A New Appraisal of Dermatologic Manifestations of Diabetes Mellitus

George Han, MD, PhD

Diabetes mellitus is a morbid and costly condition that carries a high burden of disease for patients and society as a whole. In dermatology, a great number of skin findings have been associated with diabetes, some demonstrating a stronger connection than others. Importantly, several cutaneous findings have systemic implications in patients with diabetes that should prompt the dermatologist to evaluate the conditions accordingly and, if necessary, communicate with the patient’s primary care physician and other specialists. This column will explore some of the skin conditions associated with diabetes, examine the strengths of these associations, and discuss some potential underlying pathophysiologic mechanisms to provide dermatology residents with a proper framework for approaching patients with diabetes.


Diabetes mellitus is a morbid and costly condition that carries a high burden of disease for patients (both with and without a diagnosis) and for society as a whole. The economic burden of diabetes in the United States recently was estimated at nearly $250 billion annually, and this number continues to rise. The cutaneous manifestations of diabetes are diverse and far-reaching, ranging from benign cosmetic concerns to severe dermatologic conditions. Given the wide range of etiology for diabetes mellitus and its existence on a spectrum of severity, it is perhaps not surprising that some of these entities are the subject of debate (vis-à-vis the strength of association between these skin conditions and diabetes) and can manifest in different forms. However, it is clear that the cutaneous manifestations of diabetes are equally as important to consider and manage as the systemic complications of the disease. In analyzing associations with diabetes, it is important to note that given such a high incidence of diabetes among the general population and its close association with other disease states, such as the metabolic syndrome, studies aimed at determining direct relationships to this entity must be well controlled for confounding factors, which may not even always be possible. Regardless, it is important for dermatologists and dermatology residents to recognize and understand the protean cutaneous manifestations of diabetes mellitus, and this column will explore skin findings that are characteristic of diabetes (Table 1) as well as other dermatoses with a reported but less clear association with diabetes (Table 2).

Skin Findings Characteristic of Diabetes

Diabetic Thick Skin—The association between diabetes and thick skin is well described as either a mobility-limiting affliction of the joints of the hands (cheiroarthropathy) or as an asymptomatic thickening of the skin. It has been estimated that 8% to 36% of patients with insulin-dependent diabetes develop some form of skin thickening; one series also found this association to be true for patients with non–insulin-dependent diabetes mellitus (NIDDM). Skin thickening is readily observable on clinical presentation or ultrasonography, with increasing thickness in many cases associated with long-term disease progression. This increasing thickness...
was shown histopathologically to be a direct result of activated fibroblasts and increased collagen polymerization, with some similar features to progressive systemic sclerosis. Interestingly, even clinically normal skin showed some degree of fibroblast activation in diabetic patients, but collagen fibers in each case were smaller in diameter than those found in progressive systemic sclerosis. This finding clearly has implications on quality of life, as a lack of hand mobility due to the cheiroarthropathy can be severely disabling. Underlining the need for strict glycemic control, it has been suggested that tight control of blood sugar levels can lead to improvement in diabetic thick skin; however, reports of improvement are based on a small sample population.

Huntley papules are localized to areas on the dorsum of the hands overlying the joints, demonstrating hyperkeratosis and enlarged dermal papillae. They also can be found in a minority of patients without diabetes. Interestingly, diabetic thick skin also has been associated with neurologic disorders in diabetes. Diabetic thick skin was found to be significantly (P < .05) correlated with diabetic neuropathy, independent of duration of diabetes, age, or glycosylated hemoglobin levels, though no causal or etiologic link between these entities has been proven.

**Yellow Nails**—Nail changes are well described in diabetes, ranging from periungual telangiectases to complications from infections, such as paronychia; however, a well-recognized finding, especially in elderly diabetic patients, is a characteristic yellowing of the nails, reported to affect up to 40% of patients with diabetes. The mechanism behind it likely includes accumulation of glycation end products, which also has been thought to lead to yellowing of the skin, and vascular impairment. These nails tend to exhibit slow growth, likely resulting from a nail matrix that is poorly supplied with blood, and also can be more curved than normal with longitudinal ridges (onychorrhaxis). It is important, however, not to attribute yellow nails to diabetes without considering other causes of yellow nails, such as onychomycosis, yellow nail syndrome, and yellow nails associated with lymphedema or respiratory tract disease (eg, pleural effusion, bronchiectasis).

**Diabetic Dermopathy**—Colloquially known as shin spots, diabetic dermopathy is perhaps the most common skin finding in this patient population, though it also can occur in up to 1 in 5 individuals without diabetes. Although it is very common, it is not a condition that should be overlooked, as numerous studies have shown an increase in microangiopathic complications such as retinopathy in patients with diabetic dermopathy. Although follow-up studies may be necessary to fully characterize the relationship between shin spots and diabetes, it certainly is reasonable to be more wary of diabetic patients presenting with many shin spots, as the general consensus is that these areas represent postinflammatory hyperpigmentation and cutaneous atrophy in the setting of poor vascular supply, which should prompt analysis of other areas that might be affected by poor vasculature, such as an ophthalmologic examination. Antecedent and perhaps unnoticed trauma has been implicated given a possible underlying neuropathy, but this theory has not been supported by studies.

**Bullosis Diabeticorum**—Bullosis diabeticorum is a rare but well-described occurrence of self-resolving, nonscarring blisters that arise on the extremities of diabetic patients. This entity should be distinguished from other primary autoimmune blistering disorders and from simple mechanobullous lesions. Several types of bullosis diabeticorum have been described, with the most classic form showing an

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**Table 1.**

*Characteristic Skin Findings of Diabetes Mellitus*

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<thead>
<tr>
<th>Skin Finding</th>
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<tr>
<td>Acanthosis nigricans</td>
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<td>Bullosis diabeticorum</td>
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<td>Diabetic dermopathy</td>
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<tr>
<td>Diabetic thick skin</td>
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<tr>
<td>Necrobiosis lipoidica (diabeticorum)</td>
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<td>Scleredema</td>
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<td>Yellow nails</td>
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**Table 2.**

*Dermatoses Associated With Diabetes Mellitus*

<table>
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<th>Dermatosis</th>
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<tr>
<td>Acquired perforating dermatosis</td>
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<td>Eruptive xanthomas</td>
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<td>Granuloma annulare</td>
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<tr>
<td>Lichen planus</td>
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<tr>
<td>Psoriasis</td>
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<td>Vitiligo</td>
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intraepidermal cleavage plane. These lesions tend to resolve in weeks but can be recurrent. The location of the pathology underlines its nonscarring nature, though similar lesions have been reported showing a cleavage plane in the lamina lucida of the dermoeidermal junction, which underlines the confusion in the literature surrounding diabetic bullae. Some may even use this term interchangeably with trauma or friction-induced blisters, which diabetics may be prone to develop due to peripheral neuropathy. Confounding reports have stated there is a correlation between bullosis diabeticorum and neuropathy as well as the acral location of these blisters. Although many authors cite the incidence of bullosis diabeticorum being 0.5%, no population-based studies have confirmed this figure and some have speculated that the actual incidence is higher. In the end, the term bullosis diabeticorum is probably best reserved for a rapidly appearing blister on the extremities of diabetic patients with at most minimal trauma, with a lesion containing sterile fluid and negative immunofluorescence. The mechanism for these blisters is thought to be microangiopathy, with scant blood supply to the skin causing it to be more prone to necrosis. The mechanism for these blisters consists of dark brown plaques in the flexural areas, especially the posterior neck and axillae, is a common finding in diabetic patients and is no doubt familiar to clinicians. The pathophysiology of these lesions has been well studied and is a prototype for the effects of insulin resistance in the skin. In this model, high concentrations of insulin binding to insulin-like growth factor receptor in the skin stimulate keratinocyte proliferation, leading to the clinical appearance and histologic finding of hyperkeratosis and papillomatosis, which in turn is responsible for the observed hyperpigmentation. It is an important finding, especially in those without a known history of diabetes, as it can also signal an underlying endocrinopathy (eg, Cushing syndrome, acromegaly, polycystic ovary syndrome) or malignancy (ie, adenocarcinoma of the gastrointestinal tract). Several distinct mechanisms of insulin resistance have been described, including insulin resistance due to receptor defects, such as those seen with insulin resistance in NIDDM; autoimmune processes; and postreceptor defects in insulin action. Keratolytics and topical retinoids have been used to ameliorate the appearance of these lesions.

Necrobiosis Lipoidea (Diabeticorum)—Necrobiosis lipoidea diabeticorum was first described by Urback in 1932, but reports of similar lesions were described in nondiabetic patients soon after. The dermatologic community has since come to realize that perhaps a more accurate nomenclature is necrobiosis lipoidea to fully encompass this entity. Clinically, lesions appear as erythematous papules and plaques that expand into a larger well-circumscribed plaque with a waxy yellowish atrophic center, often with telangiectases, and usually presenting in the pre-tibial area. Lesions can become ulcerated in up to one-third of cases. Necrobiosis lipoidea also is defined by characteristic histologic findings, including important features such as palisaded granulomas arranged in a tierlike fashion, necrotizing vasculitis, collagen degradation, and panniculitis. Necrobiosis lipoidea is still relatively rare, developing in approximately 0.3% of patients with diabetes, though its relationship with insulin resistance and diabetes is strong. Approximately two-thirds of patients with necrobiosis lipoidea have diabetes and an even higher number go on to develop diabetes or have a positive family history of diabetes.

Although these figures are interesting, the data are nearly a half-century-old, and it is unclear if these findings still hold true today. The etiology of necrobiosis lipoidea also remains elusive, with theories focusing on the role of microangiopathy, immunoglobulin deposition leading to vasculitis, structural abnormalities in collagen or fibroblasts, and trauma; however, the true nature of this condition is likely some combination of these factors. These lesions are difficult to treat, especially at an advanced stage. Management with topical steroids to limit the inflammatory progression of the lesions is the mainstay of therapy.

Scleredema—Scleredema adultorum (Buschke disease) refers to indurated plaques over the posterior neck and upper back. It is usually thought of as 3 distinct forms. The form that is known to occur in diabetic patients is sometimes referred to as scleredema diabeticorum; the other 2 occur as postinfectious, usually Streptococcus, or malignancy-related forms. The prevalence of scleredema diabeticorum among diabetic individuals most frequently is reported as 2.5%; however, it is worth noting that other estimates have been as high as 14%. Although there has been some correlation between poorly controlled NIDDM, treatment and tight glucose control does not seem to readily resolve these lesions with only few conflicting case studies serving as evidence for and against this premise. The lesions often are recalcitrant toward a wide variety of treatment approaches. Histopathologic analysis generally reveals a thickened dermis with large collagen bundles, with clear spaces between the collagen representing mucin and increased numbers of mast cells. Proposed
mechanisms include stimulation of collagen synthesis by fibroblasts and retarded collagen degradation, likely due to excess glucose. 31

**Dermatoses Demonstrating an Association With Diabetes**

**Granuloma Annulare**—Granuloma annulare (GA) is a dermatologic condition existing in numerous forms. The generalized form has been suggested to have some association with diabetes. The lesions of GA are classically round, flesh-colored to erythematous papules arising in the dermis that may start on the dorsal extremities where the localized form typically presents, though larger annular plaques or patches may exist in the generalized form. Histologically, GA has a characteristic granulomatous infiltrate and palisaded granulomatous dermatitis, depending on the stage of the evolution. Many studies dating back to the mid-20th century have attempted to elucidate a link between GA and diabetes, with numerous reports showing conflicting results across their study populations. 32–36 This issue is further muddled by links between generalized GA and a host of other diseases, such as malignancy, thyroid disease, hepatitis, and human immunodeficiency virus infection. The usual course of GA is spontaneous resolution, including a peculiar phenomenon noted in the literature whereby biopsy of one of the lesions led to clearance of other lesions on the body. 37 However, the generalized form may be more difficult to treat, with therapeutic approaches including topical steroids, light therapy, and systemic immunomodulators.

**Lichen Planus**—A recent small population study in Turkey demonstrated a strong relationship between lichen planus and abnormal glucose tolerance. In this study of 30 patients with lichen planus, approximately half (14/30) had abnormal glucose metabolism and a quarter (8/30) had known diabetes, but larger studies are needed to clarify this relationship. 38 Prior to this report, a link between oral lichen planus and diabetes had been shown in larger case series. 39,40 Clinically, one may see white plaques with a characteristic lacy reticular pattern in the mouth. At other cutaneous sites, lichen planus generally appears as pruritic, purple, flat-topped polygonal papules. The clinical finding of lichen planus also is linked with many other disease states, most notably hepatitis C virus, but also thymoma, liver disease, and inflammatory bowel disease, among other associations. 41

**Vitiligo**—As an autoimmune entity, it stands to reason that vitiligo may be seen more commonly associated with insulin-dependent diabetes, which has been shown to hold true in one study, while no association was found between later-onset NIDDM and vitiligo. 42 Given the nonspecific nature of this association and the relatively common presentation of vitiligo, no special consideration is likely needed when examining a patient with vitiligo, but its presence should remind the clinician that these autoimmune entities tend to travel together.

**Acquired Perforating Dermatosis**—Although the classic presentation of acquired perforating dermatosis (Kyrle disease) is linked to renal failure, diabetes also has been connected to its presentation. Extremely rare outside of the setting of chronic renal failure, acquired perforating dermatosis occurs in up to 10% of dialysis patients. 43,44 It is characterized by papules with a central keratin plug, representing transepidermal elimination of keratin, collagen, and other cellular material; its etiology has not been elucidated. The connection between acquired perforating dermatosis and diabetes is not completely clear; it would seem that renal failure is a prerequisite for its presentation. A large proportion of renal failure necessitating hemodialysis occurs in patients with diabetic nephropathy, which may explain the coincidence of diabetes, renal failure, and acquired perforating dermatosis. 45 The presentation of this cutaneous finding should not, however, affect treatment of the underlying conditions. Symptom relief in the form of topical steroids can be used as a first-line treatment of these often pruritic lesions.

**Eruptive Xanthomas**—The link between diabetes and eruptive xanthomas seems to be a rather tenuous one, hinging on the fact that many diabetic patients have abnormalities in carbohydrate and lipid metabolism. A central feature of eruptive xanthomas is an elevation in triglycerides, which can occur in diabetes. Indeed, it has been estimated that only 0.1% of diabetics will develop eruptive xanthomas, 46 and its main importance may be to prompt the physician to treat the hypertriglyceridemia and consider other concerning possibilities such as acute pancreatitis.

**Psoriasis**—Psoriasis is a common dermatologic condition that has been shown to have a far-reaching impact both on patients’ quality of life and cardiovascular risk profiles. Data have emerged linking psoriasis with diabetes as an independent risk factor; although this retrospective study had its limitations, it certainly is interesting to note that patients with psoriasis may have an increased risk for developing diabetes. Perhaps more importantly, though, this study also implied that patients with severe psoriasis may present with diabetes that is more difficult to control, evidenced by increased treatment with systemic therapies as opposed to milder forms of intervention such as diet control. 47 There almost certainly are other confounding factors and further studies would serve to reveal the strength of this association,
but it is certainly an intriguing concept. Echoing these findings, a more recent nationwide study from Denmark demonstrated that psoriasis is associated with increased incidence of new-onset diabetes, adjusting for numerous confounding factors. The relationship between psoriasis and diabetes is worth noting as evidence continues to emerge.

Conclusion
Given the diverse cutaneous manifestations of diabetes, it is important to distinguish those that are directly related to diabetes from those that suggest there may be another underlying process. For example, a new patient presenting to a primary care physician with acanthosis nigricans and yellow nails should immediately trigger a primary care physician with acanthosis nigricans. For example, a new patient presenting to that suggest there may be another underlying process. Conversely, the presence of yellow nails in a patient with chronic diabetes should not be ignored. The physician should consider onychomycosis and query the patient about possible respiratory symptoms. In the case of a multisystem disease such as diabetes, it may be challenging to reconcile seemingly disparate skin findings, but having a framework to approach the cutaneous manifestations of diabetes can help to properly identify and treat an individual patient's afflictions.

REFERENCES


