Phenotypic variation in the familial atypical multiple mole-melanoma syndrome (FAMMM)

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Summary The familial atypical multiple mole-melanoma syndrome (FAMMM) is characterised by an autosomal dominantly inherited susceptibility to multiple atypical moles which show variable colouration ranging from black to brown, tan, red, or pink, with occasional variegation. These compound naevi may be macular or papular, with regular or irregular borders, and measure 1 cm or more in size. They may be few in number or absent or may exceed 100 in a given patient. They are located predominantly on areas not exposed to the sun. Dysplastic changes in melanocytes, fibroplasia, focal chronic inflammatory cell infiltrate, and new blood vessel formation of the papillary dermis characterise their histopathology. These findings are not uniformly present. Because of these distinctive features, coupled with their propensity for transformation to cutaneous malignant melanoma, little attention has been given to the possibility of either minimal or absent cutaneous expression of the phenotype or more diverse neoplastic involvement in this disease. These latter phenomena, which we ascribe to the pleiotropic effects of the cancer-prone FAMMM genotype, were observed in a single FAMMM kindred, the subject of this report.

Variable expressivity of phenotype, including reduced penetrance of the deleterious gene, which is a cardinal feature of dominantly inherited non-cancer diseases, is also frequently observed in cancer genetics. Investigation of virtually all heritable cancer syndromes has shown cancer susceptibility to encompass an array of histological and anatomical sites in any given at risk patient. When studying cancer-prone families, a meticulous assessment of cancer of all anatomical sites is imperative. In early studies of familial breast cancer only site-specific breast cancer was recorded, since the study designs ignored tumours of other anatomical sites. However, it is now clear that genetic heterogeneity accounts for the occurrence of several breast cancer syndromes with differing tumour combinations. Other examples are common in the discipline of cancer genetics.

We postulate that the hereditary variety of malignant melanoma as observed in the FAMMM is more complex than suspected before. For example, intraocular melanoma (IOM) has now been firmly shown to be part of the phenotype of this disease and tumours of other anatomical sites are probably also associated as a result of the pleiotropic effects of the cancer-prone gene. Patients have now been observed with an absence of clinical stigmata of the FAMMM, but who, by virtue of their position in their family pedigree, must have the deleterious genotype. These facts clearly indicate the need for comprehensive clinical-genetic studies of FAMMM kindreds.

The purpose of this report is to document a FAMMM kindred with a wide ranging cutaneous and tumour phenotype.

Family study

The proband (fig 1, III.4) was a 29-year-old white female. A squamous cell carcinoma of the left tonsillar pillar was surgically resected at the age of 28 (fig 2). A synchronous nodular malignant melanoma (CMM, Clark's level II) was excised from the anterior portion of her left forearm and a split-thickness skin graft was performed. This CMM

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showed an absence of atypical moles. A single papular naevus, measuring 0.5 cm, was biopsied and pathological evidence consistent with a FAMMM mole was lacking. Laboratory studies, including chest x-ray, IVP, bone scan, and CT scan of the brain, were all interpreted as normal.

The proband’s father (fig 1, II.3) was a 60-year-old white male. He had contracted malaria during the Second World War while in Africa or Italy after discontinuing atabrine, and he may have had a recurrence of this disease in 1951. At the age of 28, he had had multiple moles (specimens lost) removed from his right shoulder and back. A lesion on his back was found to be a nodular malignant melanoma, Clark’s level IV (confirmed by us histologically).

At the age of 56, a diagnosis of adenocarcinoma of the prostate gland was established and he was treated with radium implants in this gland. He was given diethylstilbestrol and received cobalt treatment over his breasts as prophylaxis. At the age of 58, he underwent a thoracotomy with left upper lobectomy for histologically proven squamous cell carcinoma of the lung. He smoked 30 cigarettes per day from the age of 16 to 47. He drank alcohol temperately.

Physical examination revealed a well-nourished, well-developed white male. He had mild hoarseness. There was a thoracotomy scar on the left, a McBurney’s incision, and a suprapubic surgical incision secondary to the prostate operation. There was a varicocele of the left superior pole of the testis.

The patient did have several atypical naevi which had the clinical appearance of FAMMM moles. Histological findings of these naevi, however, were not characteristic of fully developed FAMMM moles.

The proband’s three children were examined by us (fig 1, IV.1, IV.2, and IV.3). None showed clinical evidence of atypical moles consistent with the FAMMM syndrome. Their history and physical findings were also unremarkable.

Patient III.1 (fig 1) was examined by a dermatologist (HSY) who had clinical experience with other FAMMM syndrome patients. He found the patient to have the classical phenotype of the FAMMM. The histological examination of one of these lesions showed a compound naevus with minimal fibroplasia and chronic inflammatory infiltrate of the papillary dermis. There was no melanocytic dysplasia.

Patient III.5 (fig 1) was examined by a dermatologist (JLS) and had clinical features of the FAMMM. She had several atypical naevi, but also showed many small macular tan-coloured lesions scattered over the trunk. She showed minimal clinical features of the more florid clinical phenotype.

was confirmed by us histologically. Her past medical history was otherwise unremarkable.

She married at the age of 20. Her first pregnancy was at the age of 21 and was followed by two more term pregnancies. Menarche occurred at 13 years of age and she has had regular menstrual cycles. She began birth control pills at the age of 19, but discontinued these after only 9 months. She had not been exposed to therapeutic radiation or significant diagnostic x-ray procedures. She has not had excessive sunlight exposure. She occasionally smokes (never more than two cigarettes per day) and drinks alcohol only temperately.

Physical examination revealed a patient with auburn hair and brown eyes who had a minimal suntanned complexion, even though she resided in the southern United States. The left side of her tongue had been partially resected. There was a surgical scar on her left anterior forearm secondary to removal of the CMM. Examination of the skin
previously reported by us.7 Two atypical naevi were biopsied and histological examination of these specimens showed compound naevi with mild dysplasia of the melanocytes at the dermal-epidermal junction. One specimen showed a mild chronic infiltrate of the papillary dermis.

Pertinent details of the cutaneous findings and pathology evaluations of moles or malignant neoplasms or both in the kindred are contained in table 1. In summary, this is a single FAMMM kindred showing marked variable expressivity in tumour and cutaneous phenotype. There were three relatives who showed variable clinical and histological cutaneous manifestations of the FAMMM. The proband, who did not have atypical moles, showed a remarkable occurrence of synchronous CMM and squamous cell carcinoma of the oral cavity. Another relative had no cutaneous naevi or associated systemic cancer other than malignant melanoma.

Discussion

Our study of a relatively small FAMMM kindred demonstrates several important nuances of this recently described disorder. Most striking is the extent of phenotypic variation. Note that the proband (fig 1, III.4) lacked the cutaneous features of the FAMMM, but developed an isolated CMM and, extraordinarily, in the absence of environmental aetiological factors, she manifested a synchronous intraoral carcinoma. In the absence of a reliable biomarker for the FAMMM genotype, we cannot rule out the possibility that these findings were fortuitous. However, when seen in context with the pedigree, it seems more likely that she has expressed the tumour phenotype of the FAMMM. This patient’s older sister (III.5) showed clinical and pathological features of the FAMMM, but has not yet expressed cancer. Their father (fig 1, II.3) showed clinical cutaneous features of the FAMMM: CMM aged 28, adenocarcinoma of the prostate aged 55, and squamous cell carcinoma of the lung aged 59. His brother (fig 1, II.1) manifested CMM aged 38. As far as we can determine, he has not shown any cutaneous features of FAMMM moles. However, this patient’s son (III.1) shows clinical-pathological features of the FAMMM phenotype.

Histological examination of these compound naevi show variation in the severity of dysplasia of the melanocytes at the dermal-epidermal junction and the amount of fibroplasia and chronic inflammation in the papillary dermis, both within and between patients. In some patients who have the FAMMM phenotype, the naevi histologically show no dysplasia of the melanocytes, but only chronic inflammation and fibroplasia of the papillary dermis. This finding has been demonstrated not only in this family, but in other families studied.8 7

From the detailed study of the FAMMM families, it seems that there are variations in the histological composition of the compound naevi. These variations may occur in different patients and they even occur in different regions of the same patient. This finding of histological variability has been described in other genodermatoses, such as neurofibromatosis and connective tissue naevi. It would not be unreasonable to expect a similar degree of variation in the histopathology of the FAMMM naevi.

A preliminary genetic analysis of this and three other kindreds strongly supports the hypothesis of autosomal dominant inheritance. The rate of penetrance was estimated to be approximately 0.86 and is surprisingly almost independent of age. CMM, atypical moles, and multiple primary cancers were, in that order, the best criteria of the affected subjects. This family shows the typical variation in expression observed in the other kindreds. It also displays what we believe to be another important aspect of the FAMMM syndrome. As a disease of the young and middle-aged adult, it is manifest by CMM or atypical moles or both. As a disease of the more elderly, it is manifest as multiple primary cancers usually, although not necessarily, with a history of CMM and atypical moles (fig 1, II.3). These cancers include intraocular melanoma (IOM) and carcinoma of the breast and lung. Proof of this progression of the natural history of the FAMMM diagnosis will require the examination of many more such families.

The study of a single kindred poses serious limitations for the delineation of genetics and phenotype since only a given number of at-risk patients can be ascertained. Further confounding this issue is the possibility of genetic heterogeneity

<table>
<thead>
<tr>
<th>Pedigree No</th>
<th>Clinical FAMMM phenotype</th>
<th>FAMMM histology</th>
<th>Malignant melanoma</th>
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<tbody>
<tr>
<td>I.1</td>
<td>-</td>
<td>-</td>
<td>+</td>
</tr>
<tr>
<td>I.2</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>I.3+</td>
<td>+</td>
<td>+</td>
<td>-</td>
</tr>
<tr>
<td>III.1</td>
<td>-</td>
<td>-</td>
<td>+</td>
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<td>III.2</td>
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<tr>
<td>III.3</td>
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<td>ND</td>
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<tr>
<td>III.4+</td>
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<tr>
<td>III.6</td>
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<td>ND</td>
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<tr>
<td>IV.1</td>
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<tr>
<td>IV.2</td>
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<td>ND</td>
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<tr>
<td>IV.3</td>
<td>-</td>
<td>ND</td>
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ND = not done.
*Adenocarcinoma of prostate, squamous cell carcinoma of lung.
†Squamous cell carcinoma of tonsil.
and the absence of reliable biomarker(s). Several questions therefore must be considered with respect to the FAMMM. (1) Is the dominant gene causing the FAMMM influenced by associated genes, chromosomal position effect, or specific environmental interactions? (2) Are there differing hereditary forms (genetic heterogeneity)? (3) What is the full range of tumour spectrum in the FAMMM? (4) Are there sporadic forms, so called phenocopies, of the FAMMM? (5) How consistent is the expression of the histological features of the FAMMM mole in these patients and is it always the precursor of malignant melanoma (that is, does CMM develop de novo in the kindred)? With respect to extracutaneous tumours, we have recently studied a family from Holland in which a patient showed cutaneous features of the FAMMM but, in addition, he manifested bilateral IOM and multiple CMMs. Cancer of diverse anatomical sites (lung, breast), including CMMs and cutaneous features of the FAMMM, occurred throughout this Dutch kindred.

Sporadic occurrences of the FAMMM have been reported. However, painstaking clinical and pathology studies of all available high risk relatives of the probands were not described. We have shown that genetic historical data are consistently missed by physicians in the appraisal of cancer-prone families. Until such data are available, we would prefer to withhold judgment on the status of so-called sporadic occurrences of the FAMMM. For example, when reassessing two allegedly sporadic cases, Elder et al. found them to have positive family histories. Our family could very well have fitted into the sporadic category had we seen the proband’s father in isolation. In turn, the proband would not have been considered to manifest the FAMMM had we not had an opportunity to study her unusual cancer occurrences together with the clinical-pathological findings in her relatives. The fact of reduced penetrance and variable expression further confounds the issue. Both parents will be unaffected 15% of the time. Consequently, one must evaluate carefully the extended kindred, including grandparents, aunts, uncles, and cousins. Indeed, experience with other dominant disorders showing similar degrees of variation in expression and penetrance has taught the geneticist to use caution in the use of the term sporadic.

It would be premature at this time to split the FAMMM into two formal sub-types, one familial and the other sporadic, and therefore to imply two different aetiologies. A priori, we would expect a proportion, if not all, of sporadic cases to be the result of new mutations at the FAMMM locus. Their familial nature may not be recognised until the next generation. The concept of a FAMMM phenoctype with different pathogenesis can only be considered a remote possibility, and proof of the phenoctype hypothesis must be supported by rigorous histopathological and family studies.

Experience in cancer genetics has clearly indicated that many cellular types and differing organ and tissue sites may be inordinately susceptible to malignant transformation. An example of variable tumour spectra in several cancer-associated genodermatoses, each of which is believed to be under the control of a single gene, is provided in table 2.

Models at the animal level, such as the Sinclair miniature swine colony, when observed, should be

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**TABLE 2** Representative tumour associations in several cancer-associated genodermatoses.

<table>
<thead>
<tr>
<th>Autosomal recessive</th>
<th>Autosomal dominant</th>
<th>X linked recessive</th>
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<tbody>
<tr>
<td>Ataxia telangiectasia (Louis-Bar syndrome)</td>
<td>Multiple nevoid basal cell carcinoma syndrome</td>
<td>Wiskott-Aldrich syndrome</td>
</tr>
<tr>
<td>Lymphocytic leukaemia, Hodgkin’s disease, reticulum cell sarcoma, lymphosarcoma and other lymphoma, possible gliomas, medulloblastoma, and gastric carcinoma</td>
<td>Multiple basal cell carcinoma, medulloblastoma, astrocytoma, ameloblastoma, ovarian carcinoma</td>
<td>Lymphoreticular malignancies, malignant lymphoma, myelogenous leukaemia, astrocytoma</td>
</tr>
<tr>
<td>Bloom’s syndrome</td>
<td>Cowden’s disease (multiple hamartoma syndrome)</td>
<td>Dyskeratosis congenita (Zinsser-Cole-Engman syndrome)</td>
</tr>
<tr>
<td>Leukaemia, lymphomas, squamous cell carcinoma of the oesophagus, adeno-carcinoma of the colon</td>
<td>Carcinoma of breast, colon, and thyroid</td>
<td>Squamous cell carcinoma of oral cavity, oesophagus, nasopharynx, skin, anus, cervix, and adeno-carcinoma of rectum</td>
</tr>
<tr>
<td>Fanconi pancytopenia (constitutional infantile pancytopeny)</td>
<td>Von Recklinghausen’s neurofibromatosis</td>
<td>Gardner’s syndrome (intestinal polyposis III)</td>
</tr>
<tr>
<td>Osteogenic sarcoma, hepatoma, adeno-carcinoma of the bile duct, breast, thyroid, malignant melanoma, squamous cell carcinoma, meningiomas</td>
<td>Sarcomas, intracranial and optic nerve gliomas, acoustic neurinoma, optic neurina, meningiomas, and phaeochromocytoma</td>
<td>Adenocarcinoma of colon, periampullary carcinoma, thyroid carcinoma, adrenal cortical carcinoma, sarcomas</td>
</tr>
</tbody>
</table>
maximally exploited.\textsuperscript{14} We have noted evidence of similarities in the histology of naevi from these swine with that from our FAMMM patients.

Experience is limited with regard to the surveillance and management of the FAMMM syndrome. Findings in our kindred, as well as those from reviewing published reports, clearly show the need for educating both the patient and the physician about the natural history of this disease. Knowledge of the marked variable expressivity of phenotype, including the risk for extracutaneous cancers, must be employed in cancer control strategy.

The risk for the development of isolated de novo malignant melanomas without pre-existing naevi, which differs from the view of Elder et al\textsuperscript{10} who feel that the atypical mole is invariably the precursor of CMM, must be studied.

The advent of new technology in television and computer image storage/retrieval should in the future enable a clinical comparative review of the entire cutaneous surface with the capability of image intensification and magnification of small areas. This type of surveillance will necessarily be restricted to centres with the capability for this technology as opposed to the typical clinical office setting. This surveillance mode will be cost effective only in at risk family members. The recognition of clinical changes would then necessitate prompt histological examination.

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References
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