Systemic Vasculitis and High IgE Level in a Patient with LPS-Responsive Beige-Like Anchor (LRBA) Deficiency

Jing Wu 1,2, Wei-Fan Wang 1, Tong-Xin Chen 1,2,** and Ji Chen 3,*

1Department of Allergy and Immunology, Shanghai Children’s Medical Center, School of Medicine, Shanghai Jiao Tong University, Shanghai, China
2Division of Immunology, Institute of Pediatric Translational Medicine, Shanghai Children’s Medical Center, School of Medicine, Shanghai Jiao Tong University, Shanghai, China
3Department of Dermatology, Shanghai Children’s Medical Center, School of Medicine, Shanghai Jiao Tong University, Shanghai, China

**Corresponding author: Department of Allergy and Immunology, Shanghai Children’s Medical Center, School of Medicine, Shanghai Jiao Tong University, 1678 Dongfang Rd., 200127 Shanghai, China. Email: tongxinc@yahoo.com

*Corresponding author: Department of Dermatology, Shanghai Children’s Medical Center, School of Medicine, Shanghai Jiao Tong University, 1678 Dongfang Rd., 200127 Shanghai, China. Email: chenji@scmc.com.cn

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Abstract

**Background:** Mutations in LPS-responsive and beige-like anchor (LRBA) gene in patients were firstly described to associate with a syndrome of immune deficiency and autoimmunity in 2012. However, there was still no LRBA deficient patient reported in China.

**Objectives:** We present a Chinese patient with heterozygous LRBA gene mutations with his clinical, immunological and genetic features.

**Methods:** Patient’s clinical data was collected and analyzed. Laboratory results included lymphocyte subsets analysis and immunoglobulin quantification. Targeted gene capture followed by next-generation sequencing was used to identify the gene mutations, and flow cytometry assay was used to analyze B cell immunophenotyping of this patient.

**Results:** The patient mainly suffered from recurrent respiratory tract infections, EBV-associated lymphoproliferative disease and systemic vasculitis. Heterozygous LRBA gene mutations were identified in the patient, which were inherited from his parents, respectively. B cell immunophenotyping revealed that he had decreased total B cells, bone marrow progenitor B cell subsets (HSC, CLP, Pro-B and Pre-B cells), immature B cells, non-switched memory B cells, switched memory B cells and B1 cell subsets. Besides, he had extremely high level of IgE.

**Conclusions:** Herein, we firstly report a patient with heterozygous LRBA gene mutations in China. LRBA might play an important role in B cell development.

**Keywords:** LRBA Deficiency, Systemic Vasculitis, Hyper IgE Level, Next-Generation Sequencing, B Cell Immunophenotyping

1. Background

LPS-responsive beige-like anchor protein (LRBA) deficiency is a rare inherited common variable immunodeficiency (CVID) caused by loss-of-function mutations in LRBA gene. Affected individuals were usually reported to suffer from immune deficiency, lymphoproliferation and various organ-specific autoimmunity, which severely affected the patients’ physical health and imposes a heavy economic burden on the patients (1-5).

LRBA gene, which is located on chromosome 4q31.3, contains 57 exons and encodes a large protein containing 2851 amino acid residues. LRBA belongs to the BEACH-WD40 protein family composed of multiple functional domains, including Concanavalin A (ConA)-like lectin binding domain, vacuolar protein sorting-27, hepatocyte growth factor-regulated tyrosine kinase substrate domain and signal transducing adaptor molecule (VHS) domain, two PKA regulatory subunit (RII) binding motifs, WD-like (WDL), and Beige and Chediak-Higashi (BEACH) domains, plekstrin homology (PH)-like domain and five WD40 repeats (6). LRBA is broadly expressed in several human tissues and involved in intracellular vesicle trafficking and signal transduction. Besides, LRBA deficiency was reported to be associated with a diminished expression of CTLA4 on the surface of Treg cells (7).

LRBA gene was firstly identified as an CVID-causing gene in 2012. Three papers were published almost at the same time to categorize LRBA deficiency as a common variable immunodeficiency (CVID), with autoimmunity and inflammation (1, 8, 9). However, its clinical spectrum has been extended with the increasing publications worldwide in the recent years, which included recurrent in-
Infections, hypogammaglobulinemia, inflammatory bowel disease, enteropathy, splenomegaly, hepatomegaly, lymphadenopathy, growth retardation, atopic skin disease, EBV infections and so on (1-4, 10). Immunologic testing showed that a majority of LRBA deficient patients had reduced total B cells and B cell subsets, as well as Treg cells (3, 10, 11). However, the broad phenotypic spectrum made the clinical diagnosis of LRBA deficiency difficult, which made genetic analysis to distinguish it from other diseases necessary.

2. Objectives

Herein, we describe a Chinese patient with compound heterozygous mutations in LRBA gene, who had decreased B cells, but extremely high level of IgE, mainly suffering from recurrent respiratory tract infections, systemic vasculitis and EBV infection. To our knowledge, this is the first Chinese LRBA deficient patient reported.

3. Methods

3.1. Patient

The affected patient with LRBA deficiency was enrolled in Shanghai Children’s Medical Center, China. Healthy control children were age matched. All study participants were recruited with written informed consent approved by the Ethics Committee of Shanghai Children’s Medical Center.

3.2. Targeted Gene Capture and Next-Generation Sequencing

DNA library was prepared according to illumine standard protocols, then captured with an immunodeficiency related gene panel, which included all PID-causing genes published in the International Union of Immunological Societies Expert Committee for Primary Immunodeficiency on 2015, by using biotinylated oligo-probes. The capture experiment was conducted according to manufacturer’s protocol (MyGenostics, Beijing, China) as previously described (12). In brief, 1 µg DNA library was mixed with Buffer BL and probes, incubated at 95°C for 7 min and 65°C for 2 min. Then, 65°C Buffer HY was added to the mixture, and incubated at 65°C for 22 hours for hybridization. Then MyOne beads (Life Technology) was used to bind the biotinylated oligo-probes. The bound DNA was eluted and amplified through PCR, and then purified using SPRI beads (Beckman Coulter). Finally, Illumina HiSeq 2000 sequencing machine was used to sequence the libraries.

3.3. Bioinformatics Analysis

High-quality reads were retrieved through eliminating low quality reads and adaptor sequences, then aligned with the human reference genome (hg19) using SOAPaligner program. The SNPs was firstly identified using the SOAPsnp program after PCR duplicates were removed by the Picard software. Then, the reads were realigned to the reference genome to detect insertions or deletions by using the GATK program. The identified SNPs and InDels were annotated by using Exome-assistant program. MagicViewer was then used to view the short read alignment and validate the candidate SNPs and InDels.

3.4. PCR and Sanger Sequencing

Genomic DNA was isolated from peripheral blood by using RelaxGene blood DNA isolation kit (Tiangen Biotech Co. Ltd.) according to the manufacturer’s instructions. PCR Products sent for Sanger sequencing. Sequence analysis software (Gene Runner) was used to identify gene mutations.

3.5. B Cell Immunophenotyping

B cell subsets were classified by flow cytometry according to the expression of surface markers as follows: hematopoietic stem cell (HSC): CD34+CD10-CD19-IgM--; common lymphoid progenitor (CLP); CD34+CD10+CD19-IgM--; Pro-B cells: CD34+CD10+CD19+IgM--; Pre-B cells: CD34-CD10+CD19+IgM--; immature B cells: CD19+CD10+CD27CD38++; memory B cells: CD19+CD27- naïve B cells: CD19+CD27-IgD--; non-switched memory B cells: CD19+CD27+IgD--; switched memory B cells : CD19+CD27+IgD--; double negative B cells: CD19+CD27- IgD--; plasma cells: CD19+CD10-CD27++; naïve B: CD20+CD27CD43--; memory B: CD20+CD27+CD43--; B: CD20+CD27+CD43+.

4. Results

4.1. Clinical Presentations

In this study, the male patient, who was born in 2005, was the first child from non-consanguineous parents and there was no history of similar disorders in their families. At 2 years of age, he got fever without obvious cause (maximum temperature, 39.6°C), paroxysmal and non-spasmodic cough and wheeze, and then was hospitalized with asthmatic suffocating pneumonia in a local hospital. After two months of treatment, His cough and wheeze was relieved, but he still had fever. In addition, enlarged cervical lymph nodes were observed in the patient.

The patient went to Beijing children’s hospital to seek treatment when 3 years old. At the time of admission,
general physical examination showed that he had congested throat and enlarged tonsils (1 degree). Some enlarged lymph nodes were also observed in his neck and groin bilaterally, which were soft, painful to touch and easy to move. Laboratory testing results showed that he had normal peripheral blood leukocytes, but raised serum C-reactive protein (CRP) levels and erythrocyte sedimentation rates (ESR), as well as positive Epstein-Barr virus (EBV) capsid antigen (VCA) IgG and EB VCA IgA, thus leading to a diagnosis of EBV-induced infectious mononucleosis complicated by bacterial infection. Anti-infective agents including ganciclovir and cefotiam were given to the patient to treat infections. During hospitalization, a progressive dilatation of coronary artery was observed in this patient by cardiac color ultrasound, while no obvious endangium lesions were detected by vascular ultrasound. Combined with his recurrent fever, relapse of raised ESR and CRP, positive lupus anticoagulant test, the patient was given the diagnosis of systemic vasculitis. Therefore, he was given a treatment with aspirin, dipyridamole and intravenous immunoglobulin. Despite these aggressive treatments, relapsing unexplained fever still occurred in this patient. Prednison was adminstrated to reduce his fever. After resuming normal body temperature, he was discharged from the hospital.

After discharge, he still had a recurrent fever. Vaccinia-like vesicles were also observed on his face and limbs, rupture of which led to ulceration (Figure 1). In November 2008, he came to our hospital to seek treatment. A clinical workup showed his body temperature was 39°C. Enlarged lymph nodes were observed in his ears, jaw, neck and right groin. Besides, he was reported to have swollen lip with obvious bleeding point and hepatosplenomegaly (at 1 cm under the ribs). Immunological tests showed that his serum IgG, IgA and IgM level were normal, while his serum IgE was up to 29400 IU/mL. Biopsies of skin vesicles showed allergic inflammation in the cutaneous and subcutaneous tissue, accompanied by focal necrosis and infiltration of eosinophils. Granulomatous lymphadenitis was also found in his right groin. The patient was suspected to have Hyper-IgE syndrome, but there were no STAT3 and DOCK8 mutations found.

After discharge, he had less fever, but lymph nodes enlargement and hepatosplenomegaly were still observed. Besides, he suffered from nasosinusitis and oral candidiasis, saprodontia, tooth loss, poor appetite and hard stools. In November 2014, he was hospitalized again because of fever. A routine blood examination showed he had decreased white blood cell (WBC), polymorphonuclear cell (PMN), lymphocyte (LYM) counts and hemoglobin (HGB). Vancomycin and fosfomycin was used to combat the infection, but the treatment was ineffective. The patient still had fever, and gradually developed severe hepatosplenomegaly, digestive hemorrhage, vomiting, hematochezia and dyspnea. Suspended red blood cells were infused to correct anemia, and omeprazole acetic acid was used for symptomatic treatment. However, these treatments were ineffective, and he died of multiple organ failure as a 9-year old in September, 2014.

4.2. Laboratory Analysis

The patient’s peripheral blood immunological data is shown in Table 1. He had normal lymphocytes and T cell subsets, but very low percentage of CD19+B lymphocytes. Interesting is that despite the low percentage of B cells, he was reported to have normal level of serum IgG, IgA, IgM, but extremely high level of IgE.

Neutrophil oxidative burst test was performed in this patient. The mean dihydrorhodamine 123 (DHR) stimulation index (SI) indicating the residual reactive oxygen intermediates (ROI) production capacity of neutrophils was 41.2 in the patient; whereas his parents’ results were 140.97 and 160.21, respectively (Table 1). These results suggested that the patient had defective NADPH oxidase activity.

4.3. Mutation Analysis

To find out the disease causing gene, targeted gene capture combined with next-generation sequencing technology was applied in this study. The results showed that this patient was a compound heterozygote for novel mutations consisting of a C to T transition at position 7016 of exon 46 (c.7016G>T, p.S2339I) and a C to A transition.
Table 1. Laboratory Parameters of the Patient with LRBA Gene Mutations

| Variables | Values |
|-----------|--------|
| **Lymphocyte counts, 10^9/L** | |
| Age at measurement, y | 3.3 | 3.4 |
| 2.55 (1.2 - 3.4) | 3.2 (1.2 - 3.4) |
| **T lymphocyte cell subsets** | |
| Age at measurement, y | 3.3 | 3.4 |
| CD3+ T cell counts, 10^9/L | 1.9 (0.7 - 2.10) | 2.1 (0.7 - 2.10) |
| CD3+, % | 73.30 (64.19 - 75.77) | 68.60 (64.19 - 75.77) |
| CD3+/CD4+, % | 46.10 (30.09 - 40.41)↑ | 41.60 (30.09 - 40.41)↑ |
| CD3+/CD8+, % | 25.70 (20.74 - 29.42) | 23.20 (20.74 - 29.42) |
| CD4+/CD8+, % |  |  |
| **B lymphocyte cell subsets** | |
| Group name | Patient | Age matched controls |
| CD19+ B cell counts, 10^9/L | 0.03↓ | 0.12 - 0.56a |
| CD19+, % | 1.132↓ | 4.3 - 18.2a |
| **Immunoglobulin levels** | |
| Age at measurement, y | 3.6 | 3.9 | 5.8 |
| IgG, g/L | 18.7 (6.94 - 16.85)↑ | 16.2 (6.94 - 16.85) | 16.40 (7.51 - 15.60)↑ |
| IgM, g/L | 1.09 (0.60 - 2.77) | 1.95 (0.60 - 2.77) | 1.49 (0.46 - 3.04) |
| IgA, g/L | 2.39 (0.68 - 3.82) | 2.54 (0.68 - 3.82) | 6.05 (0.82 - 4.53)↑ |
| IgE, IU/mL | 8600 (5.00 - 165.3)↑ | 5220 (5.00 - 165.3)↑ | 29400 (5.00 - 165.3)↑ |
| **Neutrophil oxidative burst test, SI** | |
| Patient | 41.2 | Normal range |
| Father | 140.97 |  |
| Mother | 160.2 | 127.9 (85.2 - 264.6) |

Abbreviation: ND, not determined.

4 Normal values refer to age matched controls published in van Gent R, et al. Clin Immunol 133:95 - 107.

at the fifth nucleotide of intron 45 (c.6667+5C>A, splicing) in LRBA gene (Figure 2A). The frequency of the T variant at c.7016G>T has been reported to be 4.6% in 1000 genomes, and there was no population frequency data of c.6667+5C>A. The following Sanger sequencing results showed that the compound heterozygous mutations were inherited from his parents, in which his father carried a heterozygous c.7016G>T mutation and his mother had a heterozygous c.6667+5C>A mutation (Figure 1B). Mutation damage prediction was made using standard open access software. The p.S2350I was predicted to be probably damaging with a score of 0.963 by using PolyPhen2 (http://genetics.bwh.harvard.edu/pph2); disease causing by using MutationTaster (http://www.mutationtaster.org); affecting protein function with a score of 0.00 through Sorting Intolerant from Tolerant (SIFT; http://sift.jcvi.org). The c.6667+5C>A was predicted to be disease causing by using MutationTaster.

4.4. B Cell Immunophenotyping

To understand the role of LRBA in B cell development, B cell immunophenotyping was performed in this study. Bone marrow progenitor B cell subsets in peripheral blood could be divided into HSC, CLP, Pro-B and Pre-B cells (13). Compared with age-matched healthy controls, all of the bone marrow progenitor B cell subsets (% of lymphocytes) were dramatically decreased in the patient. Besides, immature B cells were also decreased in the patient (Figure 3A and B and Table 2).

Memory B cell subsets could be sub-grouped for their expression of CD27 and IgD on their surface. CD27+IgD+ and CD27+IgD- B cells represented non-switched memory B cells and switched memory B cells, respectively (14).
Figure 2. Mutation analysis of LRBA deficiency. A, Targeted gene capture followed by next-generation sequencing technology was used to screen mutations causing disease in the patient. B, Verify LRBA gene mutations in the patient and his parents by Sanger sequencing. Compound heterozygous mutations including c.6667+5C>A and c.7016G>T were found in the patient. A heterozygous mutation of c.6667+5C>A and c.7016G>T was found in his father and his mother, respectively.

5. Discussion

LRBA deficiency was reported to have a broad and variable clinical phenotype, mainly including hypogam-
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Figure 3. B cell immunophenotyping of the patient with LRBA deficiency and an age matched healthy controls. A, bone marrow progenitor B cell subsets (HSC, CLP, pro-B and Pre-B cells); B, B cell subsets in the antigen-dependent development period (immature B, mB and plasma cells); C, memory B cell subsets (naïve B, NSM, SM, DN); D, B1 subsets (naïve B1, B1 and mB1 cells). HSC, hematopoietic stem cell; CLP, common lymphoid progenitor; Pro-B, progenitor B cells; Pre-B, precursor B cells; NSM, non-switched memory B cells; SM, switched memory B cells; DN, double negative B cells; mB1, memory B1 cells.

maglobulinemia, recurrent infection, autoimmune disorders, organomegaly and chronic diarrhea. Other clinical features included allergy and asthma, growth retardation, neurologic disease, and so on (2, 3, 5, 8). In a recent review including 32 LRBA deficient patients worldwide, 61% patients were reported to suffer from chronic diarrhea, autoimmune diseases, organomegaly and respiratory infections, and 58% patients from hypogammaglobulinemia.
Table 2. Percentages of B-Cell Subsets in the Patient and Age-Matched Healthy Controls (N = 5)

| Bone marrow progenitor B cell subsets, % of lymphocytes | Age Matched Healthy Controls | Patient in This Study |
|--------------------------------------------------------|-----------------------------|-----------------------|
| HSC                                                    | 1.53 (1.15 - 1.96)          | 0.28 ↓                |
| CLP                                                    | 0.15 (0.10 - 0.28)          | 0.01 ↓                |
| Pro-B cells                                            | 0.16 (0.13 - 0.19)          | 0.00 ↓                |
| Pre-B cells                                            | 2.01 (1.87 - 2.76)          | 0.08 ↓                |
| B cell subsets, % of CD19+ cells                       |                             |                       |
| Immature B cells                                       | 19.26 (11.61 - 30.72)       | 9.44 ↓                |
| Naïve B cells                                          | 65.44 (56.72 - 74.61)       | 75.14 ↑               |
| Memory B cell                                          | 26.6 (18.85 - 32.58)        | 7.54 ↓                |
| Non-switched memory B cells                            | 13.4 (9.48 - 19.03)         | 4.25 ↓                |
| Switched memory B cells                                | 13.7 (7.76 - 19.86)         | 3.29 ↓                |
| Double negative B cells                                | 7.03 (4.21 - 12.84)         | 16.99 ↑               |
| IgE+ B cells                                           | 0.556 (0.24 - 0.85)         | 9.63 ↑                |
| Plasma cells, % of lymphocytes                         | 0.39 (0.26 - 0.91)          | 0.13 ↓                |
| B1 cell subsets, % of lymphocytes                      |                             |                       |
| Naïve B1 cells                                         | 7.17 (5.05 - 9.00)          | 1.04 ↓                |
| B1 cells                                               | 0.16 (0.46 - 1.4)           | 0.01 ↓                |
| Memory B1 cells                                        | 2.22 (1.61 - 3.64)          | 0.07 ↓                |

Abbreviations: CLP, common lymphoid progenitor; HSC, hematopoietic stem cell; Pre-B, precursor B cells; Pre-B, progenitor B cells.

At the age of 4 years, the patient was diagnosed as EBV infection. His EBV-associated symptoms included intermittent fever, lymphadenopathy and hepatosplenomegaly, which have been reported in patients with LRBA deficiency (1, 17). However, EBV infection was not frequently observed in LRBA deficient patients. Only 1/17 and 2/31 LRBA deficient patients were reported to suffer from EBV infection in two large cohort studies, respectively (2, 3). In addition, the patient was reported to have autoimmune disease which presented as autoimmune vascular inflammation. Autoimmune disease, which is one of the most frequent clinical phenotypes in LRBA deficient patients, was reported to occur in 76.5% and 61% patient with LRBA deficiency previously (2, 3). The most common features included AIHA (autoimmune hemolytic anemia), ITP (immune thrombocytopenia), IBD (inflammatory bowel disease) and so on (4, 10). In addition, the patient also had clinical presentations which have been frequently reported in LRBA patients previously, such as granulomatous lymphadenitis, sinusitis and oral candidiasis. Azizi et al. (3) reported lymphadenopathy, sinusitis and oral candidiasis occurred in about 52.9%, 70.6% and 23.5% patients, respectively; while a lower frequency was reported in Alkhairy et al. (2).

LRBA deficiency was firstly described to associate with early-onset hypogammaglobulinemia (8). However, later reports showed that not all of the LRBA mutant patients presented with hypogammaglobulinemia (9, 10, 15). In this study, the patient did not have hypogammaglobuline-
in peripheral blood could not fully represent precursor B cell development as early as the bone marrow. Although the early B cell progenitors were still needed for a better understanding of the biological functions of LRBA gene. Further studies are still needed for a better understanding of the biological functions of LRBA gene.

LRBA deficiency becomes more feasible. NGS has many advantages, including cost saving, higher sequencing accuracy and shorter turnover time. More importantly, the data is more feasible for the bioinformatics and further clinical analysis. Thus, it provides a more robust and effective way for the rapid clinical genetic diagnosis regarding human diseases, especially monogenic disease.

As many subtypes of CVIDs, LRBA deficiency has been reported to associate with defective B cell development, but the underlying mechanism still needs to be determined. Gamez-Diaz et al. reported that most LRBA-deficient patients had low B-cell subset counts, mainly including switched memory B cells (80%) and plasmablasts (92%). A further study showed that except for switched memory B cells and plasmablasts, naïve B cells, transitional B cells and marginal zone B cells all decreased in a majority of infected individuals. Besides, T cell deficiency, especially defects in Treg cells, was also reported in a considerable LRBA-deficient patients. Azizi et al reported that, although 94.1% LRBA deficient patients in their cohort had normal T-cell counts, the hyper serum IgE in this patient. However, the underlying mechanism remains unclear.

B cells are a predominant cell population in body cavities. They produce natural antibody, which are known to participate in immune responses against bacteria, viruses and certain parasites. The patient in this study had greatly decreased B cell subsets, which indicated a defective memory B cell development in the patient. Similarly, Lopez-Herrera et al. reported LRBA deficient patients had low switched memory B-cell cells. Gamez-Diaz et al. also observed decreased switched memory B-cells in most (80%) of the LRBA deficient patients. In fact, decreased CD27 memory B cells have been reported in most CVID patients with non-LRBA deficiency. However, the underlying mechanism remains unclear.

Taken together, we reported a patient with compound heterozygous mutations in LRBA gene in China, who presented with EBV infection, systemic vasculitis, recurrent infection and hyper serum IgE. Our results indicated the importance of LRBA in B cell development. Further studies are still needed for a better understanding of the biological functions of LRBA gene.
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Footnotes

Authors’ Contribution: Ji Chen designed this study and reviewed the manuscript; Tong-Xin Chen contributed to project design; Jing Wu performed the laboratory work, analyzed the data and wrote this manuscript; Wei-Fan Wang collected and analyzed the clinical data of this patient.

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