EVALUATION THE RELATION BETWEEN AGE AT THE TIME OF DIABETES DIAGNOSIS AND GLUTAMIC ACID DECARBOXYLASE (GAD) ANTIBODY IN NON-OVERWEIGHT, OBESE DIABETIC INDIVIDUALS

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ABSTRACT
Objective: Evaluation the relation between age at the time of diabetes diagnosis and glutamic acid decarboxylase (GAD) antibody in non-overweight, obese diabetic individuals. Method: A cross-sectional study on 284 non overweight-obesity diabetic patients at Hue Central Hospital from August 2017 to August 2019. All patients were measured autoantibodies glutamic acid decarboxylase (anti-GAD). GAD antibody-positive was determined when autoantibodies to GAD concentration was higher than 5 IU/mL. Clinical data (age, sex, weight, height) were obtained. Age at the time of diabetes diagnosis was interviewed. Data were analysed by SPSS version 16.0 and Medcalc software. Results: The risk of glutamic acid decarboxylase (GAD) antibody-positive in non-overweight, obese diabetic individuals increased 2.7 time when aged at the time of diabetes diagnosis 50 and older. The cut-off of age at the time of diabetes diagnosis for detecting risk of glutamic acid decarboxylase (GAD) antibody-positive in non-overweight, obese diabetic individuals was 57. Conclusion: This study showed non-overweight, obese diabetic individuals should be screened for glutamic acid decarboxylase (GAD) antibody at aged 50 and older.

Key words: autoantibodies glutamic acid decarboxylase (anti-GAD), diabetic individual, non-overweight, obese, age at the time of diabetes diagnosis.

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1. BACKGROUND
The International Diabetes Federation (IDF) released new figures that highlight the alarming growth in the prevalence of diabetes around the world, that if there is no effective measure to prevent disease progression, by 2040, the number of people with diabetes worldwide will reach 642 million [6].

The age at the time of diabetes diagnosis is also a predictor factor of the presence of glutamic acid decarboxylase (GAD) antibody. We found that the recommended age for cut-off point for detection / diagnosis of type 2 diabetes with positive autoantibodies depending on the proportion of type 1 diabetes in the general population. For white race, the rate of autoimmune diabetes with positive autoantibodies is higher than for other races, so the cut-off point for type 2 diabetes screening has lower autoantibodies. The rate of autoimmune diabetes with positive autoantibodies among Asians is assessed to be lower than the white race, so a cut off point the age at the time of diabetes diagnosis is recommended higher[1].

Objective: Evaluation the relation between age at the time of diabetes diagnosis and glutamic acid decarboxylase (GAD) antibody in non-overweight, obese diabetic individuals.

2. SUBJECTS AND METHODS
2.1. Subjects: Non overweight-obesity
diabetic patients over 35 years old, (BMI <23 at the time of the study) at the General Internal Medicine - Geriatrics Department and Endocrinology - Neurology – Respiratory department at Hue Central Hospital, from August 2017 to August 2019.

**Selection criteria:** Patients had qualify for:
- Diabetes was diagnosed according to the standards of Vietnam Association of Diabetes and Endocrinology (2016) [5].
- BMI < 23 (non overweight-obesity).
- Age of disease detection ≥ 35 years old.

**Exclusion criteria:**
- Patients disqualified prospective subjects from inclusion in the study with:
  - Age of onset of disease < 35 years old.
  - BMI ≥ 23.
  - History of gestational diabetes mellitus

**2.2. Methods:**
A described, cross-sectional study with convenience sampling.
- All patients were carried out taking parameters of age, sex, weight, height, age of disease detection.
- Quantified anti-GAD antibody by ELISA analyzer at Department of Biochemistry, Hue Central Hospital.

<table>
<thead>
<tr>
<th>Table 2.1. Anti-GAD antibody reference value *:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anti-GAD antibody value (IU/mL)</td>
</tr>
<tr>
<td>---------------------------------</td>
</tr>
<tr>
<td>&lt;5,0</td>
</tr>
<tr>
<td>≥5,0</td>
</tr>
</tbody>
</table>

* Each laboratory should establish its own reference values.

- Calculating BMI:

\[
\text{BMI} = \frac{\text{Weight (kg)}}{(\text{height})^2 (m^2)}
\]

<table>
<thead>
<tr>
<th>Table 2.2. Classification of BMI by WHO 2000 body index (BMI), apply to Asia [5]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Classification</td>
</tr>
<tr>
<td>----------------</td>
</tr>
<tr>
<td>Underweight</td>
</tr>
<tr>
<td>Normal</td>
</tr>
<tr>
<td>At-risk of obesity</td>
</tr>
<tr>
<td>Obese 1</td>
</tr>
<tr>
<td>Obese 2</td>
</tr>
</tbody>
</table>

Data were analysed by SPSS version 16.0 and Medcalc.

**3. RESULTS**

<table>
<thead>
<tr>
<th>Table 3.1. Comparison of disease detection age between two groups of diabetic patients with positive and negative anti-GAD antibody.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Criteria</td>
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<tr>
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<tr>
<td></td>
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<tr>
<td>The age at the time of diabetes diagnosis (years)</td>
</tr>
</tbody>
</table>

*Comments:* The average age at the time of diabetes diagnosis in the group of positive anti-GAD antibodies patients was higher than the negative group: 59.84 ± 12.85 (age) compared to 52.45 ± 14.07 (years), the difference was statistically significant (p <0.05).
Table 3.2. The positive anti-GAD antibodies ratio for the age of detecting diabetes was under 50 years old

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Anti-GAD antibodies status</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Negative (n=262)</td>
<td>Positive (n=22)</td>
</tr>
<tr>
<td>&lt; 50 years (n=53)</td>
<td>45</td>
<td>8</td>
</tr>
<tr>
<td>≥ 50 years (n=231)</td>
<td>217</td>
<td>14</td>
</tr>
</tbody>
</table>

Comment: In the group of patients at the time of diabetes diagnosis under 50 years old, the rate of positive anti-GAD antibodies was 15.09%, the difference was statistically significant (p <0.05). In non overweight- obesity diabetic patients at the time of diabetes diagnosis≥ 50 years old was increased relative risk of anti-GAD antibodies is 2.7 times higher.

Figure 3.1. ROC curve of age at the time diabetes diagnosis predicts risk of positive anti-GAD antibodies in non overweight- obesity diabetic patients.

Comments: When the age at the time diabetes diagnosis was > 57, there was a risk of positive anti-GAD antibodies with an area under the curve (AUC) 0.66 (95% confidence interval: 0.602 - 0.715); sensitivity 64.5% and specificity 63.64%; p <0.01

Table 3.3. Comparison of disease detection time between two groups of diabetic patients with positive and negative anti-GAD antibodies.

<table>
<thead>
<tr>
<th>Criteria</th>
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<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Negative (n=262)</td>
<td>Positive (n=22)</td>
</tr>
<tr>
<td>The age at the time of diabetes diagnosis (years)</td>
<td>7,11±6,04</td>
<td>8,05±6,86</td>
</tr>
</tbody>
</table>

Comment: The time to detect diabetes in the group with positive anti-GAD antibodies was greater than the negative group, the difference is not statistically significant (p> 0.05).
4. DISCUSSIONS

Age of detecting diabetes is also a predictor factor of the presence of anti-GAD antibodies. In our study (Table 3.1), the average age at the time of diabetes diagnosis in positive anti-GAD positive antibodies patient groups (52.45 ± 14.07) was lower than for patients with negative anti-GAD antibodies (59.84 ± 12.85), the difference was statistically significant (p < 0.05). In Tran Quang Khanh's study [10], the average age at the time of diabetes diagnosis in positive anti-GAD antibodies (51.6 ± 12.6) was higher than that of patients with negative anti-GAD antibodies (51.2 ± 11.4), the difference was not statistically significant (p > 0.05).

The age at the time of diabetes diagnosis in group of patients with positive anti-GAD antibodies earlier than negative group was 7.39 years. This result was larger than Nguyen Thi Thu Mai’s study [11]: patients with positive anti-GAD antibodies started the disease earlier than negative group with 4.04 years, but this difference was not statistically significant. Meanwhile, Tran Quang Khanh's study noted that the difference in age of disease detection between two groups was insignificant and not statistically significant [10].

There was no consensus on the age at which autoantibodies should be screened in the population of diabetic patients. Zimmet suggested the age to detect type 2 diabetes with a positive autoantibody of 25 [17].

Many other authors had proposed the cut-off point for the detection of type 2 diabetes with positive autoantibodies was over 35 years old because it overlaps with the detection of classic type 1 diabetes in the age of 25-35 [8]. The Immunology of Diabetes Society had estimated that the minimum cut-off point for the age of detecting type 2 diabetes with positive autoantibodies varies between 25-40 years old and suggests 30 years old was the average age in detecting type 2 diabetes with positive autoantibodies [12].

In contrast, in the Asian race, Zhou noted that the age at the time of diabetes diagnosis with positive autoantibodies in China is less than 40 years old in order to avoid overlap with the classic type 2 diabetes diagnosis [16]. Similarly, Tan and Thai noted that type 2 diabetic patients with positive autoantibodies in Singapore were often found in 40 years old [13]. Kobayashi presents the age at which detection of type 2 diabetes with positive autoantibodies is recommended higher.

We found that the recommended age for cut-off point for detection / diagnosis of type 2 diabetes with positive autoantibodies depending on the proportion of type 1 diabetes in the general population. For white race, the rate of autoimmune diabetes with positive autoantibodies is higher than for other races, so the cut-off point for type 2 diabetes screening has lower autoantibodies. The rate of autoimmune diabetes with positive autoantibodies among Asians is assessed to be lower than the white race, so a cut-off point the age at the time of diabetes diagnosis is recommended higher.

According to the results of Table 3.2, among diabetes patients group was diagnosed under 50 years old, the rate of positive anti-GAD antibodies was 15.09%, the difference was statistically significant (p < 0.05) in non-overweight-obesity diabetic patients at the time of diabetes diagnosis ≥ 50 years old was increased relative risk of anti-GAD antibodies is 2.7 times higher. So, these subjects do not rule out the possibility of becoming insulin dependent in the future.

According to Figure 3.1, using ROC curve of age at the time diabetes diagnosis predicts risk of positive anti-GAD antibodies in non-overweight-obesity diabetic patients was 57 with an area under the curve (AUC) 0.66 (95% confidence interval: 0.602 - 0.715); sensitivity 64.5% and specificity 63.64%; p < 0.01.

Our results were higher than other authors, and there was discussed about the association between positive autoantibodies in type 2 diabetics and the age at the time diabetes
diagnosis:

- Fourlanos S. Author et al: a prospective study in Melbourne, Australia, when comparing between two groups of type 2 diabetes patients had positive and negative anti-GAD antibodies, Fourlanos noted the age at the timetype 2 diabetes diagnosis had positive anti-GAD antibodies was less than 50 years old and this difference was significant compared to type 2 diabetes had negative anti-GAD antibodies. This was a predictive factor for the presence of anti-GAD antibodies [4].

- Author Tran QuangKhanh [10] used cut-off point to screen for autoantibodies being over 25 years old because the unknown rate of autoimmune diabetes with positive autoantibodies in the Vietnamese population. The author also did not record the difference in age, the age at the time diabetes diagnosis and disease average time between type 2 diabetes patients group with positive and negative autoantibodies.

In Finland, the type 2 diabetic population age at the time diabetes diagnosis under 45 years had a significantly higher rate of positive anti-GAD antibodies than the population group was diagnosed after 45 years old in Toumi's cross-sectional study [14].

- Carlson suggests that age over 60 is an important risk factor for both type 2 diabetes and type 2 diabetes with positive autoantibodies [2].

- Arikan believes that Turkey type 2 diabetes with positive anti-GAD antibodies were diagnosed at younger age than patients with negative anti-GAD antibodies significantly (45.1 years of age compared to 50.8 years old) [1].

- UKPDS 25 study observed that there was a negative correlation between the rate of positive autoantibodies and the age of disease detection. The rate of positive anti-GAD antibodies decreases as patients were diagnosed diabetes at a higher age [15].

- UKPDS 70 study found that type 2 diabetes patients with positive autoantibodies were usually 4 years younger than negative anti-GAD patients [3].

In contrast, Zinman commented that there was no difference in the rate of positive anti-GAD antibodies between ages based on data from the ADOPT study [18].

In Asia, Ishii's research on the Japanese diabetes population had shown that type 2 diabetes patients with positive autoantibodies who had high anti-GAD antibodies titre were often older than type 2 diabetes patients with positive antibodies had a medium or low antibodiestitre [7].

Meanwhile, Hamaguchi did not record a difference in diagnostic age between type 2 diabetes patients with positive and negative autoantibodies. Wang noted that the rate of negative autoantibodies increases gradually as type 2 diabetes patients were diagnosed at an older age [5].

5. CONCLUSIONS

Conducting research on 284 non-overweight, obese diabetic individuals, we have some comments: The risk of positive anti-GAD antibodies in non-overweight, obese diabetic individuals increased 2.7 time when aged at the time of diabetes diagnosis 50 and older. The cut-off of age at the time of diabetes diagnosis for detecting risk of positive anti-GAD antibodies in non-overweight, obese diabetic individuals was 57.

REFERENCES


2. Carlson S., Midthjell K., et al. (2007), “Age, overweight and physical inactivity increase the risk of latent autoimmune diabetes mellitus in adults: results from the Nord-Trndelag health study”,

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10. Trần Quang Khánh (2010), “Tý lệ kháng thể kháng Glutamic Acid Decarboxylase và kháng tiểu đường tự thân bệnh nhân đạiisko đường type 2, Luận án Tiến sĩ Y học, trường Đại học Y Dược thành phố Hồ Chí Minh


