Is there any relationship between extra-pulmonary manifestations of *Mycoplasma pneumoniae* infection and atopy/respiratory allergy in children?

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Abstract

Mycoplasma pneumoniae is a common cause of respiratory infections in children, but sometimes extra-pulmonary diseases can be observed. The immunological mechanisms involved in these extra-respiratory complications are unknown. Here, we report a small case series of *Mycoplasma*-related diseases including 5 children who developed: i) aseptic meningitis; ii) urticarial rash and pericardial effusion; iii) pleural effusion with severe eosinophilia; iv) Stevens-Johnson syndrome; v) multiform erythema. Interestingly, all children were moderately to highly atopic, as a common immunologic feature.

Introduction

Mycoplasma pneumoniae (MP) is a common cause of respiratory infections in children. This infectious agent can cause bronchitis and pneumonia. However, there is evidence that MP infection is often limited to the upper airways and patients can have mild respiratory manifestations, which are often misdiagnosed as viral illnesses.¹ Indeed, the main clinical evidence of MP infection is sometimes only extra-pulmonary symptoms involving skin and joints in many cases. However, the involvement of other organs (e.g. central nervous system, heart, gastrointestinal system, eye, etc.) is well described.² Here, we report a small case series including 5 cases of extra-pulmonary diseases, which were shown to be associated to an ongoing and/or recent MP infection, after an appropriate diagnostic work-up.

Materials and Methods

In the period between October 2014 and May 2015, among all children being admitted to the pediatric ward of Melegnano Hospital (Milan,

Italy), we studied 5 consecutive patients diagnosed as having MP-related extra-respiratory diseases. Such a diagnosis was made upon the exclusion of other concomitant diseases and infections in children showing only a significant titer of specific IgM against MP (>10 AU/mL), in presence of a negative IgG specific serology (<10 AU/mL). Serum specific IgG and IgM against MP were measured by a chemiluminescent immunoassay (CLIA) performed through a Liason® XL analyzer [DiaSorin S.p.a, Saluggia (VC), Italy].³ As shown in Table 1, all children received a first-level immunologic work-up, including blood cell count, serum immunoglobulin measurement, complement screening (C3 and C4) and lymphocyte phenotype.

Case Report #1

Patient 1 was a 7-year-old boy who presented to our Emergency Pediatric Department because of high fever, persistent headache and vomiting. As neck rigor was observed, a lumbar puncture was performed: the cerebrospinal fluid resulted to be sub-limpid, showing a significant pleocytosis, but both bacterial cultures and viral genome research (with particular regard to herpesviridae) were negative. However, as the parents reported some episodes of upper airway inflammation during the previous weeks, thus serology for MP was evaluated, demonstrating a high titer of specific IgM (>27 AU/mL). Whereas the initial therapy with ceftriaxone and acyclovir did not lead to any improvement, actually the administration of intravenous clarithromycin rapidly improved patient's clinical condition.

Case Report #2

Patient 2 was a 13-year-old girl having urticaria (which responded to anti-histamines and steroids only partially), diffuse arthralgia and dactylitis-like transient fingers edema. Therefore, the patient received several diagnostic investigations including heart ultrasound, which showed a pericardial effusion greater than 0.5 cm. Eye examination did not show uveitis and autoantibody panel was completely negative; however, serum IgM against MP was present (20 AU/mL). After therapy with oral clarithromycin was started, a ready clinical improvement occurred.

Case Report #3

Patient 3 was a 10-year-old boy who was evaluated at the Pediatric Emergency Department because of acute onset of upper abdominal pain, vomiting and mild fever. Chest X-ray showed an

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interstitial pattern of lung inflammation associated with a very mild pleural effusion. Unexpectedly, the most relevant finding of the blood count was leukocytosis (WBC: 11750/mm³) with severe and absolute eosinophilia (5750/mm³). The clinical exam revealed no clinical signs of proliferative disease and blood test presented normal values of cellular turnover parameters (LDH: 456 U/mL, b2-microglobulin: 0.9 mg/mL) and of liver and kidney function. A wide diagnostic work-up for infections and autoimmunity was performed: particularly, no autoantibody was showed and stools examination did not showed intestinal viral, bacterial or parasite infections. The only significant serology was related to MP, through a very high specific IgM titer (>27 AU/mL). The treatment with clarithromycin led to a complete resolution of both symptoms and laboratory abnormalities: indeed, eosinophilia gradually decreased to normal values (500/mm³).

Case Report #4

Patients 4 was a 10-year-old boy who was admitted to the pediatric ward in order to receive intravenous nutritional and hydration support, because of the onset of severe ulcerative mucosal lesions involving lips and oral cavity, in addition to the anal area. Such a clinical picture was diagnosed as being Stevens-Johnson syndrome with prevalent mucosal involvement. After an

extensive infectious work-up, there was no other significant finding than a high titer of anti-MP IgM (15 UA/mL). The clinical condition gradually improved during the therapy with clarithromycin.

Case Report #5

Patient 5 was a 6-year-old boy complaining of the acute and sudden onset of a mildly itchy, but diffuse skin rash. That was diagnosed as being multiform erythema. Whereas viral serology, including Epstein-Barr virus and cytomegalovirus, were negative, a significant increase of anti-MP IgM (21 AU/mL) was detected. Such an acute MP infection was treated through a 2-week course of clarithromycin, leading to the resolution of the clinical picture.

All patients were Caucasian and reported a recent history of airways inflammation during the previous 2-4 weeks: however, the personal and family history resulted negative for genetic diseases and immunodeficiency. An immunologic work-up was performed for all these patients and, interestingly, we observed a significant increase of serum IgE levels for age in all of them, indicating a condition of atopy. Moreover, three of them showed an overt respiratory allergy (patient 1: house dust mite and grass pollen; patient 2: house dust mite and *Alternaria* spp.; patient 5: grass pollen).

Discussion and Conclusions

Several studies have linked respiratory MP infections to asthma exacerbations and there are some evidences that MP might elicit a Th2 immune response in the bronchial system.^{4,5} As regards MP-related extra-pulmonary complications, although those are supposed to be immune-mediated/autoimmune phenomena, the immunological mechanism –or mechanisms – is unknown.⁶ Here, through the description of our case series, a potential association between this category of reactive diseases caused by MP and atopy might be highlighted.

A tendency to IgE production during the acute phase of MP respiratory infections has been previously described and specific IgE against this microorganism could be detected.^{7,8} Stelmach *et al.* observed several antibody and cellular immune parameters in children affected with respiratory infections, but their data did not show a significant difference in total serum IgE levels between MP positive and negative patients. Actually, a trend toward a mild and gradual increase of IgE levels was described during the first year after mycoplasma infection.⁹ However, in our case series, we could observe a much greater IgE elevation in these children affected with MP-related extra-pulmonary diseases compared to the data displayed in the aforementioned study.

All these observations might raise the speculation that the predisposition to produce IgE in

some individuals, namely atopy, and/or the coexistence of respiratory allergy might predispose to (or might be involved in the pathogenesis of) extra-respiratory diseases related to MP acute infections. Of course, although the method used to measure MP serology showed a good reliability,³ serology diagnosis of MP infection should be cautiously considered and comparative studies will be required to support this preliminary observation.

References

1. Principi N, Esposito S. Emerging role of *Mycoplasma pneumoniae* and *Chlamydia pneumoniae* in paediatric respiratory tract infections. *Lancet Infect Dis* 2001;1:334-44.
2. Timitilli A, Di Rocco M, Nattero G, et al. Unusual manifestations of infections due to *Mycoplasma pneumoniae* in children. *Infez Med* 2004;2:113-7.
3. Capuano F, Grassi B, Pallavicini L, et al. New Liaison® automated immunoassays for the detection of mycoplasma pneumoniae IgG and IgM antibodies in human serum/plasma specimens. At 21st ECCMID /27th ICC 2011, Milano (Italy), 7-10 May 2011.
4. Smith-Norowitz TA, Silverberg JI, Kusonruska M, et al. Asthmatic children have increase specific anti-*Mycoplasma pneumoniae* IgM but not IgG or IgE-values independent of history of respiratory tract infection. *Pediatr Infect Dis J* 2013;32:599-603.
5. Ye Q, XU XJ, Shao WX, et al. *Mycoplasma pneumoniae* infection in children is a risk factor for developing allergic diseases. *World Sci J* 2014;986527.
6. Smith LG. *Mycoplasma pneumoniae* and its complications. *Infect Dis Clin N Am* 2010;24:57-60.
7. Nagayama Y, Sakurai N, Kojima S, Funabashi S. Total and specific IgE responses in the acute and recovery phases of respiratory infections in children. *J Asthma* 1987;24:159-66.
8. Koh YY, Park Y, Lee HJ, Kim CK. Bronchoalveolar lavage fluid from patients with micoplasma pneumonia: implication of tendency toward increate immunoglobulin E production. *Pediatrics* 2001;107:39-46.
9. Stelmach I, Podsiadlowski-Borzecka M, Grzelewski T et al. Humoral and cellular immunity in children with *Mycoplasma pneumoniae* infection: a 1 year prospective study. *Clin Diag Lab Immunol* 2005;12:1246-50.

Table 1. Immunologic parameters in patients with extra-pulmonary disease related to *Mycoplasma pneumoniae*.

Parameter	Case #1	Case #2	Case #3	Case #4	Case #5
Blood count, μ L					
Leucocytes	6100	11500	11750	9950	8200
Neutrophils	1700	5300	1700	3300	6100
Lymphocytes	3300	5300	3600	4900	1500
Monocytes	500	600	350	800	500
Eosinophils	600	300	5700	800	<100
Basophils	<100	<100	250	150	<100
Immunoglobulins					
IgA, mg/dL	143	162	155	181	93
IgG, mg/dL	810	864	1011	1030	688
IgM, mg/dL	87	104	82	108	56
IgE, UA/dL	848	168	900	431	1028
Complement, mg/dL					
C3	110	147	172	153	126
C4	31	27	39	23.8	24.5
Lymphocyte count, %					
CD3	68	55	66	62	n.a
CD4	37	36	36	31	n.a
CD8	17	23	23	21	n.a
CD19	15	25	21	17	n.a
CD16/56	10	16	10	14	n.a