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Zusammenfassung. Weichteilsarkome (WTS) und Weichteiltumoren unterschiedlicher Malignität stellen eine heterogene Gruppe von Tumoren dar, die primär in den Weichteilen entstehen und sich wahrscheinlich aus pluripotenten mesenchymalen Zellen entwickeln.

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Zusammenfassung. In diesem Kapitel erfahren Sie mehr über: Fibrosarkom, Liposarkom, Rhabdomyosarkom, Leiomyosarkom, Neurosarkom (MPNST), malignes fibröses Histiozytom (MFH), Angiosarkom, synoviales Sarkom (Synovialsarkom oder malignes Synovialom).

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Weichteilsarkome by Krebsliga Schweiz Issuu

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Weichteilsarkome Springer for Research Development

Sarkome sind bösartige Geschwülste, die ihren Ursprung von den mesenchymalen Geweben nehmen. Diese mesenchymalen Gewebe machen 80-85% der gesamten Masse des Körpergewichts aus.

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Weichteilsarkome der Extremitäten Springer for Research

Zusammenfassung. Die lokale Therapie der Weichteilsarkome hat in den vergangenen Jahrzehnten umfassende Änderungen erfahren. Primäre Amputationen sind funktionserhaltenden Therapiestrategien unter Anwendung plastisch-rekonstruktiver Maßnahmen gewichen.

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Abbatucci JS, Mandard AM, Boulier JC et al. (1990) End results in 130 cases of soft tissue sarcomas of the limbs and trunk walls, treated by a systematic radio-surgical association.

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Zusammenfassung. Bei den Weichteilsarkomen handelt es sich um eine heterogene Gruppe von malignen Tumoren, die von Muskeln, Sehnen, Gefäßen, Lymphgefäßen, Bindegewebe, Faszien und Gelenkbestandteilen abstammen oder aus primitiven mesenchymalen Zellen entstehen.

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ber uns AIO Arbeitsgemeinschaft Internistische Onkologie

The Sarcoma and GIST Working Group of the AIO is focused on the development of novel therapeutics in the field of mesenchymal tumors. Our aim is to improve the treatment of sarcoma patients, which we pursue by testing novel concepts within clinical studies.

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