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Mutations in the human SC4MOL gene encoding a methyl sterol oxidase cause psoriasiform dermatitis, microcephaly, and developmental delay

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Defects in cholesterol synthesis result in a wide variety of symptoms, from neonatal lethality to the relatively mild dysmorphic features and developmental delay found in individuals with Smith-Lemli-Opitz syndrome. We report here the identification of mutations in sterol-C4-methyl oxidase–like gene (SC4MOL) as the cause of an autosomal recessive syndrome in a human patient with psoriasiform dermatitis, arthralgias, congenital cataracts, microcephaly, and developmental delay. This gene encodes a sterol-C4-methyl oxidase (SMO), which catalyzes demethylation of C4-methylsterols in the cholesterol synthesis pathway. C4-Methylsterols are meiosis-activating sterols (MASs). They exist at high concentrations in the testis and ovary and play roles in meiosis activation. In this study, we found that an accumulation of MASs in the patient led to cell overproliferation in both skin and blood. SMO deficiency also substantially altered immunocyte phenotype and in vitro function. MASs serve as ligands for liver X receptors α and β (LXRα and LXRβ), which are important in regulating not only lipid transport in the epidermis, but also innate and adaptive immunity. Deficiency of SMO represents a biochemical defect in the cholesterol synthesis pathway, the clinical spectrum of which remains to be defined.

Introduction
Cholesterol is a key component of cell membranes and lipid rafts and is the immediate precursor of steroids, vitamin D, and bile acids. Many disorders of cholesterol synthesis share common clinical features, such as abnormal morphogenesis, growth delay, and psychomotor disabilities (1). However, there are also striking differences suggesting that reduced de novo cholesterol synthesis per se may not primarily underlie some of the symptoms, including cataracts as well as skin and immune system abnormalities. Rather, recent studies implicate the accumulation of pre-cholesterol sterols and the replacement of cholesterol with some of these sterols in lipid rafts as playing a key role in the underlying pathophysiology (2). The meiosis-activating sterols (MASs) were the first group of cholesterol biogenesis intermediates that were found to have important extrahepatic functions in mammals. These include 4,4′-dimethyl-5α-cholesta-8,24-dien-3β-ol (testis meiosis-activating sterol [T-MAS]), 4,4′-dimethyl-5α-cholesta-8,14,24-trien-3β-ol (follicular fluid meiosis-activating sterols [FF-MASs]), and zymosterol. They are found in high concentrations in testis and ovary and play roles in meiosis activation. The function of the MASs outside the reproductive organs is not well studied. FF-MAS is also a ligand for liver X receptors (LXRs) (3). LXR signaling is known to regulate crosstalk between inflammatory and cholesterol metabolism, which is important in regulating cholesterol efflux in epidermis (4). The association between the MASs and overproliferative inflammatory skin diseases is illustrated by a number of genetic defects in the pathway of processing of MASs. Six genes are expected to be involved in the metabolism of MASs (Supplemental Figure 3; supplemental material available online with this article; doi:10.1172/JCI42650DS1). Autosomal recessive defects in DHR14B/LBR lead to hydrops–ectopic calcification–“moth-eaten” (HEM) dysplasia, which is a neonatal lethal condition in humans, while mutations in this gene in mouse cause ichthyosis and neutrophilic anomalies (5). Haploinsufficiency of DHR14B/LBR leads to the Pelger-Huët anomaly of altered nuclear morphology in granulocytes (6). NSDHL and EBP are X-linked genes that lead to embryonic lethality in most males. Heterozygous mutations in NSDHL in females cause congenital hemidysplasia with ichthyosiform erythroderma and limb defects (CHILD) syndrome, and EBP mutations cause chondrodysplasia punctata 2 (CDPX2). Mutations in the murine homologs of EBP and NSDHL lead to striated and bare patches on the skin (7).

In this study, we characterize genetic defects in SC4MOL (OMIM 607545), encoding a sterol-C4-methyl oxidase (SMO). This defect represents what we believe to be the first viable autosomal recessive condition in the MAS demethylation pathway, linking MAS demethylation with cell overproliferation, cholesterol homeostasis, and immune regulation in skin and blood and providing a model to study the pathogenesis of psoriasiform skin disease in this group of disorders.
RESULTS
Clinical presentation. A 13-year-old female presented for evaluation of severe ichthyosiform erythroderma affecting her entire body but sparing the palms (Figure 1, A–C). She did not show signs of dermatitis at birth; it was first noted around her umbilicus at the age of 2 and subsequently progressed to her back, trunk, and then the remainder of her body by the age of 6. Her dermatitis worsened in the winter or when she was under stress, and once almost completely normalized. Also notable in the patient’s medical history are congenital cataracts, mild developmental delay, microcephaly (head circumference 53.5 cm at the age of 13 years, <3rd percentile; 50th percentile for a 7.5-year-old), and failure to thrive. Her height had increased along the 3rd percentile between 9 and 39 months of age, and at age 13 years, her weight was 28.6 kg (<3rd percentile; 50th percentile for a 9-year-old) and her height was 140 cm (<3rd percentile; 50th percentile for a 10.5-year-old). A skeletal survey showed delayed skeletal maturation, but bone morphology was normal. Skin histology showed psoriasiform hyperplasia, thinning of suprapapillary plate, and neutrophils in the epidermis and stratum corneum (Figure 1D). Neutrophil elastase staining of neutrophils in the stratum corneum is shown in red in Figure 1F. Although this histology was initially considered to be consistent with psoriasiform dermatitis, more detailed examination of the tissue revealed the presence of several foamy cells in the dermis, and oil red O staining (Figure 1E) revealed intracellular lipid in these cells, reminiscent of that reported in patients with CHILD syndrome (8). However, the foamy cells were CD68 negative, indicating that they were not macrophages, a distinct difference from CHILD syndrome and verruciform xanthoma. Immunohistochemistry and H&E-stained sections suggested that these cells were lipid-laden fibroblasts. Taken together, the results indicate that the histologic features are consistent with psoriasiform dermatitis with some features of a verruciform xanthoma (Figure 1, D and E). In light of the psoriasiform changes, traditional therapies for psoriasis were implemented for the patient, including topical corticosteroids, calcipotriene, cyclosporine A, etanercept, phototherapy, and oral isotretinoin. However, while her skin briefly improved with cyclosporine A, she did not exhibit a long-term response to any of the therapies. Her serum lipid profile showed a persistently low total cholesterol level (85 mg/dl; normal, 140–176), low HDL (28 mg/dl; normal, 35–75), and low LDL (49 mg/dl; normal, 70–160) but normal triglycerides (84 mg/dl; normal, 50–200) and VLDL 8 mg/dl; normal, 8–14).

Biochemical and molecular characterization of SMO deficiency. A cholesterol biosynthesis defect was suspected on the basis of clinical presentation, and plasma sterols were analyzed by gas chromatography–mass spectrometry (GC-MS). The profile showed 20- and 500-fold elevation of 4α-monomethyl sterols and 4, 4’-dimethyl sterols, respectively (Figure 2, A–D). No 4-carboxymethylsterol or 4-methylsterol was detected in either skin or plasma of the patient. To rule out the possibility that the patient’s abnormal sterol pattern developed secondarily to another primary systemic process, we determined sterol profiles in extracts from cultured diploid skin fibroblast cultures from the patient and controls under normal culture condition and in cholesterol-deficient medium (Table 1 and Supplemental Table 4). Methylsterols were indeed markedly elevated in the patient’s fibroblasts, but not any of the controls. Furthermore, the level of methylsterols in patient cells increased when they were grown in cholesterol-depleted medium. The sterol pattern in these experiments had not been previously associated with a human defect in the sterol synthesis pathway, but a review of the literature found it to be consistent with a defect in a methylsterol oxidase previously identified in yeast and plants (9, 10). Supplementation of culture medium with the addition of 1 μM miconazole, a strong inhibitor of lanosterol 14α-demethylase (CYP51, upstream of SMO) in cholesterol-deficient medium, and 5 μM simvastatin, a compound that significantly reduces cholesterologenesis by inhibiting HMG-CoA reductase, reduced methylsterol levels in patient fibroblasts from 2% of total cholesterol to 0.9% and 0.5%, respectively.

Figure 1
Severe scaling and psoriasiform dermatitis in the patient. (A) Note mild microcephaly, lusterless, fine fair hair, and blepharitis. (B and C) The ichthyosiform erythroderma covers all of the patient’s body except for the palms and soles. (D) H&E-stained section of affected skin shows hyperkeratosis (original magnification, ×10), loss of granular layer, psoriasiform hyperplasia, thinning of suprapapillary plate, and neutrophilic epidermal infiltration; these features are characteristic of psoriasis. (E) Oil red O staining of affected skin biopsy (original magnification, ×20) The arrow shows the intracellular lipid accumulation in the foamy cells in the dermis. (F) Neutrophil elastase staining (shown in red) of neutrophils in the stratum corneum of affected skin (original magnification, ×20). Photographs reproduced with signed informed consent/assent provided by the patient and her family.
These data strongly indicated a block at the step of SMO in the cholesterol synthesis pathway. Since there is no validated enzyme assay or substrate available for SMO, we sequenced the SC4MOL gene predicted to encode this enzyme. Two variations from the published sequence (NM_006745) were identified: 519T → A and 731A → G encoding H173Q and Y244C, respectively, in the SMO protein (Figure 2E). The 519T → A mutation was carried by the patient’s father, and the 731A → G mutation was present in the mother. Both positions encode amino acids that occur in highly conserved metal-binding domains in SMO (Supplemental Figure 1), and neither of the variants was identified in any of the databases for common SNPs. In addition, H173Q alters a predicted active site encompassing the second iron-binding motif of the enzyme. Both mutations are predicted to be “probably damaging” by the software package PolyPhen (http://genetics.bwh.harvard.edu/pph/), with a score of 2.5 for the Y244C mutation and 3.2 for the H173Q mutation. Neither mutation was identified in 2,876 alleles from population controls. Nor were they present in the public SNP (http://www.ncbi.nlm.nih.gov/snp) or 1000 Genomes (http://browser.1000genomes.org/index.html) databases. Thus, these variants are exceptionally rare in the human population.

Because slightly elevated methylsterols have been reported in patients with CHILD syndrome (NSDHL deficiency), we also sequenced the cDNA from NSDHL gene, and no mutations were found.

Methylsterol levels were also increased (though less so) in the plasma of the patient’s parents (Table 2), particularly the father, who carries the active site H173Q mutation, suggesting a subclinical effect in the heterozygous state. After 3 months of cholesterol supplementation, the patient’s methylsterol level decreased by about 20%, and the plasma cholesterol level was much improved, reaching a level near the low end of the normal range, but the methylsterol level decreased no further with longer treatment (Table 2).

Cell overproliferation in SMO deficiency. Previous studies of germinial cells have demonstrated meiosis-activating function of MASs, but the effect of MASs on cell cycle regulation and mitosis is unknown (11, 12). The fact that mutations in DHCR14B/LBR, SC4MOL, NSDHL, and EBP or its mouse homologs result in overproliferative skin disease prompted us to examine cell proliferation in skin fibroblasts from our patient (Figure 3, A–C). The rate of
The patient was born with a condition under which de novo cholesterol biosynthesis is stimulated (Figure 3A and ref. 13). Five different normal fibroblast lines with different passage numbers were used as controls. The S-G2-M to G0-G1 ratio in patient skin fibroblasts was 3-fold higher than in control cells (Figure 3B). This ratio peaked after 2-3 days growth in the cholesterol-restricted medium and corresponded with peaks of cellular methylsterols and total protein (Table 2). Thus, cholesterol-restricted medium induced the patient cells to proceed into S-G2-M, implicating an association between MAS production and cell division. To further examine this possibility, we cultured transformed control human lymphoblasts in the presence of a sterol methyl oxidase inhibitor, 3-amino-1,2,4-triazole (ATZ) (14, 15). Under these conditions, the S-G2-M/G0-G1 ratio increased 3-fold in treated lymphoblasts, while neither simvastatin (an inhibitor of HMG-CoA reductase) nor fluconazole (a weak inhibitor of lanosterol 14α demethylase; CYP51) had a significant effect on cell cycle progression (Figure 3C). The stronger inhibitor miconazole was not used here due to its adverse effects on cell growth. These data strongly suggest that alterations in sterol methyl oxidase affect cell cycle activation.

**Immune dysregulation in SMO deficiency.** The patient was born with normal skin, and widespread skin lesions did not develop until she was 6 years old, suggesting that primary barrier dysfunction was unlikely to be the sole cause of her inflammatory skin condition. Many of the defects of cholesterologenesis are associated with immunological manifestations, and our patient’s skin briefly improved with treatment with cyclosporine A; however, she subsequently became resistant to treatment, suggesting that her immune response was atypical compared with other cholesterol pathway disorders and common psoriasis. We therefore examined a panel of immunologic parameters in the patient and her father as described in Methods and Supplemental Table 1. Blood samples from 20 normal individuals were studied as controls. Selected multicolor flow cytometry profiles of granulocytes and T cells are shown in Figure 4A, and the geometric mean fluorescence intensity (GMFI) and percentage of cells for each measurement are shown in Supplemental Table 2. We found that activated CD16+ granulocytes (identified by CD25+CD69+ and CD86+HLA-DR+ subsets) were increased 30- and 20-fold in the patient and her father, respectively, compared with healthy controls (Figure 4A and Supplemental Table 2). Downregulation of CD8 and CD28 and accumulation of CD86+ T cells are indicative of pervasive immune activation in the setting of chronic inflammatory disease or aging (19). It is surprising that the patient’s father, who has normal skin, showed an immunologic profile similar to that of the patient, and that both are strikingly different from all the controls in this study and those seen with other inflammatory diseases (19).

Because psoriasis and psoriatic arthritis are known to be driven by the production of certain cytokines, including the TNF-α and IL-6, the serum cytokine profiles of the patient and father were investigated by multiplex Luminex assay (Supplemental Table 3). Consistent with the flow cytometry data, the levels of a number of proinflammatory and chemokines associated with granulocyte and monocyte

Table 1

<table>
<thead>
<tr>
<th></th>
<th>10% FBS</th>
<th>Cholesterol-depleted</th>
<th>3 d</th>
<th>10% FBS</th>
<th>Cholesterol-depleted</th>
<th>3 d</th>
</tr>
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<tbody>
<tr>
<td>Methylsterols (μg/mg protein)</td>
<td>2.3</td>
<td>1.4</td>
<td></td>
<td>12.3</td>
<td>24</td>
<td></td>
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<tr>
<td>Total cholesterol (%)</td>
<td>0.08 and 0.10A</td>
<td>0.13 and 0.11</td>
<td></td>
<td>0.48 and 0.29</td>
<td>2.0 and 2.0</td>
<td></td>
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<tr>
<td>Total protein (mg)</td>
<td>22</td>
<td>9.2</td>
<td></td>
<td>23</td>
<td>51</td>
<td></td>
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</tbody>
</table>

*Data are from 1 experiment representative of 2 independent experiments. The total protein and cholesterol varied moderately between the 2 experiments, while the methylsterol in percent total cholesterol was consistent between replicates. The average methylsterol in control and patient cells was 0.12% and 2.0%, respectively, in cholesterol-depleted medium; 0.09% and 0.4% in the regular medium.

Table 2

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<tr>
<th></th>
<th>Methylerols (mg/ml)</th>
<th>Total cholesterol (mg/dl)</th>
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<tbody>
<tr>
<td>Normal</td>
<td>2.8–3.2</td>
<td>140–176</td>
</tr>
<tr>
<td>Patient</td>
<td>41 and 42A</td>
<td>85 and 93A</td>
</tr>
<tr>
<td>Patient + cholesterol</td>
<td>32 ± 1.7B</td>
<td>117 ± 12</td>
</tr>
<tr>
<td>Mother</td>
<td>6.4</td>
<td>188</td>
</tr>
<tr>
<td>Father</td>
<td>18.5</td>
<td>231</td>
</tr>
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</table>

*Baseline methylsterol and cholesterol levels in patient are from measurements at 2 time points that were 1 month apart. AMean ± SD of measurement at three time points: 1, 3, and 5 months after start of treatment.
activation were increased in the patient, and to a lesser extent in her father, including GM-CSF, a cytokine reported to trigger psoriatic eruption in humans (20). IL-6 and IL-8 were also elevated in the patient’s serum (Supplemental Table 3). Elevation of IL-6 and IL-8 has been demonstrated in psoriatic skin and fibroblasts and is considered an amplifier of skin inflammation (21). The classical pro-inflammatory cytokine TNF-α was normal in the patient, consistent with the failure of the dermatitis to respond to treatment with TNF-α inhibitor etanercept. Cultures of patient skin fibroblasts showed constitutive production of IL-6 compared with control fibroblasts regardless of passage number (Figure 4D), and IL-6 concentration was further increased with the addition of recombinant TNF-α, a known inducer of IL-6. In agreement with these findings, statins have been reported to suppress Th1-biased cytokine production in both cultured microglia and a mouse model (22).

To investigate a possible association between the defect in cholesterologenesis and cytokine production, we treated control and patient fibroblasts with simvastatin, which reduces cholesterol biosynthesis intermediates including methylsterols (23, 24). After 24 hours, IL-6 production by patient fibroblasts was significantly decreased even in the presence of exogenous TNF-α (Figure 4D), suggesting that the abnormal cytokine production is likely associated with cholesterologenesis.

Reduction of TLR-4 expression induced by SMO inhibition. Activation of TLRs such as TLR4 by microbial ligands inhibits cholesterol efflux by regulating ABCA1 (4), promoting immunocyte proliferation through LXR signaling (25). Of note, ABCA1 is an important transporter for controlling lipid and cholesterol composition in epidermis, critical to maintenance of normal skin barrier function (26). Therefore, we next examined expression of TLR receptors by direct inhibition of SMO in normal leukocytes. Strikingly, by inhibiting SMO in normal leukocytes with ATZ, we found a greater than 6-fold increase in the TLR-2+ TLR-4– granulocyte population (Figure 4C) and 2-fold decrease in CD8 expression in normal lymphocytes (Table 3). Half of the TLR-2+ TLR-4+ and 40% of TLR-2+ TLR-4+ normal granulocytes lost their TLR-4 expression, and the majority of them became TLR-2 positive, suggesting a direct regulating effect of sterol methyl oxidase on TLR receptor expression. TLRs are the connection between innate and adaptive immunities in humans and also the connection between bacterial infection and lipid metabolism in skin.

Discussion
Sterols are important components of cell membranes and lipid rafts, and serve as precursors of many bioactive compounds. SC4MOL is expressed almost ubiquitously in human tissues according to EST database profiles (http://www.ncbi.nlm.nih.gov/UniGene/ESTProfileViewer.cgi?uglist=Hs.105269) and the gene atlas of the human protein–encoding transcriptome in 79 different human tissues from a variety of sources in the GEO database (GDS596; http://www.ncbi.nlm.nih.gov/sites/GDSbrowser). The expression of this gene is highest in liver, brain, adrenal gland, lymphoblast, skin, testis, and vascular tissues. Expression data of SC4MOL in cultured epidermal keratinocytes (GDS2611, GDS2903) and normal leukocytes (GDS3207) can also be found in the GEO database.

A C4 demethylation reaction is found in plants, animals, and yeast, although the structure of sterols and their biosynthetic pathways vary in each group. Previous studies have demonstrated that
only one SMO in yeast is involved in demethylation of both 4,4′-dimethyl and 4α-monomethyl sterols, while two different SMOs exist in plants (9). Since there is only one known SMO in humans (encoded by the SC4MOL gene), we speculate that human SMO will function similarly to its yeast homolog. The sterol intermediates detected in our SMO-deficient patient for the first time to our knowledge provide direct evidence of the role of human SMO in demethylation of 4,4′-dimethyl and 4α-monomethyl sterols. Besides the accumulation of 4,4′-dimethylsterol, the patient accumulated a significantly increased amount of 4α-monomethylsterols, which suggests that both 4,4′-dimethylsterols and 4α-monomethylsterols are substrates for human SMO. No 3-oxo-4α-methylsterols were detected in either plasma or skin of the patient, suggesting a functional difference between human and yeast SMO (27). Thus, it is unlikely that human SMO utilizes 3-oxomethylsterol as a substrate. Rather, in contrast to yeast, 3-oxomethylsterol is reduced to 4α-monomethylsterol by 3-ketosteroid reductase (HSD17B7) before it enters the second round of demethylation in humans (Supplementary Figure 3 and ref. 28). In addition, although 4,4′-dimethylsterol was more abundant than 4α-monomethylsterols in the samples from our patient, only 4α-monomethyl-7-en-sterol accumulated, with no trace of 4,4′-dimethyl-7-en-sterol. These findings suggest that 4α-monomethylsterol, but not the 4,4′-dimethylsterols, can be utilized downstream by Δ8-Δ7 isomerase. Taken together, this pattern indicates that 4α-monomethylsterol may have physiological functions differing from those of 4,4′-dimethylsterol.

Skin cell hyperproliferation causes the flaking skin and characteristic histopathology seen in psoriasiform dermatitis. All known genetic disorders that cause increased levels of MASs present with ichthyosiform or psoriasiform dermatitis in humans and mice, including defects in DHCR14/LBR, NSDHL, and EBP (Supplementary Figure 3 and refs. 1, 5, 7). However, none of the defects adjacent or distal to the MAS demethylation complex, including CYP51, SC5D, or DHCR7, result in skin lesions (24, 29, 30). The present cell proliferation studies revealed a perturbation of cell cycle in cells from our patient, both in cholesterol-restricted medium (which induces increased de novo cholesterol biosynthesis) and with specific inhibition of SMO by ATZ. Interestingly, patients with genetic

**Figure 4**

Immunocyte abnormalities in SMO deficiency. (A and B) Leukocyte populations were analyzed by multicolor flow cytometry; subsets identified by forward/scatter profiles, with mature granulocytes, monocytes, and lymphocytes falling in gates A, B, and C, respectively. The control data shown here represent blood samples donated from 20 normal individuals. (A) Cytometric profiles for CD25, CD69, HLA-DR, TLR-2, and TLR-4 (gate i, and fluorescence histograms of CD16b) and (B) for mature granulocytes/neutrophils. Also shown in A are the CD4, CD8 profiles of CD3+ T cells; and CD28nullCD56+ in CD8dim T cells (gate iv). No significant differences were observed in the monocyte population (data not shown). (C) Overlay of treated and untreated cytometric profiles. Note that 50% of control granulocytes migrated to TLR-2+TLR-4- after the ATZ treatment. (D) IL-6 production by skin fibroblasts from healthy control and patient upon treatment of TNF-α, and TNF-α plus simvastatin, or in medium alone. *P < 0.01. Data are presented as mean ± SD.
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Table 3
Analysis of granulocyte and T cell phenotypes of normal leukocytes treated with ATZ by multicolor flow cytometry

| Cell population | Untreated | ATZ-treated
<table>
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<tr>
<td>Granulocytes</td>
<td></td>
<td></td>
</tr>
<tr>
<td>% CD16(−) (GMFI)</td>
<td>80.6 (57)</td>
<td>60.7 (55.6)</td>
</tr>
<tr>
<td>% CD16(+) (GMFI)</td>
<td>89.4 (48.2)</td>
<td>61.2 (40.3)</td>
</tr>
<tr>
<td>% TLR-2(+) (GMFI)</td>
<td>45.2 (129)</td>
<td>68.0 (397)</td>
</tr>
<tr>
<td>% TLR-4(+) (GMFI)</td>
<td>85.0 (67.8)</td>
<td>46.0 (35.8)</td>
</tr>
<tr>
<td>CD3(+) T cells</td>
<td></td>
<td></td>
</tr>
<tr>
<td>% CDB(+) (GMFI)</td>
<td>26.7 (38.9)</td>
<td>27.7 (26.1)</td>
</tr>
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</table>

*aATZ treatment had no significant effect on cell viability as determined by trypan blue exclusion. *Granulocyte gate iiii in Figure 4D; CD3-CD20-CD16(−) cells. CGMF1 of the indicated antigenic marker; all GMFI values shown have robust coefficients of variation less than 1%. *Lymphocyte gate iii in Figure 4D, and gated for CD3-CD20-CD16(−) cells.

defects in NSDHL, which encodes the enzyme immediately downstream of SMO, occasionally present with clinical features nearly identical to those in individuals with defects in EBP; however, they differ in the histopathology of their skin lesions (1). Outside of the newborn period, NSDHL-deficient patients have psoriasisiform dermatitis, and sometimes verruciform xanthoma–like lesions secondary to the presence of lipid-laden macrophages in the dermis. EBP-deficient individuals have early resolution of their skin disease, with a typical appearance of atrophy but not inflammation on skin biopsy. We have previously reported the presence of elevated methylsterols in skin flanks from patients with common psoriasis (31). Our patient and NSDHL-deficient patients accumulate 4,4′-dimethylsterols in the skin, but EBP-deficient patients do not. This implies that although the MASs may play a role in overproliferation of skin cells and disruption of skin barrier function, 4,4′-dimethylsterols are unique in contributing to perturbation of immune function.

Almost all of the genetic diseases in cholesterol biosynthesis present with some degree of immune dysfunction, including Smith-Lemli-Opitz syndrome (SLOS) and mevalonate kinase (MVK) deficiency (32, 33). Patients with mild mutations in the MVK gene present with hyper-IgD syndrome, periodic fever, and high levels of IgA. Skin lesions, arthritis, lymphadenopathy, and splenomegaly are common in these patients. Previous studies suggest that anomalies in the immune system of SLOS and MVK patients may relate to the accumulation of mevalonate and associated isoprenoid synthesis (34) or abnormal cholesterol composition of lipid rafts in immunocytes (35). However, these studies did not report comprehensive immunological parameters or immunocyte profiles. Thus, our results provide the first evidence to our knowledge that the immunologic abnormalities seen in patients with defects in cholesterologenesis are distinct from other disorders. Since epidermal lipid composition is important for skin barrier function, a defect in de novo cholesterol synthesis that affects epidermal lipid composition could impair barrier function and contribute to dysregulation of immune function. However, the unique immunocyte profile seen in our patient and her father is strikingly different from that usually seen with skin barrier dysfunction, suggesting they are unlikely solely secondary abnormalities. In addition, methylsterols are potent ligands for LXR, while most other sterols are not, and activated LXR binds to TLR-4 promoter through an LXRE site to regulate TLR-4 expression (36). Thus, the reduction of TLR-4 expression in SMO deficiency may be related to inhibition of the LXR by T-MAS, which is structural similar to FF-MAS. Since TLR-4 plays an inhibitory role in both immune response and sterol efflux by ABCA1, the reduction in TLR4 expression could lead to increased lipid transport and cytokine response in the skin in response to bacterial infection, ultimately resulting in the severe psoriasiform dermatitis seen in our patient. Interestingly, the promoter region of ABCA1 also has an LXRE site, which is strongly inhibited by TLR-4 signaling (4). These potential connections imply the unique functions of sterol methyl oxidase in immune function regulation in skin and blood.

Of note, the most profound immunocyte abnormalities in the patient and, to a lesser degree, in her father were found in the granulocytes. Defects in forming mature granulocyte nuclei are also seen in individuals who carry a C14 sterol reductase (lamin B receptor) mutation (37). The appearance of similar granulocyte abnormalities in individuals with defects in adjacent steps of the cholesterol synthesis pathway could be coincidental. However, granulocyte/neutrophil-rich pathology is also characteristic of psoriasis and psoriatic arthritis, and our findings suggest a direct association between demethylayation of MAs and innate immunity (38, 39). It is also intriguing to speculate that the mild elevation of 4,4′-dimethylsterols seen in the patient’s father could be relevant to his history of early-onset inflammatory joint disease. In support of this, two other paternal relatives with histories of early-onset arthritis also carry the 519T→A mutation found in the father (Supplemental Figure 2). Interestingly, it has become evident in recent years that statins suppress several key functions of the immune system and influence the development of autoimmune disease independent of cholesterol reduction (40). Rather, its immune-modulating effects are likely related to the reduction in cholesterologenesis intermediates (41).

In summary, we have identified the first patient to our knowledge deficient in SMO, a novel enzyme in the C4 methylsterol demethylase complex. She had congenital cataracts and developed psoriasiform dermatitis with the presence of lipid-laden cells in the dermis in a pattern reminiscent of CHILD syndrome. Importantly, our patient also had an aberrant TLR expression profile likely induced by the abnormal metabolites accumulating due to the SMO deficiency. Our studies provide evidence that SC4MOL gene function not only is important in de novo sterol synthesis and thus critical for skin barrier function (42), but also regulates cell proliferation and immune regulation. Taken together, the results indicate that defects in SC4MOL gene affect most of the major processes that are critical to the pathogenesis of psoriasiform dermatitis.

Methods
This study was conducted according to an IRB protocol approved by the University of Pittsburgh. Written informed consent was received from participants in the study.

Genomic DNA and cDNA amplification and sequencing. Total RNA and genomic DNA (gDNA) were extracted from patient fibroblasts for reverse transcription, PCR amplification, and sequencing as described previously (43). Primers 5′-CCTTTACCTGGAATCCTCGTG-3′ (forward) and 5′-GCCAGGATTCTCAGGTAAAAG-3′ (reverse) were used for cDNA amplification covering the entire coding region. Primers 5′-ACTTTCATATCTCAGTTGACGG-3′ (forward) and 5′-GAATGAGGAAAGAGCAGTT-3′ (reverse) were used for the PCR amplification of exon 6 to confirm the 731A→G (Y224C) variant, and primer 5′-GGTCAACCTGGAATGTTG-3′ (forward) and primer 5′-TGGTTGTGTGTTGTTATTG-3′ (reverse) were used for the PCR amplification of exon 4 to confirm the 519T→A (H173Q) variant.
Lymphoblasts were cultured with cholesterol-depleted medium as described before. For cell proliferation assays, primary fibroblasts or EBV-transformed human lymphocytes were cultured in DMEM supplemented with 10% FBS, 100 U/ml penicillin, and 100 μg/ml streptomycin; GlutaMAX (Invitrogen), TNF-α (10 ng/ml), and simvastatin (5 μM) were added to the medium before the cells were subjected to the proliferation assay.

Cell proliferation and cell cycle stage analysis. Skin fibroblasts or EBV-transformed lymphoblasts (2 x 10⁵ cells/ml) were labeled with 5- (and 6-)CFSE (0.25 μM) (Molecular Probes, Invitrogen) using established procedures (19). At different days of culture, cells were harvested and stained with DAPI. The fluorescence intensity of both CFSE and DAPI was analyzed by multicolor flow cytometry. The number of cell divisions was determined by the number of CFSE fluorescence peaks. The number of cells in G₁-G₄ and G₂-M phases of the cell cycle was proportional to the amount of linear DAPI fluorescence, G₂-G₄ cells having half the amount of DAPI staining as the S-G₂-M cells.

Statistics. For all experiments with error bars, standard deviation was calculated to indicate the variation within each experiment. Mean differences between groups were examined by pooled Student’s t test (2-tailed). P values less than 0.05 were considered statistically significant.

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4. Liu XY, et al. The gene mutated in bare patches and mevalonate kinase deficiency: differential response to lipid supplied by serum, and skin flakes obtained from patients and controls by gas chromatography and selected-ion mass spectrometry as described previously (7, 44).
5. Flow cytometry on peripheral blood cells. Flow cytometry on cells from patients and controls was carried out and analyzed as described previously (19). In the sterile oxidase inhibition studies, 10 ml red blood cell–depleted whole blood from a healthy 16-year-old girl were incubated with 5 mM ATZ for 60 minutes at 37°C with 5% CO2. Both ATZ-treated and untreated total leukocytes were then isolated and stained as described in Supplemental Table 2.
6. Cell culture, proliferation, and ELISA. Primary dermal fibroblasts were cultured in DMEM supplemented with 10% FBS, 100 U/ml penicillin, and 100 mg/ml streptomycin, GlutaMAX (Invitrogen), TNF-α (10 ng/ml), and simvastatin (5 μM) were added to fibroblasts at 80% confluence; and culture media were refreshed biweekly. The sterol oxidase inhibition studies, 10 ml red blood cell–depleted whole blood from a healthy 16-year-old girl were incubated with 5 mM ATZ for 60 minutes at 37°C with 5% CO2. Upon completion of PCR, plates were read on an ABI PRISM 7900 HT sequence detector. Genotypes were called successfully for 1,438 samples at both variants using Allelic Discrimination Sequence Detection Software (Applied Biosystems) version 2.3 (SDS v2.3).
7. Sterol analyses. Sterol distribution was measured in cultured skin fibroblasts, transformed lymphocytes, serum, and skin flakes obtained from patients and controls by gas chromatography and selected-ion mass spectrometry as described previously (45, 46).
8. Flow cytometry on peripheral blood cells. Flow cytometry on cells from patients and controls was carried out and analyzed as described previously (19). In the sterile oxidase inhibition studies, 10 ml red blood cell–depleted whole blood from a healthy 16-year-old girl were incubated with 5 mM ATZ for 60 minutes at 37°C with 5% CO2. Both ATZ-treated and untreated total leukocytes were then isolated and stained as described in Supplemental Table 2.
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