Vaginal rhabdomyosarcoma in a patient with Noonan syndrome

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Abstract
This is the first report of a Noonan syndrome patient who has had a vaginal rhabdomyosarcoma. Recent reports of Noonan syndrome patients with leukaemia have prompted speculation that there may be a slightly increased malignancy risk associated with this syndrome.

Noonan syndrome (NS), as first described, comprises a specific group of dysmorphic features without any demonstrable chromosomal alteration. It can be inherited as an autosomal dominant disorder, a gene locus having recently been mapped to chromosome 12.

Recent evidence for a Noonan syndrome patient with leukaemia, prompted speculation that there may be a slightly increased malignancy risk associated with this syndrome. (J Med Genet 1995;32:743-745)

Discussion
Since the original description of Noonan syndrome, there have been occasional reports of NS patients developing malignant disease including malignant schwannoma, phaeochromocytoma, ganglioneuroma, and acute lymphoblastic leukaemia. Some of these malignancies occur with increased frequency in neurofibromatosis type 1 (NF-1), and there are some NF-1 patients who have a Noonan phenotype (so-called NF 1X, neurofibromatosis/Noonan syndrome). Furthermore, patients with Watson syndrome, which is allelic to NF-1, can have some features of NS, notably pulmonary stenosis. However, none of the reports of NS patients who have had malignancies mentions any clinical features suggestive of NF-1 or Watson syndrome.

We report a patient with NS who presented with a vaginal rhabdomyosarcoma which is a previously unreported association and we speculate on the nature of the Noonan syndrome gene as a result.

Case report
A 20 month old girl presented with a three month history of intermittent vaginal bleeding. Four weeks before presentation, the vaginal bleeding became worse and her mother noticed a grape-like mass protruding from the introitus. An abdominal ultrasound scan showed a vaginal mass measuring 1.3 cm x 1.5 cm x 5 cm. CT scanning was consistent with a diagnosis of rhabdomyosarcoma without evidence of metastatic spread. Histology of a biopsy specimen confirmed a botryoid rhabdomyosarcoma. She was treated with chemotherapy to which she had a good response and a repeat biopsy of the vagina after completion of treatment showed no evidence of tumour cells.

At the time of admission, the child was noted to have several dysmorphic features including low set ears, proptosis, hypertelorism, widely spaced nipples, pectus excavatum, and a cardiac murmur owing to mild pulmonary valve stenosis. Her facial appearance is shown in the figure. The features were diagnostic of Noonan syndrome. Neither parent had any signs of NS. She had no cutaneous nor ophthalmological signs of NF-1, although at this age a diagnosis of NF-1 cannot be excluded. Chromosome analysis in the patient was normal. Constitutional and tumour DNA from the patient was analysed for a gross rearrangement within the NF-1 gene using four intragenic polymorphisms in introns 27, 38, 39, and 41. No rearrangement was detected and there was no loss of heterozygosity in the tumour DNA.

Another possibility is that this patient has NF-1 and has yet to develop any cutaneous manifestations. However, we think it more
probable that the diagnosis is Noonan syndrome. Some patients with the NF-1-Noonan syndrome phenotype have had large deletions or rearrangements within the NF-1 gene and we have not been able to find any such change in this patient.

If it is subsequently proven that there is a small increased malignancy risk in Noonan syndrome, this may provide a clue to the function of the Noonan gene. There is evidence that the NF-1 gene, neurofibromin, acts as a tumour suppressor, although the actual contribution of NF-1 mutations to both NF-1 specific and unrelated sporadic tumours still has to be determined. It is possible that one of the small genes embedded within the NF-1 gene may play a role in addition to the tumour suppressor function of neurofibromin itself. Neurofibromin has extensive sequence homology with a family of GTPase activating protein (GAP) related genes which play an intimate role in the regulation of Ras activity. Neurofibromin is likely to accelerate the conversion of active p21^ras^ GTP to inactive p21^ras^ GDP. If there are similarities between NF-1 and NS in the type, if not the frequency, of malignancies reported, it is tempting to speculate that the gene responsible for NS could have similar characteristics, that is, have a tumour suppressor function and be involved in a signal transduction pathway.

It is of considerable interest that the recently isolated gene (FGD-1) for Aarskog syndrome, a disorder that can be confused with NS, proves to be a member of the rho/rac GEF (guanine nucleotide exchange factor) family of proto-oncogenes, which also play a role in cell signal transduction, and that the initial studies indicate that the clinical phenotype of Aarskog syndrome results from loss of function of the FGD-1 gene. If the NS gene also represented a gene involved in a cell signal transduction pathway, this might be consistent with the observed similarities in phenotype with Aarskog syndrome and NF-1.

No increased incidence in malignancy has been reported in Aarskog syndrome and it is uncertain whether there is an increased risk of malignancy in NS although this and recent reports suggest that this is a possibility. Even if subsequently proven, this increased tumour risk would seem to be small given the absence of malignancies reported in various studies of NS patients, including a recent large survey of 151 patients.

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