Tipo	Periódico
Título	Rare perianal extramammary Paget disease successfully treated using topical Imiquimod therapy
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Programa/Curso (s)	Programa de Pós-Graduação Stricto Sensu em Ciências da Saúde
DOI	10.1186/s12885-018-4815-6
Assunto (palavras chaves)	Perianal Paget's disease; Histological markers; Differential diagnosis; Topical Imiquimod therapy
Idioma	Inglês
Fonte	Título do periódico: BMC Cancer ISSN: 1471-2407 Volume/Número/Paginação/Ano: v. 18, p. 921, 2018
Data da publicação	25 September 2018
Formato da produção	Vários https://doi.org/10.1186/s12885-018-4815-6
Resumo	Background: Perianal Paget's disease (PPD) is a rare intraepithelial adenocarcinoma of the anal margin. Primary PPD likely represents intra-epithelial neoplasm from an apocrine source, whereas secondary disease may represent "pagetoid" spread from an anorectal malignancy. Case presentation: Histologic CDX-2 and CK20 are hallmark markers for colorectal-derived Paget's cells. Interestingly, our primary PPD patient presented both positive and no internal malignancy was identified. In addition, a negative CK7 marker was observed in our case in contrast with previously reported. Surgical excision is the standard treatment; however, previous studies have demonstrated good response with Imiquimod 5% cream in patients with vulval extramammary Paget disease (EMPD). The efficiency of Imiquimod treatment for PPD has not been well described. Our PPD patient was successfully treated using Imiquimod 5% cream. Conclusions: This study describes a primary cutaneous PPD patient CDX-2+/CK20+/CK7- without invasion of the dermis and no associated colorectal carcinoma effectively treated using topical Imiquimod therapy, suggesting that Imiquimod might potentially be considered as a first-line treatment for PPD.
Fomento	