

A mutation in the STAT1 DNA-binding domain associated with hemophagocytic lymphohistocytosis

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ABSTRACT

Introduction: The transcription factor Signal Transducer and Activator of Transcription 1 (STAT1) is a key element in many of the signalling cascades involved in immune system function. Different mutations in STAT1 are associated with heterogeneous clinical phenotypes that range from early fatality due to overwhelming infection to limited involvement of the mucus membrane with recurrent *Candida* infections. Multiple genes related to immune function have been associated with the development of hemophagocytic lymphohistiocytosis (HLH), but the association between *STAT1* mutation and HLH has not been described in detail.

Methods: We report the genetic background of a patient with chronic mucocutaneous candidiasis (CMC) as well as an unusual clinical course.

Results: In this study we describe a patient with a mutation in the *STAT1* DNA-binding domain and a history of CMC who developed a refractory and fatal case of HLH despite having bone marrow transplantation.

Conclusion: We describe a patient with refractory and fatal HLH who was found to have a mutation in the DNAbinding domain of *STAT1*.

Statement of novelty: The association of chronic mucocutaneous candidiasis with HLH.

Introduction

The DNA-binding protein Signal Transducer and Activator of Transcription 1 (STAT1) is a member of a family of transcription factors that plays a major role in different and diverse functions of the cell. Following activation by interferon-gamma or interferon-alpha, STAT1 protein is phosphorylated and forms dimers. It then translocates into the nucleus, binds to DNA, and initiates its effector function (Darnell et al. 1994; Stark et al. 1998; Levy and Darnell 2002). Different genetic mutations in *STAT1* resulting in different clinical phenotypes have been described (Dupuis et al. 2001, 2003; Chapgier et al. 2006a, 2006b). Patients with complete loss-of-function of the STAT1 protein usually die in childhood from overwhelming viral and mycobacterial infections (Dupuis et al. 2003). Other patients, who have the autosomal dominant *STAT1* loss-of-function form, usually present with a predisposition to low pathogenic mycobacterial infections (Tsumura et al. 2012). Different mutations, mostly in the coiled–coiled domain, are believed to lead to gain of function of the

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STAT1 protein and cause an autosomal dominant form of chronic mucocutaneous candidiasis (CMC) (Liu et al. 2011; van de Veerdonk et al. 2011). Recently, a severe and fatal form of combined immunodeficiency was found to be associated with a declining immunity during childhood, caused by predominantly de-novo mutations in the *STAT1* DNA-binding domain (DBD) (Sharfe et al. 2014).

Hemophagocytic lymphohistiocytosis (HLH) is a rare, life-threatening inflammatory process that is characterized by uncontrolled activation of macrophages and histiocytes (Favara et al. 1997). Traditionally, HLH was divided into a genetic form (also known as the familial or primary type) and an acquired form (secondary HLH) (Writing Group of the Histiocyte Society 1987; Henter et al. 1991).

The genetic form usually presents in a young infant with parental consanguinity or a history of other affected family members. Patients in this group have a genetic defect in genes that are responsible for the cytotoxic function of T cells and NK cells. Most commonly, mutations in the gene *PFR1* that encodes for Perforin1, are detected in this group of patients; however, other genes, such as UNC13D and STX11 that encode for cytolytic granule trafficking and exocytosis may also be associated with the aberrant inflammation (Jordan et al. 2011). Although some of the primary immunodeficiency disorders that are associated with HLH have abnormal cytotoxic function of T cells and NK cells (for example Chédiak-Higashi syndrome and Griscelli syndrome type 2) others such as Wiskott-Aldrich and DiGeorge syndrome do not, and the mechanism leading to HLH in these conditions is not fully understood (Introne et al. 1999; Ménasché et al. 2000; Cesaro et al. 2003; Pasic et al. 2003).

In contrast to the genetic form of HLH, the acquired forms seem not to be associated with a known genetic abnormality or immunodeficiency syndrome. As in some of the genetic forms, the pathogenesis is not completely understood, and it seems that the disease appears in association with infection, malignancy, rheumatic conditions, metabolic disorders, or as a consequence of drug toxicity (Henter et al. 1991; Janka et al. 1998). Nevertheless, it is conceivable that modifier genes may predispose individuals to developing secondary HLH. The symptoms and signs of HLH are nonspecific; fever, hepatosplenomegaly, lymphadenopathy, cytopenia, morbiliformic rash, and in some cases involvement of the central nervous system (CNS) with seizures or encephalopathy (Henter et al. 2007).

Without treatment, the prognosis of HLH is poor, with less than 5% survival 1 year post diagnosis (Janka 1983). Even with appropriate treatment, the mortality rate can be as high as 45% (Henter et al. 2002). The main goal of the treatment is to suppress the exaggerated activation of the immune system with immunosuppressive agents such as dexamethasone, etoposide, and cyclosporine, and when needed and indicated a bone marrow transplantation (Henter et al. 2002).

We report here an occurrence of fatal HLH in a patient with CMC due to STAT1 deficiency.

Methods

Flow cytometry

Peripheral blood mononuclear cells were obtained by Ficoll–Hypaque density gradient centrifugation, and surface phenotypes were determined by flow cytometry on a Coulter EPICS V flow cytometer (Beckman Coulter, Brea, Calif.), with a single argon laser, which analyzes up to 3 colours simultaneously. Single colour and (or) isotype antibody controls were both used for multicolour staining.

Proliferation assay

Lymphocyte proliferative responses to mitogens (including phytohemagglutinin and anti-CD3 antibodies) and to a panel of recall antigens (including *Candida*, tetanus, *Herpes zoster*, and cytomegalovirus) were determined by thymidine incorporation. All assays were performed in triplicate and were compared with simultaneously stimulated random normal controls.

DNA sequencing

Genomic DNA was extracted from peripheral blood lymphocytes using the Geneaid Genomic DNA Mini Kit. Genomic DNA was amplified by polymerase chain reaction (PCR) with specific primers for exons 1–23 of the coding sequence of *STAT1* (NM_007315.3) (available upon request). Exons and their flanking intronic regions were sequenced with the GenomeLab Dye Terminator Cycle Sequencing Quick Start Kit (Beckman Coulter) and analyzed on a CEQ 8000 Genetic Analysis System (Beckman Coulter).

Results

Case presentation

The patient, a female of nonconsanguineous East Indian origin, presented at 5 months of age with recurrent episodes of skin infection and persistent diaper rash and oral thrush. Her past medical history was unremarkable and her family history was negative for immunodeficiency, malignancy, or autoimmune disorders. The laboratory findings were unremarkable with normal levels of leukocytes and immunoglobulins (Table 1). She responded to Nystatin but continued to have periodic episodes of oral thrush. At the age of 3 years she was diagnosed with hypothyroidism and began suffering repeated episodes of pneumonia as well as asthma. At the age of 9 years she was found to have oral thrush and herpetic vesicles on her chest without hepatosplenomegaly or lymphadenopathy. She responded well to treatment with oral Fluconazole and Acyclovir. She subsequently developed Coombs-positive hemolytic anemia.

Evaluation of the immune system

Evaluation of the immune system revealed normal numbers of circulating white blood cells and lymphocytes. Flow cytometry analysis showed a steady decline of CD3⁺

Table 1: Evaluation of immunity over time.

circulating lymphocytes over time. Predominantly CD4⁺ T cells diminished over time down to 440 cells/ μ L. In addition, the number of NK cells was also declining to 77 cells/ μ L at the age of 10 years. These changes are consistent with the recently documented decline in immunity in patients with *STAT1* mutations in the DBD (Sharfe et al. 2014).

The number of circulating B cells appears to have been normal with little decline. Serum immunoglobulin levels increased over time particularly for IgG and IgA (Table 1), and specific antibodies to vaccination appeared to be preserved. In-vitro response to mitogens and antigens were still preserved in this patient.

Genetic analysis

Genetic analysis of the *STAT1* gene (Figure 1) revealed a mutation in the DBD of the molecule. The mutation was a de novo mutation, as it was not present in the parents or in the brother of the patient.

Course of HLH

The patient presented at the age of 10 years in the emergency department with a high fever, cough, and pancytopenia. An abdominal ultrasound and abdominal

		Age in years		
	1.5	9	10	Reference values
White blood cells	9.1	7.8	6.2	4–10 × 10 ⁹ /L
Markers				
CD3	1742	1128	1061	800–3500 cell/µL
CD4	939	520	440	400–2100 cell/µL
CD8	634	456	504	200–1200 cell/µL
CD20/19	1968	486	837	200–600 cell/µL
CD56	160	160	77	70–1200 cell/µL
Immunoglobulins				
lgG	11	16.5	12.4	6.6–15.3 g/L
lgM	1	1.3	1.1	0.5–1.9 g/L
IgA	0.7	3	4.6	0.5–2.2 g/L
lgE	27			<60 IU/mL
Specific antibodies				
Tetanus	1.96 IU/mL	0.26 IU/mL	0.19 IU/mL	> 0.01
Mumps		Positive (44 RU/L)	Borderline	Positive
Measles		Positive (495 IU/L)	Positive (465 IU/L)	Positive
Rubella		Positive (199 IU/L)	Positive (200 IU/L)	Positive
Varicella		Negative	Positive	Positive
Iseohemmagglutinin		1:32		1:8



Figure 1: Electropherogram showing the wild type vs patient sequence with heterozygous mutation c.(1189A > G)];(=) in *STAT1* (NM_007315.3) resulting in amino acid substitution p.(Asn397Asp);(=)(A). Human *STAT1* showing the N terminal domain, coiled-coil domain, DNA-binding domain, linker domain, SH2 domain, tail segment domain, and transactivation domain, their boundaries and the location of the N397D mutation.

CT (taken later), showed findings suggestive of a microabscess in the kidneys and spleen. An extensive infection workup that included blood cultures, serology, and PCR testing for bacteria, mycobacteria, fungi, and different viruses (*Herpes* family, respiratory viruses, and parvovirus) identified Epstein–Barr Virus viremia with a viral load of 1127 copies/mL in the blood (but not in the plasma). In addition, *Herpes zoster* virus was isolated from a skin vesicle.

As the fever continued, an initial blood work was sent to assess for HLH. The diagnostic criteria for HLH, according to the HLH-2004 protocol by the Histiocyte Society, include a molecular diagnosis consistent with HLH or 5 out of 8 clinical and laboratory criteria as described in Table 2 (Henter et al. 2007).

Although the initial blood work didn't fully support the diagnosis of HLH, the repeated blood work that was done on the second week of her admission confirmed the diagnosis (Table 3).

The patient was treated with broad spectrum antibiotics, anti-fungal, and anti-viral medications. After HLH was diagnosed, she started treatment with dexamethasone but showed only partial response. A day following Table 2: Criteria for the diagnosis of HLH.

Clinical criteria					
Fever					
Splenomegaly					
Laboratory criteria					
Cytopenia (affecting >2 of 3 lineage in peripheral blood)					
Hypertriglyceridemia or hypofibrinogenemia					
Low or absent NK cell activity					
Hyperferritinemia >500 µgram/L					
Increased levels of sCD25 >2400 U/mL					
Histopathological criteria					
Hemophagocytosis in the bone marrow, spleen, or lymph nodes					
No evidence of malignancy					

initial steroid treatment she developed seizures and her brain MRI showed bilateral multifocal abnormal signals in the cortex and subcortical white matter associated with leptomeningeal enhancement, findings that are consistent but not specific to CNS involvement of HLH (Figure 2).

Despite receiving appropriate antibiotic treatment, she did not improve and continued to be febrile with increased laboratory markers of HLH. The treatment for HLH was escalated to include etoposide, cyclosporine, and intrathecal methotrexate similar to the HLH-2004 protocol recommendation. The patient had a temporary response to the treatment as fevers subsided and repeated head imaging showed resolution of the abnormal MRI findings (Figures 3 and 4). However, she failed to get into full remission as her laboratory markers persistently showed cytopenia, increased ferritin, and elevated sCD25 levels.

After 8 weeks of treatment and following the HLH-2004 protocol recommendation for severe cases of HLH, a bone marrow transplant (BMT) was performed using an HLA matched unrelated donor in the absence of a matched sibling. The patient received a full myeloablative conditioning that included Busulfan (3.6 mg/kg) on days -8 to -5, etoposide (30 mg/kg/dose) on day -4, and cyclophosphamide (60 mg/kg) on days -3 and -2. For graft versus host disease prophylaxis she received steroids, cyclosporine, and methotrexate. The patient fully engrafted on day +8 with 100% chimerism, but she spiked a fever and once again suffered seizures within a week of transplant. A repeat MRI of her head again showed multiple cortical and white matter lesions, similar to the lesions described in Figure 2. The laboratory markers for HLH, which are sCD25, sCD163, and ferritin peaked to 4063 U/mL, 8178 ng/mL, and 949 510 µg/L, respectively. She subsequently developed severe complications including gastrointestinal bleeding, pulmonary hemorrhage, toxic epidermal necrosis, and renal failure. In addition, alemtuzumab was added to the regimen with no response. The patient deteriorated gradually until her death from multi-organ failure.

Discussion

In the last few years, different mutations in the *STAT1* gene have been reported (Liu et al. 2011; van de

Marker	Initial workup	Workup at diagnosis	Workup post bone marrow transplantation	Reference values
White blood cells	0.9	1.5	5.7	4–10 × 10 ⁹ /L
Hemoglobin	90	82	97*	120–160 g/L
Platelets	78	74	144*	150–400 × 10 ⁹ /L
Ferritin	3156	41 950	949 510	30–177 μg/L
Triglycerides	3.1	4.2	16	0.4–1.3 mmol/L
Fibrinogen	1.5	1.8	3.5	1.9–4.3 g/L
sIL2 receptor (sCD25)	1770	3101	4063	340–1500 U/mL
Bone marrow aspiration	No hemophagocytosis present	Hemophagocytosis present	_	_
Perforin expression by flow cytometery	Normal	—	—	—
Perforin gene sequencing	Normal	_		_
NK cells degranulation assay	Normal	—	—	—

*Post blood/platelets transfusion.

Table 3: Markers of HLH over time.



Figure 2: Axial fluid attenuated inversion recovery (A), axial diffusion weighted image (B), and contrast-enhanced coronal T1 images (C) show an abnormal signal in the cortex and subjacent white matter bilaterally. Subtle diffusion restriction is seen within these abnormal areas. In addition, there is enhancement in the leptomeninges, which is more pronounced in the sulci associated with those areas of an abnormal signal.



Figure 3: Axial fluid attenuated inversion recovery (A), axial diffusion weighted image (B), and contrast enhanced coronal T1 images (C) on the follow-up MRI show that the previously noted signal changes improved significantly with residual abnormality in the left parietal and occipital parietal lobes. No definite diffusion restriction is noted. The post-gadolinium image shows that the previously noted leptomeningeal enhancement is no longer visualized.



Figure 4: Axial fluid attenuated inversion recovery (A), axial diffusion weighted image (B), and contrast enhanced coronal T1 images (C) on another follow-up MRI a week after the MRI shown in Figure 3. There is further improvement with complete resolution of the abnormal cerebral signal changes and the abnormal leptomeningeal enhancement. The ventricles and pericerebral spaces are prominent bilaterally and have mildly increased in size compared with the previous studies with no midline shift. These findings are in keeping with volume loss.

Veerdonk et al. 2011; Tsumura et al. 2012). Monoallelic mutations in the DBD of STAT1 seem to cause a severe and frequently fatal immunodeficiency (Sharfe et al. 2014). In this study we described a mutation in the DBD of the molecule. The fact that these mutations are de novo mutations, support the hypothesis that changes in the DBD of the STAT1 gene are lethal at a young age before the carrier has a chance to produce offspring. The patient we described here had clinical symptoms of CMC and declining immunity that specifically displayed a decline in CD4⁺ T cells and NK cells. The diminished number and function of NK cells were previously reported to be associated with HLH. In this case, it is also conceivable that the decline in NK cell numbers underscored at least in part the development of HLH, although the NK cell degranulation assay was normal.

CNS involvement is common in HLH patients, and symptoms can range from abnormal findings in the cerebrospinal fluid to altered mental status, seizures, and coma. Active HLH, infection, and posterior reversible encephalopathy syndrome (PRES) are in the differential diagnosis in HLH patients with CNS involvement (Haddad et al. 1997; Goo and Weon 2007; Horne et al. 2008; Lee et al. 2013).

The overall survival of children with HLH who were treated with the HLH-94 protocol is around 55% and two-thirds of those who survived underwent BMT. It appears that survival after BMT in HLH cases is fair, regardless of whether a HLA matched sibling or an unrelated match donor are used (Henter et al. 2007). Although there are some reports that support reduced intensity over full myeloablative conditioning (Cooper et al. 2006; Marsh et al. 2010), at present there is insufficient evidence to support the use of one protocol over another. Out of the few reports of BMT in patients with STAT1 mutation, only 2 patients have been reported with long-term survival after receiving a BMT from a match sibling donor (Deeg et al. 1986; Hoh et al. 1996), whereas 2 other patients died after the procedure (Aldave et al. 2013).

In conclusion, we have shown an association between a monoallelic *STAT1* mutation and a severe and refractory case of HLH. A thorough immunologic workup that includes *STAT1* genetic analysis is needed in severe and refractory cases of HLH to identify those who have this uncommon genetic abnormality. More studies are in order to define optimal treatment options for patients with *STAT1* mutation and HLH.

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