













Dyskeratosis follicularis

| Dyskeratosis follicularis = 000 000000 000000 |
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| Dyskeratosis follicu1aris is a genetic disorder characterized by pathogenetic changes of keratinization. |
| byskeratosis follow rans is a genetic disorder characterized by pathogenetic changes of keratifization. |
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| Dyskeratosis follicularis (Darier's disease) is a ge-netic disorder characterized by the loss of adhesion by |
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| Dyskeratosis follicularis is an autosomal dominant disease caused by mutation in the ATP2A2 gene at |
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| Dyskeratosis follicularis clinically manifests as keratotic papules and histologically as dyskeratosis (4). |
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| testicular agenesis, psoriasis, and multiple cafe-au-lait maculae (1,5,6,11-14) can also be found |
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| in patients with dyskeratosis follicularis. Stress, ultraviolet light expo-sure, heat, sweat, friction, |
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| and oral contraceptives may exacerbate the disease symptoms (2). A case compli-cated by |
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| harman standard to the color harman and | |
| herpes simplex virus has also been reported | |
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| A 28-year-old man from Zagreb, Croatia, was ad-mitted to the Department of Dermatovenerology at the |
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| On his second visit, he had high fever (41 °C) and nu-merous keratotic papules and erosions of the skin |
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| macerations The | toenails were hy | nerkeratotic and | d vel-lowish | |
| maderations. The | tochano were ny | perioratorio ani | a yer lewien . | |
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| Staphylococcus aureus was isolated in hemoculfure and the culture of sklir), slowe atots a specific positions of sklir), slowe atots a specific position of sklir), slowe atots a specific position of sklir). |
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| The patient was treated (caenotielas typpo) th saline and fighneous in ith feathons, tantile and identical in the cate of the |
| The patient was treated patients as a gradual initial as a gradual as a |
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| The case we presented the symbol of the substitution of the substi | á b ureko |
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| One ofthe possible explanations for the infection pro-gressing to sepsis in our patient could lie in impaire |
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| It would be interesting to search for specific muta-tions of the ATP2A2 gene in our patient to see wheth |
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