Schamberg's Disease

Synonyms: progressive pigmented purpuric dermatitis, Gougerot-Blum capillaritis, itching purpura, pigmented purpuric eruption

Schamberg's disease was first described by Schamberg in 1901 in a 15-year-old boy. It represents benign dermatoses with purpura due to leaking from capillaries close to the skin surface. It most often affects the legs and spreads slowly. The discolouration is brown/orange due to haemosiderin deposition in the skin.

Pigmented purpuric reactions have six disease types:[1]

- Progressive pigmentary purpura or Schamberg's disease.
- Pigmented purpuric lichenoid dermatitis of Gougerot and Blum - red/brown papules and plaques in men - which responds to psoralen combined with ultraviolet A (PUVA) treatment.
- Purpura annularis telangiectodes - rare, with a preponderance in young females and manifests as annular erythematous plaques and patches.
- Eczematoid-like purpura of Doucas and Kapetanakis - occurs in men, with bilateral intensely itchy lesions on legs.
- Lichen aureus - a localised persistent form of pigmented purpuric dermatitis.
- Itching purpura of Lowenthal (disseminated pruriginous angiodermatitis) - rare (like Schamberg's disease) but accompanied by itching.

There is clinical and histological overlap between these and they may actually represent variable presentations of the same disease process.

Epidemiology

The condition is rare and there are few epidemiological studies in the literature. One study of patients attending an Indian outpatient clinic over an 18-month period found that there were 100 cases of pigmented purpuric dermatosis in a total of 55,323 patients (0.18%). 95 of these patients had Schamberg's disease. The male-to-female ratio was 3.8:1. The age range was from 11-66 years with a mean of 34 years.[2] It is a common cause of petechiae in children.[3] There has been a case report of four family members with Schamberg's disease, suggesting a possible genetic link.[4]

Aetiology[1, 5, 6]

The underlying cause is not known. However, the following have been postulated:

- Recent viral infection.
- Hypersensitivity to an unknown causal agent.
- Aberrant cell-mediated immunity (perivascular infiltrate has specific types of CD cells only).
- Associated with certain medications - thiamine, aspirin, chlordiazepoxide and paracetamol. It has also been reported with bezafibrate and amlodipine.[7]

Presentation[6]

There are no symptoms apart from itching and patients note their skin looks blotchy. For some this is enough to cause psychological distress. However, some patients have reported pains in their limbs - which may be coincidental.

Lesions are most commonly on the lower limbs bilaterally but can occur anywhere or be unilateral. A case involving the genitals has been reported.

The lesions consist of:

- Asymmetrical brown/orange patches.
- Non-blanchable purpura.
- Petechiae called 'cayenne pepper' spots (develop at the edge of old lesions).

Patterns can vary - eg, annular, linear. There may also be associated lichenification, scaling and pruritic marks.

Differential diagnosis[5, 8]

Other causes of purpura:

- Vasculitis - eg, leukocytoclastic vasculitis.
- T-cell lymphoma (especially if presenting in young males).
Investigations

- Drug eruption.
- Trauma.
- Self-induced purpura.
- Mycosis fungoides.[6]
- Primary benign hypergammaglobulinemic purpura of Waldenström.[10]

Associated diseases[1, 5]

- Diabetes mellitus.
- Rheumatoid arthritis.
- Systemic lupus erythematosus.
- Thyroid abnormalities.
- Hepatic disease, including hepatitis B.
- Porphyria.
- Malignancies.
- Dyslipidaemias.

Management[1, 6]

Any suspected precipitants should be withdrawn.

- Itching - treat with mild topical corticosteroid or antihistamines.
- Good results have been obtained with narrow-band ultraviolet light.
- One study reported the successful use of aminaphtone, a drug normally used in other venous conditions such as chronic venous congestion of the lower limbs.[13]
- Superimposed infection - will need antibiotics.
- Systemic steroids provide some benefits but these are outweighed by the risks of systemic side-effects.
- Advanced fluorescent technology has produced some promising cosmetic results.[14]

Other tried treatments include vitamin C supplements, laser therapy and wearing support hosiery to prevent venous stasis. There is no evidence of definite benefit of the former two. Immunosuppressants have also been used.[5] PUVA therapy has been used and the results look promising.[15]

Prognosis

Schamberg's disease usually runs a chronic course with frequent exacerbations and remissions. The rash may be present for many years with slow extension. Lesions may occasionally disappear spontaneously.

The development of T-cell lymphoma in patients with Schamberg's disease has been reported.[16]

Further reading & references

- Schamberg Disease Photograph; Dermnet Skin Disease Atlas, 2011
- Capillaritis; DermNet NZ
- Klopkov U; DermoScopy and Trichoscopy in Diseases of the Brown Skin, 2013.


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