

[PICTURES IN CLINICAL MEDICINE]

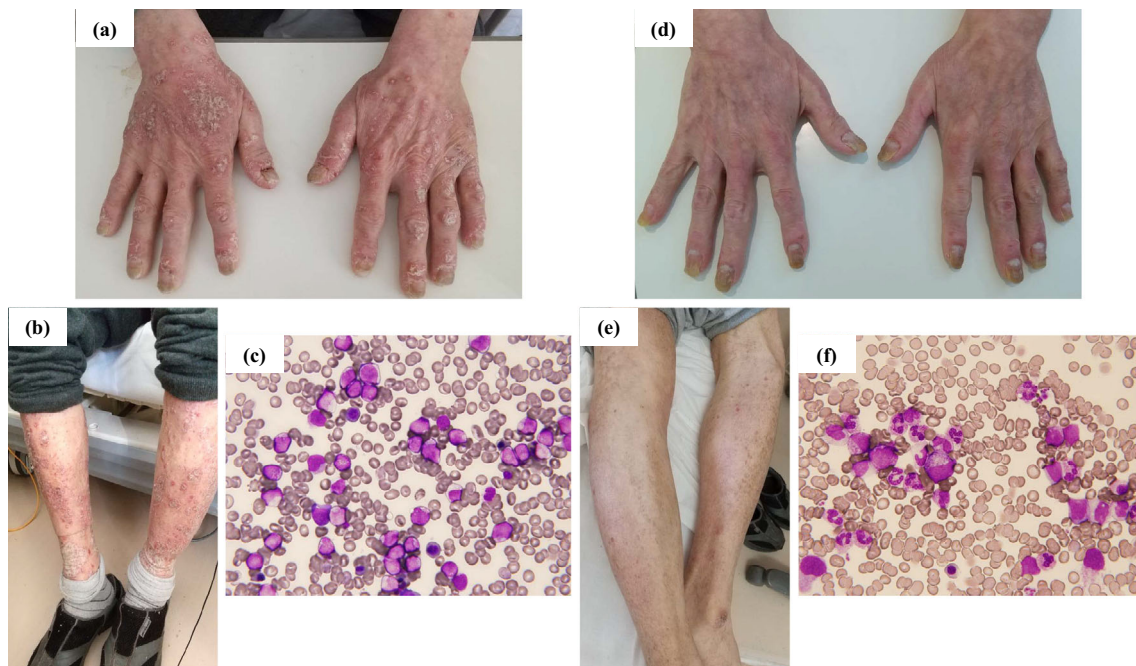
Bazex Syndrome in Acute Myeloid Leukemia

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Key words: Bazex syndrome, psoriasis, acute myeloid leukemia

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Picture.

A 72-year-old man was hospitalized with itching, discoloration, hardening, and exfoliation involving the hands, feet, and back that was refractory to common topical treatment for five months (Picture a, b). A complete blood count test performed at the time of initial presentation showed anemia (hemoglobin 10.3 g/dL). Skin biopsy revealed focal parakeratosis and lymphohistiocytic infiltration of the upper dermis without malignant cells. Laboratory tests revealed leukocytosis with blast cells (29%), anemia, and thrombocytopenia. A bone marrow examination showed >20% blasts with multilineage dysplasia (Picture c). The patient was diagnosed with acute myeloid leukemia with myelodysplasia-related changes and was treated with induction therapy (cytarabine and idarubicin). After induction therapy, the skin le-

sions markedly regressed (Picture d, e) and a bone marrow examination showed complete remission (Picture f). Acrokeratosis paraneoplastica (Bazex syndrome) is a paraneoplastic syndrome. It is most commonly associated with squamous cell carcinoma. Cases associated with acute myeloid leukemia (AML) are rare (1). Cutaneous paraneoplastic lesions usually precede tumor symptoms (2). In the present case, the hyperkeratotic eruption was considered to have occurred in association with the development of AML. When we encounter patients with hyperkeratotic eruptions that are difficult to treat, Bazex syndrome should be considered.

The patient detailed in this case report gave his consent for

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