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**Management Of
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Why a book dedicated to rare cancer management? Given the large variety of rare cancers, they take up a significant part of our daily practice.

Therefore, we owe it to our patients with rare tumours that we use the knowledge gained in the treatment of the more common cancers

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to provide them with a better future.

Management of rare adult tumours: 9782287922459: Medicine ...

Management of rare adult tumours A reference book on rare adult cancer management with the contribution of 70 experts. The knowledge comes from recent developments in molecular

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diagnostics,... Authors have analysed the literature and recommend treatments for rare cancers.

Management of rare adult tumours | Romain Bosc | Springer

René-Olivier Mirimanoff is Head of the Department of Radiation Oncology at the Centre Hospitalier Universitaire Vaudois (CHUV) and the

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University of Lausanne, Switzerland. He is also the founder of the Rare Cancer Network.

(www.rarecancer.net).

His main research interests are brain tumours and rare cancers.

Management of Rare Adult Tumours | SpringerLink

Suprasellar germ cell tumours are rare, and there are few series of patients outlining the

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problems in diagnosis and management, and providing clear guidelines for optimal therapy.

Management of Rare Adult Tumours | Request PDF

Management of Rare Tumours of the Adult is meant to provide a concise summary of the state of the art of rare tumor management for clinicians and medical

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scientists interested in uncommon malignancies. Arranged on the basis of anatomic site of the primary tumours, the authors address the classification, diagnosis, and therapy of 61 rare tumours.

Management of rare adult tumours | Yazid Belkacemi, René ...

Management of rare adult tumours. Why a

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book dedicated to rare cancer management? Given the large variety of rare cancers, they take up a significant part of our daily practice. Therefore, we owe it to our patients with rare tumours that we use the knowledge gained in the treatment of the more common cancers to provide them with a better future.

Management of rare

Page 10/24

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(eBook, 2010)

[WorldCat.org]

Tumours of the vagina
and tumours of the
vulva, C. Haie-Meder
SECTION 6. BREAST1.
Invasive breast cancer
and ductal carcinoma
in situ in young
women, G Gruber, S.
Aebi2.Tubular
carcinoma of the
breast, T. Sullivan, A.
Taghian 3.

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[WorldCat.org]

Rare Tumors Studied
by MyPART. Rare
tumors can form
anywhere in the body.
MyPART is studying
tumors in several
different body systems.
Rare cancers are those
that affect fewer than
40,000 people per year
in the U.S. As a group,
they make up just over
a quarter of all
cancers.

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Rare Tumors Studied by MyPART - National Cancer Institute

management of pineal area tumours in adults. The brain tumour IOG recommended that national tumour groups for rare CNS tumours should be established to coordinate the approach to care; this should include developing protocols for the investigation,

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management,
registration and clinical
research into rare

Rare Brain and CNS Tumours Guidelines

Adult central nervous
system tumor
treatment may include
surgery, radiosurgery,
radiation therapy,
chemotherapy,
surveillance, and
targeted therapy.
Treatment depends on
the tumor type. Learn
more about brain and

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spinal tumor treatment
In this expert-reviewed
summary.

Adult Central Nervous System Tumors Treatment (PDQ ...

Primary cardiac
tumours are rare, with
a necropsy incidence of
0.05%.¹ Secondary
deposits are seen more
frequently, in 1% of
postmortem
examinations, but
usually in the setting of

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widely disseminated malignancy.² The relative incidence of presentation is shown in table 1, and demonstrates that atrial myxoma is by far the most common primary cardiac tumour in adults, and rhabdomyosarcoma is the most common in children.

**Cardiac tumours:
diagnosis and
management | Heart**

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subgroup of primary malignant central nervous system tumours, the commonest malignant CNS tumour in childhood are primitive neuroectodermal tumours (PNETs). Adult PNET is rare and accounts for under 1% of primary CNS malignancy. Whilst exceptional cases of supratentorial PNET have been reported the vast majority of adult

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PNETs are (like the

Rare Brain and CNS Tumours Guidelines

Salivary gland tumours are rare and have very wide histological heterogeneity, thus making it difficult to generate high level evidence. This paper provides recommendations on the assessment and management of patients with cancer originating from the

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salivary glands in the
head and neck.

Management of Salivary Gland Tumours: United Kingdom ...

Granulosa cell tumors
are of two types based
on clinical and
histopathological
characteristics:
Juvenile and adult type.
Juvenile type account
for 90% of the
granulosa cell tumors
that occur in

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prepubertal girls and in women younger than 30 years of age. In the current study, three patients were of juvenile type and all were below 30 years of age.

Ovarian Granulosa Cell Tumor: Clinical Features, Treatment

...

For such rare tumors, randomized trials are hardly feasible and indeed require data

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collection over decades. This important homogeneous series focusing on this rare tumor of the young adult [11, 27., 28.] thus has limitations including missing data for old patient charts and various chemotherapy protocols. It shows, however, a number of ...

Osteosarcomas of
Page 21/24

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**the mandible:
multidisciplinary ...**

For these tumours alternative study designs, robust collection of data through national registries and audits could lead to improvements in the treatment of rare tumours. In addition, concentrating the care of patients with rare tumours into a limited number of centres will help to build expertise,

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facilitate trials and
improve outcomes.

Rare ovarian tumours: Epidemiology, treatment challenges ...

Brain tumours can affect people of any age, including children, although they tend to be more common in older adults. More than 11,000 people are diagnosed with a primary brain tumour

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in the UK each year, of which about half are cancerous.

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