Familial testicular cancer: a report of the UK family register, estimation of risk and an HLA Class I sib-pair analysis

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Summary Forty-two families with two or more cases of testicular cancer have been reported to the UK Register for Familial Testicular Cancer, comprising two pairs of identical twins, 27 sets of other brothers (25 pairs, two triples), nine father-son pairs, two pairs of first cousins and two uncle-nephew pairs. In total 91 testicular tumours are described in 86 individuals (42 (46%) pure seminoma, 49 (54%) other germ cell tumours). The median age at diagnosis in these patients was significantly younger than that in a comparable series of non-familial patients (29 cf. 32.5 years, P<0.01). In a case-control comparison of 794 testicular cancer patients, eight patients (1.0%) had a brother and four patients (0.5%) had a father with a previous diagnosis of testicular cancer at the time of their own diagnosis (and these families are all included in this report). Two out of 794 controls (0.3%) had a first degree relative with testicular cancer. The cumulative risk to a brother of a patient for developing testicular cancer by the age of 50 years was estimated to be 2.2% (95% C.I. 0.6-3.8%) which results in a relative risk of 9.8 (95% C.I. 2.8-16.7) in comparison to the general population. HLA Class I typing of 21 affected sib-pairs demonstrated four (19%) sharing two haplotypes, 13 pairs (62%) sharing one and four pairs (19%) sharing none. This did not differ significantly from the expected proportions of 25%/50%/25%. It is unlikely, therefore, that there is a major gene associated with testicular cancer predisposition within or closely linked to the major histocompatibility gene complex on chromosome 6.

There are numerous reports in the literature of families where two or more first degree relatives have been diagnosed with germ cell tumours of the testis. To date at least 21 pairs of identical twins, 82 sets of brothers (other than identical twins) and 31 father-son pairs have been reported (Weissbach & Widman, 1986; Dieckmann et al., 1987 and references therein; Dieckmann & Keyserlingk, 1989 and references therein; Goss & Bulbul, 1990; Patel et al., 1990). Only one attempt has been made to estimate the risk of testicular cancer in men who have an affected first degree relative. Tollerud et al. (1985), using data derived from a study of 225 men with testicular cancer, calculated that having a first degree relative with testicular cancer was associated with a 6-fold elevated risk in comparison with the general population. There has been relatively little research into whether the excess in familial cases occurs as a result of a genetic predisposition, common environment or both (Gedde-Dahl et al., 1985; Dieckmann et al., 1987; Forman, 1989; Oliver, 1990).

We have established a UK-based register for familial testicular cancer to provide a means for the systematic documentation of new cases, including histological verification, and for obtaining standardised lymphocyte-DNA samples from affected and unaffected family members for subsequent genetic linkage analysis. In this paper we describe data on the first 42 families reported to the register for which confirmation of the diagnosis has been obtained. A sub-set of these families were identified from interviews about family history with men diagnosed as having testicular cancer for whom an age-matched control was also interviewed. Using these families, it was possible to estimate the risk of developing testicular cancer if a first-degree relative had been previously affected.

There have been a number of studies (De Wolf et al., 1979; Majsy, 1979; Pollack et al., 1982; Oliver et al., 1986a,b; Dieckmann & Keyserlingk, 1988; Kratzik et al., 1989) which show that patients with testicular cancer have a different distribution of histocompatibility antigens (HLA) compared with control populations. Because of this, it has been suggested that disease susceptibility may be partly influenced by a gene within, or closely linked to, the HLA complex on chromosome 6 (Oliver et al., 1986a,b). We have, therefore, conducted a sib-pair analysis (Haseeman & Elston, 1972; Thomson & Bodmer, 1976) for HLA haplotype association on those families from whom appropriate blood samples could be obtained. If there was a linked gene in the HLA region, a much stronger association should be observed within families than occurs when sporadic cases are typed (Bodmer, 1982).

Methods

Case ascertainment

Familial cases of testicular cancer were mainly identified as a result of a general mailing to consultants in oncology and radiotherapy throughout the UK. Subsequently, other, further families were notified to the register on an ad-hoc basis. Another source of cases arose from structured interviews with men participating in a national population-based case-control study of testicular cancer (full details of which will be published elsewhere). These men had been diagnosed with testicular cancer between 1984 and 1986 and were aged 15 to 49 years at diagnosis. All men within this age range who were resident in any of eight predefined geographical areas (each within a different Regional Health Authority) were eligible for entry into the study, with the exception of those in non-white Caucasian racial groups and those mentally unable to participate in an interview. In total 863 men were eligible for entry, of whom 794 (92%) completed an interview. One part of the interview was concerned with
diagnosis of cancer in male relatives.

When a potential testicular cancer family had been identified, full details were recorded about the cases and a family pedigree was constructed. Usually this process involved interviews with the case and with other family members. Confirmation of the diagnosis in all suspected cases was accomplished by obtaining copies of histopathology reports from the hospitals where treatment took place. As it was not intended, because of small numbers, to look at histological sub-types of testicular cancer in fine detail, a secondary review of pathology was not undertaken. In a few cases, full histopathology reports were unavailable and in these instances supplementary clinical details were obtained to confirm the diagnosis. Having established the diagnosis and obtained other relevant details, efforts were then made to obtain blood samples from the affected family members and then from other informative relatives in the pedigree.

Relative risk estimation

The prevalence of testicular cancer among first-degree relatives (i.e. fathers and brothers) of the 794 cases participating in the case-control study was used to estimate the size of familial risk. An initial comparison was made with the prevalence of testicular cancer among the 794 control men included in the same study. These men were selected at random from the list of the general practitioner with whom the case was registered and were matched (± 1 year) for age. Any reports of relatives with testicular cancer were followed up and had to be confirmed by a histopathology report in order to be included in the comparison. A matched case-control analysis (Breslow & Day, 1980) to estimate relative risk was then carried out. Case-control analysis is, however, extremely imprecise. This is partly because of the small number of affected relatives, especially in the control group (see Results), and partly because it does not fully take account of the number of brothers at risk or the length of time for which they are at risk. An additional estimate of the risk to brothers was, therefore, obtained by actuarial analysis (Peto et al., 1976). Person years were calculated for all brothers of cases from the date of birth to the date of interview of the case or to prior death of the brother. This then enabled the calculation of a cumulative risk for a brother developing testicular cancer by a given age. This risk was then compared with the equivalent population risk calculated from England and Wales National Cancer Registration Scheme data for the year 1985 (OPCS, 1990).

Blood preparation and HLA typing

Blood (25 ml) was taken into bottles containing 0.33% Trisodium Citrate and 0.5 μM Mercaptoethanol, in RPMI 1640 Hepes buffered medium. Lymphocytes for HLA typing were prepared by centrifugation over a Lymphoprep (Nyegaard) density gradient, and HLA typed according to the method of Bodmer and Bodmer (1979), using a panel of locally available antisera.

HLA types were assigned by HLA typing of parents and sibs and looking at the patterns of antigen inheritance. This allowed for the determination of haplotype sharing between affected sibs. In some cases it was not possible to estimate unambiguously the number of shared haplotypes because either one parent was homozygous or one parent was not tested and could potentially have been homozygous. The observed distribution of haplotype sharing was compared with that expected on the basis of mendelian segregation and tested with a chi-square statistic (Thomson & Bodmer, 1976). Where there was ambiguity about the number of shared haplotypes a proportion of maximum sharing was made solely for the purposes of this analysis. One sibship, for which the HLA typing has been reported previously (and partly gave rise to the hypothesis under consideration), was excluded from the initial analysis.

Reported identical twins were checked for monozygosity by southern blotting with mini-satellite probes (R. Hawkins, personal communication).

Results

Description of families

In total, 42 families with two or more members diagnosed with testicular cancer had been reported to us or identified by us by March 1991. Details of the cases in these families are listed in Table I. Three of these families (nos. 10, 39, 42) have been previously reported (Oliver et al., 1986b). Table I shows, for each case, the age at diagnosis, the side and histological classification of the tumour and whether or not the patient had a history of undescended testis (UDT). Twelve families that were identified through the national case-control study, and were used in the risk estimates, are also indicated.

In Table I, there are two pairs of identical twins, 27 sets of other brothers (including one pair of half brothers), nine father-son pairs, two pairs of first cousins and two uncle-nephew pairs. There are four families where one brother of a pair has had bilateral testicular cancer (family nos. 1, 2, 19, 20) and there is one such father (family no. 31). There are also two sibships of three affected brothers (family nos. 7, 10).

There are 91 tumours described in Table I present in 86 individuals in the 42 families, five patients having had bilateral disease. Of the 91 tumours, 42 have been categorised by local pathologists as pure seminoma, two as teratoma differentiated, 20 as malignant teratoma intermediate, 16 as malignant teratoma undifferentiated, seven have combined seminoma and teratoma histologies and four could only be classified as malignant teratoma without further specification. Altogether, there were 45 right-sided and 41 left-sided tumours and five unilateral tumours where the side was not recorded.

Table II summarises the histological classification into two groups, pure seminoma and other germ cell tumours, for the 86 patients and shows the median age at diagnosis for each group. For comparison purposes, similar data are presented for the 781 patients in the case-control study without a family history. The median age at initial diagnosis for the 86 familial cases was 29 years (range 16–51 years). For patients with seminoma, the median age was 32.5 years (range 20–51 years), while for patients with other germ cell tumours it was 26 years (range 16–47 years).

The median age at diagnosis of the 781 non-familial patients, who were all aged between 15 and 49 years, was 32.5 years overall and 35.5 and 28.5 years for patients with seminoma and other germ cell tumours respectively. For both histological categories, therefore, the median age at diagnosis was younger in the familial patients. This difference was statistically significant for those with seminoma and for all histologies combined (P < 0.05 and <0.01 respectively, Mann-Whitney test).

The median age at diagnosis of the first tumour in the five familial patients with bilateral cancer was 26.5 years (range 20–33 years) which was younger than in those with unilateral cancer (29.5 years) although the difference was not statistically significant. Two of these five patients with an initial seminoma were diagnosed at 32 and 33 years while three with a teratoma were diagnosed at 20, 22 and 26 years.

In the father-son pairs, the median age at diagnosis in the fathers (34.3 years) was significantly (P < 0.01, Mann-Whitney test) older than in the sons (24.5 years) with a mean difference of 13.2 years (range 3–22 years). There were six seminomas in the fathers compared to five in the sons. In the non-twin brother pairs there was no tendency for the elder brother to have a later age at diagnosis than the younger brother and the mean difference in age at diagnosis between all sibships (including twins) was 5.5 years (range 1–20 years).

Information about a prior history of UDT was available for 79 individuals and, of these, eight (10.1%) had such a diagnosis which was bilateral in three cases. In four of the five unilateral cases the cancer was diagnosed in the ipsilateral testis. In no family was there more than one case of UDT. All the unilateral cases of UDT underwent orchido-
Table I  Summary of confirmed familial cases of testicular cancer on UK register

| No. | Age at diagnosis | Side | Histology | UDT* (Age at orchidectomy) | Age at diagnosis | Side | Histology | UDT* (Age at orchidectomy) | Notes |
|-----|------------------|------|-----------|-----------------------------|------------------|------|-----------|-----------------------------|-------|
|     |                  |      |           |                             |                  |      |           |                             |       |
| **Identical twins** |     |                  |           |                             |                  |      |           |                             |       |
| 1   | 22               | R    | MTI       | L & R (untreated)           | 20 & 22          | R & L | MTU & MT  |                             | Case 1 & 2: minor congenital abnormalities |
| 2a  | 32               | R    | MTU (+YS) |                             | 26 & 33          | L & R | MTI & S   |                             | Case 1: adult hernia |
|     |                  |      |           |                             |                  |      |           |                             |       |
| **Brothers (other than identical twins)** |     |                  |           |                             |                  |      |           |                             |       |
| 3   | 23               | L    | S         |                             | 26               | R    | MTU       | L (3 years)                 | Mother: breast cancer (51 years) |
|     |                  |      |           |                             |                  |      |           |                             | Father: sarcoma (44 years) |
|     |                  |      |           |                             |                  |      |           |                             | Father and paternal grandfather: adult hernia |
|     |                  |      |           |                             |                  |      |           |                             | Case 1: infantile hernia/hydrocoele |
|     |                  |      |           |                             |                  |      |           |                             | Mother: oesophageal cancer (66 years) |
|     |                  |      |           |                             |                  |      |           |                             | Father: 'facial' cancer (40 years) |
|     |                  |      |           |                             |                  |      |           |                             | Brother and uncle: adult hernia |
|     |                  |      |           |                             |                  |      |           |                             | Uncle: hydrocoele |
| 5a  | 20               | L    | MTI       |                             | 23               | R    | MTU (+YS) |                             | Father: stomach cancer (63 years) |
| 6   | 34               | L    | S         |                             | 30               | L    | S         |                             | 2nd cousin: testicular cancer (age n/k) |
| 7a  | 25               | R    | MTI       |                             | 28               | R    | MTU (+YS) |                             | Father: hydrocoele, multiple myeloma (50 yrs), glioma (48 yrs) |
| 8a  | 26               | R    | S         |                             | 24               | L    | S         |                             | Case 2: infantile hernia |
| 9a  | 19               | R    | MTU       |                             | 22               | R    | MTI       |                             | 4th brother: hypogonadal |
| 10a | 29               | R    | MTI       |                             | 20               | L    | S         |                             | 5th brother: infertile |
|     |                  |      |           |                             |                  |      |           |                             | Grandfather: hernia (age n/k) |
|     |                  |      |           |                             |                  |      |           |                             | Case 1: adult hernia |
|     |                  |      |           |                             |                  |      |           |                             | Father: adult hernia |
| 11  | 23               | R    | MTI       |                             | 18               | L    | MTI (+YS) |                             | Mother: breast cancer (64 years) |
|     |                  |      |           |                             |                  |      |           |                             | Sister: lung cancer (44 years) |
| 12  | 31               | R    | MTI (+CC/YS) |                         | 26               | R    | S         |                             | Case 2: infantile hernia |
| 13a | 31               | R    | MTU       |                             | 37               | R    | S         |                             | Father: mumps orchitis |
| 14  | 40               | L    | S/MTI     |                             | 25               | L    | MTU       | L (12 yrs)                 | Uncle: adult hernia |
| 15  | 39               | L    | S         |                             | 36               | L    | S         |                             | Mother: colon cancer (89 yrs) |
|     |                  |      |           |                             |                  |      |           |                             | Sister: Down's syndrome |
|     |                  |      |           |                             |                  |      |           |                             | Case 2's son: testicular torsion |
| 16  | 37               | L    | S/MTI     | L (15 yrs)                 | 35               | L    | MTI       |                             | Father: adult hernia |
| 17  | 24               | L    | MTI (+YS) |                             | 23               | R    | MTI       |                             | Mother: melanoma (66 years) |
| 18  | 23               | L    | MTU (+YS) |                             | 25               | L    | S         |                             | Case 2: mumps orchitis |
| 19  | 32 & 35          | L & R | S & S |                             | 24               | L    | TD        |                             | Mother: breast cancer (57 years) |
| 20  | 22 & 23          | L & R | TD & MTI |                             | 39               | L    | S         |                             | Case 1's son: hydrocoele |
|     |                  |      |           |                             |                  |      |           |                             | Uncle: testicular cancer (seminoma - 41 yrs) |
|     |                  |      |           |                             |                  |      |           |                             | Case 1's son: hydrocoele |
| 21  | 24               | L    | MTU       |                             | 32               | R    | S         | R (7 yrs)                  | Case 1: infantile |
| 22  | 35               | R    | S         | R (10 yrs)                 | 25               | R    | S         |                             | Case 2: adult hernia |
| 23a | 27               | n/k  | MT        | n/k                        | 47               | L    | S         |                             | Mother: breast cancer (57 years) |
| 24  | 33               | L    | S         | n/k                        | 32               | R    | S/MTI (+YS) | n/k                      | Twins-non identical |
| 25  | 42               | L    | S/MTI     | n/k                        | 31               | L    | MTI       | n/k                       | - |

(continued overleaf)
| No. | Age at diagnosis | Side | Histology | UDT (Age at orchidectomy) | Age at diagnosis | Side | Histology | UDT (Age at orchidectomy) | Notes |
|-----|-----------------|------|-----------|--------------------------|-----------------|------|-----------|--------------------------|-------|
| 26  | 26              | L    | MTU       | –                        | 29              | R    | MTU       | –                        | Father: stomach cancer (48 years) |
| 27  | 21              | R    | MTU (+YS) | –                        | 23              | R    | MTI       | –                        | Cousin: UDT |
| 28  | 25              | R    | S         | –                        | 26              | L    | S         | –                        | (Maternally related – half brothers) |
| 29  | 38              | R    | MTU       | –                        | 30              | L    | S         | –                        | Father: kidney cancer (60 years) |

**Father and son**

|   |   |   |   |   |   |   |   |   |   |
|---|---|---|---|---|---|---|---|---|---|
| 30 | 34 | L | S | – | 28 | R | S | – | – |
| 31 | 33 & 40 | R & L | S & S/MTU | – | 24 | R | S | – | – |
| 32 | 46 | R | S | – | 16 | L | MTU | – | – |
| 33 | 25 | R | MTI (+Rhabdo) | – | 22 | R | S | – | – |
| 34 | 40 | R | S | – | 24 | L | MTI | – | – |
| 35 | 33 | L | S | – | 24 | R | S | – | – |
| 36 | 51 | n/k | S | – | 29 | n/k | MT | – | – |
| 37 | 33 | n/k | MT | n/k | 22 | n/k | MT | – | – |
| 38 | 47 | L | MTI | – | 34 | R | S | – | – |

**Cousins**

|   |   |   |   |   |   |   |   |   |   |
|---|---|---|---|---|---|---|---|---|---|
| 39 | 40 | R | MTU | – | 29 | L | S/MTI | – | – |

**Uncle and nephew**

|   |   |   |   |   |   |   |   |   |   |
|---|---|---|---|---|---|---|---|---|---|
| 41 | 32 | L | MTI | L & R (untreated) | 41 | R | S | – | Case 2’s son: UDT |
| 42 | 29 | R | S | – | 22 | L | S/MTI | – | – |

*L = left side; R = right side. *Histological classification according to British Testicular Tumour Panel (Pugh, 1976) with subsidiary elements, according to WHO (Mostofi & Sobin, 1977), in parentheses. Histology of bilateral tumours distinguished by ‘&’; components of combined tumours distinguished by ‘/’. Abbreviations: S = Seminoma, TD = Teratoma Differentiated, MTI = Malignant Teratoma Intermediate, MTU = Malignant Teratoma Undifferentiated, MT = Malignant Teratoma not otherwise specified, YS = Yolk Sac elements, CC = Chorio Carcinoma elements, Rhabdo = Rhabdomyosarcoma. *Family identified through national case-control study. *Family reported, in part, previously (Oliver et al., 1986b).
pexy in childhood while none of the bilateral cases had a corrective operation.

Table I documents reports of testicular cancer in other family members, reports of cancer at other sites in first degree family members, urogenital abnormalities in the cases and other family members and other miscellaneous information. The two reports of testicular cancer (a second cousin in family no. 6 and an uncle in family no. 20) were confirmed from histology but the reports of other cancers were based solely on the information supplied by members of the family. There were 15 such reports in other family members, seven in mothers of cases (family nos. 3, 4, 13, 16, 19, 23, 40), six in fathers (family nos. 3, 4, 6, 7, 26, 29), one in a sister (family no. 13), and one in a second son of a father-son pair (family no. 31). One father (family no. 7) had two primary cancers and one case (family no. 39) developed a colon cancer after his testicular tumour. A range of urogenital abnormalities were reported in the cases and their relatives including inguinal hernia, hydroscele, and testicular torsion. The most frequent was inguinal hernia which was reported in six of the 86 cases (7%), three (3.5%) being infantile.

Risk estimation

In the case-control study of 794 patients with testicular cancer, one patient reported an identical twin with a previous diagnosis of testicular cancer (family no. 2), seven patients reported a non-twin brother with such a diagnosis (family nos. 4, 5, 7 (two brothers), eight (twice), 9, 13, 23) and four patients reported a father (family nos. 30, 31, 34, 38). In total, therefore, 12 out of the 794 patients (1.5%) reported a first degree relative with a history of testicular cancer (excluding the first case in family no. 8). * There was one diagnosis of testicular cancer among the fathers of the 794 controls and one among the brothers i.e. two cases out of 794 (0.3%). All these diagnoses were histologically confirmed. These figures result in relative risks of 8.0 (95% C.I. 1.1–355.0) for developing testicular cancer if a brother has been previously affected and 4.0 (95% C.I. 0.4–197.0) if a father has been previously affected. Each testicular cancer case reported a mean of 1.18 brothers and each control a mean of 1.26 brothers (12 and eight twin brothers, 923 and 995 non-twin brothers reported by cases and controls respectively). Restricting the case-control analysis to those cases and controls with brothers did not substantively alter the relative risk.

Using actuarial analysis, the estimated cumulative risk for brothers of cases developing testicular cancer was 2.2% (95% C.I. 0.6–3.8%) by the age of 50 years. This compares with a risk of 0.23% based on 1985 England and Wales cancer registrations and results in a relative risk of 9.8 (95% C.I. 2.8–16.7).

HLA haplotype analysis

The HLA haplotype analysis for the 26 families where blood samples were taken is shown in Table III. A comparison of the extent of haplotype sharing with that expected as a result of random mendelian segregation was carried out for the 21 sib-pairs that have not been previously reported. Assuming maximum sharing of haplotypes in those families which could not be unambiguously characterised, there were four pairs of brothers where both haplotypes were shared, 13 pairs where one was shared, and four pairs where no haplo-

Discussion

The families reported here add substantially to the literature increasing the total number of reports of familial cases by about 30%. Our best estimate of the proportion of cases that are familial, derived from the case-control study, is 1.5% which is similar to other recent estimates (Tollerud et al., 1985; Dieckmann et al., 1987). This would indicate that most familial cases are not reported in the literature and that each year there should be about 15 such cases in the UK and 90 in the USA alone (assuming 1,000 and 6,000 new diagnoses per annum in the UK and USA respectively (Cancer Research Campaign, 1990; American Cancer Society, 1990).

As with other forms of cancer (Anderson, 1975), familial cases were diagnosed at younger ages than non-familial cases. The non-familial cases from the case-control study were an appropriate group to use for these comparisons because similar standards of histological review were adopted. The case-control study did not, however, recruit cases over the age of 49 years and, as approximately 14% of testicular cancers are diagnosed after this age (OPCS, 1990), differences between familial and non-familial cases in age at diagnosis are likely to be greater than those reported. Indeed the 13 familial cases from the case-control study had a median age at diagnosis of 28.5 years (28.5 years for nine cases with seminoma and 24 years for four cases with other germ cell tumours) i.e. for each category slightly younger, although not significantly so, than the median age for all familial cases presented in Table II.

The difference in age at diagnosis between familial and non-familial cases is unlikely to be due to increased surveillance of cases after the first cases had been identified. The median age at diagnosis of the first diagnosed members (who would be free of any surveillance bias) of the 42 families was 26.5 years overall (30.5 years for 17 cases with seminoma and 25.5 years for 25 cases with other germ cell tumours) i.e. not substantially different from the medians of all family cases presented in Table II.

There was a substantial difference in age of diagnosis between fathers and sons among the nine pairs in this study and this could indicate ‘genetic anticipation’ as suggested by Raghavan et al. (1980) i.e. the earlier appearance of a genetic condition with increased severity in successive generations. It is, however, likely that selection due to death or infertility of young cases in the paternal generation before they could produce children, could entirely explain this difference.

The presence of bilateral tumours in five of the 86 individ-

Table II Number and median age at diagnosis of testicular cancer familial cases on UK register and non-familial patients in national case-control study by histological classification

| Histology        | Familial cases* Median age at diagnosis (years) | Non-familial cases Median age at diagnosis (years) | P  |
|------------------|-----------------------------------------------|--------------------------------------------------|----|
|                  | No. (%) | No. (%) |                  |       |
| Pure seminoma    | 40 (46.5) | 32.5 | 391 (50.0) | 35.5 | 0.02 |
| Other germ cell  | 46 (53.5) | 26 | 390 (50.0) | 28.5 | 0.21 |
| All              | 86      | 29 | 781      | 32.5 | 0.007 |
| Chi-square*      | 0.39, 1 d.f., n.s. |

*Data refer to initial tumour for five bilateral cases. *Mann-Whitney test for comparison of median ages at diagnosis. *Chi-square test to compare distribution by histology.
Table III A HLA haplotypes* for 22 (non-identical twin) sets of
brothers with testicular cancer

| Family no. | Case 1 | Case 2 | Number of haplotypes shared* |
|------------|--------|--------|-----------------------------|
| (from Table I) | A | B | A | B | |
| 3 | 29 - 44 | 31 | w10 | w60 | 0 |
| 2 | w6 w57 | 2 | w6 | w57 |
| 4 | 2 - 7 | 2 or 3 - 7 | 1 |
| 3 | 7 | 2 or 3 w4 | 35 |
| 5 | 1 - 38 | 29 | w10 | w62 | 0 |
| 2 | w2 w27 | 31 | w10 | w60 |
| 6 | 1 w3 w62 | 1 | w3 | w62 |
| 32 | w5 w4 | 2 | w3 | w62 |
| 7 | 2 - 8 | (3) | 1 - 51 | 1 |
| 3 | w1 w51 | 2 - 8 |
| 8 | 1 - 8 | 1 | w7 | 8 |
| 24 | - 18 | 2 | w6 | 37 |
| 9 | 1 w6 w57 | 1 | w6 | w57 |
| 1 | w6 w57 | 2 - 8 |
| 10* | 29 - 7 | (2) | 2 - 13 | 1 & 2 = 1 |
| 32 | w2 | 32 | w2 | 27 | 1 & 3 = 1 |
| | (3) | 2 - 13 | 2 & 3 = 2 |
| | | 32 | w2 | 27 |
| 11 | 1 | w7 | 8 | 1 | w7 |
| 26 | - | 38 | - 38 |
| 12 | 1 | w7 | 8 | 1 | w7 |
| 2 | - | 44 | 30 | w6 | 13 |
| 13 | 31 | w2 | 27 | 31 | w2 | 27 |
| | 26 | w60 | 26 | w6 | 13 |
| 14 | 2 | w10 | w60 | 2 | w10 | w60 |
| | 3 | - | 51 | 3 | - | 51 |
| 15 | 2 | w9 | w62 | 2 | w9 | w62 |
| | 29 | - | 44 | w68 | w4 | w53 |
| 16 | 1 | w7 | 8 | 2 | w5 | 44 |
| | 2 | w10 | w62 | 3 | w7 | 7 |
| 17 | 26 | w4 | 35 | 26 | w4 | 35 |
| | 29 | - | 44 | 29 | - | 44 |
| 18 | 24 | - | 39 | 11 | - | 51 |
| | 32 | - | 44 | 26 | w10 | w60 |
| 19 | 2 | w7 | 7 | 2 | w7 | 7 |
| | 2 | w6 | 13 | 3 | w7 | 7 |
| 20 | 3 | w1 | 51 | 3 | - | 44 |
| | 3 | - | 44 | 26 | w8 | w64 |
| 21 | 3 | w7 | 7 | 3 | w7 | 7 |
| | w68 | w7 | 44 | w68 | w7 | 44 |
| 26 | 2 | w1 | w22 | 1 | w6 | w57 |
| | 3 | w7 | 7 | 3 | w7 | 7 |
| 27 | 29 | - | 44 | 2 | w4 | 35 |
| | w68 | w2 | 27 | w68 | w2 | 27 |
| 29 | 25 | - | 44 | 25 | - | 44 |
| | 2 | w10 | w60 | 33 | - | 18 |

Table III (cont'd)
C Sib-pair analysis for 21 pairs of non-identical twin brothers

| No. haplotypes | Shared | Observed | Expected |
|----------------|--------|----------|----------|
| 0              | 4      | 5.25     |          |
| 1              | 13     | 10.5     |          |
| 2              | 4      | 5.25     |          |

*A w prefix is conventionally assigned to antigens which are not yet fully defined. Cw antigens are often not serologically detectable and lack of a Cw characterisation does not indicate antigen absence. A assessment of the number of shared haplotypes was based on the HLA class I characterisation of the affected cases and of other family members (results for the latter not shown). For seven families (nos 11, 12, 15, 19, 20, 26, 42), it was not possible to assess unambiguously the degree of haplotype sharing because parental haplotypes were not available to assess the affecteds could have been inherited from different parental chromosomes. In one family (no. 41) the Aw 68 Cw 4 Bw 53 haplotype is so rare that sharing could be assumed despite the absence of parental material. 'Family HLA typing reported previously (Oliver et al., 1986b) (family no 39 now differs from original report of the haplotype and the latter estimate was considerably greater, reflected in the much narrower confidence intervals, and the analysis on which it was based adequately takes into account the period of time for which each brother was at risk, the case with two affected brothers (family no. 7) and the double-ascertained family (no. 8). A case-control comparison could also be biased in that controls may be less likely to know about or to remember cancer diagnoses in their brothers. A problem with using cancer registration rates for comparison of the actuarial risk is that there may be some routine under-reporting to cancer registries (Swedlow, 1986). For this reason we only made use of the latest available rates (for 1985) rather than earlier years for which under-reporting would be more extensive. Also, under-reporting is likely to occur less frequently in young and middle-aged men with testicular cancer who are of intense clinical interest. For these reasons, we believe that the extent of under-reporting would be no more than 10%. Under-report-
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