

Changes in the Expression of Insulin Pathway, Neutrophil Elastase and Alpha 1 Antitrypsin Genes from Leukocytes of Young Individuals with Insulin Resistance

Walter David Cruz-Pineda¹, Olga Lilia Garibay-Cerdenares^{2,3}, Hugo Alberto Rodríguez-Ruíz^{1,3}, Inés Matia-García¹, Linda Anahí Marino-Ortega¹, Mónica Espinoza-Rojo⁴, Zyanya Reyes-Castillo⁵, Natividad Castro-Alarcón⁶, Eduardo Castañeda-Saucedo⁷, Berenice Illades-Aguilar³, Icela Parra-Rojas¹

¹Laboratorio de Investigación en Obesidad y Diabetes, Facultad de Ciencias Químico-Biológicas, Universidad Autónoma de Guerrero, Chilpancingo, Guerrero, México; ²CONACyT-Universidad Autónoma de Guerrero, Chilpancingo, Guerrero, México; ³Laboratorio de Biomedicina Molecular, Facultad de Ciencias Químico-Biológicas, Universidad Autónoma de Guerrero, Chilpancingo, Guerrero, México; ⁴Laboratorio de Biología Molecular y Genómica, Facultad de Ciencias Químico-Biológicas, Universidad Autónoma de Guerrero, Chilpancingo, Guerrero, México; ⁵Instituto de Investigaciones en Comportamiento Alimentario y Nutrición, Centro Universitario del Sur, Universidad de Guadalajara, Ciudad Guzmán, Jalisco, México; ⁶Laboratorio de Investigación en Microbiología, Facultad de Ciencias Químico-Biológicas, Universidad Autónoma de Guerrero, Chilpancingo, Guerrero, México; ⁷Laboratorio de Investigación en Biología Celular del Cáncer, Facultad de Ciencias Químico-Biológicas, Universidad Autónoma de Guerrero, Chilpancingo, Guerrero, México

Correspondence: Icela Parra-Rojas, Laboratorio de Investigación en Obesidad y Diabetes, Facultad de Ciencias Químico Biológicas, Universidad Autónoma de Guerrero, Avenida Lázaro Cárdenas S/N, Ciudad Universitaria, Chilpancingo, Guerrero, CP 39090, México, Tel/Fax +52 7474719310, Email iprojas@yahoo.com; Olga Lilia Garibay-Cerdenares, CONACyT-Universidad Autónoma de Guerrero, Avenida Lázaro Cárdenas S/N, Ciudad Universitaria, Chilpancingo, Guerrero, CP 39090, México, Tel/Fax +52 7474710901, Email olgaribayce@conacyt.mx

Background: Chronic hyperinsulinemia is a hallmark of insulin resistance that affects a diversity of cells, including leukocytes modifying the expression of some genes involved in insulin signaling.

Purpose: The aim of this study was to evaluate how hyperinsulinemia affects the expression of genes involved in the proximal insulin signaling pathway in leukocytes from 45 young individuals grouped: normal weight with not insulin resistance (NIR), with insulin resistance (IR) and with obesity (OB-IR).

Methods: qPCR was performed to analyze the expression of insulin receptor (INSR), insulin receptor substrate 1 and 2 (IRS-1 and IRS-2), neutrophil elastase (NE), alpha 1 antitrypsin (A1AT), glucose transporters 1, 3 and 4 (GLUT-1, GLUT-3 and GLUT-4) by the $2^{-\Delta C_t}$ method, and the correlation between the genes was determined by Spearman's test.

Results: The mRNA expression analysis of all genes between NIR and IR individuals revealed no differences. However, when comparing NIR and IR individuals with OB-IR, an increase in NE and A1AT expression and a clear trend towards a decrease in IRS-2 expression was observed, whereas the comparison of IR and OB-IR showed a decrease in GLUT-3 expression. Overall, the correlation analysis showed that in the IR group there was a positive correlation only between NE with IRS-1 ($r = 0.72$, $p = 0.003$), while in the OB-IR group, there was a positive correlation between the NE and A1AT with INSR ($r = 0.62$, $p = 0.01$ and $r = 0.74$, $p = 0.002$, respectively) and with IRS-2 ($r = 0.74$, $p = 0.002$ and $r = 0.76$, $p = 0.001$, respectively).

Conclusion: These results suggest that hyperinsulinemia and obesity are associated with changes in the expression of genes in leukocytes involved in the insulin pathway that are related to NE and A1AT.

Keywords: insulin resistance, obesity, insulin signaling, leukocytes, neutrophil elastase, alpha 1 antitrypsin

Introduction

Insulin resistance is a condition in which cells, especially those of insulin-dependent metabolic tissues, cease to respond to the physiological stimuli of insulin.¹ It is considered a condition that is associated with type 2 diabetes,² obesity,^{3,4} and

nonalcoholic hepatic steatosis,⁵ among others, and the hallmark metabolic alteration is hyperinsulinemia.⁶ The causes of insulin resistance are diverse and include inflammation,⁷ oxidative stress,⁸ and lipotoxicity.^{9,10} Insulin resistance is an important pathophysiological mechanism in the development of metabolic syndrome, which includes cardiometabolic alterations such as central obesity, hypertension, and dyslipidemia, and affects approximately one-quarter of the world population.¹¹ More than 80 million people in the United States alone are affected,^{12,13} and in Mexico, a prevalence of 41% has been reported.¹⁴ Furthermore, insulin resistance is considered to be a factor that increases the risk of developing type 2 diabetes.^{15,16}

Obesity is the predominant disease associated with insulin resistance,¹⁷ as the frequency of this condition in individuals with obesity is as high as 76%.¹⁸ In addition, at least two phenotypes have been identified: metabolically healthy individuals with obesity who have a high body mass index (BMI) ($\text{BMI} \geq 30 \text{ kg/m}^2$) but lack metabolic syndrome and show insulin sensitivity¹⁹ and normal weight individuals who have a normal BMI ($\text{BMI} < 25 \text{ kg/m}^2$) but present with a metabolism associated with obesity and insulin resistance.²⁰ The prevalence of insulin resistance in normal weight individuals is 16 to 46.5%,^{18,21} and the molecular mechanisms involved in the development of this condition are not fully understood.

Insulin is produced by pancreatic β -cells, and it is the hormone critical for stimulating plasma glucose uptake in adipose and muscle tissue²² and leukocytes^{23,24} and suppresses endogenous glucose production in the liver.¹ The signaling pathway is initiated when insulin binds to its receptor and induces signaling through adaptor proteins such as Src homology and collagen (Shc) and insulin receptor substrates (IRS), which are both crucial for regulating mitogenic processes and glucose uptake.²⁵ The IRS-1 and IRS-2 isoforms are the main proteins that transmit signals in cells to coordinate glucose homeostasis. However, although they are homologous, these isoforms have distinct and, in some cases, complementary functions, and they are both regulated in a tissue-dependent manner.²⁶ Insulin resistance can be induced by aberrant IRS degradation through the involvement of neutrophil elastase (NE), a serine protease stored in leukocyte azurophilic granules, which is released, in response to inflammation, into the extracellular space where it has immunomodulatory and tissue remodeling effects.²⁷

The effects of NE are intrinsically regulated by its main inhibitor, alpha 1 antitrypsin (A1AT),²⁸ which is an acute-phase protein synthesized mainly in the liver; however, leukocytes also synthesize A1AT through a mechanism of local regulation of inflammation.²⁹

Insulin resistance impacts leukocytes, among other cells, by affecting their glucose uptake as well as their metabolism. This influence leads to the deterioration of their functions, which is associated with the development of other chronic degenerative complications, such as metabolic syndrome and cardiovascular diseases.³⁰⁻³²

The final result of the insulin signaling pathway in glucose uptake is the mobilization of GLUT transporters to the membrane: GLUT-1 is a transporter considered to be independent of insulin signaling,³³ whereas the main insulin-dependent transporter is GLUT-4,³⁴ although insulin has also been shown to promote GLUT-3 translocation.³⁵

Insulin exerts a regulatory role on the abundance and post-translational modifications of proteins,³⁶ however, it also shows effects on the regulation of gene expression of proteins involved in its signaling pathway on classical target cells (hepatic, muscle and adipose), as well as in other tissues that are not commonly associated with metabolic functions.^{37,38}

Dysregulation in the abundance of NE and A1AT proteins is implicated in processes such as inflammation, development of obesity and hepatic steatosis, in addition to the fact that NE has direct proteolytic activity on IRS-1 and IRS-2 when its abundance is higher than that of A1AT, leading to insulin resistance.^{39,40} However, it has not been explored whether insulin resistance by itself induces changes in the gene expression of molecules involved in the proximal insulin signaling pathway in leukocytes, especially in those individuals with hyperinsulinemia with normal weight or obesity. Therefore, the aim of this study was to analyze the expression of these genes and to detect changes among young individuals with normal weight or obesity who present insulin resistance. This work is shown the changes in the expression of these genes and their association with NE and A1AT expression, indicating a probable relationship with the insulin signaling pathway.

Materials and Methods

Individuals

A total of 45 unrelated individuals (14 males and 31 females) in an age range of 18 to 26 years were included and assigned to three groups: 15 with normal weight ($\text{BMI} \leq 24.9 \text{ kg/m}^2$) without insulin resistance ($\text{HOMA-IR} < 3.2$) termed the NIR group; 15 with normal weight ($\text{BMI} \leq 24.9 \text{ kg/m}^2$) with insulin resistance ($\text{HOMA-IR} \geq 3.2$) termed the IR group; and 15 with obesity ($\text{BMI} \geq 30 \text{ kg/m}^2$) and with insulin resistance ($\text{HOMA-IR} \geq 3.2$) termed the OB-IR group. All participants are originally from the State of Guerrero, Mexico. Individuals with signs of infectious disease or receiving any treatment that might influence biochemical or hematological parameters (previous diagnosis of diabetes, pregnant women, polycystic ovary syndrome, liver or kidney disease) were excluded from the study. Written informed consent was obtained from all the participants according to the ethical guidelines of the Declaration of Helsinki 2008, and the study was approved by the Research Ethics Committee of the Autonomous University of Guerrero.

Clinical and Anthropometric Measurements

Weight, BMI and fat mass were determined with a body composition analyzer (Tanita BC-553, Arlington, USA) with participants wearing lightweight clothing and wearing no shoes. Body circumferences were measured with anthropometric tape (Seca, Hamburg, Germany). Assignment of participants into the normal weight and obesity groups was made based on a $\text{BMI} \leq 24.9 \text{ kg/m}^2$ for the normal weight group and a $\text{BMI} \geq 30 \text{ kg/m}^2$ for the obesity group. Blood pressure was measured by placing the individual seated in a chair with support using a blood pressure monitor (HEM-7120, Omron Healthcare Inc, Hoffman Estates, IL, USA) on the left arm after a 10-minute rest. Systolic blood pressure (SBP) and diastolic blood pressure (DBP) were the mean of two readings, and they were taken at least 10 minutes apart.

Laboratory Measurements

A 5 mL blood sample was obtained from each participant by antecubital venipuncture under fasting conditions. Serum glucose, cholesterol, triglycerides, HDL-C and LDL-C levels were analyzed with semiautomatic equipment (Mindray BS-200, Shenzhen, China) using commercially available kits (Spinreact, Barcelona, Spain), and insulin levels were measured by enzyme-linked immunosorbent assay (GenWay INS-EASIA kit). To determine insulin resistance, a homeostasis model assessment (HOMA-IR) was performed according to the protocol described by Matthews et al 1985, using the formula: $\text{fasting serum insulin } (\mu\text{U/mL}) \times \text{fasting plasma glucose } (\text{mmol/L}) / 22.5$.⁴¹ Insulin resistance was defined as a HOMA-IR value equal to or above the 90th percentile (≥ 3.2) as the cutoff value, obtained from analysis of apparently healthy individuals with normal weight and serum glucose levels.

Gene Expression Analysis

Total RNA extraction was performed with leukocytes isolated from peripheral blood with TRIzol reagent (Thermo Fisher Scientific, Waltham, MA, USA) and according to the method of Chomczynski and Sacchi.⁴² Synthesis of cDNA was performed using 1 μg of RNA with a commercial kit following the manufacturer's specifications (Promega Corporation, USA) and stored at -80°C until use. Gene expression analysis was performed by real-time PCR. NE (ELANE, Hs00236952_m1), A1AT (SERPINA1, Hs00165475_m1), INSR (Hs00961558_m1), IRS-1 (Hs00178563_m1), IRS-2 (Hs00275843_s1), GLUT-1 (SLC2A1, Hs00892681_m1), GLUT-3 (SLC2A3, Hs00359840_m1) and GLUT-4 (SLC2A4, Hs0016168966_m1) levels were analyzed in duplicate using TaqMan probes (Thermo Fisher Scientific, Waltham, MA, USA) according to the manufacturer's instructions with a CFX96 Touch Real-Time PCR Detection System Analyzer (BIO-RAD, California, USA). Relative expression of each gene was performed using the $2^{-\Delta\text{Ct}}$ method, and the endogenous β -actin (ACTB, Hs03023943_g1) was used as a reference. To validate the absence of GLUT-4 gene expression in the leukocytes of the individuals analyzed, human adipose tissue was used as the positive control.

Statistical Analysis

The analysis was performed with STATA (V.9.2) and GraphPad Prism 5 software. Differences in variables between groups were evaluated by Fisher's exact test, ANOVA or Kruskal–Wallis according to the distribution of the data. Post hoc analysis between groups was done using the Tukey or Dunn test, as appropriate. To evaluate the association between genes in the insulin signaling pathway, NE and A1AT, Spearman's correlation test was performed. Values of $p < 0.05$ were considered significant.

Results

Characteristics of the Study Population

A total of 45 nondiabetic, nonhypertensive, apparently healthy individuals participated in the study. The clinical and anthropometric characteristics of the NIR, IR and OB-IR groups are summarized in Table 1. The age range was 18–26 years, and differences were observed between groups with respect to weight, BMI, fat mass, waist circumference, hip circumference, insulin and HOMA-IR ($p < 0.001$) as well as in age, waist-to-hip ratio (WHR), systolic blood pressure (SBP), diastolic blood pressure (DBP), cholesterol and triglyceride levels ($p < 0.05$). The post hoc analysis confirms the differences between the OB-IR group with respect to NIR and between the IR and OB-IR groups.

Hyperinsulinemia Induces Changes in the Expression of Genes Involved in the Insulin Signaling Pathway in IR and OB-IR Individuals

The expression of genes involved in the insulin signaling pathway, NE and A1AT was analyzed in leukocytes. Figure 1 shows the mRNA expression levels of each gene in the study groups. For insulin receptor (INSR), a low mRNA expression level was observed in the IR and OB-IR groups, but differences were observed only between the OB-IR and NIR groups ($p = 0.0440$). The mRNA expression of IRS-1 was lower in the IR group, and mRNA expression of IRS-2 was decreased in the IR and OB-IR groups. However, a comparison among groups did not reveal significant differences in the mRNA expression of either IRS isoform. In the OB-IR group, mRNA expression of NE was significantly higher than that in the NIR ($p = 0.0314$) and IR ($p = 0.0391$) groups, similar results were observed for mRNA expression of A1AT with NIR ($p = 0.0012$) and IR ($p = 0.0045$) groups. Finally, mRNA expression of GLUT-1 did not differ in any of the groups; however, mRNA expression of GLUT-3 was significantly decreased in the OB-IR group compared to the IR group ($p = 0.0432$). Interestingly, GLUT-4 was not expressed in the leukocytes in the three groups studied.

The NE and A1AT Gene Expression Correlates with the Insulin Signaling Pathway

Table 2 shows an overall and between-group correlation analysis of the expression levels of certain genes involved in the insulin signaling pathway in leukocytes. For all individuals, there was a correlation between NE and IRS-1 ($r = 0.43$; $p = 0.003$), whereas A1AT correlated with INSR ($r = 0.31$; $p = 0.04$) and IRS-2 ($r = 0.31$; $p = 0.04$). When performing a correlation analysis by group, NIR showed a correlation between A1AT and IRS-1 ($r = 0.80$; $p = 0.0003$). For the IR group, a correlation between NE and IRS-1 ($r = 0.72$; $p = 0.003$) was found. In the OB-IR group, correlations of NE and A1AT ($r = 0.80$; $p = 0.0003$), both NE and A1AT with INSR ($r = 0.62$; $p = 0.01$, $r = 0.74$; $p = 0.002$, respectively); and with IRS-2 ($r = 0.74$; $p = 0.002$, $r = 0.76$; $p = 0.001$, respectively) were found.

Table 3 shows the correlation between NE and A1AT gene expression and anthropometric and metabolic parameters. NE correlated with BMI ($r = 0.30$; $p = 0.048$) and WHR ($r = 0.36$; $p = 0.02$), while A1AT demonstrated a negative correlation with the following parameters: weight ($r = -0.52$; $p = 0.0003$), BMI ($r = -0.51$; $p = 0.0003$), fat mass (%) ($r = -0.52$; $p = 0.0003$), fat mass (kg) ($r = -0.51$; $p = 0.0003$), waist circumference ($r = -0.41$; $p = 0.005$), hip circumference ($r = -0.53$; $p = 0.0002$), insulin ($r = -0.41$; $p = 0.005$), HOMA-IR ($r = -0.42$; $p = 0.004$) and triglyceride levels ($r = -0.33$; $p = 0.03$).

Discussion

Chronic hyperinsulinemia is a characteristic of insulin resistance; therefore, every cell in the body is influenced by the microenvironment created by this condition.^{43,44} It is clear that the transient increase in metabolic mediators, such as

Table 1 Anthropometric and Biochemical Parameters by Study Group

Variables	NIR (n = 15)	IR (n = 15)	OB-IR (n = 15)	p value	p value
Age (years) ^c	18(18–20)	19(18–21)	21(19–22)	0.02	0.06* 0.004** 0.14***
Sex n (%) ^a				0.61	
Women	12(80)	10(67)	9(60)		
Men	3(20)	5(33)	6(40)		
Height (cm) ^c	156(150–162)	156(152–169)	162(154–167)	0.22	
Weight (kg) ^c	48.1(46–59.5)	50.9(46–58.3)	82.6(79.2–87.2)	0.0001	0.39* <0.001** <0.001***
BMI (kg/m ²) ^c	20.3(19.4–22.5)	20.9(19.4–23.1)	31.3(30.7–32.7)	0.0001	0.30* <0.001** <0.001***
Fat mass (%) ^b	24.6 ± 5.9	23.8 ± 6.1	34.4 ± 5.1	<0.001	0.92* <0.001** <0.001***
Fat mass (kg) ^c	11.2(9–18.9)	12(11.1–16.9)	29.7(25.9–33.3)	0.0001	0.28* <0.001** <0.001***
Waist (cm) ^c	72(65–77)	70(64–80.7)	93(87.5–105)	0.0001	0.28* <0.001** 0.0001***
Hip (cm) ^c	90.5(85–94)	89(88–84)	110(105.5–113)	0.0001	0.47* <0.001** <0.001***
WHR ^c	0.77(0.75–0.79)	0.78(0.74–0.87)	0.85(0.78–0.93)	0.048	0.23* 0.008** 0.045***
SBP (mmHg) ^c	101(91–104)	107(101–114)	111(108–116)	0.003	0.02* 0.0004** 0.11***
DBP (mmHg) ^b	61 ± 9.0	66.4 ± 10.2	71.6 ± 7.7	0.01	0.25* 0.007** 0.27***
Glucose (mg/dL) ^b	74 ± 7.3	80.2 ± 9.3	82 ± 12.1	0.07	
Insulin (μU/mL) ^c	10.9(8.36–13.2)	23.5(20.1–27.5)	39.2(30.5–46.9)	0.0001	0.0001* <0.001** 0.02***
HOMA-IR ^c	2.02(1.28–2.52)	4.74(3.82–5.41)	7.93(5.79–8.34)	0.0001	0.0001* <0.001** 0.02***

(Continued)

Table 1 (Continued).

Variables	NIR (n = 15)	IR (n = 15)	OB-IR (n = 15)	p value	p value
Cholesterol (mg/dL) ^c	166(149–181)	167(135–172)	188(163–211)	0.04	0.32* 0.03** 0.008***
Triglycerides (mg/dL) ^c	70(58–90)	79(64–100)	124(100–141)	0.01	0.24* 0.002** 0.01***
LDL-C (mg/dL) ^b	101.8 ± 22.4	104.2 ± 25.8	101.8 ± 38.4	0.97	
HDL-C (mg/dL) ^b	39.5 ± 3.7	39.1 ± 3.6	37.4 ± 3.4	0.23	

Notes: ^aData presented as n and percentage (%); Fisher's exact test. ^bData presented as mean ± SD; ANOVA test followed by Tukey's post hoc multiple comparisons test. ^cData presented as median and 25th and 75th percentile; Kruskal–Wallis test followed by Kruskal–Wallis with multiple comparisons (post hoc) showing differences between: *NIR vs IR, **NIR vs OB-IR, ***IR vs OB-IR. p values <0.05 were considered significant. Bold numbers highlight statistical significance.

Abbreviations: WHR, waist to hip ratio; SBP, systolic blood pressure; DBP, diastolic blood pressure; LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol.

insulin after food intake, is favorable for resolving postprandial glycemia, but its chronic elevation, even when minimal, can generate toxicity.⁴⁵ In this study, expression levels of genes in the insulin signaling pathway, NE and A1AT were evaluated in leukocytes obtained from apparently healthy individuals and from insulin-resistant individuals with normal weight or obesity. Even though individuals from the 3 groups analyzed showed normal blood glucose, cholesterol and triglyceride levels, individuals in the NIR group had the lowest levels of these three parameters, including insulin and HOMA-IR levels.

INSR is present in all cells of an organism; however, its expression levels vary depending on tissue type and in response to microenvironmental signals.^{46–49} In the IR group, serum insulin and HOMA-IR levels were observed to be elevated twofold; however, in the OB-IR group, these values were elevated almost fourfold compared to those of the NIR group, with a slight trend toward increased glucose levels. In INSR expression, it was significantly decreased in the OB-IR group. One of the mechanisms of regulation in INSR expression is insulin signaling itself, where a greater availability of nutrients, in this case a greater amount of circulating insulin, generates negative feedback that limits the expression of the receptor itself and IRS-2 by inhibiting the activity of the transcription factor FoxO1 to modulate signal intensity,⁵⁰ although this mechanism has not been evaluated in leukocytes. The results showed that INSR expression in the OB-IR group was positively correlated with all the genes evaluated in this study. These results suggest that impairment of the proximal pathway of insulin signaling in leukocytes from individuals with obesity and insulin resistance may impact intracellular signaling, which could affect leukocyte metabolic and immune functions.

No studies have evaluated the expression of IRS in leukocytes of individuals with insulin resistance and obesity, but it is clear that it plays an important role in the transduction of signals that regulate leukocyte functions. Although the role of IRS in leukocyte signaling is not known, in this work, the changes in IRS-1 and IRS-2 expression were not significant between groups. It is known that in monocytes from individuals susceptible to developing type 2 diabetes, IRS-2 expression levels are decreased because of insulin resistance.⁵¹ In obesity, IRS-2 gene expression is decreased in macrophages, leading to increased production of proinflammatory cytokines and preventing their polarization toward the M2 phenotype.⁵² However, a trend toward decreased IRS-2 expression was observed in both the IR and OB-IR groups that may be related to the presence of insulin resistance, probably associated with chronic low-grade inflammation and immune cell dysfunction in obesity conditions.^{53,54}

One of the recently described mechanisms in insulin resistance is the dysregulation of NE and its main inhibitor, A1AT. In the present study, no differences in NE expression were observed between individuals with normal weight with or without insulin resistance, although a trend toward an increase was observed in the IR group. Moreover, in these individuals, NE correlated with the expression of IRS-1, which is its degradation target that may lead to insulin resistance. NE is known to be synthesized in leukocytes such as neutrophils, monocytes and lymphocytes and stored

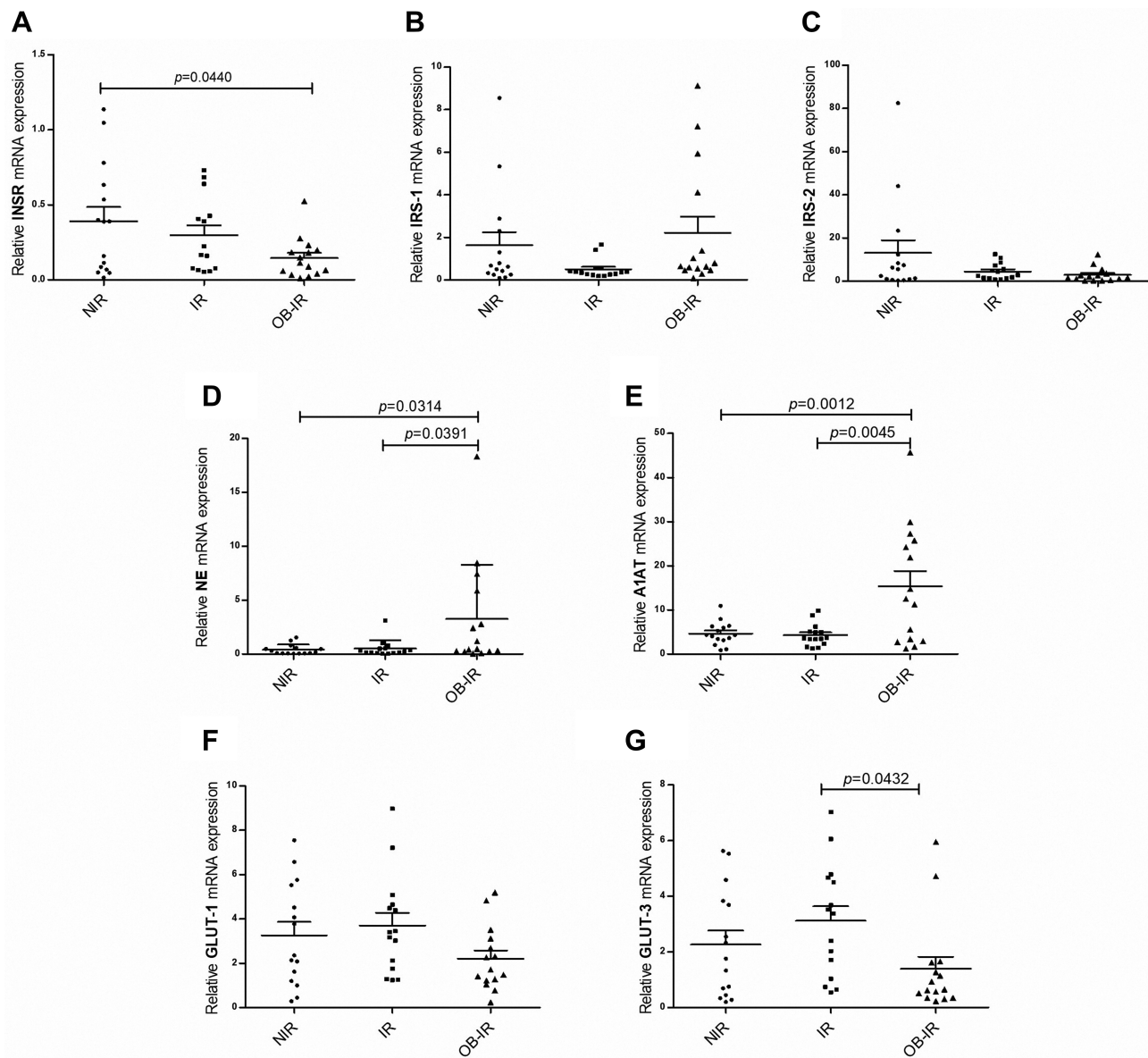


Figure 1 Relative expression of genes of the insulin signaling pathway, NE and A1AT. Relative mRNA expression of INSR (A), IRS-1 (B), IRS-2 (C), NE (D), A1AT (E), GLUT-1 (F) and GLUT-3 (G) genes. The analysis was performed using the $2^{-\Delta\Delta Ct}$ method and β -actin as reference gene. The *p* value was calculated using ANOVA and Tukey's tests.

in azurophilic granules, in response to inflammatory signals. In the OB-IR group, NE correlated with A1AT and additionally with IRS-2 expression and probably has a regulatory effect on these proteins in the insulin signaling pathway. These results that show an increase of NE expression in OB-IR group were similar to Ali et al, 2018, who found that NE levels in peripheral blood leukocytes were higher in individuals with overweight or obesity and insulin-resistance than in subjects with normal weight.⁵⁵ Elevation in A1AT expression may inhibit the proinflammatory reactions associated with NE, possibly acting as a local anti-inflammatory mechanism.⁵⁶ The results suggest that increased NE expression in peripheral blood leukocytes may be associated with the chronic low-grade inflammation that accompanies obesity but not with hyperinsulinemia per se because the NE expression levels did not correlate with insulin or HOMA-IR levels, although A1AT did (Table 3). Ferdous et al, 2020 reported decreased NE levels in individuals with newly diagnosed type 2 diabetes,⁵⁷ these individuals showed no significant differences in

Table 2 Correlations Between the Expression of Insulin Signaling Pathway, NE and AIAT Genes

Variables	NE		AIAT		IRS-1		IRS-2		GLUT-1		GLUT-3	
	r	p	r	p	r	p	r	p	r	p	r	p
Total												
INSR	0.06	0.69	0.31	0.04	0.07	0.64	0.75	<0.001	0.75	<0.001	0.66	<0.001
IRS-1	0.43	0.003	0.18	0.25	-	-	-	-	-	-	-	-
IRS-2	0.11	0.48	0.31	0.04	0.09	0.56	-	-	-	-	-	-
NE	-	-	-	-	-	-	-	-	-	-	-	-
AIAT	0.15	0.31	-	-	-	-	-	-	-	-	-	-
GLUT-1	0.13	0.40	0.36	0.02	0.18	0.24	0.64	<0.001	-	-	-	-
GLUT3	0.09	0.54	0.37	0.01	0.06	0.69	0.68	<0.001	0.64	<0.001	-	-
NIR												
INSR	-0.33	0.23	-0.02	0.95	0.10	0.71	0.91	<0.001	0.84	0.0001	0.61	0.01
IRS-1	0.30	0.28	0.80	0.0003	-	-	-	-	-	-	-	-
IRS-2	-0.29	0.29	-0.004	0.99	0.14	0.63	-	-	-	-	-	-
NE	-	-	-	-	-	-	-	-	-	-	-	-
AIAT	0.48	0.07	-	-	-	-	-	-	-	-	-	-
GLUT-1	-0.33	0.23	0.06	0.83	0.21	0.44	0.91	<0.001	-	-	-	-
GLUT3	-0.07	0.81	0.07	0.81	0.13	0.66	0.60	0.02	0.58	0.02	-	-
IR												
INSR	0.26	0.37	0.02	0.96	0.12	0.68	0.31	0.29	0.49	0.08	0.69	0.007
IRS-1	0.72	0.003	0.05	0.86	-	-	-	-	-	-	-	-
IRS-2	0.32	0.25	0.17	0.54	0.14	0.63	-	-	-	-	-	-
NE	-	-	-	-	-	-	-	-	-	-	-	-
AIAT	0.40	0.14	-	-	-	-	-	-	-	-	-	-
GLUT-1	0.49	0.06	0.19	0.49	0.21	0.44	0.91	<0.001	-	-	-	-
GLUT3	0.29	0.29	0.17	0.55	0.13	0.66	0.60	0.02	0.58	0.02	-	-
OB-IR												
INSR	0.62	0.01	0.74	0.002	0.58	0.02	0.75	0.001	0.71	0.003	0.81	0.0003
IRS-1	0.31	0.25	0.40	0.14	-	-	-	-	-	-	-	-
IRS-2	0.74	0.002	0.76	0.001	0.25	0.36	-	-	-	-	-	-
NE	-	-	-	-	-	-	-	-	-	-	-	-
AIAT	0.80	0.0003	-	-	-	-	-	-	-	-	-	-
GLUT-1	0.65	0.009	0.82	0.0002	0.45	0.10	0.70	0.004	-	-	-	-
GLUT3	0.58	0.02	0.76	0.001	0.54	0.04	0.86	<0.001	0.71	0.003	-	-

Notes: Spearman's correlation coefficient (r). p values <0.05 were considered significant. Bold numbers highlight statistical significance.

blood insulin levels but significant differences in hyperglycemia, which correlated negatively with NE expression levels.

Leukocytes require constant energy production to fulfill the energetic demands involved in their activation during host defense, utilizing glucose that enters cells through GLUT transporters. GLUT-1 expression levels were not different among the groups. GLUT-1 transporter expression levels have not been evaluated in leukocytes of individuals with insulin resistance; however, increased GLUT-1 expression has been found in the muscle of individuals with this condition.⁵⁸

GLUT-3 transporter levels were found to be different only between the IR and OB-IR groups. In the skeletal muscle of insulin-resistant individuals, decreased GLUT-3 expression levels have been reported.⁵⁸ Insulin treatment has been shown to increase the expression of both GLUT-1 and GLUT-3, indicating that their biosynthesis is insulin-dependent.⁵⁹

Table 3 Correlation Between NE and A1AT Gene Expression Among Anthropometric and Metabolic Parameters

Variables	NE		A1AT	
	r	p	r	p
Age	0.19	0.20	-0.03	0.83
Weight	0.17	0.26	-0.52	0.0003
Height	0.02	0.88	-0.07	0.63
BMI	0.30	0.048	-0.51	0.0003
Fat mass (%)	0.05	0.74	-0.52	0.0003
Fat mass (kg)	0.21	0.16	-0.51	0.0003
Waist	0.25	0.10	-0.41	0.005
Hip	0.18	0.23	-0.53	0.0002
WHR	0.36	0.02	-0.08	0.63
SBP	0.22	0.14	-0.30	0.047
DBP	0.17	0.26	-0.24	0.11
Glucose	0.20	0.20	-0.05	0.76
Insulin	0.28	0.07	-0.41	0.005
HOMA-IR	0.29	0.06	-0.42	0.004
Triglycerides	0.19	0.20	-0.33	0.03
Cholesterol	0.09	0.54	-0.18	0.23
LDL-C	0.07	0.64	0.07	0.63
HDL-C	-0.21	0.16	0.14	0.36

Notes: Spearman's correlation coefficient (r). p values <0.05 were considered significant. Bold numbers highlight statistical significance.

Abbreviations: WHR, waist-to-hip ratio; SBP, systolic blood pressure; DBP, diastolic blood pressure; LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol.

These results may suggest that the insulin signaling pathway regulates the expression of these transporters to maintain glucose uptake, which is dysregulated in obesity.

Although the presence of the three GLUT isoforms has been reported in peripheral blood leukocytes,⁶⁰ in this study, GLUT-4 expression was not observed in any of the groups; therefore, their capacity to store large amounts of energy, compared to classical target cells in metabolic tissues, is limited. Whereas leukocytes are not cells that could be used as classical models of insulin signaling, they are cells that depend on glucose metabolism for their survival,⁶¹ differentiation,⁶² activation⁶³ and immune functions,^{64,65} so that tight regulation of insulin-induced translocation of GLUT-3 transporters to the membrane is required to modulate glucose uptake,²³ due that deterioration of insulin signaling could promote excessive glucose uptake and generate immune hyperactivation and even immune pathology.

There are certain limitations in the present study, due to the expression of the components in the insulin signaling pathway at the protein level, as well as the phosphorylation of molecules such as INSR or IRS-1/2, which are necessary for signal transduction, were not evaluated.

Conclusion

In the present study, the expression of certain genes involved in the insulin signaling pathway, NE and A1AT in peripheral blood leukocytes was analyzed, and a differential expression pattern was found among the three groups studied. The expression of insulin signaling pathway genes was significantly altered, and these changes were associated with the increased expression of NE and A1AT, suggesting that the hyperinsulinemia that occurs in obesity may be related to the expression of these genes. A positive correlation between NE and A1AT expression and INSR, IRS-2, GLUT-1 and GLUT-3 was found in the OB-IR group. Future studies are needed to elucidate the molecular mechanisms involved in insulin resistance in leukocytes, where NE and A1AT may play relevant role.

Data Sharing Statement

The authors indicate that all supporting data are included within the main article. The data that support the findings of this study are available from the corresponding author upon reasonable request.

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Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

Disclosure

The authors declare no conflicts of interest.

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