A mutation in the immunoproteasome subunit PSMB8 causes autoinflammation and lipodystrophy in humans

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Proteasomes are multisubunit proteases that play a critical role in maintaining cellular function through the selective degradation of ubiquitinated proteins. When 3 additional β subunits, expression of which is induced by IFN-γ, are substituted for their constitutively expressed counterparts, the structure is converted to an immunoproteasome. However, the underlying roles of immunoproteasomes in human diseases are poorly understood. Using exome analysis, we found a homozygous missense mutation (G197V) in immunoproteasome subunit β type 8 (PSMB8), which encodes one of the β subunits induced by IFN-γ in patients from 2 consanguineous families. Patients bearing this mutation suffered from autoinflammatory responses that included recurrent fever and nodular erythema together with lipodystrophy. This mutation increased assembly intermediates of immunoproteasomes, resulting in decreased proteasome function and ubiquitin-coupled protein accumulation in the patient’s tissues. In the patient’s skin and B cells, IL-6 was highly expressed, and there was reduced expression of PSMB8. Downregulation of PSMB8 inhibited the differentiation of murine and human adipocytes in vitro, and injection of siRNA against Psmb8 in mouse skin reduced adipocyte tissue volume. These findings identify PSMB8 as an essential component and regulator not only of inflammation, but also of adipocyte differentiation, and indicate that immunoproteasomes have pleiotropic functions in maintaining the homeostasis of a variety of cell types.

Introduction

Autoinflammatory syndrome is characterized by autoimmunity and inflammatory responses in the absence of infection (1, 2). Recent studies have revealed several genes that are etiologic in many autoinflammatory diseases (3). Initially, inherited missense mutations in TNF receptor superfamily, member 1A (TNFRSF1A), were found in several families afflicted with prolonged fevers and severe localized inflammation, now known as TNF receptor-associated periodic syndrome (4). Subsequently, mutations in the gene NLR family, pyrin domain containing 3 (NLRP3) were reported to cause familial periodic fever syndromes, including Muckle-Wells syndrome (also known as familial cold-induced autoinflammatory syndrome) and neonatal-onset multisystem inflammatory disease.

The proteasome is a multisubunit protease that degrades the majority of nonlysosomal proteins in eukaryotic cells (5). The proteasome degrades ubiquitinated proteins for rapid turnover to homeostasis in a variety of human tissues. In the patient’s skin and B cells, IL-6 was highly expressed, and there was reduced expression of PSMB8. There are 3 additional catalytic β subunits, β1i (also known as immunoproteasome subunit, β type 9 [PSMB9]), β2i (PSMB10), and β5i (PSMB8; or large multifunctional peptidase 7 [LMP7]), whose expression is induced by IFN-γ. The immunoproteasome is assembled in a stepwise manner (6). For instance, β1i first enters the assembly pathway of the immunoproteasome and forms an assembly intermediate containing an α ring, β1i, β2i, β3, and β4 (7). β5i (referred to herein as PSMB8) is incorporated preferentially over β3 into intermediates that contain β1i and β2i and is required for the processing of β1i and β2i (8, 9). PSMB8 deficiency retards immunoproteasome assembly and leads to the accumulation of proteasome precursors containing unprocessed β1i and β2i (10). PSMB8-deficient mice do not exhibit any gross changes, although they have reduced MHC class I expression (11). Although a recent paper indicated that a PSMB8 inhibitor was able to suppress autoimmune responses in mice (12), PSMB8-deficient mice have a high susceptibility for experimental autoimmune encephalomyelitis, which suggests that the loss of PSMB8 in mice is involved in the progression of inflammatory responses (13). Recently, Agarwal et al. demonstrated a missense mutation in PSMB8 in patients with a variety of symptoms, including lipodystrophy and joint contractures (14), although they did not address the mechanism by which the mutated PSMB8 contributed to each symptom. Therefore, it remains unclear how the mutation in immunoproteasomes contributes to human diseases and how immunoproteasomes contribute to homeostasis in a variety of human tissues.
We examined candidate genes of 3 patients, from 2 consanguineous families, with an autoinflammatory syndrome characterized by recurrent fever, nodular erythema, and hypergammaglobulinemia together with a loss of adipose tissue in the upper part of the body (15, 16). Clinically similar cases have been categorized as Nakajo syndrome (OMIM 256040; ref. 17), and other cases with similar clinical manifestations were previously reported in Japan (18). These manifestations, characteristic of a newly recognized type of Japanese autoinflammatory syndrome with lipodystrophy (JASL), suggest that the causative genes for JASL contribute to both immunity and adipocyte metabolism. Here, we identified a missense mutation in PSMB8 in patients with JASL, which we believe to be novel. The identified PSMB8 mutation increased assembly intermediates of immunoproteasomes, resulting in reduced expression of PSMB8. Low PSMB8 expression in humans caused increased p38 phosphorylation, which resulted in increased IL-6 production. Surprisingly, siRNA-mediated downregulation of PSMB8 disturbed adipocyte differentiation both in vitro and in vivo. These data indicate that a mutation in immunoproteasome complexes causes autoinflammation through increased IL-6 production and lipodystrophy through disturbed adipocyte differentiation. Moreover, our results illustrate the crucial functions of immunoproteasomes in cellular homeostasis and provide molecular insights into immunoproteasome-mediated human diseases.

**Results**

**Clinical phenotypes of JASL patients.** We identified 3 patients with JASL from 2 Japanese consanguineous families. The parents of the families were consanguineous, and the 2 families were not known to be related (Figure 1A). The clinical characteristics of the 3 patients were summarized in Supplemental Table 1 (supplemental material available online with this article; doi:10.1172/JCI58414DS1). Clinical and laboratory findings of our JASL patients and previously reported Nakajo syndrome or Nakajo syndrome-like diseases are matched by the same clinical and laboratory features.
summarized in Supplemental Table 2. Patients began to present recurrent high fever (>40°C) with nodular erythema (Figure 1B) at 1 month to 3 years of age. The recurrent high fever without infectious episodes and nodular erythema on the entire body are common clinical features among JASL patients. Muscle weakness, deformities of hands (Figure 1B), and frostbitten hands were also seen in all patients. The patients began to develop partial lipodystrophy from 6 to 12 years of age (Supplemental Tables 1 and 2). The lipodystrophy was particularly prominent in the face, fingers, and upper limbs (Figure 1B), while nodular erythema developed on the skin of the entire body. Biopsy of skin lesions from patient 2-2B (from family 2) showed a massive infiltration of neutrophils and lymphocytes in the dermis (Figure 1C). The clinical and laboratory findings for JASL and other types of primary or secondary lipodystrophy are compared in Supplemental Table 3. Serum C-reactive protein, IgG, and IgA levels were very high. However, autoantibodies (including anti-nuclear antibody, dsDNA, SS-A, SS-B, and Jo-1 antibodies) were not detected, and blood glucose, total cholesterol, triglyceride, and complement levels were normal (Supplemental Table 1). Patients were treated with moderate doses of corticosteroids (10–15 mg/d), which controlled nodular erythema and high fever and reduced acute-phase reactants, but never inhibited the progression of lipodystrophy. Female patient 1-3PT (from family 1) has 1 daughter who is healthy at the age of 35 years. All patients lived over 47 years of age and died of respiratory or cardiac problems.

Genetic analysis of JASL. Family 2 has been previously reported (15, 16). To identify candidate loci for this disease, homozygosity mapping was performed in both families. We genotyped 3 affected patients in families 1 and 2 and 1 unaffected sibling in family 1. Multipoint linkage analysis using Merlin software revealed a maximum LOD score of 3.435 at the chromosome 6p region (chr6, 27,268,074–40,724,868; Figure 1D). We then searched for consecutive homozygous SNPs by HomozygosityMapper (19) in order to identify candidate regions of homozygosity in affected but not unaffected offspring, which showed only 6p (chr6, 26,476,155–41,414,550) as a candidate region (Figure 1E) that nearly overlapped the region obtained from the linkage analysis. To further narrow these candidate regions, we identified shared haplotypes of common homozygous SNPs within and across families. Based on these approaches, we identified a single 6.8-Mb region of homozygosity on 6p22.2–6p21.31 (between rs9283878 and rs210120) shared by all patients (Figure 1F). This interval was found to contain 465 gene entries, according to the NCBI database (http://www.ncbi.nlm.nih.gov/).

In order to identify the causative genes in the candidate region on chromosome 6, we focused on exonic and flanking intronic variants within this region. Alignment of the sequences in this region with exome analysis revealed that targeted exons (90%) had greater than 10 high-quality reads. The targeted exons with fewer than 10 high-quality reads were further analyzed by Sanger sequencing. The 684 homozygous variants in both patients were identified in this region; of these, 633 appeared in dbSNP130, and the remaining 51 were novel variants (Table 1). We found 2 variants in STK19 or PSMB8 to cause nonsynonymous substitutions; others were located in introns (not splicing-related regions) or were synonymous substitutions. In order to examine whether the 2 amino acid–altering mutations were previously identified variants in the Japanese population, we genotyped each in 312 healthy Japanese controls. The frequency of the missense mutation in STK19 was seen in 4 of 100 control alleles, which suggests that it is a benign polymorphism. In contrast, G197V of PSMB8 was not present in the 624 alleles of the 312 healthy controls (Table 2).

Functional analysis of mutant PSMB8. Whereas 1-3PT carried the homozygous mutation, the mother of family 1 (1-1M) and the affected patient’s healthy sibling (1-2ES) carried the heterozygous mutation (Figure 2A and data not shown). PSMB8 has 2 isoforms, and the missense mutation changed an amino acid from G to V at position 197 or 201 in the different isoforms (Figure 2B). The corresponding region was conserved among species (Figure 2C). G197 occupies a surface-exposed position in a β hairpin turn (Supplemental Figure 1, A–D). From the homology model, we can deduce that a substitution for a larger residue (V) would create some crowding, but, more significantly, would restrict the flexibility of the backbone in the region corresponding to the hairpin turn. We modeled mouse subunits β1i, β2i, and β5i (i.e., PSMB8) with the corresponding constitutive subunits and found that position 197 sat outside the subunit-subunit interface (Figure 3A) and S1 pocket (Figure 3B). We also found that G197 sat outside the β-ring–β-ring interface (Figure 3C).

The amino acid sequence around G197 of PSMB8 was aligned with 83 sequence homologs obtained by running BLAST (http://blast.ncbi.nlm.nih.gov/) against the protein data bank with known structure, which indicated that G197 was the most conserved residue in PSMB8, at 100% conservation among 84 proteins (Figure 3D). The structure modeling and alignment data suggest that the G197 region of PSMB8 is important for stability or folding of the protein.

Peripheral blood B cells from 1-3PT, 1-2ES, and 1-1M and a healthy unrelated control were transformed with EBV. The mRNA level of PSMB8 in 1-3PT B cells was relatively higher than those of 1-2ES or 1-1M (Figure 4A), whereas expression of mature PSMB8 protein was much lower in 1-3PT B cells than in those from 1-2ES, 1-1M, and the unrelated control (Figure 4B). The higher level of PSMB8 mRNA expression in 1-3PT B cells relative to control cells might be due to the presence of compensa-
tory mechanisms for low PSMB8 protein expression. Treatment of 1-3PT B cells with a proteasome inhibitor did not increase PSMB8 expression (Supplemental Figure 2A). The constitutively expressed proteasome components β1, β2, and β5 were comparable between 1-3PT B cells and those of 1-1M and 1-2ES (Supplemental Figure 2B). To examine whether the missense mutation reduced the stability of immature PSMB8, 293T cells transfected with wild-type or mutant immature PSMB8 were labeled with [35S]-methionine and cysteine, and the stability of PSMB8 was examined. The speed of degradation was comparable between the 2 PSMB8 groups (Supplemental Figure 2C).

We then determined whether the mutation affected assembly of immunoproteasomes by immunoblotting lysates from transduced B cells using anti-β1i, anti-α6, and anti-UMP1 antibodies (20). In each blot, we found an increased proportion of an assembly intermediate (an intermediate stage of immunoproteasome before reaching full maturation; ref. 6) in cell lysates from 1-3PT compared with control B cells (Figure 4C). The lysates from 1-1M and 1-2ES, with 1 allele of mutant PSMB8, also had more assembly intermediates than did the unrelated control (Figure 4C).

Cell lysates from control or 1-3PT B cells were separated by glycerol gradient centrifugation to determine which mature or immature immunoproteasome subunits were incorporated in assembly intermediates or in 20S or 26S proteasomes. The assembly intermediates containing immature β1i and α6 (fraction 10) were increased in 1-3PT compared with control B cells (Figure 4D). Immature PSMB8 was observed in the 20S (fraction 16) and the 26S (fraction 22) immunoproteasomes only in 1-3PT B cells (Figure 4D). Furthermore, β1i, α6, and PSMB8 in the 20S fraction were decreased in 1-3PT compared with control B cells, which suggests that the total amount of mature immunoproteasomes decreased in 1-3PT cells. Taken together, these data indicate that the mutation in PSMB8 disturbs assembly of immunoproteasomes, resulting in low levels of their expression.

We then compared the chymotrypsin-like activities of proteasomes from 1-3PT, 1-1M, 1-2ES, and the unrelated control. We found that 1-3PT cells, with mutant PSMB8 protein, possessed reduced chymotrypsin-like activity compared with control cells (Figure 5A). Furthermore, total proteasome activity was reduced in B cells from 1-3PT compared with that of the healthy control, as determined by ornithine decarboxylase (ODC) degradation (Figure 5B). ODC is degraded by proteasomes independent of ubiquitin. Because a previous paper showed that PSMB8 is expressed in the skin (21), we next compared the expression level of PSMB8 in skin from a control and from 2-2B. Whereas PSMB8 was expressed in control skin, localized in both the nucleus and the cytoplasm, its expression was much lower in 2-2B (Figure 5C). Taken together, these data suggest that the missense mutation in PSMB8 in JASL patients reduces both protein expression and proteasomal functions.

Accumulation of ubiquitinated proteins in cells carrying the mutant PSMB8. PSMB8 is one of the proteasomal components contributing to the degradation of ubiquitinated proteins. Based on the low expression of PSMB8 in B cells from 1-3PT, we first test-
ed the expression of ubiquitin in transformed B cells from the unrelated control, 1-1M, and 1-3PT. Ubiquitin expression level was higher in B cells from 1-3PT than in those from the 2 controls (Figure 6A). We then examined by immunohistochemistry whether ubiquitin accumulates in 1-3PT cells as a result of impaired PSMB8 function. Staining of skin sections with an anti-ubiquitin antibody showed little expression of ubiquitin in healthy control skin (Figure 6, B and C). In contrast, accumulated ubiquitin was observed in the nucleus or the cytoplasm of epidermal cells from 2-2B (Figure 6D). Furthermore, a dense accumulation of ubiquitin with a punctate pattern and irregular shapes was detected in the cytoplasm of sebaceous cells of sweat glands and hair follicular cells (Figure 6, E and F). Taken together, our data regarding accumulation of ubiquitinated protein in transformed B cells and skin of JASL patients suggest that increased ubiquitin protein accumulation results from reduced expression of PSMB8 altering proteasomal activity, rather than from inflammation.

Increased IL-6 expression in cells carrying mutant PSMB8 occurs via p38. Hyperactivation of NLRP3 is involved in the pathogenesis of hereditary autoinflammatory syndromes (22). The activation of NLRP3 cleaves pro–caspase-1, which generates p10 caspase-1 (22). We tested whether NLRP3 inflammasomes contributed to autoinflammatory symptoms in JASL patients by measuring caspase-1 activation. B cells from the unrelated control, 1-1M, and 1-3PT did not show any activation of caspase-1 without stimulation with Alum (Supplemental Figure 3). The 3 subjects' cells showed comparable expression of p10 of caspase-1 by stimulation with Alum (Supplemental Figure 3), which suggests that NLRP3 is not involved in autoinflammatory symptoms in JASL.

Kasagi et al. reported that serum IL-6, a crucial mediator of inflammation, was elevated in a JASL patient (17). Since the 3 JASL patients in the 2 families were deceased, we assessed expression of IL6 mRNA in skin from 2-2B, and found it to be much higher than that in control skin (Figure 7A). Next, we evaluated whether PSMB8 was involved in IL-6 regulation. Using transformed...
B cells from 1-3PT, we examined IL-6 expression with or without transduction by a retrovirus carrying wild-type PSMB8. The 1-3PT B cells infected with PSMB8-encoding retrovirus (referred to herein as 1-3PT–PSMB8 cells) expressed levels of PSMB8 comparable to those of control 1-3PT B cells (1-3PT–EV; Figure 7B). Expression of IL6 by PMA- and ionomycin-stimulated 1-3PT–EV cells was reduced in 1-3PT–PSMB8 cells, but still higher than that in control B cells from the unrelated healthy control or 1-1M (Figure 7B). This incomplete reduction of IL6, even by overexpressing wild-type PSMB8, might be due to the secondary effect on transformed B cells induced by long-lasting impaired immunoproteasomes functions. Since a MAP kinase pathway controls IL-6 expression (23, 24), we tested various MAP kinase inhibitors to determine whether they could suppress IL6 expression in the B cells of a JASL patient. Transformed 1-3PT B cells were cultured in the presence of inhibitors of MEK1/2, p38, or JNK. The addition of an inhibitor for p38 suppressed IL6 expression; in contrast, MEK1/2 and JNK inhibitors could not significantly suppress IL-6 expression, although the JNK inhibitor demonstrated partial suppression (Figure 7C). We next tested whether PSMB8 affects p38 phosphorylation in 1-3PT B cells stimulated with PMA and ionomycin. 1-3PT–EV and 1-3PT–PSMB8 cells expressed comparable levels of total p38 (Figure 7D). Phosphorylation of p38 in 1-3PT–EV cells was greater after stimulation with PMA and ionomycin than in 1-3PT–PSMB8 cells, although the latter was still higher than that in control B cells (Figure 7E). Again, this incomplete correction of high phospho-p38 expression, even by overexpressing wild-type PSMB8 in patient B cells, might be due to the secondary effects on transformed B cells induced by long-lasting impaired immunoproteasomes functions, although it might be difficult to compare the expression in different types of transformed B cells. Taken together, these data indicated that low PSMB8 expression in 1-3PT resulted in the upregulation of IL-6, at least partially through a p38-mediated pathway.

PSMB8 regulated adipocyte differentiation. Patients with JASL gradually develop partial lipodystrophy. Therefore, we next examined the possibility that PSMB8 regulates adipocyte differentiation. The murine preadipocyte cell line 3T3-L1 highly expressed PSMB8 (Supplemental Figure 4A). In order to evaluate whether PSMB8 is involved in preadipocyte maturation, we tested the change of PSMB8 expression during adipocyte differentiation. We found that PSMB8 expression did not change during differentiation toward adipocytes (Supplemental Figure 4A). PSMB8 expression was then downregulated using a siRNA targeting PSMB8. The PSMB8 siRNA transfection efficiently downregulated PSMB8 (Figure 8A) and did not affect the viability of 3T3-L1 cells (Figure 8B). Transfected 3T3-L1
cells were allowed to differentiate into adipocytes, and cytoplasmic lipid droplets were evaluated by Oil Red staining 6–7 days after transfection. The downregulation of PSMB8 in 3T3-L1 cells inhibited adipocyte differentiation, as evidenced by the reduced number and amount of Oil Red–positive areas (Figure 8C). Downregulation of the β2i subunit did not affect adipocyte differentiation (Supplemental Figure 4B), which indicates that PSMB8-mediated proteasomal activity is required for adipocyte differentiation. We also examined the expression of PSMB8 in human preadipocytes and found high expression of PSMB8 (Figure 8D). Transfection of siRNA against PSMB8 in human preadipocytes efficiently downregulated PSMB8 (Figure 8E), which disturbed their differentiation into mature adipocytes, as evaluated by Oil Red staining (Figure 8F). Although to our knowledge, it has not been reported that animals deficient in PSMB8 exhibit lipodystrophy, we next inhibited PSMB8 expression in the skin of adult BALB/c mice by subcutaneously injecting PSMB8 siRNA. Treatment of mice with PSMB8 siRNA efficiently downregulated Psmb8, as evaluated by PCR and histological analysis (Supplemental Figure 4, C and D). Histological analysis of skin specimens showed atrophic subcutaneous adipose tissue and a decreased number of hair follicles in mice that received PSMB8 siRNA compared with control mice (Figure 8G). A smaller number of adipocytes was observed in mice that received PSMB8 siRNA, as determined by staining cells with anti-FABP4 antibody (Supplemental Figure 4E). Taken together, these data indicate that PSMB8 is critically involved in the differentiation of preadipocytes into adipocytes, which could account for the progressive lipodystrophy observed in JASL patients.

Discussion
We found patients from 2 consanguineous families characterized by nodular erythema, recurrent fever, and lipodystrophy and investigated the gene causing JASL. We found a PSMB8 homozygous missense mutation — which we believe to be novel — that increased an intermediate form of immunoproteasomes, resulting in reduced proteasomal function. The discovery of the mutation in human PSMB8 allowed us to evaluate the physiological and pathological roles of immunoproteasomes. A distinct missense mutation in PSMB8 was previously reported in patients suffering from lipodystrophy, joint contractures, and muscle atrophy (14). However, there has been no understanding of the molecular mechanisms underlying the low function of immunoproteasomes and the variety of associated symptoms. Therefore, in this study, we focused on evaluating the mechanisms underlying the 2 major symptoms of JASL, autoinflammatory responses and lipodystrophy. We found that B cells and skin from JASL patients expressed high levels of IL-6, and that JASL B cells upregulated IL-6 through the p38 pathway. High levels of IL-6 production were partially reversed by overexpression of wild-type PSMB8. We also found that downregulation of PSMB8 in mouse or human preadipocytes disturbed their differentiation toward mature cells. Furthermore, injection of siRNA targeted against PSMB8 in mouse skin reduced the number of adipocytes in their skin. These data provide mechanistic insights into autoinflammation and lipodystrophy of JASL and, furthermore, demonstrate physiological roles of immunoproteasomes in regulating both inflammation and adipocyte differentiation.
The NLRP3-mediated autoimmune inflammatory syndrome depends on increased production of active IL-1 through caspase-1 activation (22, 25). JASL patient B cells did not show increased production of active caspase-1 compared with control B cells, which suggests that PSMB8 deficiency does not activate the NLRP3 pathway. In contrast, patients with JASL exhibited high expression of IL-6, which exerted pleiotropic effects in promoting inflammatory responses in the skin (present study) and in serum (17). Indeed, a previous paper indicating that increased expression of IL-6 causes skin inflammation (26) supports the link between increased IL-6 expression and autoinflammation in JASL patients. As for the mechanism of high IL-6 production in cells of JASL patients, our data indicated that overproduction was associated with increased phosphorylation of p38 in B cells, although it remains unclear which p38-related molecules actually promote IL-6 production. Furthermore, treatment of JASL patient B cells with a p38 inhibitor did not completely abrogate IL-6 production, which suggests that a p38-independent pathway is also involved in elevated IL-6 production in JASL. Additionally, a recent paper reported that targeted deletion of Psmmb8 in mice exacerbated experimental autoimmune encephalomyelitis with an increase in oxidized proteins (13), which suggests that PSMB8 is also involved in the progression of inflammatory responses through increased accumulation of oxidized protein. It should be noted that treatment with a PSMB8 inhibitor suppressed rheumatoid arthritis in mice (12). Therefore, careful analysis of PSMB8 inhibitors could elucidate the roles of PSMB8 in distinct types of inflammatory responses. We are currently developing a knockin mouse that has the same mutation as 3 JASL patients in our study. Those mice will broaden our understanding of the roles of immunoproteasomes in the pathogenesis of autoinflammation and other inflammatory diseases.

The missense mutation in PSMB8 decreased expression of mature PSMB8, although it did not affect the stability of immature forms of the protein. We found increased levels of assembly intermediates containing immature β1i and reduced expression of 20S immunoproteasomes in JASL patient B cells. As β1i is incorporated into immunoproteasomes before incorporation of PSMB8 (7), these data suggest that the mutation in PSMB8 disturbs incorporation of PSMB8 into immunoproteasomes or facilitates separation of PSMB8 from mature immunoproteasomes. In addition, the increased levels of immature forms of PSMB8 in 20S and 26S immunoproteasomes in JASL patient B cells indicated that the mutation in PSMB8 affects its processing. Together, these data suggest that the mutation in PSMB8 leads to abnormal assembly of immunoproteasomes and reduced levels of mature immunoproteasomes. Indeed, JASL patient B cells exhibited low immunoproteasomal function compared with control cells, as evidenced by their reduced chymotrypsin-like activity and ODC degradation, although the former was more profound than the latter. Given that PSMB8 possesses chymotrypsin-like activity (6), assessing that activity might be a more sensitive measure of PSMB8 function than the ODC degradation assay, which provides a measure of total protease activity.

In our study of immunoproteasome functions in JASL patients, we detected homogenous accumulation of ubiquitin or a dense accumulation with irregular shapes in the nucleus and cytoplasm of keratinocytes, sweat gland cells, or hair follicular cells together within transformed JASL patient B cells. Although we do not have evidence that the accumulation of ubiquitin-coupled protein in JASL patients is associated with autoinflammation, the characteristic accumulation patterns might disturb cellular functions or trigger inflammatory responses. We previously reported that

Figure 6
Ubiquitinated proteins accumulate in JASL cells. (A) Expression of ubiquitin in transformed B cells from healthy control, 1-1M, and 1-3PT, evaluated by Western blotting with anti-ubiquitin and β-actin mAbs. Data are representative of 5 experiments. (B–F) Skin sections of a healthy control (B and C) and 2-2B (D–F) stained with anti-ubiquitin mAb together with hematoxylin. Original magnification, ×20 (B and D); ×40 (C and E). The boxed region in E is shown enlarged in F. Histological data are representative of 4 independent staining experiments using a biopsy sample from each patient. Scale bars: 100 μm (B and D); 50 μm (C and E); 10 μm (F).
DNase1 deficiency can cause systemic lupus erythematosus as a result of accumulated DNA-related products (27–29). Such a defective clearance of intracellular products might be a primary cause for other immune-associated diseases. If this is the case, it would be reasonable to develop strategies for reducing or inactivating accumulated molecules, rather than suppressing the immune system in order to treat such diseases. In addition, immunoproteasomes contribute to antigen processing for MHC class I. Therefore, it will be important to analyze the contributions of mutant PSMB8 to this pathway in future studies.

We demonstrated herein that PSMB8 is involved in the differentiation of preadipocytes to adipocytes, which might contribute to lipodystrophy in JASL patients. The finding that corticosteroid therapy could completely suppress autoinflammatory responses, but not lipodystrophy progression, also suggests that the lipodystrophy in JASL patients is caused by the primary effect of impaired PSMB8 function on adipocyte differentiation, rather than a secondary effect of autoinflammation. We examined the amount of ubiquitin-coupled protein in 3T3-L1 cells in which PSMB8 was downregulated and did not observe gross accumulation of ubiquitin-coupled protein (data not shown). Therefore, we think it unlikely that nonspecific accumulation of large amounts of ubiquitin-coupled protein in preadipocytes (in which PSMB8 expression is reduced) disturbs the differentiation of preadipocytes to adipocytes. Rather, altered turnover of specific proteins that regulate adipocyte differentiation might be affected by downregulation of PSMB8. The identification of such molecules might provide new targets for therapeutic strategies in the treatment of metabolic diseases.
PSMB8-deficient mice do not spontaneously develop autoinflammatory symptoms and lipodystrophy. One possibility to explain the discrepancy between the distinct phenotypes of JASL patients and PSMB8-deficient mice is that the mutant PSMB8-mediated assembly defect in immunoproteasomes might lead to unexpected immunoproteasome activity. Reduced functions of immunoproteasomes with altered activity in JASL might contribute to some aspects of JASL phenotypes. We expect to clarify those discrepancies by establishing knockin mice that carry the JASL mutation.

Agarwal and colleagues independently published results about PSMB8 mutation in JASL-like syndrome (14), although they did not address how the mutation in PSMB8 was involved in the symptoms of JASL-like syndrome. In the present report, we characterized an autosomal-recessive mutation in JASL, which we believe to be novel, and JASL-like syndrome. In the present report, we characterized an autosomal-recessive mutation in JASL, which we believe to be novel, and JASL-like syndrome.

Methods

Further information can be found in Supplemental Methods.

DNA samples. Genomic DNA was extracted from peripheral blood and autopsy samples obtained from 2 families after obtaining informed consent. SNP homozygosity mapping and linkage analysis. Both affected and unaffected siblings from 2 consanguineous families were studied with the Illumina Human 610 Quad. Sample processing and labeling were performed in accordance with the manufacturer’s instructions. An average call rate greater than 99% was obtained. Homozygous regions were identified using HomozygosityMapper (19). Our laboratory has developed a graphic interface that allows for easy comparison of common homozygous regions among 3 patients from different families. For linkage analysis, multipoint LOD scores were obtained using Merlin software (30).

Flow cytometry. Cells were resuspended in staining buffer at a density of 2 × 10^6 cells/ml. Cells were stained with rabbit IgG (10500; Caltag Laboratories) or polyclonal anti–phospho-p38 (4511; Cell Signaling Technology) or anti–phospho-p38 antibody, we did not observe phospho-p38–positive cells in the p38-deficient KOP cell line (Riken Cell Bank) or in KOP cells transfected with cDNA for human p38 (provided by S. Matsuda, Kansei Medical University, Moriguchi, Japan) or constitutively active MKK6 (provided by Y. Gotoh, University of Tokyo, Tokyo, Japan; ref. 31). However, we observed phospho-p38–positive cells in KOP cells transfected with both human p38 and constitutively active MKK6 (Supplemental Figure 5). The fluorescence intensity of 10^6 cells was examined using a FACSCalibur or FACS Canto II cytometer (BD Biosciences).

Assay of proteasome activity. Transformed B cells from 1-1M, 1-2ES, 1-3PT, or a healthy unrelated control were lysed and subjected to 8%–32% glycerol gradient centrifugation. Subsequently, peptidase activity was measured using a fluorescent peptide substrate, succinyl-Leu-Leu-Vl-Tyr-7-amido-4-methylcoumarin (Suc-LLVY-MCA), for chymotrypsin-like activity as previously reported (32).

Structure modeling. A structural model of PSMB8 was built using Spanner (http://sysimm.ifrec.osaka-u.ac.jp/cgi-bin/spanner), which uses a fragment assembly algorithm to produce a gapless alignment for the template. Multiple sequence alignments were computed using MAFFT (33). Electrostatic surfaces were prepared using the program eSurf (34).
Protein extracts, immunological analysis, and glycerol gradient analysis. Cells were lysed in ice-cold lysis buffer (50 mM Tris–HCl, pH 7.5; 0.5% [v/v] NP-40; 1 mM dithiothreitol; 2 mM ATP; and 5 mM MgCl₂), and the extracts were clarified by centrifugation at 20,000 g for 15 minutes at 41°C. The supernatants were subjected to native PAGE (7% Tris–acetate gel; Invitrogen) according to the instructions provided by the manufacturer. The separated proteins were transferred onto a polyvinylidene difluoride membrane and reacted with the indicated antibody. Anti-Ump1, anti-α6, and anti-β61 polyclonal antibodies were described previously (20, 35). ODC degradation activity was assayed as described previously (36). For glycerol gradient analysis, cell lysates were fractionated by linear glycerol density gradient centrifugation (22 hours, 100,000 g) as described previously (36).

Statistics. Unless otherwise indicated, data are shown as mean ± SD. For all experiments, the significance of differences between groups was calculated using the Mann-Whitney U test for unpaired data. Differences were considered significant for P values less than 0.05.

Study approval. All genetic studies were conducted after Ethical Commit tee approval of human genome research of the University of Tokushima and informed consent was obtained from patients, or from their relatives if patients were deceased. The study was conducted in accordance with the principles of the Declaration of Helsinki. All animal work was approved by the Animal research committee of the University of Tokushima.

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