



POSTER PRESENTATION

Open Access

Is it worth allowing the presence of morning stiffness in the definition of inactive disease in juvenile idiopathic arthritis?

Maddalena Allegra¹, Maria Chiara Gallo¹, Sara Verazza¹, Serena Calandra¹, Sara Dalprà¹, Federica Mongelli¹, Alberto Martini^{1,2}, Angelo Ravelli^{2,3}, Alessandro Consolaro^{1*}

From 21st European Pediatric Rheumatology (PReS) Congress
Belgrade, Serbia. 17-21 September 2014

Introduction

Morning stiffness is a major symptom of juvenile idiopathic arthritis (JIA) and it is usually associated with active disease. However, it is common view that children with disease quiescence may have some degrees of residual morning stiffness. The 2004 preliminary criteria for inactive disease (ID) in JIA did not include the assessment of morning stiffness, whereas the 2011 revision of the criteria has allowed the presence of morning stiffness lasting ≤ 15 minutes. However, it is still unknown whether the disease status of children with ID who have or do not have morning stiffness is comparable.

Objectives

To compare the disease status of children with JIA who meet the 2011 revised criteria for ID and have or do not have a morning stiffness lasting ≤ 15 minutes.

Methods

A database at the study center including 785 patients who had undergone a total of 2957 visits, which included a parent report of the presence and duration of morning stiffness, was analyzed to identify all visits in which patients met the criteria for ID. In each visit, the duration of morning stiffness was categorized as follows:

Table 1

	Patients meeting 2004 ID criteria			p-value
	Patients meeting 2011 ID criteria			
	No MS N = 390	MS ≤ 15 min N = 41	MS > 15 min N = 29	
Median (IQR) disease duration	3.8 (1.8; 7.3)	2.8 (1.4; 6)	5.7 (2.6; 7.9)	0.30
Functional ability (JAFA score) >0, N (%)	64 (16.4)	18 (43.9)	23 (79.3)	< 0.001
Physical health (PRQL PhS) >0, N (%)	173 (44.4)	32 (78)	28 (96.6)	< 0.001
Psychosocial health (PRQL PhS) >0, N (%)	188 (48.2)	29 (70.7)	24 (82.8)	< 0.001
VAS well-being >0, N (%)	155 (40.4)	32 (82.1)	28 (100)	< 0.001
VAS pain >0, N (%)	102 (26.8)	28 (68.3)	21 (87.5)	< 0.001
Acceptable symptom state, N (%)	366 (95.6)	33 (80.5)	16 (57.1)	< 0.001

MS: morning stiffness; IQR: interquartile range

¹Pediatria II, Istituto Giannina Gaslini, Genova, Italy
Full list of author information is available at the end of the article

≤15 min, 15-30 min, 30-60 min, 1-2 hr, >2 hr. Clinical assessments included demographic features, and parent-reported outcomes. In case a patient met the ID criteria in more than 1 visit, only the first visit was retained.

Results

A total of 460 visits in which the patient met the criteria for ID were evaluated. Absence of morning stiffness was reported in 390 (84.8%) visits, whereas in 70 visits (15.2%) there was morning stiffness. Among the visits with morning stiffness, in 41 (8.9%) duration was ≤15 min, and in 29 (6.3%) duration was >15 min. Table 1 shows the comparison of disease duration and parent-reported outcomes between patients with absence or presence of morning stiffness.

Conclusion

Among patients who met the 2011 criteria for ID, those with morning stiffness ≤15 min had worse parent-reported outcomes than those without morning stiffness. This finding suggests that parents may not perceive their child's disease state as true remission when lower degrees of morning stiffness are present. Notably, a sizeable proportion (6.3%) of children meeting the 2004 ID criteria had morning stiffness lasting > 15 min. The change of the criterion "Duration of morning stiffness of ≤ 15 minutes" to "Absence of morning stiffness" in the definition for ID should be considered.

Disclosure of interest

None declared

Authors' details

¹Pediatria II, Istituto Giannina Gaslini, Genova, Italy. ²Pediatrics, Università degli Studi di Genova, Italy. ³Pediatrics, Istituto Giannina Gaslini, Genova, Italy.

Published: 17 September 2014

doi:10.1186/1546-0096-12-S1-P174

Cite this article as: Allegra *et al.*: Is it worth allowing the presence of morning stiffness in the definition of inactive disease in juvenile idiopathic arthritis? *Pediatric Rheumatology* 2014 **12**(Suppl 1):P174.

**Submit your next manuscript to BioMed Central
and take full advantage of:**

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

