Kindler Syndrome

First described in 1954 by Theresa Kindler, Kindler syndrome is a rare autosomal recessive genodermatosis characterized by congenital acral skin blistering, photosensitivity, progressive poikiloderma, and diffuse cutaneous atrophy. The syndrome is a combination of features of inherited blistering skin disorders (eg, dystrophic epidermolysis bullosa) and congenital poikilodermas (eg, Rothmund-Thompson syndrome). Kindler syndrome is identified as entry 173650 in the Online Mendelian Inheritance of Man database.

The Medscape Genomic Medicine Resource Center may be of interest.

In 2003, Siegel et al mapped the disease locus to band 20p12.3 by using linkage and homozygosity analysis in an isolated cohort of patients with Kindler syndrome. Loss-of-function
Kindler syndrome = congenital acral skin blistering, photosensitivity, progressive poikiloderma, and diffuse cutaneous atrophy

mutations were identified in the candidate gene

FLJ20116,

which was renamed

KIND1

. This gene encodes a 677–amino acid protein, kindlin-1, which is thought to play a regulatory role in inhibiting oversecretion of basement membrane components by basal keratinocytes at the dermoepidermal junction.

Kindlin-1 is a human homolog of the Caenorhabditis elegans protein UNC-112, a membrane-associated structural/signaling protein that had been implicated in linking the actin cytoskeleton to the extracellular matrix (ECM). Kindler syndrome is the first genodermatosis caused by a defect in actin-ECM linkage rather than keratin-ECM linkage, underlying the pathology of other inherited skin fragility disorders such as epidermolysis bullosa.

Since the first description in 1954 by Theresa Kindler, more than 100 cases of Kindler syndrome have been reported worldwide. A cluster of 26 patients with the syndrome has been identified within a tribe in the Bocas del Toro province on the northwestern Caribbean coast of Panama.4

Morbidity and mortality are mostly related to secondary infections arising from cutaneous bullae and to cosmetic disfigurement.

Persons of any race can be affected.

No sex predilection has been documented.

Patients usually present with the initial skin manifestations during the first year of life.

- The hallmark of Kindler syndrome is congenital blistering and photosensitivity, combined
with progressive poikiloderma and diffuse cutaneous atrophy.
- Both blistering and photosensitivity begin in infancy or early childhood and improve significantly with age. The poikiloderma appears gradually and becomes more prominent later in life. Some patients develop sclerodermoid changes of the fingers and nails.
- Recurrent trauma-induced blister formation occurs primarily on hands and feet, which may prompt an incorrect diagnosis of epidermolysis bullosa.
- Photosensitivity can manifest as increased susceptibility to sunburn.
- Patients with Kindler syndrome frequently have poor dental hygiene and dental problems.

- Diffuse poikiloderma (reticular telangiectasia, patchy hypopigmentation and hyperpigmentation, epidermal atrophy), skin fragility, and atrophic changes (cigarette paper–like wrinkled appearance of the skin) are most prominent in sun-exposed areas, most commonly on the dorsal surfaces of the hands and feet.
- During the neonatal period and early childhood, acral blisters and bullae are seen in trauma-prone areas.
- Dental abnormalities occur commonly in affected persons and include advanced periodontal bone loss, mild-to-severe gingivitis, dental caries, and leukokeratosis of buccal mucosa. In one study, 13 of 18 patients with the syndrome had marked periodontal disease.

- Less common and variable features of the syndrome include pitted or punctate palmoplantar hyperkeratosis (≤65% of individuals), webbing of fingers and toes, nail dystrophy, and, in one case report, abnormal skeletal maturation.
- Mucosal involvement is frequent and leads to urethral, anal, and esophageal stenosis.
- Ophthalmic abnormalities have also been described in some patients and include ectropion, keratoconjunctivitis, and conjunctival scarring.
- Early development of actinic keratoses may occur. Squamous cell carcinoma of the lower lip and transitional cell carcinoma of the bladder have also been reported in one patient with Kindler syndrome.

Kindler syndrome has been shown to result from mutations in the \textit{KIND1} gene on band 20p12.3 (see Pathophysiology). An autosomal recessive pattern of transmission has been reported, but sporadic cases are common, with many originating in consanguineous families. Variable expressivity within families has also been documented. Mutations in the gene encoding type VII collagen (\textit{COLA7A1}) have been excluded, distinguishing Kindler syndrome from dystrophic epidermolysis bullosa.
Treatment is mainly symptomatic and preventative in nature. Patients should be advised to avoid trauma, which helps prevent blister formation. Sun avoidance and photoprotection may prevent or slow the progression of poikiloderma. Good wound care includes the use of topical and systemic antibiotics for infected bullous lesions, which may reduce morbidity.

Telangiectasias can be treated with pulsed-dye laser therapy. Surgical correction of urethral, anal, or esophageal stenosis may be needed.

**Consultations**

- Dermatologist
- Geneticist
- Dentist
- Psychological counselor
- Ophthalmologist and/or surgeon depending on related complications

The role of pharmacotherapy is supportive in nature. Emollients and good blister care reduce morbidity and help prevent complications.