

Bilateral Klippel-Trenaunay Weber syndrome intricate with digestive bleeding

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Summary

Klippel Trenaunay Weber syndrome or nevus vasculosus hipertrophicus is a vascular malformation characterized by a triad of extensive capillary malformation, underlying venous varicosities, and underlying soft-tissue or bony hypertrophy.

We report a case less commonly of these syndrome in a 20-year old woman with extensive bilateral port-wine stains, which involved preferentially the right-half body, with the hypertrophy of the both right limbs, syndactily and hematochezia suggestive for gastrointestinal bleeding.

Key words: *Klippel-Trenaunay Weber syndrome, digestive bleeding*