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LETTER TO THE EDITOR

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ZEB2, a new candidate gene for asplenia

Linda Pons¹, Sophie Dupuis-Girod^{1*}, Marie-Pierre Cordier¹, Patrick Ederly^{1,2} and Massimiliano Rossi^{1,2}

Abstract

Primary asplenia is a rare condition with poorly known etiology. Mowat-Wilson syndrome (MWS) is characterized by typical facial dysmorphisms, intellectual disability, microcephaly, epilepsy and the possible presence of internal organ malformations. It is caused by heterozygous mutations or deletions in the *ZEB2* gene. Nearly 180 patients have been reported to date, but only one with asplenia. We report here spleen hypo/aplasia in 4 out of 6 MWS patients, with severe infectious complications for 3 of them. Our report shows that spleen hypo/aplasia is part of the MWS phenotype and makes *ZEB2* a possible candidate gene for primary asplenia.

Keywords: Mowat-Wilson syndrome, *ZEB2*, Asplenia

Letters to the editor

Primary congenital asplenia is a rare condition with poorly known genetic bases. It can be part of multiple congenital abnormalities syndromes or it can be isolated, which is extremely rare with about 70 patients reported to date [1]. Recently Bolze et al. identified heterozygous mutations in the *RPSA* gene in more than half the patients studied with isolated congenital asplenia [2]. Mowat-Wilson syndrome (MWS, OMIM #235730) is characterized by typical facial features (large medially sparse eyebrows, hypertelorism, deep set eyes, uplifted ear lobes with central depression, saddle nose and a pointed chin), intellectual disability, microcephaly, epilepsy and congenital malformations including Hirschsprung disease, genito-urinary abnormalities, cardiac defects, corpus callosum agenesis and ocular anomalies [3]. This syndrome was first described in 1998 [4] and is caused by heterozygous mutations or deletions in the Zinc finger E-box-binding homeobox 2 gene (*ZEB2*) [5,6]. Nearly 180 patients have been reported to date. Asplenia was reported in only one case [7]. Interestingly, it has been shown that *ZEB2* has a diffuse expression in several mouse and human organs, including the spleen [8,9].

We report here spleen hypo/aplasia in 4 out of 6 unrelated MWS patients referred to our genetic department, with severe infectious complications for 3 of them.

The first patient is a female born to unrelated parents, presenting with a typical facies, microcephaly, postnatal short stature, developmental delay, corpus callosum agenesis, ventricular septal defect, strabismus and left dimmed vision. A *de novo* c.2083C > T heterozygous mutation of the *ZEB2* gene was identified, thus confirming the diagnosis. At the age of 8 months, she suffered from purpura fulminans related to a severe *Streptococcus Pneumoniae* infection (serotype 12 F) with severe necrosis sequelae including the loss of 5 toes and the right heel requiring a skin graft. Asplenia was diagnosed on ultrasound scan (USS).

The second patient is a female born to unrelated parents. Typical dysmorphic features, microcephaly, developmental delay, epilepsy, corpus callosum agenesis, ventricular septal defect and patent ductus arteriosus, club foot and strabismus were consistent with the diagnosis of MWS. A *de novo* c.600_640dup heterozygous mutation of the *ZEB2* gene was identified. At the age of 1 year, she developed meningitis related to *Streptococcus Pneumoniae* infection (serotype 17 F on cerebrospinal fluid culture) complicated by moderate intracranial hypertension. USS revealed asplenia.

The third patient is a female born to unrelated parents. She had typical facial dysmorphisms, severe intellectual disability, microcephaly, epilepsy, postnatal short stature, ventricular septal defect, vesico-ureteric reflux, club foot and unilateral choanal atresia. Molecular analysis of the *ZEB2* gene showed a *de novo* c.1762G > T heterozygous mutation. She had two pneumococcal septicemias at the ages of 2 and 3 years. Abdominal USS revealed severe splenic hypoplasia (main axis: 34 mm, average for weight: 80 mm, range 78–87 mm). Howell-Jolly bodies were absent.

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The fourth patient is a female born to non-related parents, presenting with a typical facies, microcephaly, post-natal short stature, developmental delay, epilepsy, atrial septal defect and patent ductus arteriosus, strabismus. A *de novo* c.2761C > T heterozygous mutation of the *ZEB2* gene was identified confirming the diagnosis of MWS. She had no severe infections. USS showed moderate splenic hypoplasia (main axis: 46 mm, average for weight: 78 mm, range 76–78 mm).

The 2 other MWS patients followed in our genetic department had a normal spleen on USS and did not have severe infections.

On one hand, our report shows that spleen hypo/aplasia is a part of the phenotype of MWS. USS should be systematically performed on MWS patients in order to rule out spleen hypo/aplasia because of potential complications' severity. Prevention of severe infections in cases of asplenia or severe hypoplasia, effectively requires appropriate antibiotic prophylaxis and vaccination that can restore the pool of memory B cells [10].

As our cohort only includes 6 MWS subjects, we think that other studies should be done to confirm if asplenia/ spleen hypoplasia is a new feature of this multisystem disorder.

Indeed, the case of asplenia previously reported has a *de novo* c.696C > G heterozygous mutation of *ZEB2*, so a different one from our patients. Furthermore, no spleen hypo/aplasia has been described in previously reported patients with *de novo* c.2083C > T or c.2761C > T heterozygous mutations of *ZEB2* found in patients 1 and 4 [11].

On the other hand, this report emphasizes the potential role of *ZEB2* in human spleen development. This association makes this gene a possible new candidate for isolated congenital asplenia. Indeed all cases cannot be attributed to *RPSA* gene, suggesting a possible genetic heterogeneity. Although mutations of *ZEB2* are generally associated with much more complex conditions, it cannot be excluded that some missense mutations could be responsible for apparently isolated asplenia [2].

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References

1. Mahlaoui N, Minard-Colin V, Picard C, Bolze A, Ku CL, Tournilhac O, et al: Isolated congenital asplenia: a French nationwide retrospective survey of 20 cases. *J Pediatr* 2011, **158**(1):142–148. 8 e1.

2. Bolze A, Mahlaoui N, Byun M, Turner B, Trede N, Ellis SR, et al: Ribosomal protein SA haploinsufficiency in humans with isolated congenital asplenia. *Science* 2013, **340**(6135):976–978.
3. Garavelli L, Mainardi PC: Mowat-Wilson syndrome. *Orphanet J Rare Dis* 2007, **2**:42.
4. Mowat DR, Croaker GD, Cass DT, Kerr BA, Chaitow J, Ades LC, et al: Hirschsprung disease, microcephaly, mental retardation, and characteristic facial features: delineation of a new syndrome and identification of a locus at chromosome 2q22-q23. *J Med Genet* 1998, **35**(8):617–623.
5. Garavelli L, Donadio A, Zanacca C, Banchini G, Della Giustina E, Bertani G, et al: Hirschsprung disease, mental retardation, characteristic facial features, and mutation in the gene ZFH1B (SIP1): confirmation of the Mowat-Wilson syndrome. *Am J Med Genet Part A* 2003, **116A**(4):385–388.
6. Verstappen G, van Grunsven LA, Michiels C, Van de Putte T, Souopgui J, Van Damme J, et al: Atypical Mowat-Wilson patient confirms the importance of the novel association between ZFH1B/SIP1 and NuRD corepressor complex. *Hum Mol Genet* 2008, **17**(8):1175–1183.
7. Zweier C, Thiel CT, Dufke A, Crow YJ, Meinecke P, Suri M, et al: Clinical and mutational spectrum of Mowat-Wilson syndrome. *Eur J Med Genet* 2005, **48**(2):97–111.
8. Bassez G, Camand OJ, Cacheux V, Kobetz A, Dastot-Le Moal F, Marchant D, et al: Pleiotropic and diverse expression of ZFH1B gene transcripts during mouse and human development supports the various clinical manifestations of the "Mowat-Wilson" syndrome. *Neurobiol Dis* 2004, **15**(2):240–250.
9. Mowat DR, Wilson MJ, Goossens M: Mowat-Wilson syndrome. *J Med Genet* 2003, **40**(5):305–310.
10. Rosado MM, Gesualdo F, Marcellini V, Di Sabatino A, Corazza GR, Smacchia MP, et al: Preserved antibody levels and loss of memory B cells against pneumococcus and tetanus after splenectomy: tailoring better vaccination strategies. *Eur J Immunol* 2013, **43**(10):2659–2670.
11. Dastot-Le Moal F, Wilson M, Mowat D, Collot N, Niel F, Goossens M: ZFH1B mutations in patients with Mowat-Wilson syndrome. *Hum Mutat* 2007, **28**(4):313–321.

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