylated Haemoglobin was determined by ion exchange chromatography as described by Goldstein et al, using ion exchange chromatography kits.⁸

SERUM^{1,2} Insulin levels were done using solid phase two site direct sandwich enzyme immunoassay. Results were read using ELISA reader.

Serum CORTISOL^{3,4} levels were measured using DRG Cortisol ELISA kit – a solid phase enzyme linked immunosorbent assay, based on the principle of competitive binding.

Serum hormone levels were measured in a biochemistry laboratory and pathology blood transfusion laboratory by chemiluminescence and Elisa techniques using first-thawed specimens from the 2011 to 2013 venipuncture during 2011-2013. Free cortisol and insulin levels were thus determined.

Data were analyzed using simple statistical techniques. Analyses were performed using mean values and bar diagrams. Unadjusted associations between hormone levels and diabetes were evaluated using Student's *t* test and χ^2 test and calculation of *p* values.

RESULTS

Baseline Characteristics and Diabetes

Mean fasting blood glucose levels (149.46±29.28 mg/dL) were significantly higher in men ($P < 0.001$) and women ($P < 0.01$) with diabetes compared with persons without diabetes (95.72±6.21 mg/dL). (Table 1)

Table 1: Fasting Blood Glucose and Glycosylated Haemoglobin Levels in Cases and Controls

Parameter	Cases	Controls	p Value
Fasting Blood Glucose (mg/dl)	149.46± 29.28	95.72±6.21	<0.001
HbA1c (%)	9.32±2.85	4.37±0.845	<0.001

The mean levels of glycosylated haemoglobin in diabetic and control group were 9.32±2.85% and 4.37±0.845% respectively, and the difference was statistically highly significant ($p < 0.001$). (Table 1)

No differences were observed for age and sex (Table 2)

Table 2: Age and Sex Wise Distribution of Cases and Controls

	Cases (n=100)	Controls (n=100)
Mean age	53.73±11.30	51.43±14.11
Range	31-78	24-80
Male	50 (50%)	50(50%)
Female	50 (50%)	50 (50%)

Diabetes had significantly higher mean waist circumference, BMI (Table 3), triglycerides. (Table 4) and HbA1c (Table 2) Mean total cortisol (Table 6), insulin (Table 5) and HDL-cholesterol levels were lower

Table 3: Body Mass Index (BMI) and Waist Hip Ratio (W/H R) in Cases and Controls (All Values are in Mean±SD)

	Cases	Controls	p Value
BMI (kg/m ²)	29.17±6.50	25.66±5.07	<0.001
W/H Ratio (Waist Hip Ratio)	0.951±0.022	0.934±0.073	<0.001

Table 4: Lipid Profile in Cases and Controls (Mean±SD)

	Cases	Controls	p-Value
TG (mg/dl)	170.74±44.18	151.09±83.91	<0.001
HDL-C (mg/dl)	42.73±18.24	47.78±5.40	<0.001
VLDL-C (mg/dl)	34.14±8.83	30.21±16.78	<0.001
LDL-C (mg/dl)	111.86±48.42	101.21±32.03	<0.05

Table 5: Fasting Blood Glucose, Post Prandial Blood Glucose, Glycosylated Haemoglobin and Serum Insulin Levels in Cases and Controls

Parameter	Cases	Controls	p value
Fasting Blood Glucose(mg/dl)	149.46± 29.28	95.72±6.21	<0.001
HbA1c (%)	9.32±2.85	4.37±0.845	<0.001
Serum Insulin (µIU/ml)	13.56±9.99	6.00±3.9	<0.001

The subjects in the diabetic group had meant fasting blood glucose levels of 149.46±29.28 mg/dL whereas in the control group it was 95.72±6.21 mg/dL. The difference in levels of FBG in diabetic and control groups was statistically highly significant (p<0.001). The mean levels of glycosylated haemoglobin in diabetic and control group were 9.32±2.85% and 4.37±0.845% respectively, and the difference was statistically highly significant (p<0.001). The mean levels of serum insulin in diabetic and control group were 13.56±9.99 µIU/mL and 6.0±3.9µIU/mL respectively, and the difference was statistically highly significant (p<0.001)(Table 6).

Table 6: Mean Cortisol Levels in Cases and Controls

	Cases	Controls	p Value
Cortisol Levels (sng/ml)	106.07±46.57	84.32±54.84	<0.001

Cortisol was significantly higher in diabetic cases as compared to controls, (p =0.000) which is statistically significant.

CONCLUSIONS

In present study, Increased insulin levels were observed in the study group (13.56 ± 9.99 $\mu\text{IU/ml}$ vs. 6.0 ± 3.9 $\mu\text{IU/ml}$) ($p < 0.001$) (Table 6). These observations are in conformation with previous studies which state that hyperinsulinemia is a feature of type 2 diabetes. Elevated fasting insulin levels are described as a marker of insulin resistance which may be due to poor compliance and various other factors. In insulin-resistant states accompanied by hyperinsulinemia, such as occur in type II diabetes, there is resistance to the anti-lipolytic effect of insulin; this resistance results in increased serum levels of free fatty acids and glycerol. All of these effects tend to worsen hyperglycemia.⁹

In our study cortisol levels were significantly higher in cases as compared to controls ($p < 0.001$). This is in accordance with literature which reveals that cortisol directly plays role in glucose metabolism and affects insulin as well. Cortisol counteracts insulin, contributes to hyperglycemia-causing hepatic gluconeogenesis and inhibits the peripheral utilization of glucose (insulin resistance) by decreasing the translocation of glucose transporters (especially GLUT4) to the cell membrane.¹⁰

Our finding is further supported by previous studies carried out by Oltmanns, Chiodini and Reynolds which suggest that high cortisol levels in diabetes type 2 are positively associated with deranged glycosylated haemoglobin, blood pressure changes, relative abdominal mass, severity of clinical features and complications.^{6,7,11} Moreover, it has been found to affect cognitive function in elderly DM 2 patients and thus strategies targeted at maintaining its level would help in improving health profile of DM 2 patients.¹²

To conclude, patients with DM type 2 have abnormalities in various hormone levels. These associations may be considered in the pathogenesis of the disease and should be taken into account for the treatment of patients of DM type 2.

Moreover, from a clinical perspective, the consistent findings among both men and women of significant associations for cortisol and insulin, suggest possible clinical applications of hormone biomarkers in potentially adding predictive risk information above and beyond obesity. More prospective investigations are needed to better define risk levels.

Furthermore, the hormone therapy may have a role to play. The role of various strategies targeted at maintaining cortisol levels in improving health profile of DM 2 patients, might be considered.

To conclude, patients with DM type 2 have abnormalities in various hormone levels. These associations may be considered in the pathogenesis of the disease and should be taken into account for the treatment of patients of DM type 2.

SUMMARY AND CONCLUSIONS

Mean insulin and cortisol levels were significantly higher in diabetics. In conclusion, patients with DM type 2 have abnormalities in lipid profile, cortisol and insulin hormone levels. These associations may be considered in the pathogenesis of the disease and should be taken into account for the treatment of patients of DM type 2.

The role of various strategies targeted at maintaining cortisol levels in improving health profile of DM 2 patients, might be considered.

REFERENCES

1. Ashby, J. And Frier, B.: Circulating C-Peptide: Measurement and Clinical Applications. *Annals of clinical Biochemistry*. 18:125, 1981
2. Beischer, W.: Proinsulin and C-Peptide in Humans. *Hormones in Normal and Abnormal Human Tissues*. Volume 3K, Fotherby and Pal, S., ed. (Berlin: Walter DeGruyter). Pp. 1-43, 1983.
3. Tietz NW. *Clinical guide to Laboratory Tests*. 2nd edition. Philadelphia, USA. WB Saunders; 1990. p. 554-6.
4. L. Thomas, *Labor and Diagnose*, 4. Auflage, 1992.
5. Piroli GG, Grillo CA, Reznikov LR, Adams S, McEwen BS, Charron MJ, et al. Corticosterone Impairs Insulin-Stimulated Translocation of GLUT4 in the Rat Hippocampus. *Neuroendocrinol* 2007; 85:71–80.
6. Chiodini I, Adda G, Morelli V, Lembo S, Epaminonda P, Masserini B. Cortisol Secretion in Patients With Type 2 Diabetes. *Diabetes Care* 2007; 30:83-8.
7. Reynolds RM, Labad J, Strachan MWJ, Brarm A, Lee AJ, Frier BM et al. Morning cortisol levels and cognitive abilities in people with Type 2 Diabetes. *Diabetes Care* 2010; 33:714-20.
8. Goldstein DE, Little RR, Wedmayer HM. Glycated haemoglobin: methodology and clinical application. *Clin Chem* 1986; 32:64-70.
9. Poretsky L. On the paradox of insulin induced hyperandrogenism in insulin resistant states. *Endocr Rev* 1989; 12:3-13.
10. De Quervain DJ, Roozendaal B, Nitsch RM, McGaugh JL, Hock C. Acute cortisone administration impairs retrieval of long-term declarative memory in humans. *Nat Neurosci* 2000; 3:313–4.
11. Kerstin MO, Baerbel D, Bernd S, Hans HR, Ulrich S, Jan B. Glucocorticoid-induced diabetes and adrenal suppression: How to detect and manage them. *Cleve Clin J Med* 2011; 78:748-56.
12. De Quervain DJ, Roozendaal B, McGaugh JL. Stress and glucocorticoids impair retrieval of long-term spatial memory. *Nature* 1998; 394:787–90.