Case 7

A 35-year-old Thai woman, from Ubonrajchathani

Chief Compliant: Relapsing pruritic vesicles and bullae on both forearms and lower legs for

10 years

Present illness: She had a 10-year history of relapsing pruritic vesicles and bullae which

localized on both forearms and shins. When the lesions resolved, they gradually turned

into hyperkeratotic brownish papules

Past History: Nil Family History: Nil

Physical examination: A healthy looking woman, not pale, no jaundice

HEENT: WNL Heart & Lung: WNL

Abdomen: No hepatosplenomegaly

Skin: Few tense hemorrhagic vesicles with multiple closely-set brownish hyperkeratotic papules

intermingled with hypopigmented macules located on both shins

Investigation

CBC: Hct 38% WBC 7,710/mm3 N 71% L 36% M7% EO 5% PLT 292,000/cumm.

Urine Bence-Jones protein: negative

BUN 12 mg/dl, Cr 0.6 mg/dl

LFT: WNL

Protein electrophoresis: albumin 3.59 g% alpha-1 0.3 g% alpha-2 1.3% Beta 1.3g% gamma 0.2g%

CXR: normal



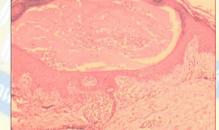




Fig. 7.1

Fig. 7.2 Fig. 7.3

Histopathology (slide no. 41-2762)

- Subepidermal blister with reepithelization
- Small globular deposit of homogeneous amphophilic material at the broadened dermal papillae
- Stallate fibroblasts, melanophages and blood vessels associated with the globules

Diagnosis: Bullous dyschromic amyloidosis

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References

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- 3. Bieber T, Ruzicka T, Linke RP, et al. Hemorrhagic bullous amyloidosis, Arch Dermatol 1988;124:1683-6

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