

Session S29 - Pediatric Multiple Sclerosis

○ Itinerary

S29.002 - Pediatric Radiologically Isolated Syndrome: Clinical and Radiological Outcomes

 April 19, 2016, 6:45 - 7:00 AM


Authors

Naila Makhani,¹Christine Lebrun Frenay,²Aksel Siva,³Clarisse Carra-Dalliere,⁴Jerome De Seze,⁵Francoise Durand Dubief,⁶Orhun Kantarci,⁷Megan Langille,⁸Jean Pelletier,⁹Juan Ignacio Rojas,¹⁰Teri Schreiner,¹¹Mar Tintore,¹²Ugur Uygunoglu,³Darin Okuda,¹³Daniel Pelletier,⁸Observatoire Francophone de la Sclérose en Plaques (OFSEP), Société Francophone de la Sclérose en Plaques (SFSEP), Radiologically Isolated Syndrome Consortium (RISC)
¹New Haven, CT, USA, ²Nice, France, ³Istanbul, Turkey, ⁴Montpellier, France, ⁵Strasbourg, France, ⁶Lyon, France, ⁷Rochester, MN, USA, ⁸Los Angeles, CA, USA, ⁹Marseille, Cedex 5, France, ¹⁰Buenos Aires, Argentina, ¹¹Aurora, CO, USA, ¹²Barcelona, Spain, ¹³Dallas, TX, USA

Disclosures

N. Makhani: None. **C. Lebrun Frenay:** ; Dr Lebrun received personal compensation from Bayer schering, biogen idec, merck serono, Genazyme, Almirall, Allergan, Novartis and sanofi for speaking in symposia. **A. Siva:** ; Dr Siva also reports receiving honoraria or consultation fees from Biogen Idec & Gen Pharma of Turkey, Novartis and Teva Turkey. He had received travel and registration coverage for attending, several national or international congresses or symposia, from Merck Serono, Biogen Idec/Gen Pharma of Turkey, Novartis and Teva.. **C. Carra-Dalliere:** None. **J. De Seze:** None. **F. Durand Dubief:** None. **O. Kantarci:** None. **M. Langille:** None. **J. Pelletier:** None. **J. Rojas:** None. **T. Schreiner:** ; Dr. Schreiner will receive honorarium from Biogen and MSAA for consulting services.. **M. Tintore:** None. **U. Uygunoglu:** ; Dr. Uygunoglu received personal compensation from Merck Serono and Biogen Idec. **D. Okuda:** ; Dr. Okuda received lecture fees from Acorda Therapeutics, Genzyme, and TEVA Neuroscience, consulting and advisory board fees from Genzyme, Novartis, and TEVA Neuroscience., Dr. Okuda received research support from Biogen for a multi-center clinical trial. **D. Pelletier:** ; Dr. Pelletier received consulting fees from CNS Imaging Consultant, LLC, Dr. Pelletier on behalf of Yale receives research grant funding from Biogen-Idec, Genzyme, and Hoffman-LaRoche..

Abstract

Objective:

To describe the natural history of children displaying incidental imaging findings suggestive of MS.

Background:

The term radiologically isolated syndrome (RIS) has been applied to asymptomatic individuals with incidental MR imaging findings highly suggestive of MS. RIS in adults carries a 34% risk of a first clinical event within 5 years. Such risk in individuals under 18 years of age (children) is unknown.

Methods:

Children demonstrating DIS on MRI were retrospectively analyzed from databases in 13 sites from 5 different countries using standardized variables. We used Fisher's exact test (categorical variables) or Mann-Whitney U test (continuous variables) for univariate analyses.

Results:

We identified 26 children with RIS (18F, 8M) detected at a median age of 15.3 years (range 8.9-17.8). Median follow-up time after RIS diagnosis was 4.31 years (range 0.6-17.4). The most common reason for initial imaging was headache (12/26, 46.2%). A first clinical event or MS diagnosis occurred in 11/26 (42.3%) children after a median of 2.6 years of follow-up (range 0.3-17.1). Radiological evolution occurred in 15/26 (57.7%) children after a median of 1.6 years (range 0.1-7.0). There were no differences in any of the baseline demographic variables (reason for imaging, age, sex, presence of CSF oligoclonal bands, presence of contrast-enhancing lesions, presence of spinal cord lesions) between children with and without either clinical or radiological evolution, but children with new MRI activity had lower serum 25-hydroxyvitamin D levels as compared to those with stable MRI scans (mean 17.6±7.2 vs. 47.7±12.7 ng/mL, p=0.012).

Conclusions:

We describe the first international cohort of children with RIS. RIS in childhood is associated with a high risk of a first clinical event, MS diagnosis and subsequent MRI lesions. Our preliminary findings suggest that low serum vitamin D level may increase risk of future MRI activity. Further study is warranted.