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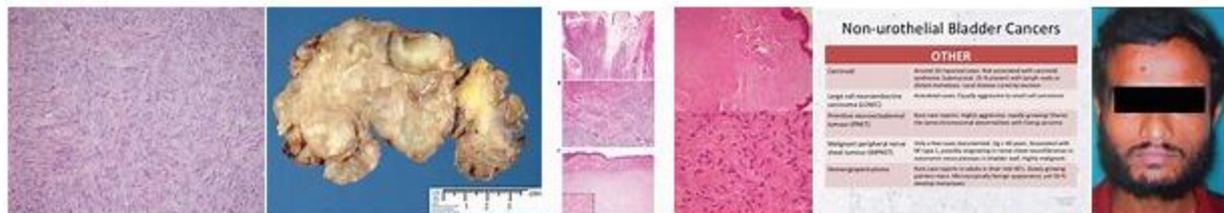
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Published in: IOSR Journal of Dental and Medical Sciences - 2014

Authors: Swapnil Moghe · Moni Gupta · Mrinal Sathpathy · Ajay Pillai · Saksham Nahar

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SPINDLE-CELL SARCOMA OF THE THIGH: A CASE REPORT SAMUEL W. VERNICK, M.D. Assistant Visiting Physician, City Hospital, New York AND JAMES R. LISA, M.D. Pathologist, City Hospital, New York The following case of highly malignant, large **spindle-cell**

Published in: American Journal of Cancer - 1932

Authors: Samuel W Vernick · James R Lisa

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Type I neurofibromatosis with spindle cell sarcoma: A case report

Yu Zhang, Jiao-Jiao Chao, Xiu-Feng Liu, Shu-Kui Qing

Abstract

BACKGROUND

Neurofibromatosis type I is the most frequent subtype of neurofibromatosis. It is ¹related tumor-suppressor syndromes characterized by a predisposition to multiple tumor types and other disorder presentations, what's more, its incidence of tumors is much higher than normal people. However, the reports on it are very few at home and abroad, then we present a case of Neurofibromatosis type I with spindle cell sarcoma.

CASE SUMMARY

A 50-year-old male was found a right axillary mass for 20 years. Specialist examination: coffee milk spots can be seen in many parts of the skin, rounded nodules in the skin; bulge in the right armpit, touching a lump (10 cm × 6 cm), hard, unclear boundary, poor mobility, local tenderness; right The anterior side of the thigh feels weakened on the opposite side; the right groin can touch a swollen lymph node, hard, clear border, good mobility, local tenderness. According to the results of PET/CT, Puncture pathology and immunohistochemistry, Genetic testing, Neurofibromatosis type I with spindle cell sarcoma was confirmed. According to the Genetic testing result, the patient was given a targeted treatment with crizotinib.

CONCLUSION

Surgery, chemotherapy and radiotherapy are the main treatment methods of Neurofibromatosis type I. However, with the continuous progress of molecular biology research, molecular targeted therapy may bring benefits for patients.

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Neurofibromatosis type I (von Recklinghausen's disease): A ...

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3491339>

The term **neurofibromatosis** (NF) is used for a group of genetic disorders that primarily affect the cell growth of neural tissues. **Neurofibromatosis type 1** (NF1), also known as von Recklinghausen's disease, is the most common **type** of NF, and accounts for about 90% of all cases.

Spindle Cell Sarcoma of Thyroid: A Case Report

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4354824>

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Cited by: 2 Author: A. K. Abdul Siyad, Praveen Gopi, N. P. J...

Publish Year: 2014

Metaplastic spindle cell carcinoma of the breast in a ...

<https://onlinelibrary.wiley.com/doi/full/10.1111/tbj.12957>

We report a unique case of a 67-year-old woman with **neurofibromatosis type 1**, who was also diagnosed with **metaplastic** spindle cell carcinoma of the left breast. She had many **neurofibromatosis** lesions on her body, as well as the mass in the left breast.

Cited by: 2 Author: Han Shin Lee, Eun Jung Jung, Ju Yeon Ki...

Publish Year: 2018

Left atrial spindle cell sarcoma: a case report | European ...

<https://academic.oup.com/ehjcr/article/3/1/ytz005/5308399>

Feb 06, 2019 · This case report describes a case of cardiac spindle cell sarcoma diagnosed after surgery, where the initial diagnostic possibilities included an intramural thrombus and a cardiac myxoma. A 57-year-old woman was referred to our hospital for evaluation of ...

Author: Natalia J Braams, Selma E Kaffka Ge... Publish Year: 2019

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Neurofibr...

Medical Condition

Genetic disorder of the



A disorder of the development of ne

Rare (Fewer year in US)

Requires lab

Treatments no known cu

Can last sev

Tumors can form system including nerves. It is caused that are either inherited spontaneously. D symptoms include freckles on the ar loss, loss of vision It cannot be cured control it and mal

Symptoms