

Retinal Microperimetry: A Useful Tool for Detecting Insulin Resistance-Related Cognitive Impairment in Morbid Obesity

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Abstract: Background: There is clear association between type 2 diabetes (T2D) and cognitive decline. Retinal microperimetry is a useful tool for detecting cognitive impairment in T2D. Morbid obesity (MO) has been associated with cognitive impairment. Insulin resistance (IR) seems a major determinant, but the data are unclear. The aim of this study was to evaluate the cognitive impairment in MO as well as the utility of retinal microperimetry in identifying these alterations. Methods: In total, 50 consecutive patients with MO were matched by age and gender with 30 healthy controls. All patients underwent cognitive evaluation (Montreal Cognitive Assessment Test-MoCA) and retinal microperimetry, using MAIA microperimeter 3rd generation. Retinal sensitivity and gaze fixation parameters were used for the evaluation of the analysis. Results: MO patients showed a significantly lower neurocognitive performance than the controls: MoCA score 24.94 $\pm$ 2.74 vs. 28.95 $\pm$ 1.05, $p < 0.001$. Cognitive function inversely correlated with the HOMA-IR ($r = 0.402$, $p = 0.007$). The AUROC for cognitive impairment using microperimetry was 0.807, CI 95% (0.592–0.947), $p = 0.017$. Conclusions: (1) Systemic insulin resistance is a major underlying mechanism accounting for the higher prevalence of cognitive impairment detected in young MO subjects. (2) Retinal microperimetry is a useful tool for identifying MO patients with cognitive impairment.

Keywords: morbid obesity; cognitive impairment; retinal microperimetry; insulin resistance

1. Introduction

Obesity represents a major public health problem and is associated with significant economic burden in the health systems of developed countries. This is mainly due to associated co-morbidities, type 2 diabetes (T2D) being the most important. At present, there is a clear relationship between type 2 diabetes and dementia [1,2]. Furthermore, T2D acts as an accelerator for cognitive decline [3] and

shares with Alzheimer's disease (AD) common pathophysiological pathways, such as inflammation, oxidative stress, and insulin resistance (IR) [4,5].

T2D has been associated with lower grey matter volumes and white matter disruption [6,7], as well as with impairment of perceptual speed, attention, and primary memory [8]. At present, there are no reported phenotypic indicators or reliable cost-effective screening tests for identifying T2D patients with cognitive impairment. Moreover, the diagnosis is based on neuropsychological tests whose complexity makes their incorporation into current standards of care for the T2D population unfeasible.

Recent evidence has shown that retinal neurodegeneration is an early event in the pathogenesis of diabetic retinopathy (DR) [9,10]. The retina is ontogenically a brain-derived tissue, and it has been suggested that it may provide an easily accessible and noninvasive way of examining the pathology of the brain [11,12]. Therefore, it seems reasonable to propose that the evaluation of retinal parameters related to neurodegeneration, such as retinal function, could be useful for identifying those patients with type 2 diabetes who have a high risk of developing AD. Several methods can be used to measure retinal function. Among them, fundus-driven microperimetry has emerged as a simple, noninvasive, and rapid test that can be used in clinical practice [13,14]. Microperimetry measures retinal sensitivity in terms of the minimum light intensity that patients can perceive when spots of light stimulate specific areas of the retina. Some recent studies have suggested that microperimetry is even more sensitive than multifocal electroretinography in detecting early functional changes of the retina [15]. Our group showed that retinal sensitivity assessed by fundus-driven microperimetry was related to brain neurodegeneration and could be a useful biomarker for identifying those patients at risk of developing AD [16]. Additionally, gaze fixation parameters evaluated during microperimetry improved the predictive value of retinal sensitivity in identifying cognitive impairment in subjects with T2D [17].

Obese patients with metabolic syndrome without hyperglycemia are more prone to showing global cognitive decline [18,19]. More recently, it has been demonstrated that cognitive decline even occurs at younger age, with the executive function and the perceptual speed related to visuospatial task being especially altered [20,21]. Several studies showed small brain volumes and white matter disruption in obese patients in comparison with normo-weighted counterparts [22–24]. Furthermore, similar brain structure findings related to AD (example cortical glyosis, hyppocampal atrophy, beta-amyloid deposits) were found in obese murine models [25,26]. Since similar alterations in brain structures are found in both T2D and obesity, it can be postulated that IR is a major underlying mechanism of the increased prevalence of AD reported in these populations. However, further clinical and basic research to support this concept is needed.

The aim of the present study was to explore, by means of a pilot study, whether retinal sensitivity and gaze fixation assessed by microperimetry could be useful for identifying obese patients with cognitive impairment and their relationship with IR.

2. Material and Methods

Between September 2018 and December 2018, a total of 50 consecutive patients who were attending the obesity outpatient clinic of Vall Hebron University Hospital for bariatric surgery (BS) evaluation were included in a case-control study as a part of the PREDIBAR trial- NCT 03784508. The inclusion criteria (cases) were: (a) Age between 18 and 60 years; (b) fulfill the criteria for BS; (c) previous accomplishment of the preoperative protocol for BS at our site; (d) ability to read and understand the specific questionnaires; and (e) written informed consent form. The exclusion criteria were: (a) Any form of diagnosed cognitive impairment; and (b) neurodegenerative diseases of the brain or retina (i.e., glaucoma) or cerebrovascular diseases. The control group comprised healthy persons with normal weight matched by age. Most of them were family members of patients included in the study from the same catchment area.

The study was approved by the local ethics committee and carried out in accordance with the Declaration of Helsinki.

All patients underwent complete physical, biochemical, and psychiatric evaluations as part of the preoperative protocol for bariatric surgery at our site. Cognitive function was evaluated using the Spanish version of Minimental State Evaluation (MMSE) [27] and the Montreal Cognitive Assessment Test (MoCA) [28]. The MoCA test evaluates several domains of cognitive function: Visual spatial executive (5 points), naming (3 points), attention (6 points), language (3 points), abstraction (2 points), delayed recall (5 points), and orientation (6 points). The required administration time is approximately 10 minutes. The maximum score is 30. The score was adjusted by age and education level (if the education level was <12 years, 1 point was added). We used a cut-off of 26 to define mild cognitive impairment as previously described [28].

Retinal sensitivity was evaluated by fundus-driven microperimetry MAIA 3rd generation after a previous pupillary dilation of a minimum of 4 mm. The standard MAIA test covers a 10-diameter area with 37 measurement points, and a red 1 radius circle was used as the fixation target. We employed an expert exam 4–2 strategy (Goldmann III size stimulus, background luminance of 4 asb, and maximum luminance of 1000 asb, with a 25 decibel (dB) dynamic range). For the evaluation of fixation, the MAIA microperimeter uses high-speed eye trackers (25 Hz) for the control of fixation losses and to register the fixation pattern. All fixation positions during the examination are used by the instrument to calculate the fixation indexes, P1 and P2, which represent the percentage of fixation points inside a circle of 2 and 4 degrees of diameter, respectively. Subjects were classified as having stable fixation if P1 was more than 75%, relatively unstable fixation if P1 was less than 75% and P2 more than 75%, and unstable fixation if both P1 and P2 were below 75%. Microperimetry also provides a more accurate estimation of the fixation pattern using the bivariate contour ellipse area (BCEA). This area is calculated as an ellipse that covers fixation eye positions and takes into account 1 or 2 times the standard deviation, including 63% (BCEA63) and 95% (BCEA95) of the fixation points. The area of this ellipse is calculated through major and minor axes, which are 2 orthogonal diameters, describing the extent of the fixation points (horizontal and vertical diameters) [29]. For retinal sensibility and fixation, data corresponding to the right eye was used, which was the first eye explored in all subjects.

Statistical Analysis

To assess differences between groups, the Chi-square test for qualitative variables and ANOVA followed by DMS post-hoc tests for quantitative variables were used. To evaluate the correlations, Spearman's correlation test and regression analysis were performed. Significance was accepted at the level of $p < 0.05$. The Bonferroni correction was used for multiple comparisons. Logistic regression was used to predict the ROC curves and the Chi-squared test for ROC area comparison. Statistical analyses were performed with SPSS 24.0 statistical package.

3. Results

The baseline demographical and laboratory data as well as the neurocognitive and psychological evaluation of obese patients and healthy controls included in the study are shown in Table 1. It should be noted that as per the protocol requirements, we did not find any difference in age.

We did not find differences in the results of MMSE between obese and control subjects. By contrast, a significant difference in the global cognitive function evaluated by MoCA was observed (Table 1). The main cognitive domains affected in the obese group were the executive function, attention, and recent memory (delayed recall) (Table 2).

Regarding microperimetry assessment, we found no significant differences between the obese patients and controls in terms of retinal sensitivity. By contrast, we found significant differences in all the parameters related to gaze fixation: P1, P2, BCEA63, and BCEA95 (Table 3).

The BCEA95 parameter, which is the most integrative parameter of gaze fixation, was significantly correlated to the MoCA score ($R = 0.512$, $r = 0.510$, $p < 0.001$) (Figure 1).

MoCA Domains	Obese Patients	Controls	p
Visuo Spatial Executive Function	4.0 ± 0.791	4.93 ± 0.26	0.001
Naming	2.94 ± 0.243	3.00 ± 0.00	n.s.
Abstraction	1.71 ± 0.772	1.96 ± 1.92	n.s.
Delayed Recall	3.14 ± 1.581	4.11 ± 1.31	0.015
Orientation	6.00 ± 0.00	6.00 ± 0.00	n.s.

Table 1. The baseline characteristics of the subjects included in the study.

Characteristics	Total	Obese Patients	Controls	p
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Gaze fixation assessment	N but not retinal sensitivity	50	is impaired	30 in subjects with NA	morbid obesity and
Gender (females %)		71.12		70.45	n.s.
Age (years)		45.88	10.23	41.17	12.33
BMI (kg/m ²)		44.54	5.53	22.99	3.51
Regarding microperimetry	HOMA-IR assessment	6.43	we 2.64	found no	1. significant
HbA1c (%Hb DCCT)		5.96	1.15	5.24	0.27
T2D duration (month)		249.1		NA	NA

all the parameters related MoCA to score gaze fixation: 24.94 P1, P2, 74 BCEA63, 28.95 and 1. BCEA95 (Table <0.0013).

MMSE	28.24	1.14	29	0.62	n.s.
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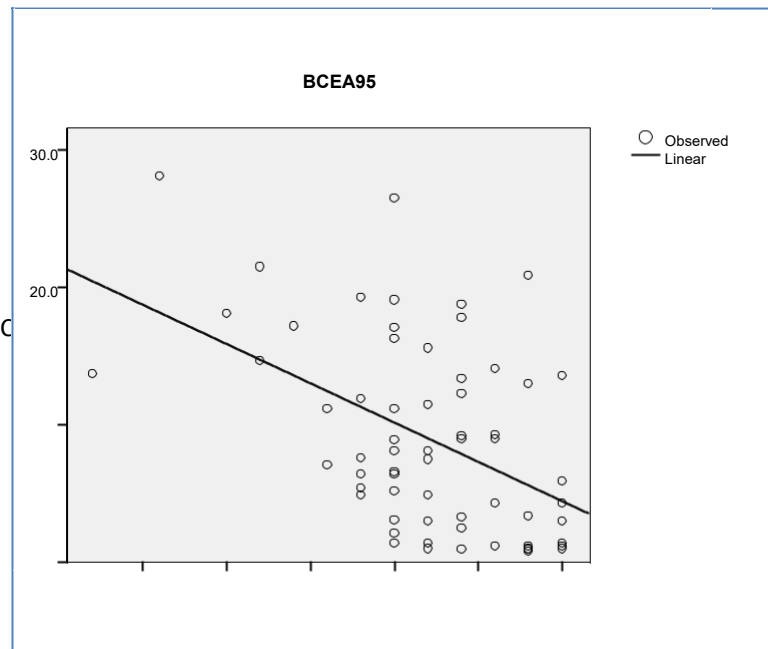
Table 3. Microperimetry parameters between obese patients and control.

Microperimetry Table 2. The scores Parameters of the different Obese domains Patients evaluated Controls by the MoCA test p.

MoCA Domains	Obese Patients	Controls	p
Visuo Spatial Executive Function	4.0 0.791	4.93 0.26	0.001
Naming	2.94 0.243	3.00 0.00	0.001
Attention	4.41 1.73	5.67 0.62	0.001
Language	2.71 0.588	2.91 0.32	n.s.
Abstraction	1.71 0.772	1.96 1.92	n.s.
Reliability index	93.52 ± 11.75	95.45 ± .07	0.015
Orientation	6.00 0.00	6.00 0.00	n.s.
Total	24.94 2.74	28.95 1.05	<0.001

The BCEA95 parameter, which is the most integrative parameter of gaze fixation, was

significantly correlated with the MoCA Assessment Score (R = 0.512, β = -0.510, p < 0.001) (Figure 1).



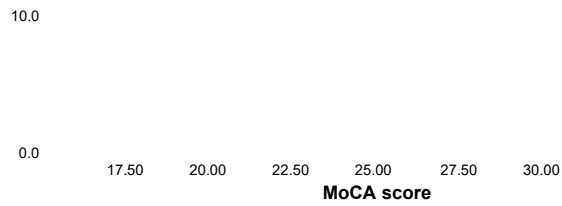


Figure 1 Correlation between BECA95 and MoCA. BECA95: BECA95; 95% bivariate confidence ellipse area. MoCA: Montreal Cognitive Assessment test.

MoCA: Montreal Cognitive Assessment test.

Additionally, the BECA95 parameter was able to predict the cognitive status of the patients evaluated in the study by means of the MoCA test, showing an AUROC of 0.807 (CI 95% (0.592–0.947), $p = 0.017$) (Figure 2).

Table 3. Microperimetry parameters between obese patients and control.

Microperimetry Parameters	Obese Patients	Controls	p
N	50	30	NA
Sensitivity (dB)	27.6 3.81	29.28 1.39	n.s.
Fixation P1 (%)	78.48 26.16	97.18 3.2	0.001
Fixation P2 (%)	90.31 17.84	98.40 4.39	0.001
BCEA63	1.50 1.32	0.40 0.34	0.001
BCEA95	8.33 4.66	3.01 2.01	0.021
Reliability index	93.52 11.75	95.45 15.07	n.s.

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0.947), $p = 0.017$) (Figure 2).

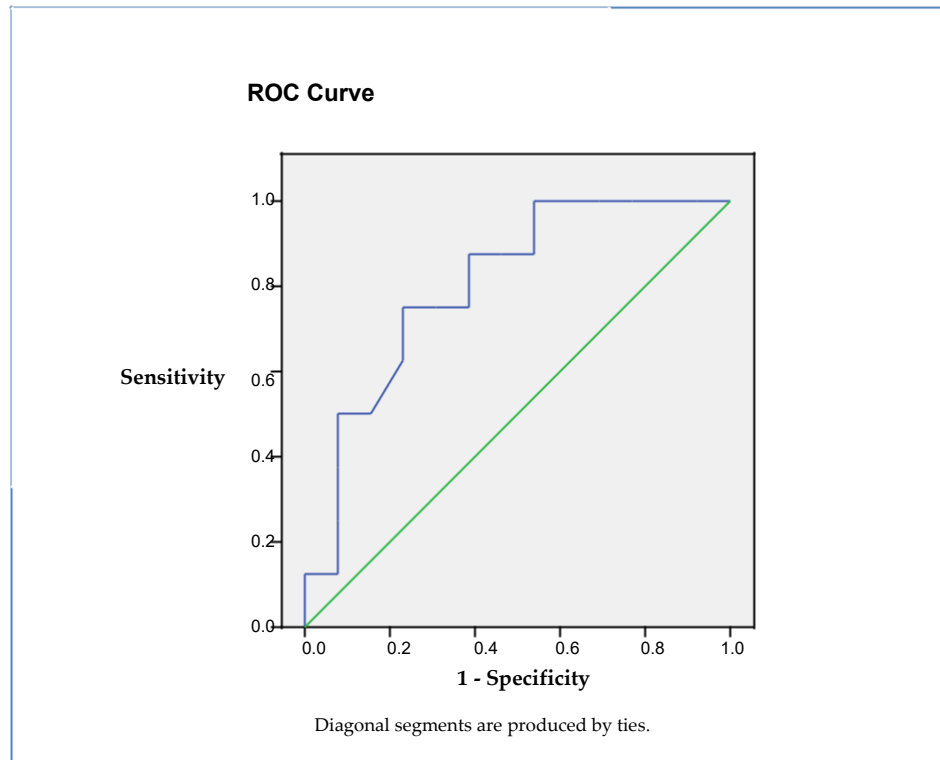


Figure 2. AUROC of BCEA95 and MoCA. MoCA: Montreal Cognitive Assessment test.

Insulin Resistance (IR) as an Independent Predictor of Cognitive Status

The characteristics of obese subjects regarding the presence of diabetes are displayed in Table 4.

4. We did not find significant differences in the demographic characteristics, cognitive function, retinal sensitivity, or gaze fixation between groups. As expected, HOMA-IR and HbA1C levels

retinal sensitivity, or gaze fixation between groups. As expected, HOMA-IR and HbA1C levels were significantly higher in obese subjects with diabetes.

Regarding antidiabetic treatment, 35% were on a diet only, 52% of the patients received one

Regarding antidiabetic treatment, 35% were on a diet only, 52% of the patients received one drug (metformin), and 13% received two drugs (metformin plus GLP-1 receptor agonists). No patient was on insulin treatment.

on insulin treatment.

Since obese patients Table 4. Comparison between cognitive MoCA patient scores with using T2D and MoCA, without T2D, wanted to explore the variables accounting for this finding. For this purpose, a multiple regression analysis, including as Characteristics Obese Patients with T2D Obese Patients Without T2D *p*

variables age, gender, T2D status, HbA1c levels, BMI, and HOMA-IR, was performed. We found that

HOMA-IR Gender was the (female only independent %) predictor 70.37 of cognitive impairment 71.42 as reflected by n Table.s. 5.

	N	24	26	NA
Age (years)	47.63 ± 8.62	44.33 ± 11.41	n.s.	
BMI (kg/m ²)	44.36 ± 5.5	44.70 ± 5.66	n.s.	
HOMA-IR	8.34 ± 2.08	4.71 ± 2.047	0.001	
HbA1c (%Hb DCCT)	6.62 ± 1.3	5.33 ± 0.28	0.001	
T2D duration (month)	24 ± 9.1	NA	NA	
MoCA score	25.5 ± 2.61	24.44 ± 2.92	n.s.	
MMSE	28.1 ± 1.53	28.09 ± 1.78	n.s.	
Retinal sensitivity (dB)	27.76 ± 2.45	27.59 ± 4.58	n.s.	

Table 4. Comparison between MO patients with T2D and without T2D.

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Characteristics	Obese Patients with T2D	Obese Patients Without T2D	p
Fixation P1 (%)	82.48 ± 23.59	78.19 ± 27.6	n.s.
N	24	26	NA
FixationGender(femaleP2(%))	92.0070±.3717.21	89.5471.42±18.04	n.sn..s.
Age (years)	47.63 8.62	44.33 11.41	n.s.
BCEA63	44.36 5.4	44.70 5.66	n.s.
BCEA95	8.14 ± 4.5	8.46 ± 4.85	n.s.
HOMA-IR	8.34 2.08	4.71 2.047	0.001
Reliability index	91.52 ± .75	92.65 ± 12.96	n.s.
HbA1c (%Hb DCCT)	6.62 14.3	5.33 0.28	0.001
T2DEducationdurationlevel(month)	7.0824± 09..331	7.16NA± 0.28	NA n.s.
MoCA score	25.5 2.61	24.44 2.92	n.s.
MMSE	28.1 1.53	28.09 1.78	n.s.
Retinal sensitivity (dB)	Since obese patients presented poor cognitive scores using MoCA, we wanted to explore the	27.59 4.58	n.s.

variables accounting for this finding. For this purpose, a multiple regression analysis, including.s as variables age,Fixationgender,P2(%)T2D status, HbA1c92.00 levels,17.21 BMI, and HOMA89-.54IR, was18.04 performed.nWe.s. found

Coefficients(a)									
Model	Unstandardized Coefficients		Standardized Coefficients		Sig.	95% Confidence Interval		Collinearity Statistics	
	B	Std. Error	Beta	Coe cients(a)		Lower Bound	Upper Bound	Tolerance	VIF
(Constant)	25.200	3.734		6.749	0.000	17.684	32.715		
Gender	-.503	0.776	-.084	-.0648	0.520	-2.065	1.060	0.869	1.151
Age	-.063	0.034	-.242	-1.823	0.075	-0.132	0.007	0.831	1.203
Type 2									
Diabetes	-.169	0.286	-.076	0.589	0.559	-0.745	0.408	0.888	1.127
(Constant)	25.200	3.734		6.749	0.000	17.684	32.715		
HbA1c levels	0.233	0.375	0.081	0.620	0.538	0.523	0.989	0.869	1.151
Gender	0.233	0.503	0.776	0.084	0.648	0.520	2.065	0.869	1.151
BMI	0.078	0.050	0.209	1.574	0.122	0.022	0.179	0.836	1.196
Age	0.063	0.034	0.034	0.242	1.823	0.075	0.132	0.831	1.203
HOMA-IR	-0.149	0.053	-0.375	-2.836	0.007	0.559	-0.255	0.842	1.188
HbA1c levels	0.233	0.375	0.081	0.620	0.538	0.523	0.989	0.869	1.151
BMI	0.078	0.050	0.209	1.574	0.122	0.022	0.179	0.836	1.196
HOMA-IR	0.149	0.053	0.375	2.836	0.007	0.255	0.043	0.842	1.188

In fact, the cognitive function evaluated by the MoCA score was correlated with the insulin resistance state evaluated by HOMA -IR (R = 0.402, $p < 0.0001$) as reflected in Figure 3.

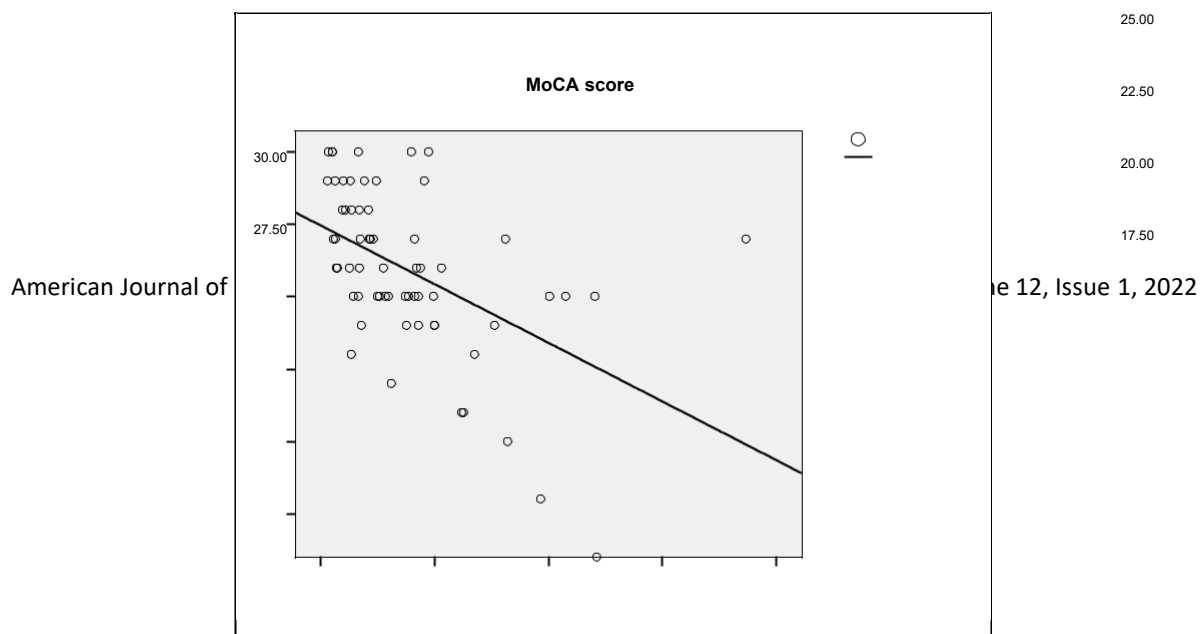




Figure 3. Correlation between MoCA score and HOMA-IR. MoCA: Montreal Cognitive Assessment test. HOMA-IR: Homeostatic model assessment for insulin-resistance.

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Finally, we found a significant correlation between BCEA95 and HOMA-IR ($R = 0.411$, $\beta = 0.470$; $p < 0.001$) (Figure 4).

$p < 0.001$) (Figure 4).

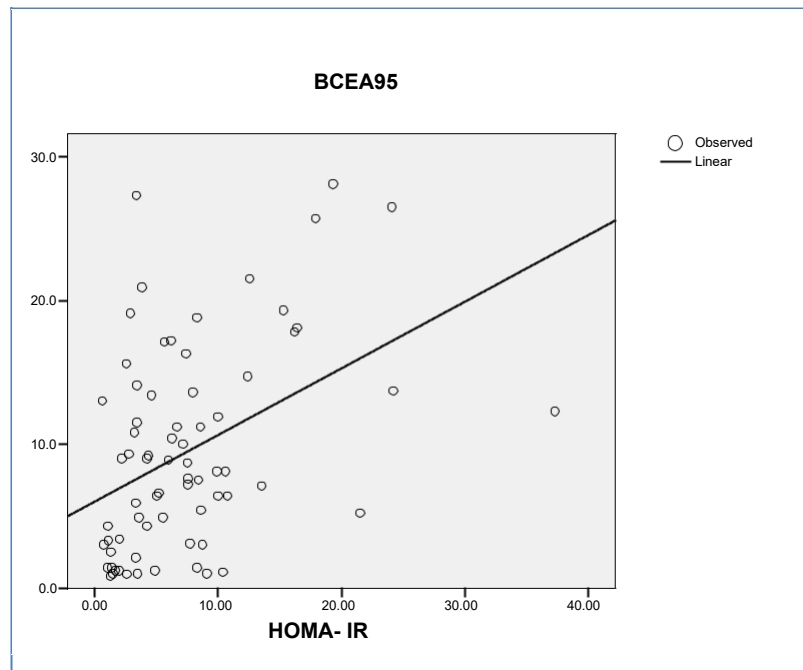


Figure 4. Correlation between BCEA95 and HOMA-IR.. BCEA95: 95% bivariate contour ellipse area.. HOMA-IR: Homeostatic model assessment for insulin-resistance..

4. Discussion

4. Discussion

In this study, we found a significant reduction in the cognitive function in young morbidly obese patients, when compared to their counterparts with normal weight, regardless of the presence of obese patients, when compared to their counterparts with normal weight, regardless of the diabetes. In addition, we found that cognitive impairment assessed by the MoCA score adjusted by presence of diabetes. In addition, we found that cognitive impairment assessed by the MoCA score age correlated with the insulin resistance state evaluated by HOMA-IR. Furthermore, the assessment adjusted by age correlated with the insulin resistance state evaluated by HOMA-IR. Furthermore, of gaze fixation by microperimetry was significantly correlated to the MoCA score and the HOMA-IR. the assessment of gaze fixation by microperimetry was significantly correlated to the MoCA score

Overall, our results suggest that IR could be a primary underlying mechanism accounting for both and the HOMA-IR. Overall, our results suggest that IR could be a primary underlying mechanism accounting for both cognitive impairment related to obesity and gaze fixation abnormalities.

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The latter can be very useful for early detection of mild cognitive impairment and dementia [28] and The latter can be very useful for early detection of mild cognitive impairment and dementia [28] we found a significantly lower score in obese patients in comparison with controls. It should be noted and we found a significantly lower score in obese patients in comparison with controls. It should be that the MMSE score was within the normal range in all cases and no differences were found in our noted that the MMSE score was within the normal range in all cases and no differences were found

study in the MMSE score between cases and controls. This finding suggests that MMSE is not useful in our study in the MMSE score between cases and controls. This finding suggests that MMSE is not for detecting early cognitive impairment in young obese patients. Regarding the cognitive domains useful for detecting early cognitive impairment in young obese patients. Regarding the cognitive explored by MoCA, we found significant reductions in the obese patient group in the visual spatial domains explored by MoCA, we found significant reductions in the obese patient group in the executive function, attention, and delayed

There is a clear relationship between T2D and cognitive impairment [3,4]. Clinical studies showed that T2D is associated with smaller brain volumes in grey matter [32]. These structural network abnormalities were related to the slowing of the information-processing speed [33]. Interestingly, brain damage and white matter disruption have been reported in the IR stage without overt diabetes [34]. These structural abnormalities could be involved in early cognitive impairment, in particular in perceptual speed, attention, and primary memory [34]. These studies were performed in middle-aged or older adults whereas we found similar results in terms of cognitive impairment in younger obese subjects. It should be noted that we did not observe differences in the cognitive status between obese subjects with and without diabetes, but IR was an independent predictor of cognitive decline. Taken together, these findings strongly suggest that the pathways related to insulin signaling rather than blood glucose levels play a key role in the development of cognitive impairment in the natural history of diabetes.

The impairment of insulin signaling (leading to insulin resistance) in the brain plays an essential role in the pathogenesis of dementia and precedes cognitive dysfunction and pathological alterations even by decades. Insulin signaling is essential for neuronal survival, as well as for learning and memorizing. In fact, in animal models, impaired insulin signaling precedes A β accumulation [35,36]. Once formed, A β oligomers deplete insulin receptors from the surface membrane of neurons, thereby contributing to insulin resistance [37], and produce an aberrant phosphorylation of insulin receptor substrate (IRS). This reduces the normal pro-survival signaling in neurons and promotes their death by apoptosis [38]. It should be noted that most of these studies focus on brain insulin resistance, rather than systemic IR. By contrast, in the present study, HOMA-IR (a measurement of systemic IR) was independently associated with the MoCA score. This raises the question of whether insulin-sensitizer drugs could be useful in preventing or arresting cognitive decline. In this regard, Luchsinger et al. [39] showed a slight arrest in cognitive decline related to lowered fasting insulin levels in middle-aged (65 years) non-diabetic patients with obesity (BMI 31 kg/m²) after 12 months of treatment with metformin. Additionally, Alosco et al. [40] showed a significant improvement in the cognitive function in young MO at 12 months follow-up after BS.

Another interesting finding in our study was that the gaze fixation parameters (ex. BECA 95) were correlated with both the HOMA-IR and cognitive function. In fact, the AUROC of BECA95 for identifying young morbidly patients with cognitive impairment was 0.807 (CI 95%: 0.592–0.947). We previously reported that retinal sensitivity evaluated by retinal microperimetry was able to identify T2D subjects with cognitive impairment [16]. It should be noted that in this previous study, the T2D patients with Alzheimer's disease and with mild cognitive Impairment were not obese but were insulin resistant. The patients presented a BMI of 27.72 $\pm$ 3.79 kg/m² and 28.82 $\pm$ 4.48 kg/m², respectively. Furthermore, the predictive value in identifying those T2D subjects with mild cognitive impairment was significantly increased by adding fixation parameters to the retinal sensibility [17]. This finding could be attributed to the fact that the brain areas involved in retinal fixation are not the

same as in retinal sensitivity. Therefore, the combined test (retinal sensitivity + gaze fixation) results in an improvement of the efficiency of microperimetry in detecting MCI due to the examination of different neural circuits. In this regard, retinal sensibility relies on the retina and the neural pathway of vision. The first station of the optic tract is the lateral geniculate body of the thalamus, which is a major visual processing region in the brain and plays a crucial role in relaying information from the retinal ganglion cells to the primary visual cortex [41]. By contrast, the superior colliculus and the parietal and frontal cortex play an essential role in gaze fixation. There is evidence that both the geniculate body and the superior colliculus are affected by AD [42–44].

We have found that gaze fixation is impaired earlier than retinal sensitivity in obese patients. There are several reasons that could explain this finding, but perhaps the fact that gaze fixation depends on white matter complex connectivity networks, which are impaired earlier in subjects with IR, could be one of the most important [45,46].

To the best of our knowledge, this is the first study that provides a clinical link between systemic insulin resistance in obese subjects, cognitive impairment, and abnormal gaze fixation assessed by microperimetry. Our results suggest that retinal microperimetry could be a useful tool in identifying early cognitive impairment in MO patients. Further studies are needed to validate our results and also to evaluate the possible role of retinal microperimetry in monitoring the evolution of cognitive impairment and its relationship with IR after BS or pharmacological interventions.

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References

1. Biessels, G.J.; Strachan, M.W.J.; Visseren, F.L.J.; Kappelle, L.J.; Whitmer, R.A. Dementia and cognitive decline in type 2 diabetes and prediabetic stages: Towards targeted interventions. *Lancet Diabetes Endocrinol.* **2014**, *2*, 246–255. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/24622755> (accessed on 1 October 2019). [CrossRef]
2. Exalto, L.G.; Whitmer, R.A.; Kappelle, L.J.; Biessels, G.J. An update on type 2 diabetes, vascular dementia and Alzheimer's disease. *Exp. Gerontol.* **2012**, *47*, 858–864. [CrossRef]
3. Ciudin, A.; Espinosa, A.; Simó-Servat, O.; Ruiz, A.; Alegret, M.; Hernández, C.; Boada, M.; Simo, R. Type 2 diabetes is an independent risk factor for dementia conversion in patients with mild cognitive impairment. *J. Diabetes Complicat.* **2017**, *31*, 1272–1274. [CrossRef] [PubMed]
4. Simó, R.; Ciudin, A.; Simó-Servat, O.; Hernández, C. Cognitive impairment and dementia: A new emerging complication of type 2 diabetes—The diabetologist's perspective. *Acta Diabetol.* **2017**. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/28210868> (accessed on 1 October 2019).
5. Biessels, G.J.; Reagan, L.P. Hippocampal insulin resistance and cognitive dysfunction. *Nat. Rev. Neurosci.* **2015**, *16*, 660–671. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/26462756> (accessed on 1 October 2019).
6. Moran, C.; Beare, R.; Phan, T.; Srikanth, V. Brain, imaging in type 2 diabetes. In *Type 2 Diabetes and Dementia*; Academic Press: Cambridge, MA, USA, 2019; Volume 92, pp. 823–830. [CrossRef]
7. Reijmer, Y.D.; Brundel, M.; de Bresser, J.; Kappelle, L.J.; Leemans, A.; Biessels, G.J. Microstructural white matter abnormalities and cognitive functioning in type 2 diabetes: A diffusion tensor imaging study. *Diabetes Care* **2013**, *36*, 137–144. [CrossRef] [PubMed]

8. Marseglia, A.; Fratiglioni, L.; Laukka, E.J.; Santoni, G.; Pedersen, N.L.; Bäckman, L.; Xu, W. Early cognitive deficits in type 2 diabetes: A population-based study. *J. Alzheimer's Dis.* **2016**, *53*, 1069–1078. [[CrossRef](#)]
9. Simó, R.; Hernández, C. Novel approaches for treating diabetic retinopathy based on recent pathogenic evidence. *Prog. Retin. Eye Res.* **2015**, *48*, 160–180. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/25936649> (accessed on 1 October 2019).
10. Simó, R.; Hernández, C.; European Consortium for the Early Treatment of Diabetic Retinopathy (EUROCONDOR). Neurodegeneration in the diabetic eye: New insights and therapeutic perspectives. *Trends Endocrinol. Metab.* **2014**, *25*, 23–33. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/24183659> (accessed on 1 October 2019).
11. Cheung, C.Y.; Ikram, M.K.; Chen, C.; Wong, T.Y. Imaging retina to study dementia and stroke. *Prog. Retin. Eye Res.* **2017**, *57*, 89–107. [[CrossRef](#)]
12. Liao, H.; Zhu, Z.; Peng, Y. Potential utility of retinal imaging for Alzheimer's Disease: A review. *Front. Aging Neurosci.* **2018**, *10*, 188. [[CrossRef](#)]

13. Acton, J.H.; Greenstein, V.C. Fundus-driven perimetry (microperimetry) compared to conventional static automated perimetry: Similarities, differences, and clinical applications. *Can. J. Ophthalmol.* **2013**, *48*, 358–363. [[CrossRef](#)] [[PubMed](#)]
14. Rohrschneider, K.; Bültmann, S.; Springer, C. Use of fundus perimetry (microperimetry) to quantify macular sensitivity. *Prog. Retin. Eye Res.* **2008**, *27*, 536–548. [[CrossRef](#)] [[PubMed](#)]
15. Wu, Z.; Ayton, L.N.; Guymer, R.H.; Luu, C.D. Comparison between multifocal electroretinography and microperimetry in age-related macular degeneration. *Investig. Ophthalmol. Vis. Sci.* **2014**, *55*, 6431–6439. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/25159206> (accessed on 1 October 2019). [[CrossRef](#)] [[PubMed](#)]
16. Ciudin, A.; Simó-Servat, O.; Hernández, C.; Arcos, G.; Diego, S.; Sanabria, Á.; Sotolongo, Ó.; Hernández, I.; Boada, M.; Simó, R. Retinal microperimetry: A new tool for identifying patients with type 2 diabetes at risk for developing Alzheimer disease. *Diabetes* **2017**, *66*, 3098–3104. [[CrossRef](#)]
17. Simó-Servat, O.; Ciudin, A.; Ortiz-Zúñiga, Á.M.; Hernández, C.; Simó, R. Usefulness of eye fixation assessment for identifying type 2 diabetic subjects at risk of dementia. *J. Clin. Med.* **2019**, *8*, 59. [[CrossRef](#)]
18. Profenno, L.A.; Porsteinsson, A.P.; Faraone, S.V. Meta-analysis of Alzheimer's disease risk with obesity, diabetes, and related disorders. *Biol. Psychiatry* **2010**, *67*, 505–512. [[CrossRef](#)]
19. Feng, L.; Chong, M.S.; Lim, W.S.; Lee, T.S.; Collinson, S.L.; Yap, P.; Ng, T.P. Metabolic syndrome and amnesic mild cognitive impairment: Singapore Longitudinal Ageing Study-2 findings. *J. Alzheimer's Dis.* **2013**, *34*, 649–657. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/23246920> (accessed on 1 October 2019). [[CrossRef](#)]
20. Tsai, C.L.; Huang, T.H.; Tsai, M.C. Neurocognitive performances of visuospatial attention and the correlations with metabolic and inflammatory biomarkers in adults with obesity. *Exp. Physiol.* **2017**, *102*, 1683–1699. [[CrossRef](#)]
21. Tsai, C.L.; Chen, F.C.; Pan, C.Y.; Tseng, Y.T. The neurocognitive performance of visuospatial attention in children with obesity. *Front. Psychol.* **2016**, *7*, 1033. [[CrossRef](#)]
22. Ho, A.J.; Raji, C.A.; Becker, J.T.; Lopez, O.L.; Kuller, L.H.; Hua, X.; Lee, S.; Hibar, D.; Dinov, I.D.; Stein, J.L.; et al. Obesity is linked with lower brain volume in 700 AD and MCI patients. *Neurobiol. Aging* **2010**, *31*, 1326–1339. [[CrossRef](#)]
23. Yokum, S.; Ng, J.; Stice, E. Relation of regional gray and white matter volumes to current BMI and future increases in BMI: A prospective MRI study. *Int. J. Obes.* **2012**, *6*, 656–664. [[CrossRef](#)] [[PubMed](#)]
24. Dekkers, I.A.; Jansen, P.R.; Lamb, H.J. Obesity, brain volume, and white matter microstructure at MRI: A cross-sectional UK biobank study. *Radiology* **2019**, *292*, 270. [[CrossRef](#)] [[PubMed](#)]
25. Chunchai, T.; Chattipakorn, N.; Chattipakorn, S.C. The possible factors affecting microglial activation in cases of obesity with cognitive dysfunction. *Metab. Brain Dis.* **2018**, *3*, 615–635. [[CrossRef](#)] [[PubMed](#)]
26. Puig, K.L.; Floden, A.M.; Adhikari, R.; Golovko, M.Y.; Combs, C.K. Amyloid precursor protein and proinflammatory changes are regulated in brain and adipose tissue in a murine model of high fat diet-induced obesity. *PLoS ONE* **2012**, *7*, e30378. [[CrossRef](#)] [[PubMed](#)]
27. Folstein, M.F.; Folstein, S.E.; McHugh, P.R. "Mini-mental state". A practical method for grading the cognitive state of patients for the clinician. *J. Psychiatr. Res.* **1975**, *12*, 189–198. [[CrossRef](#)]

28. Nasreddine, Z.S.; Phillips, N.A.; Bédirian, V.; Charbonneau, S.; Whitehead, V.; Collin, I.; Cummings, J.L.; Chertkow, H. The Montreal Cognitive Assessment, MoCA: A brief screening tool for mild cognitive impairment. *J. Am. Geriatr. Soc.* **2005**, *53*, 695–699. [[CrossRef](#)]
29. Schrijvers, E.M.C.; Witteman, J.C.M.; Sijbrands, E.J.G.; Hofman, A.; Koudstaal, P.J.; Breteler, M.M.B. Insulin metabolism and the risk of Alzheimer disease: Rotterdam Study. *Neurology* **2010**, *75*, 1982–1987. [[CrossRef](#)]
30. Rasgon, N.L.; Kenna, H.A.; Wroolie, T.E.; Kelley, R.; Silverman, D.; Brooks, J.; Williams, K.E.; Powers, B.N.; Hallmayer, J.; Reiss, A. Insulin resistance and hippocampal volume in women at risk for Alzheimer's disease. *Neurobiol. Aging* **2011**, *2*, 1942–1948. [[CrossRef](#)]
31. Ursache, A.; Wedin, W.; Tirsi, A.; Convit, A. Preliminary evidence for obesity and elevations in fasting insulin mediating associations between cortisol awakening response and hippocampal volumes and frontal atrophy. *Psychoneuroendocrinology* **2012**, *37*, 1270–1276. [[CrossRef](#)]

32. Espeland, M.A.; Bryan, R.N.; Goveas, J.S.; Robinson, J.G.; Siddiqui, M.S.; Liu, S.; Hogan, P.E.; Casanova, R.; Coker, L.H.; Yaffe, K.; et al. Influence of type 2 diabetes on brain volumes and changes in brain volumes: Results from the Women's Health Initiative Magnetic Resonance Imaging studies. *Diabetes Care* **2013**, *36*, 90–97. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/22933440> (accessed on 1 October 2019). [[CrossRef](#)]
33. Reijmer, Y.D.; Leemans, A.; Brundel, M.; Kappelle, L.J.; Biessels, G.J.; Utrecht Vascular Cognitive Impairment Study Group. Disruption of the cerebral white matter network is related to slowing of information processing speed in patients with type 2 diabetes. *Diabetes* **2013**, *62*, 2112–2115. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/23349494> (accessed on 1 October 2019). [[CrossRef](#)] [[PubMed](#)]
34. Willette, A.A.; Xu, G.; Johnson, S.C.; Birdsill, A.C.; Jonaitis, E.M.; Sager, M.A.; Hermann, B.P.; La Rue, A.; Asthana, S.; Bendlin, B.B. Insulin resistance, brain atrophy, and cognitive performance in late middle-aged adults. *Diabetes Care* **2013**, *36*, 443–449. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/23069842> (accessed on 1 October 2019). [[CrossRef](#)] [[PubMed](#)]
35. Chua, L.-M.; Lim, M.-L.; Chong, P.-R.; Hu, Z.P.; Cheung, N.S.; Wong, B.-S. Impaired neuronal insulin signaling precedes A β 42 accumulation in female APPSW/PS1DE9 mice. *J. Alzheimer's Dis.* **2012**, *29*, 783–791. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/22337827> (accessed on 1 October 2019). [[CrossRef](#)] [[PubMed](#)]
36. Macklin, L.; Grieth, C.M.; Cai, Y.; Rose, G.M.; Yan, X.-X.; Patrylo, P.R. Glucose tolerance and insulin sensitivity are impaired in APP/PS1 transgenic mice prior to amyloid plaque pathogenesis and cognitive decline. *Exp. Gerontol.* **2017**, *88*, 9–18. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/28025127> (accessed on 1 October 2019). [[CrossRef](#)] [[PubMed](#)]
37. de Felice, F.G. Alzheimer's disease and insulin resistance: Translating basic science into clinical applications. *J. Clin. Investig.* **2013**, *123*, 531–539. [[CrossRef](#)] [[PubMed](#)]
38. Bomfim, T.R.; Forny-Germano, L.; Sathler, L.B.; Brito-Moreira, J.; Houzel, J.C.; Decker, H.; Silverman, M.A.; Kazi, H.; Melo, H.M.; McClean, P.L.; et al. An anti-diabetes agent protects the mouse brain from defective insulin signaling caused by Alzheimer's disease-associated A β oligomers. *J. Clin. Investig.* **2012**, *122*, 1339–1353. [[CrossRef](#)] [[PubMed](#)]
39. Luchsinger, J.A.; Perez, T.; Chang, H.; Mehta, P.; Steiner, J.; Pradabhan, G.; Ichise, M.; Manly, J.; Devanand, D.P.; Bagiella, E. Metformin in amnesic mild cognitive impairment: Results of a pilot randomized placebo controlled clinical trial. *J. Alzheimer's Dis.* **2016**, *51*, 501–514. [[CrossRef](#)]
40. Alosco, M.L.; Spitznagel, M.B.; Strain, G.; Devlin, M.; Cohen, R.; Paul, R.; Crosby, R.D.; Mitchell, J.E.; Gunstad, J. Improved memory function two years after bariatric surgery. *Obesity* **2014**, *22*, 32–38. [[CrossRef](#)]
41. Jones, E.G. A new view of specific and nonspecific thalamocortical connections. *Adv. Neurol.* **1998**, *77*, 49–71. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/9709817> (accessed on 1 October 2019).
42. Dugger, B.N.; Tu, M.; Murray, M.E.; Dickson, D.W. Disease specificity and pathologic progression of tau pathology in brainstem nuclei of Alzheimer's disease and progressive supranuclear palsy. *Neurosci. Lett.* **2011**, *491*, 122–126. [[CrossRef](#)]
43. Parvizi, J.; Van Hoesen, G.W.; Damasio, A. The selective vulnerability of brainstem nuclei to Alzheimer's disease. *Ann. Neurol.* **2001**, *49*, 53–66. [[CrossRef](#)]

44. Erskine, D.; Taylor, J.P.; Firbank, M.J.; Patterson, L.; Onofrj, M.; O'Brien, J.T.; McKeith, I.G.; Attems, J.; Thomas, A.J.; Morris, C.M.; et al. Changes to the lateral geniculate nucleus in Alzheimer's disease but not dementia with Lewy bodies. *Neuropathol. Appl. Neurobiol.* **2016**, *42*, 366–376. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/25967384> (accessed on 1 October 2019). [[CrossRef](#)] [[PubMed](#)]
45. Ellis, E.M.; Gauvain, G.; Sivyer, B.; Murphy, G.J. Shared and distinct retinal input to the mouse superior colliculus and dorsal lateral geniculate nucleus. *J. Neurophysiol.* **2016**, *116*, 602–610. [[CrossRef](#)] [[PubMed](#)]
46. Siegel, M.; Donner, T.H.; Oostenveld, R.; Fries, P.; Engel, A.K. Neuronal synchronization along the dorsal visual pathway reflects the focus of spatial attention. *Neuron* **2008**, *60*, 709–719. Available online: <http://www.ncbi.nlm.nih.gov/pubmed/19038226> (accessed on 1 October 2019). [[CrossRef](#)]