WEGENER'S GRANULOMATOSIS: CASE OF EARLY DIAGNOSTICS

I. Yu. Vatanskaya, O. S. Olenitskaya, T. I. Stepanenko, I. P. Strecozova, Yu. O. Vatanskaya

Abstract

Wegener's granulomatosis (WG) — a rare systemic disease of unknown etiology. Histopathologically the disease is characterized by granulomatous necrotizing systemic vasculitis, most often involving the upper and lower airways with further lung damage. Clinical debut of WG in most cases is nonspecific. Timely diagnostics is very complex, leading to a delay of treatment and poor prognosis. The case of GW, which caused considerable difficulties in diagnosis at an early stage, was presented in current report. The diagnosis was established based on the presence of two out of four criteria of American college of rheumatologists (1990): presence of dense infiltrate in the lung and haematuria, as well as positive proteinase-3 anti-neutrophilic cytoplasmatic antibodies (ANCA) test.

Key words: systemic vasculitis, Wegener's granulomatosis, diagnostics.

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Iryna Yu. Vatanska
State Institution "Lugansk State Medical University"
Chair of Internal Medicine
PhD, Assistant Professor
Block of Oborony Luganska, 1≥, 91045, Lugansk, Ukraine
Tel/fax: 380642580979, cell: +380509324645, doktor vatanskaya@ukr.net