[PICTURES IN CLINICAL MEDICINE]

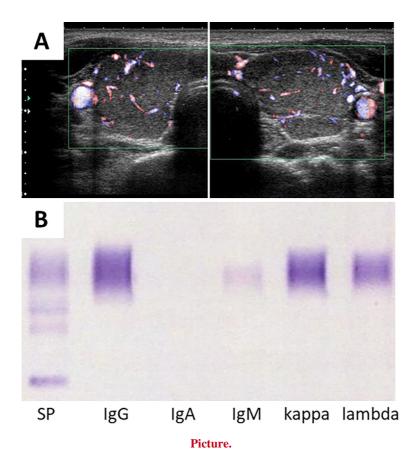
Hyperthyroidism with Selective Immunoglobulin A Deficiency

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Key words: Graves' disease, hyperthyroidism and selective immunoglobulin A deficiency

(Intern Med 58: 145-146, 2019)

(DOI: 10.2169/internalmedicine.1215-18)



A 38-year-old Japanese woman with a history of Graves' disease (GD) presented with a chief complaint of fatigue. The results of examinations showed recurrent empyema, ton-sillitis and respiratory infections. She had no family history of immunodeficiency. Her thyroid was slightly enlarged with increased blood flow by ultrasound (Picture A). Laboratory tests revealed selective IgA deficiency (SIgAD) with recurrent GD as follows: IgA, 2.6 mg/dL (normal: 110-410); IgG, 1,676 mg/dL (861-1747); IgM, 173.9 mg/dL (50-269); IgE,

<10 IU/mL (<180); thyrotropin, 0.005 μ IU/mL; free T3, 12.73 pg/mL; free T4, 3.99 ng/dL; and TSAb, 283% (<120). SIgAD was confirmed by immunofixation electrophoresis (Picture B). The prevalence of SIgAD in Japanese is very low, but patients with SIgAD are at an increased risk for the development of autoimmune diseases, including GD (1). It is possible that SIgAD and GD share predisposing genetic factors or that increased infection due to SIgAD precipitates autoimmune disorders, such as GD (2).

The authors state that they have no Conflict of Interest (COI).

Wang N, Shen N, Vyse TJ, et al. Selective IgA deficiency in autoimmune diseases. Mol Med 17: 1383-1396, 2011.

References

 Abolhassani H, Gharib B, Shahinpour S, et al. Autoimmunity in patients with selective IgA deficiency. J Investig Allergol Clin Immunol 25: 112-119, 2015. The Internal Medicine is an Open Access journal distributed under the Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License. To view the details of this license, please visit (https://creativecommons.org/licenses/by-nc-nd/4.0/).

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