IgG subclasses serum concentrations in a population of children with Down syndrome: Comparative study with siblings and general population

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SUMMARY

Serum total IgG and subclasses were determined in three different groups of children: with Down syndrome, their siblings and general pediatric population. Several cases of IgG2 and IgG4 deficiency were identified, predominantly in children with Down syndrome. The differences, considering three age groups, were statistically significant for both groups in relation to the general population group, with an increase of IgG1 and IgG3 and a decrease in serum concentrations of IgG2 and IgG4. Down syndrome children and their siblings tend to have a similar variation of the IgG4 serum concentration levels (P < 0.05). The mechanisms of this concordance are not well understood. The results point out that an adequate strategy to improve the immune status of Down syndrome children could have a positive manifestation in the immune profile of their brothers.

Key words: Down syndrome. Immunoglobulins. IgG subclasses deficiency.

MATERIAL AND METHODS

In our study were included 206 caucasian children, divided in three different groups. One group included children with T21, another formed by their siblings and a third with otherwise considered normal children. In the first group were included children with T21 followed in the Development Consultation of the University Clinic of Pediatrics. These children are ac-
complishing for a follow-up protocol with incidence in all aspects of their medical and daily life, being well integrated regarding familiar, scholar and social aspects. The T21 group was constituted by 79 individuals, 39 girls and 40 boys, with an average age (SD) of 4.6 (3.0) years and a variation between 1 and 14 years. The group of the siblings (FT21) included 73 individuals, 36 girls and 37 boys, with an average age (SD) of 7.5 (4.8) years, variation between 1 and 18 years. Fifty-four normal children formed a third group, 31 girls and 23 boys with an average age (SD) of 3.9 (3.5) years, variation between 1 and 12 years. Exclusion criteria involved all children suffering from acute or chronic systemic or respiratory infectious diseases. Samples from normal children were collect from those undergoing minor surgery procedures or followed in the outpatient clinic after hospital discharge. Informed consent was obtained in all cases. The blood samples were drawn by venopuncture according to standard procedures, centrifuged and the serum stored at –20 °C until submitted to laboratorial analysis. Serum samples processing took place in the Pediatrics Laboratory of the University School of Medicine of Hospital Santa Maria.

The IgG concentration was determined by nephelometry (Behring Nephelometer 100 Analyzer, Messer Greisheim GmbH, Frankfurt, Germany), and IgG subclasses levels by radial immunodifusion using the Radial Immunodiffusion Kit BINDARID 3 plate kits (The Binding Site Limited) according to protocol procedure defined by the manufacturer. Those values under two standard deviation below of the mean for normal children were considered as low serum concentrations.

Three age sub-groups in the groups of SD and FTSD were considered, in function of the homogeneity of the variation of serum concentrations in each one of them: below four years old, between four and eight years and above eight years old. The normality of the distribution of the values of IgG subclasses serum concentrations was verified in each group with the test of Kolmogorov-Smirnov. If normal distribution was verified then comparative matching between the diverse age sub-groups using the test t-Student was made. In alternative, the Mann-Whithney test was applied.

The results were expressed graphically using clustered boxplot showing median and quarter centile values after logarithmic transformation of data. Statistical calculations and graphic analysis were made with SPSS version 10.

RESULTS

Only in two cases, both in the group of control, lower serum concentrations of total IgG (under –2 SD) were verified, both with reduction of the IgG4. In the group of T21 we did not identify any cases of total IgG deficiency, despite the fact some individual showed lower values in several subclasses (IgG2 and IgG4), mostly due to compensatory higher values of IgG1 and IgG3 subclasses. We compared the number of the individual IgG subclass deficiency for each group (table I). Only IgG2 and IgG4 demonstrated a relevant number of deficiency cases. None of the pairs of brothers T21-FT21 showed concordance in the variation of IgG2 serum concentration towards deficiency. Among T21 group, 32/77 children evidenced low values of IgG4, and in 16 pairs of brothers T21-FT21 that we analyzed, 10/16 had low serum concentrations of IgG4 in both brothers (table II).

<table>
<thead>
<tr>
<th>Variation</th>
<th>IgG</th>
<th>IgG1</th>
<th>IgG2</th>
<th>IgG3</th>
<th>IgG4</th>
</tr>
</thead>
<tbody>
<tr>
<td>T21</td>
<td>FT21</td>
<td>Normal</td>
<td>T21</td>
<td>FT21</td>
<td>Normal</td>
</tr>
<tr>
<td>Lower</td>
<td>0</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Normal</td>
<td>73</td>
<td>74</td>
<td>54</td>
<td>73</td>
<td>75</td>
</tr>
</tbody>
</table>

Table I

Number of individuals with total IgG and subclass deficiency. There is a significant difference in the number of cases of Down syndrome with low levels of IgG4 to the other two groups ($p < 0.05$)

Table II

Qualitative alterations of serum concentrations in the cases of matching pairs of brothers T21-FT21. More than half of the pairs brothers studied had simultaneously lower values of IgG4 in serum

<table>
<thead>
<tr>
<th>Matching pairs T21-FT21</th>
<th>Number of cases</th>
</tr>
</thead>
<tbody>
<tr>
<td>↓ FT21, ↔ T21</td>
<td>1</td>
</tr>
<tr>
<td>↔ FT21, ↓ T21</td>
<td>5</td>
</tr>
<tr>
<td>↓ FT21, ↓ T21</td>
<td>10</td>
</tr>
<tr>
<td>Total</td>
<td>16</td>
</tr>
</tbody>
</table>

FT21, sibling; T21, Down syndrome; ↓, low values; ↔, normal values.
Considering the different age sub-groups, statistical significant differences between the three groups were found. Down syndrome children showed tendency to present higher values of total IgG, with statistical significance above four years old in relation to the other two groups (P < 0.05). In the oldest group of siblings, there also a significant difference to the normal population (P < 0.01). In relation to IgG1, Down syndrome children and their siblings show statistically higher values to the normal population in all age groups. Above four years old, T21 group has a significant different to the rest (P < 0.01). Down syndrome children present statistically lower values of IgG2 to the normal population below eight years old (P < 0.05) and to their siblings in the oldest groups (P < 0.01). Between 4-8 years, FT21 group was identified having also lower serum concentration, comparing to the normal population. Down syndrome children present statistically higher values of IgG3 to the normal population above four years old and to their siblings groups between 4 and 8 years old (P < 0.01); the siblings group has higher values than normal population in the oldest group (P < 0.05). In relation to IgG4, there is no difference between normal and siblings group. Down syndrome group revealed statistically lower values, compared with normal population below 8 years old (P < 0.01), and their siblings in the group between 4 and 8 years old (P < 0.05) (figs. 1-5).

**DISCUSSION**

IgG subclasses have biological properties with intrinsic variances. The complement fixation, placenta crossing, receptor binding, antigen binding and functional valence are different in the four different subtypes of IgG (1). Several studies have established an association between the deficiency of certain IgG subclasses and the predisposition for recurrent infections of the respiratory airways (6). The antibodies for virus are specially IgG1 and IgG3 associated subclasses. The response to bacterial polysaccharide antigens is predominantly an IgG2 and IgG4 reaction (2).

More than to define structural differences, it is important to consider the biological dynamics of each subclass and how this deficiency is clinically expressed. In fact, the deficiency of one or several subclasses can be conditioning factor of an immunological system incapable to answer adequately to pathogenic aggressions. In the other hand, there are several reports of individuals with IgG subclass-deficient profile without abnormal number of infections (4). This is due...
probably to compensatory immunological mechanisms present in these subjects not yet well studied. Recent studies emphasized that there is a lack of correlation between the level of IgG subclasses and the individual susceptibility to infections, claiming more importance for the antibody response to certain antigens (4).

However, several studies referred that those patients with respiratory *Haemophilus influenzae* infectious processes are more prone to have IgG deficiencies than individuals with other types of infectious diseases (1, 4, 7, 8). The IgG4 deficiency is the most common in this group. Several other studies tried to establish a correlation between the occurrence of specific IgG subclass and infectious pathology. In any case, the reported incidence of IgG subclass deficiencies vary considerably between the published studies, often limited by different methodological approaches (4).

The subclass IgG2 and IgG4 antibodies have been associated with the protection against infections by capsulated bacterial microorganisms, such as *Streptococcus pneumoniae* and *Haemophilus influenzae* (2). Particular emphasis has been placed in the role played by IgG4 (4). Diverse authors have enhanced the importance of the IgG4 in the lung and mucosal immunological defenses. Although IgG4 is

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Figure 3.—Comparison of total serum IgG2 levels (mg/dl) in three different groups, according to age; data in clustered boxplot show the median values for each case. Down syndrome children present statistically lower values to the normal population below 8 years old and to their siblings in the oldest groups (p < 0.05).

Figure 4.—Comparison of total serum IgG3 levels (mg/dl) in three different groups, according to age; data in clustered boxplot show the median values for each case. Down syndrome children present statistically higher values to the normal population above 4 years old and to their siblings groups between 4 and 8 years old; the siblings group has higher values than normal population in the oldest group (p < 0.05).

Figure 5.—Comparison of total serum IgG4 levels (mg/dl) in three different groups, according to age; data in clustered boxplot show the median values for each case. There is no difference between normal and siblings groups. Down syndrome children present statistically lower values to the normal population below 8 years old and to their siblings in the group between 4 and 8 years old (p < 0.05).
RESUMEN

Se determinaron el suero total IgG y subclases en tres grupos distintos de niños: con síndrome de Down, sus hermanos y población pediátrica general. Se identificaron varios casos de deficiencia en suero IgG2 e IgG4, principalmente en niños con síndrome de Down. Las diferencias, considerando tres grupos de edad, resultaban estadísticamente significativas en ambos grupos en relación con el grupo de población general, con un aumento de IgG1 e IgG3 y un descenso de las concentraciones de suero de IgG2 e IgG4. Los niños con síndrome de Down y sus hermanos tienen tendencia a presentar una variación similar de los niveles de concentración de suero IgG4 (p < 0,05). Los mecanismos de esta concordancia no se entienden demasiado bien. Los resultados indican que una estrategia adecuada para mejorar las condiciones inmunológicas de los niños con síndrome de Down podría tener una manifestación positiva en el perfil inmunológico de sus hermanos.

Palabras clave: Síndrome de Down. Deficiencia de subclasses de IgG.

REFERENCES


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