B-cell lymphoma in a tubular adenoma with high-grade dysplasia: a rare extramuraly manifestation of high-grade diffuse large B-cell lymphoma

B-cell lymphomas represent about 90% of all non-Hodgkin lymphomas (NHLs). Almost all express pan-B-cell antigens including CD19, CD20, and PAX5. Most patients with NHL present persistent painless peripheral lymphadenopathy. Staging tests do not include colonoscopy. Here we present a rare case of synchronous occurrence of a high-grade diffuse large B-cell lymphoma (DLBCL) initially presenting in a tubular adenoma of the large bowel and subsequently in the bone marrow. A 73-year-old woman was referred to our hospital because of leukopenia (minimum 0.6 × 10^9/L), anemia (hemoglobin 83 g/L), dyspepsia, and B symptoms. Previously, 27 years earlier, the patient suffered from a pluriform differentiated mucinous breast carcinoma (pT3 pN0 M0) treated with radiation (46 Gy high dose) after mastectomy.

A computed tomography (CT) scan of the abdomen detected few enhanced paraaortal lymph nodes, and routine laboratory analysis showed elevated values for lactate dehydrogenase (maximum 2892 U/L), C-reactive protein (312 mg/L), liver enzymes, ferritin (6334 ng/mL), and CA 12-5 (160 U/mL). Colonoscopy revealed a 2-cm tubular adenomatous sigmoid polyp, which was completely removed by endoscopic resection (Fig. 1).

Histological examination and appropriate immunohistochemical staining showed high-grade intraepithelial neoplasia and focal infiltration of the stroma by highly proliferative lymphoid blasts positive for CD20 and PAX5. Bone marrow immunocytochemistry displayed a monoclonal B-cell lymphocytosis. Subsequent bone marrow biopsy revealed an interstitial and diffuse infiltration by a DLBCL with positivity for CD20, bcl2, and PAX5, negativity for TdT, CD10, and CD34, and a proliferation index of 80% (MIB1 staining). Infiltration by the previously diagnosed breast carcinoma was excluded by negativity for the pan-keratin markers A/E1–3 and MNF116.

Finally, molecular genetic analysis detecting the rearrangement of the FR3a region of the immunoglobulin heavy chain was performed, and identical monoclonal amplificates of approximately 248 base pairs were detected in both manifestations, demonstrating malignancy and clonal association of the lymphoma infiltrates in the adenoma and the bone marrow (Fig. 2).

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Fig. 2  Histochemical and molecular genetic analysis of lymphoma infiltrates in: a, c, e the colonic adenoma; b, d, f bone marrow biopsy. Lymphoid blast infiltrates in the adenomatous stroma (a, circle and inset) and the bone marrow (b) are strongly positive for the B-cell marker CD20 (c and d) and show proliferation rates $>80\%$ with the MIB1 stain (insets c and d). Molecular genetic analysis detecting the rearrangement of the FR3a region of the immunoglobulin heavy chain (IgH) demonstrates monoclonal amplificates for both lymphoma infiltrates, showing monoclonal peaks (e and f) with a so-called slight polyclonal background in the colonic adenoma (e), representing the reactive non-neoplastic B-cells in the surrounding tissue.