

POSTER PRESENTATION

Open Access

Brazilian multicentre study of Takayasu's arteritis in children and adolescents – preliminary results of a clinical, imaging and therapeutic study

Maria Teresa Terreri^{1*}, Gleice Clemente¹, Sheila K Oliveira², Marise Lessa², Clóvis Artur Silva³, Adriana Sallum³, Lúcia Campos³, Flávio Sztajnbok⁴, Rosana Gasparello⁴, André Cavalcanti⁵, Virginia Ferriani⁶, Teresa Robazzi⁷, Silvana Sacchetti⁸, Maria Odete Hilário¹

From 18th Pediatric Rheumatology European Society (PReS) Congress Bruges, Belgium. 14-18 September 2011

Background

Takayasu's arteritis (TA) is a chronic inflammatory disease that affects the aorta and its branches. Although it is the third most common vasculitis in childhood, reports with large number of patients are lacking.

Aim

To evaluate the clinical features and outcome of TA in children and adolescents from Brazil.

Methods

We retrospectively evaluated 55 patients with Takayasu's arteritis from 8 Pediatric Rheumatology centers in Brazil. All patients fulfilled the specific classification criteria for TA that have most recently been developed for the pediatric age group. Clinical, data were collected at two time points (at disease onset and at the last available visit).

Results

The majority of patients were girls (84%), with mean age at disease onset of 9.3 years, mean age at diagnosis of 10.6 years and a mean follow-up period of 5.6 years. The most common initial clinical manifestations included headache (56.3%), malaise (56.3%), weight loss (45.4%), dyspnea (41.8%) and limb pain (41.8%). The most common cardiovascular finding was hypertension (85.4%). The most frequent laboratory abnormality was an increased erythrocyte sedimentation rate (70.9%). The Mantoux test was positive in 38% of patients. The most

common alteration was narrowing or stenosis. The most involved vessels were abdominal aorta (71.8%) followed by left (56.2%) and right renal artery (53.1%). At the last visit the most common clinical manifestations were: headache (22%), inferior limb claudication (14%) and arthralgias (12%); Cardiovascular finding included the absence or decrease of the inferior limb pulses (52%). The treatment included imunossupressors other than corticosteroids in 83.6% and anti-TNF antagonists in 7.2% of patients. Seventeen (30.9%) patients underwent surgical intervention and 5 (9%) died of complication from treatment or from disease related complications.

Conclusion

In this large survey of children and adolescents with Takayasu's arteritis from Brazil a delay in the diagnosis and a high rate of Mantoux test positivity were observed. A death rate similar to the described in literature was found.

Author details

¹Universidade Federal de São Paulo/ Escola Paulista de Medicina, Brazil.
²Universidade Federal do Rio de Janeiro, Brazil.
³Universidade de São Paulo, Brazil.
⁴Universidade do Estado do Rio de Janeiro, Brazil.
⁵Universidade Federal de Pernambuco, Brazil.
⁶Universidade Estadual Paulista, Brazil.
⁷Universidade Federal da Bahia, Brazil.
⁸Santa Casa de Misericórdia de São Paulo, Brazil.

Published: 14 September 2011

doi:10.1186/1546-0096-9-S1-P89

Cite this article as: Terreri et al.: Brazilian multicentre study of Takayasu's arteritis in children and adolescents – preliminary results of a clinical, imaging and therapeutic study. *Pediatric Rheumatology* 2011 9 (Suppl 1):P89.

¹Universidade Federal de São Paulo/ Escola Paulista de Medicina, Brazil Full list of author information is available at the end of the article



^{*} Correspondence: teterreri@terra.com.br