Ichthyosis prematurity syndrome (IPS) is a rare autosomal recessive disorder characterized by prematurity, a thick caseous scale at birth and lifelong atopic diathesis. Here, we describe the first Japanese case of IPS and report novel compound heterozygous mutations (p.C403Y and p.R510H) in fatty acid transport protein 4 (FATP4). She is the first reported patient of Asian origin, entirely distinct from the Scandinavian population, in whom the heterozygote carrier frequency is very high.

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Figure 1. Clinical skin features seen at birth.
eosinophilia. These features are typically indicative of atopic diathesis.

Here, we present the case of a 14-year-old female with atopic dermatitis, increased total IgE and eosinophilia. This patient represents the first Japanese case of genetically diagnosed IPS. Using a next-generation sequencing strategy, we identified compound heterozygosity of novel missense mutations in the fatty acid transporter member 4 (FATP4) gene (p.C403Y and p.R510H).

The proposita was born at the Tosei General Hospital of nonconsanguineous parents. The birth occurred at 32 weeks, 2 days of gestation via spontaneous vaginal delivery. The birth weight was 2,186 g (> 90 percentile). The patient had no family history of ichthyosis. Apgar scores were 2 and 4 at 1 and 5 min, respectively. Tracheal intubation was performed immediately. The patient was placed on a respirator with high inspiratory pressure and she received surfactant replacement therapy. Respiratory assistance was discontinued on day 3 with no further need of oxygen or positive-pressure breathing.

At birth, the patient was coated in a thick layer of white, vernix, caseosa-like material (Figure 1). The entire body, including the face, showed thickening and scaling of the skin, which became bright red during crying. Histological analysis of skin biopsy samples showed a diffuse hyperkeratosis with intracorneal pustule and psoriasiform hyperplasia foci. No evidence of epidermolytic hyperkeratosis was recognized, resulting in a tentative diagnosis of nonbullous congenital ichthyosiform erythroderma. Transient eosinophilia up to 8,400/μl was noted during hospitalization, but this level gradually normalized. After discharge, the eruption improved into a mild flexural dermatitis with generalized ichthyosis under moisturizing-agent treatment.

![Figure 2.](image)

(a) Conservation of the FATP4 protein near two SNVs. (b) Chromatogram derived from targeted capillary sequencing of the patient and her parents for mutations in FATP4, c.1208G>A and c.1529G>A. (c) Schematic drawing of the FATP4 protein and a summary of the mutations reported previously (black arrow) and in this report (red arrow). Functional domains include the transmembrane region (TM), an ER localization signal (ERx) and ATP/AMP (ATP/AMP) and very long-chain acyl-CoA synthetases/fatty acid transport proteins (VLACS/FATP) (FATP) motifs.
When the patient was 14 years old, she visited a nearby clinic because of abdominal pain. Blood examination revealed eosinophilia and elevated serum IgE levels. She was referred to the Tosei General Hospital for further examination. Thickenings of the skin of the patient’s forehead, trunk and limbs was observed at the first visit. This thickening was accompanied by fine desquamation and mild pigmentation. The patient’s limbs were also slightly reddish. Blood examination revealed eosinophilia (2.068/μl), elevated serum IgE (18,000 IU/ml) and multiple sensitizations to respiratory and food allergens (specific IgE to house dust, 84.9 UA/ml; Dermatophagoides pteronyssinus, 93.5 UA/ml; cat dander 2.51 UA/ml; Japanese cedar, 2.21 UA/ml; egg white, 1.06 UA/ml; and ovomucoid, 0.91 UA/ml). As she had no past history of recurrent infections suggesting hyper IgE syndrome, atopic diathesis due to skin barrier defects was speculated.

As the list of differential diagnosis included a variety of nonbullous congenital ichthyosiform erythrodermas,2 we resorted to next-generation sequencing analysis. Targeted exome sequencing was performed using a Trusight Exome sequencing panel on a MiSeq platform (Illumina, San Diego, CA, USA). This kit enables enrichment and final analysis of a panel of ~2,700 genes. The patient and her parents gave written informed consent for genome analysis.

Of the nonsynonymous single-nucleotide variants (SNVs) detected in FATP4, two (c.G1208A and c.G1529A) were not found in the 1000 Genomes database (1000g2012feb), the Exome Variant Server database from 13,006 chromosomes (NHDLGI GO Exome Sequencing Project)3 or the Japanese genetic variation database (Human Genetic Variation Database).4 These variations were strictly conserved among multiple species (Figure 2a) and were predicted to damage protein function by PolyPhen2 (scores 0.996 and 1, respectively) and AVSIFT (score 0 for both SNVs). Mutation validation was performed using PCR and Sanger sequencing of the candidate gene region. This analysis revealed that c.G1208A and c.G1529A were of paternal and maternal origins, respectively (Figure 2b).

The FATP4 gene, consisting of 13 exons, produces a transmembrane protein that transports exogenous fatty acids into cells. The gene also functions as an acyl-CoA synthetase. FATP4 contains a N-terminal transmembrane region, an ER localization signal and a conserved VLACS/FATP motif, which is strictly conserved among multiple species (Figure 2a) and is important for fatty acid binding (VLACS; Figure 2c). IPS is very rare, except in Scandinavia with an estimated heterozygote carrier frequency of 1 in 50.1 Although 13 FATP4 mutations (2 nonsense, 9 missense and 2 splice site) have been reported to date (Figure 2c),5–11 all except two (one from the Middle East and the other from North Africa) are of European origin. Thus, p.C403Y and p.R510H are the first mutations reported in individuals of Asian origins. Clinical features in these individuals were not found to be substantially different from those of Scandinavian origin. Although IPS has not been reported in Japan, a lack of clinician experience may lead to potential cases going undiagnosed. Genetic testing can help to provide a correct diagnosis. Parents can then be informed of the prognostically favorable nature of IPS and its mode of inheritance. This information is necessary for genetic counseling.

Common loss-of-function mutations of the epidermal barrier protein filaggrin, the causative gene of ichthyosis vulgaris, are the major predisposing factors for atopic diathesis.11 Further study of IPS will contribute toEXT-Supported Program for the Strategic Research Foundation at Private Universities from the Ministry of Education, Culture, Sports, Science and Technology (MEXT).

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COMPETING INTERESTS

The authors declare no conflict of interest.

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