

Fibromatosis agresiva en hueso poplíteo. A propósito de un caso

Aggressive Fibromatosis. Case report

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Resumen. Se presenta un caso de una masa indolora en región posterior de la pierna en un varón de 35 años de crecimiento rápido, consistencia firme y mal delimitada. La resonancia magnética mostraba infiltración de músculos y tejido celular subcutáneo, siendo sugestiva de fibromatosis, tumor neurogénico o sarcoma de partes blandas. La biopsia incisional confirmó el diagnóstico de fibromatosis agresiva. La resección amplia es el tratamiento aconsejado por la mayoría de autores, aunque en algunos casos y debido al carácter infiltrativo de la lesión es difícil resecarla ampliamente. La radioterapia postoperatoria se aconseja en los casos de resección marginal y en los casos de recidiva.

Summary. We present a case of a 35-year-old man who presented a fast growing painless mass localized in the posterior leg, which had at physical examination poorly defined contour and firm consistency. Magnetic resonance imaging showed local infiltration of muscle and subcutaneous cellular tissue, being suggestive of fibromatosis, neurogenic tumor, or soft tissue sarcoma. The incision biopsy confirmed the diagnosis of aggressive fibromatosis. Wide resection is the treatment recommended by most authors, although in some cases and due to the infiltration tendency of the tumor, it is difficult to achieve wide margins. Post-operative radiotherapy is recommended in cases of marginal resection and in case of recurrence.

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Introducción. La fibromatosis agresiva o también llamada desmoide extra-abdominal, se considera un proceso neoplásico benigno de partes blandas, pero localmente agresivo - estadio 3 de Enneking (1,2) - y con tendencia a recidivar después de su resección quirúrgica (2-6). Constituye cerca del 3,5% de todos los tumores de tejidos fibrosos y el 0,03% de todas las neoplasias (7).

Se origina principalmente en el tejido conectivo del músculo y la fascia o aponeurosis suprayacente (fibromatosis musculoa-poneurótica) y afecta principalmente a los músculos de la cintura escapular y pelviana y los muslos en adolescentes y adultos jóvenes.

Este tipo de tumores continua siendo un problema en cuanto al reconocimiento y manejo, debido a la notable discrepancia entre su aspecto microscópico engañosamente inocuo y su potencial de lograr un gran tamaño, recidivar e infiltrar tejidos vecinos en la forma de un fibrosarcoma.

Teniendo en cuenta su escasa frecuencia, consideramos de interés la presentación de un caso tratado por nuestro servicio.

Caso clínico. Paciente de 35 años de edad que acude a consulta ambulatoria de traumatología por presentar una tumoración en cara posterior de pierna derecha, de crecimiento rápido en los últimos tres meses e indolora. En la anamnesis no existía antecedente traumático ni patología previa de interés.

En la exploración se palpaba una masa de consistencia dura, adherida a planos profundos, mal delimitada y de aproximadamente unos 10x4 cm, localizada en hueso poplíteo y cara posterior de pierna derecha. La exploración por aparatos fue normal y la radiografía simple de pierna y rodilla no mostró lesiones óseas ni calcificación intratumoral. Los estudios analíticos no mostraron alteraciones significativas en ninguno de los parámetros evaluados (hemograma, VSG, bioquímica completa).

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Table of Contents Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1

1. Understanding the eBook Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - The Rise of Digital Reading Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Advantages of eBooks Over Traditional Books
2. Identifying Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - User-Friendly Interface
4. Exploring eBook Recommendations from Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Personalized Recommendations
 - Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1 User Reviews and Ratings
 - Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1 and Bestseller Lists
5. Accessing Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1 Free and Paid eBooks
 - Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1 Public Domain eBooks
 - Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1 eBook Subscription Services

- Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1 Budget-Friendly Options
- 6. Navigating Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1 eBook Formats
 - ePub, PDF, MOBI, and More
 - Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1 Compatibility with Devices
 - Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1 Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Highlighting and Note-Taking Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Interactive Elements Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
- 8. Staying Engaged with Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
- 9. Balancing eBooks and Physical Books Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Setting Reading Goals Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Fact-Checking eBook Content of Manejo De La Fibromatosis Musculoaponeur Tica Agresiva 1
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
- 14. Embracing eBook Trends

- Integration of Multimedia Elements
- Interactive and Gamified eBooks

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