Reduced life expectancy seen in hereditary diseases which predispose to early-onset tumors

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Abstract: There are several hereditary diseases that are a predisposition to early-onset tumors. These include syndromic conditions like neurofibromatosis 1 and 2, von Hippel–Lindau syndrome, Gorlin syndrome, multiple endocrine neoplasia, and familial adenomatous polyposis; and conditions which are usually not possible to diagnose clinically in a single individual, such as Lynch syndrome and BRCA1/2. Understanding of the mortality in hereditary cancer predisposing diseases is important for developing effective disease treatment programs. A number of studies have been undertaken to investigate the genetic predictors, prevalence and incidence, and treatment outcomes of these diseases; however, the majority examine only the most common of these diseases (eg, neurofibromatosis or BRCA), or look into postoperative survival. The mortality of individuals who are diagnosed with one of these hereditary diseases remains an area for investigation. This review is the first to attempt identification of studies investigating life expectancy in hereditary diseases which predispose to early-onset tumors.

Keywords: mortality, survival, life expectancy, early-onset, tumors

Introduction

Contemporary studies have shown the hereditary diseases of neurofibromatosis 1 (NF1), neurofibromatosis 2 (NF2), familial adenomatous polyposis (FAP), von Hippel–Lindau syndrome (VHL), Gorlin syndrome (GS), and Lynch syndrome (LS) to be major causes of early-onset tumors in adults, often leading to reduced life expectancy and a greater dependence on health services.¹ ²

At a population level, these hereditary diseases are rare; however, in those diagnosed, they can have dramatic consequences with regards to quality of life and mortality.¹ To date, a number of studies have examined the impact of these diseases (mainly focusing on the most common of these conditions [eg, neurofibromatosis]) and postsurgical survival.¹ ³ ⁴ The number of studies which investigate the full effect of hereditary diseases that predispose to early-onset tumors upon life expectancy remains limited.

The purpose of this review is to summarize the current knowledge within the published literature regarding hereditary cancer predisposing diseases. This information is essential for etiological research and for assessment of existing treatments. Such information aids the development of guidelines, like those produced by the National Health Institute of Clinical Excellence. This review is limited to the well-known autosomal dominant conditions (Table 1).
Table 1: Chromosomal location and implications of various dominant tumor predisposing conditions

<table>
<thead>
<tr>
<th>Disease</th>
<th>Location</th>
<th>Tumors</th>
<th>Age of tumor onset</th>
<th>Risk</th>
<th>Birth incidence</th>
<th>Life expectancy</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>FAP</td>
<td>5q</td>
<td>Adenomas</td>
<td>1st year</td>
<td>100%</td>
<td>1 in 8,600</td>
<td>63–70 years</td>
<td>1,18</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Colorectal</td>
<td>4–7 years</td>
<td>99%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NF1</td>
<td>17q</td>
<td>Neurofibroma</td>
<td>1st year</td>
<td>100%</td>
<td>1 in 2,600</td>
<td>54–72 years</td>
<td>2,4,5,7,11</td>
</tr>
<tr>
<td>NF2</td>
<td>22q</td>
<td>Schwannomas</td>
<td>1st year</td>
<td>100%</td>
<td>1 in 33,000</td>
<td>62–69 years</td>
<td>1,15,17</td>
</tr>
<tr>
<td>VHL</td>
<td>3p</td>
<td>Hemangioblastomas</td>
<td>1–2 years</td>
<td>90%</td>
<td>1 in 40,000</td>
<td>49–53 years</td>
<td>1,22–24</td>
</tr>
<tr>
<td>MEN1</td>
<td>11q</td>
<td>Parathyroid</td>
<td>5 years</td>
<td>95%</td>
<td>1 in 35,000</td>
<td>Close to normal</td>
<td>30,31</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Insulinoma</td>
<td>5 years</td>
<td>95%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>MEN2a</td>
<td>10q</td>
<td>Medullary thyroid</td>
<td>3 years</td>
<td>80%</td>
<td>1 in 40,000</td>
<td>Reduced</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Parathyroid</td>
<td>3 years</td>
<td>80%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Pheochromocytoma</td>
<td>3 years</td>
<td>80%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>MEN2b</td>
<td>10q</td>
<td>Medullary thyroid</td>
<td>1st year</td>
<td>100%</td>
<td>1 in 1,000,000</td>
<td>Reduced</td>
<td></td>
</tr>
<tr>
<td>Gorlin</td>
<td>9q</td>
<td>Basal cell carcinoma</td>
<td>5 years</td>
<td>90%</td>
<td>1 in 18,000</td>
<td>73.4 years</td>
<td>1</td>
</tr>
<tr>
<td>syndrome</td>
<td></td>
<td>Medulloblastoma</td>
<td>1st year</td>
<td>5%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cowden</td>
<td>10q</td>
<td>Breast cancer</td>
<td>30 years</td>
<td>60%</td>
<td>1 in</td>
<td>Reduced</td>
<td>41,43,46</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Thyroid</td>
<td>30 years</td>
<td>40%</td>
<td>200,000–250,000</td>
<td>in women</td>
<td></td>
</tr>
<tr>
<td>PJS</td>
<td>19p</td>
<td>GI malignancy</td>
<td>20 years</td>
<td>60%</td>
<td>1 in 25,000</td>
<td>58 years</td>
<td>33,34</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Breast/cervix</td>
<td>20 years</td>
<td>40%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>JPS</td>
<td>18q, 10q</td>
<td>GI malignancy</td>
<td>20 years</td>
<td>40%</td>
<td>1 in 100,000</td>
<td>Reduced</td>
<td>37,40</td>
</tr>
<tr>
<td>LFS</td>
<td>17p</td>
<td>Sarcoma</td>
<td>1st year</td>
<td>80%</td>
<td>1 in 30,000</td>
<td>Severely reduced</td>
<td>4,7,15</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Glommas</td>
<td>1st year</td>
<td>20%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HDGC</td>
<td>16q</td>
<td>Gastric</td>
<td>&lt;40 years</td>
<td>70%–80%</td>
<td>Very rare</td>
<td>Reduced</td>
<td>55,57,60,61</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Breast (women)</td>
<td>&gt;50 years</td>
<td>20%–40%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lynch</td>
<td>2p, 3p, 2q, 7p</td>
<td>Colorectal</td>
<td>&gt;16 years</td>
<td>50%–80%</td>
<td>1 in 1,000–4,000</td>
<td>Reduced</td>
<td>50–54</td>
</tr>
<tr>
<td>syndrome</td>
<td></td>
<td>Endometrium</td>
<td>&gt;20 years</td>
<td>60%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Ovary/ureter</td>
<td>&gt;20 years</td>
<td>10%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Gastric</td>
<td>&gt;20 years</td>
<td>10%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Pancreas</td>
<td>&gt;30 years</td>
<td>5%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BRCA1</td>
<td>17q</td>
<td>Breast/ovary (women)</td>
<td>&gt;16 years</td>
<td>60%–90%</td>
<td>1 in 1000</td>
<td>62 years</td>
<td></td>
</tr>
<tr>
<td>BRCA2</td>
<td>13q</td>
<td>Breast/ovary (women)</td>
<td>&gt;16 years</td>
<td>40%–90%</td>
<td>1 in 800</td>
<td>68 years</td>
<td>66</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Prostate</td>
<td>&gt;30 years</td>
<td>20%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Pancreas</td>
<td>&gt;30 years</td>
<td>5%</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: FAP, familial adenomatous polyposis; NF, neurofibromatosis; VHL, von Hippel–Lindau syndrome; MEN, multiple endocrine neoplasia; PJS, Peutz–Jeghers syndrome; JPS, juvenile polyposis syndrome; LFS, Li–Fraumeni syndrome; HDGC, hereditary diffuse gastric cancer; GI, gastrointestinal.

Types of studies

There are generally two types of study assessing mortality in tumor predisposing syndromes. The first assesses survival in cohorts of ascertained patients. The second assesses mortality based on death certificate notification. The latter relies on the notification of the syndrome on the death certificate, which could only reliably occur in conditions that are easily diagnosed in a single individual. NF1 can be identified easily by cutaneous examination and documentation of café au lait patches and neurofibromas; NF2 by the presence of bilateral vestibular schwannomas; and FAP by the presence of >100 adenomas. However, this would not be possible in either LS or BRCA1/2. We used the following search terms to identify the articles required for this review: mortality, survival, and life expectancy by each of the conditions studied.

Neurofibromatosis (type I)

The neurofibromatoses are the most well-documented of the cancer predisposing hereditary diseases and can
be divided into three genetic types: NF1, NF2, and schwannomatosis. NF1 is an autosomal dominant dis-

ease caused by mutations in the NF1 gene at 17q11.2 on 
chromosome 17 that normally codes for the cytoplasmic 

protein neurofibromin. The pathogenic mutations lead 

to a wide variety of symptomatic presentations, though 

the distinctive phenotypic expressions include >6 café 

au lait patches, >2 Lisch nodules of the iris, as well as 

central nervous system tumors, and vasculopathy. The 

overwhelming characteristic feature of NF1 is peripheral 

nerve sheath tumors (neurofibromas, malignant peripheral 

nerve sheath tumors [MPNST]), which form as a result of 

the loss of the tumor suppressor protein neurofibromin. 


MPNST are a major cause of reduced life expectancy in 

NF1 patients. 

Epidemiological data shows NF1 to be a relatively 

common autosomal dominant disease with birth incidence 
estimates ranging from 1 in 2,500 to 1 in 3,000 live births. Evans et al conducted a study looking into the prevalence and incidence of tumor-prone syndromes in North West England and found an estimated birth incidence for NF1 of 1 in 2,712, based on the highest incidence decade values.

A number of studies have looked at the genetic and phenotypic expression of NF1; however, relatively few have looked into the effect on mortality. Recently reported estimates of reduced life expectancy vary from 8 years to 15 years. An Italian death certificate study by Masocco et al reported a dramatic increase in mortality between adolescence and 40 years, with a further increase after 50 years of age. They also found a mean reduction in life expectancy of 20 years in comparison to the general population. Another death certificate study in the United States assessed cause of death in 3,770 NF1 patients, who died between 1983–1997, and found a reduced life expectancy of 15.7 years (95% confidence interval [CI] 15.0–16.3) compared to the general population. Mean age at death was recorded as 54.4 years in NF1 patients and 70.1 years in the general US population.

A 12-year follow-up study conducted by Zöller et al on 70 Swedish NF1 adults looked into life expectancy, mortality, and cause of death. Compared to the general Swedish population, the mean age of death (61.6 years) was 15 years younger. Compared to the 5.1 deaths expected in the general population, 22 deaths were found to have occurred in the NF1 group ($P < 0.001$). A Danish nationwide cohort of 212 NF1 patients were followed by Sørensen et al over a period of 42 years. As with all of the subsequent studies, Sørensen et al found significantly reduced survival rates in both those diagnosed with neurofibromatosis and their relatives compared to the general population.

Two more recent cohort studies have again confirmed reduced life expectancy in patients with NF1. A population-based study in the UK showed an 8-year reduction in life expectancy, mainly due to deaths from MPNST and glioma, although there was an increased PMR (proportionate mortality rate) from cardiovascular disease in men (relative risk 4.2; 95% CI 2.6–6.2). The other recent cohort study was from France; consecutive NF1 patients referred to the National French Referral Center for neurofibromatosis were included. Between 1980 and 2006, 1,895 NF1 patients were followed up for a median of 6.8 years (range, 0.4–20.6 years). Vital status was available for 1,226 (65%) patients, of whom 1,159 (94.5%) survived and 67 (5.5%) died. Overall mortality was significantly increased in the NF1 cohort (PMR, 2.02; CI, 1.6–2.6; $P < 10^{-4}$). The excess mortality occurred among patients aged 10–20 years (PMR, 5.2; CI, 2.6–9.3; $P < 10^{-4}$), and 20–40 years (PMR, 4.1; 2.8–5.8; $P < 10^{-4}$). Significant excess mortality was found in both males and females. In the 10–20 year age group, females had a significant increase in mortality compared to males (PMR, 12.6; CI, 5.7–23.9; and PMR, 1.8; CI, 0.2–6.4, respectively). The cause of death was reported for 58 (86.6%) patients; MPNST was the most frequent cause of death (60%). They found significantly increased PMRs, indicating excess mortality in NF1 patients compared to the general population. Overall mortality was significantly increased in NF1 patients aged 10–40 years and tended to be higher in females than in males. There was no excess mortality >40 years of age, which may have been due to limited numbers in the older age groups.

Cohort studies may be biased by ascertaining more severely affected cases, although this would be ameliorated by a highly ascertained cohort. Although death certificate studies might be expected to overcome the ascertainment bias of cohort studies, they are dependent on NF1 being accurately recorded on the death certificate. In the Evans et al 2011 study, only 35% of NF1 patients with available death certification had NF1 as a contributing cause on the death certificate. The use of death certificates by Rasmussen et al in 2001 and Masocco et al in 2011, therefore, needs to be assessed against the likelihood that all deaths associated with NF1 were recorded. The Rasmussen et al study found that NF1 was only recorded
on 1 in 8,679 deaths. In the Masocco et al study, this was even lower at 1 in 10,685.

In summary, while there are clearly excess deaths from malignancy in NF1 in the first 40–50 years of life and overall life expectancy is reduced, it is likely that life expectancy overall is not reduced by the 15–20 years suggested in many studies.

**Neurofibromatosis (type 2)**

NF2 is a chronic autosomal dominant disease, which is genotypically expressed by the inactivation of the NF2 tumor suppressor gene on chromosome 22q12. Phenotypic expression of this genotypic loss includes spinal tumors, peripheral nerve tumors, and brain tumors, with vestibular schwannomas being the hallmark manifestation. Indeed, an evaluation study into the predictors of risk by Mautner et al revealed almost 95% of individuals diagnosed with NF2 presented vestibular schwannomas, with 90% presenting with spinal tumors. Upon examination, the mortality rate of NF2 can be related to the development of these benign tumors, which can develop in uncontrolled numbers, with (until recently) surgery and sometimes radiotherapy the only effective treatments.

With an incidence of 1 in 33,000, NF2 is a rare disease affecting only a small number of the general population. Despite its uncommon occurrence, NF2 is a chronic condition that substantially reduces life expectancy in those reported as sufferers. However, as with NF1, although life expectancy is shortened in comparison to the general population, improvement in treatments shows mean actuarial survival to be 62 years. Conversely, this same study by Evans et al looked into symptoms and survival in 150 NF2 patients and found a mean age of death of 36.25 years for 40 cases, highlighting the presence of intra- and interfamily differences in disease presentation. Advancements in surgical and pharmacological treatments continue to improve life expectancy in those diagnosed with NF2. A more recent assessment of NF2 mortality in a population-based sample showed a life expectancy of 69 years, with an improvement of 14.7 years since the inception of a genetic register in 1990. Predictors of mortality include mutation type (truncation is worst), and the presence of meningiomas; importantly, treatment in a specialist center was associated with improved mortality.

**Familial adenomatous polyposis**

FAP is an autosomal dominant disease caused by mutations in the APC gene on chromosome 5q. Characteristics include a predisposition to benign and malignant colorectal and duodenal tumors, as well as rare malignant brain tumors, thyroid cancer, and hepatoblastoma. If unscreened, >95% of FAP patients will develop colorectal cancer, which is therefore the main recorded cause of death. Deaths may also occur due to duodenal cancer and infiltrative desmoid disease. Reduced life expectancy can be seen in FAP-affected individuals, with current estimates from the UK of 70.4 years. However, life expectancy has improved dramatically by regular screening and surgical intervention to remove the at-risk colon and rectum. Under these assumptions, current life expectancy appears to be similar to those diagnosed with NF1.

**von Hippel–Lindau syndrome**

Three studies have reported survival for patients with VHL, with two studies classifying the average survival as less than 50 years, and median survival measured as 49 years. In comparison to the other hereditary diseases that predispose to early-onset tumors, VHL is found to have the lowest life expectancy. Survival can often be limited in patients with this disease by the development of benign tumors in highly vascular areas, such as hemangioblastomas of the central nervous system. It is thought that the recurrence and number of tumors that develop within these vascular areas lead to the early death of VHL patients.

**Gorlin syndrome**

GS is considered to have the highest life expectancy from among the tumor predisposing syndromes. Although the numbers of patients are small, with an estimated birth incidence of 1 in 18,000, GS is one of the more common hereditary early-onset tumor diseases. In a 2011 study, Wilding et al demonstrated a reduced life expectancy of 73.4 years. Only 19% of deaths appeared to be related to GS, a much lower percentage than for the four conditions described above (55%–73%). We were unable to find any other papers assessing life expectancy. A potential reason for the relatively normal life expectancy in GS patients could be that the main complications of multiple basal cell carcinomas and benign jaw cysts only rarely metastasize, and even medulloblastomas in GS have relatively high survival rates.

**Multiple endocrine neoplasia**

Multiple endocrine neoplasia (MEN) encompasses several distinct syndromes (Table 1) featuring tumors of endocrine glands, each with its own characteristic pattern. In MEN1, the tumors are predominantly benign, while in MEN2, tumors are often malignant, with medullary thyroid cancer being the
main problem. A study assessing mortality in MEN1 found that 17/60 (28%) individuals died of MEN1-related causes, most commonly metastatic islet cell tumors (ten patients). The remaining patients died of causes unrelated to MEN1. The overall 20-year survival of MEN1 patients in this series was 64% (95% CI 56%–72%) and that of an age- and gender-matched population was 81% (P < 0.001). This suggested an increased risk of premature death in MEN1. However, a population-based study in Finland showed that over a 30-year period the spouses of MEN1-affected individuals lived the same length of time.

While the life expectancy in MEN1 may not be dramatically affected, this may not be the case in MEN2, particularly for those born with de novo MEN2b mutations that can cause medullary thyroid cancers in the first 5 years of life. There is nonetheless very little information regarding life expectancy in MEN2 in the literature. One 20-year follow-up study showed 10-year survival of only 68% in medullary thyroid cancer patients. However, the identification of the RET oncogene as the cause of MEN2a and MEN2b and early thyroidectomy before cancer diagnosis almost certainly normalizes life expectancy.

**Peutz-Jeghers syndrome**

Peutz–Jeghers syndrome (PJS) is characterized by a typical benign polyp that can occur throughout the gastrointestinal tract and mucocutaneous pigmentation. The majority of patients that meet the clinical diagnostic criteria have a causative mutation in the STK11 gene at 19p13.3; however, it is the high risk of gastrointestinal malignancy, as well as breast and a rare form of cervical cancer that can affect life expectancy. Among 72 patients with PJS at St Mark’s Hospital (London, UK), malignant tumors developed in 16 (22%), causing death in all but one. There were nine gastrointestinal and seven nongastrointestinal tumors. The relative risks of death from gastrointestinal cancer and all cancers were 13 (95% CI 2.7–38.1) and 9 (95% CI 4.2–17.3), respectively. The chance of dying of cancer by the age of 57 was 48%.

In a more recent study of 133 PJS patients, 49 cancers were diagnosed in 42 patients (32%) in 5,004 person-years of follow-up, including 25 gastrointestinal cancers. The median age at first cancer diagnosis was 45 years. The cumulative cancer risk was 20% at age 40, increasing to 76% at age 70. Cumulative cancer risks were higher for females than for males, reflecting the malignancies of breast and cervix (P = 0.005). The relative cancer risk was higher in PJS patients than in the general population (hazard ratio [HR] 8.96; 95% CI 6.46–12.42) and higher among female (HR 20.40; 95% CI 13.43–30.99) than among male patients (HR 4.76; 95% CI 2.82–8.04). Forty-two patients died at a median age of 45 years, including 28 cancer-related deaths (67%). Mortality was increased in this Dutch cohort compared to the general Dutch population (HR 3.50; 95% CI 2.57–4.75).

**Juvenile polyposis syndrome**

First defined in 1964 by McColl and followed by several case reports in the late 1960s (eg, Johnston et al), juvenile polyposis syndrome (JPS) is a familial condition that affects the upper and lower gastrointestinal tract. JPS is a relatively rare condition, affecting approximately 1 in 100,000 children. Deletion and/or mutation is seen in the tumor-suppressor genes SMAD4 and BMPR1A on the chromosomes 18q and 10q, respectively. At initial presentation, symptoms include multiple polyps, mucosal lesions, rectal bleeding, diarrhea, and exudative enteropathy. Such phenotypic expression, particularly in the gastrointestinal tract, greatly reduces the life expectancy of the affected children. A more recent study in 2007 by Brosens et al addressing colorectal cancer has shown a cumulative lifetime risk of 38.7% in JPS, with a mean diagnosis age of 43.9 years. We found that very few studies have looked into JPS and even fewer have reported on mortality.

**Cowden’s syndrome**

Cowden’s syndrome is a rare familial autosomal disease that has been localized through several genetic studies to chromosome 10q23 and the tumor-suppressor gene PTEN. The phenotypic hallmarks of this disease include multiple facial papules, trichilemmomas, oral papillomas, and macrocephaly, though there are varied expressions of the disease even among families. Breast cancers are the most frequent malignancies reported to be associated with this syndrome, though various organ systems can be involved, including gastrointestinal and urogenital.

Originally estimated as having an incidence of 1 in 1,000,000, a more recent investigation into the novel genotype–phenotype correlations of Cowden’s syndrome revised the figure by post-genetic identification to a prevalence of 1 in 200,000–250,000 within a Dutch population. Further investigations have examined potential chemoprevention and treatment of Cowden’s disease in mouse models, with the intent to increase life expectancy. However, as with many of the other diseases, a large body of work can be found on the genetics of the syndrome, though few articles discuss the mortality and life expectancy of Cowden’s patients.
Li–Fraumeni syndrome

It is likely that life expectancy is substantially reduced in Li–Fraumeni syndrome, which is an autosomal dominant disorder caused by germ-line mutations in the TP53 gene. Mutation carriers have a 20% chance of developing malignancy in childhood (particularly soft tissue and osteosarcoma, glioma, and adrenal carcinoma) and a very high chance of sarcoma and brain tumors in adulthood. Women have close to a 100% lifetime risk of malignancy, as the risk of breast cancer at <50 years is higher than that for BRCA1/2. There have been no studies that have directly addressed life expectancy in a large series of mutation carriers, but the very early onset of malignancy and poor survival means that life expectancy is likely to be below 40 years of age on average. A recent study using multi-modal screening, including whole body MRI scans, found tumors in seven of 15 screened mutation carriers. All 15 were alive after a mean follow-up time of 47 months (compared with two of ten [20%] mutation carriers in whom 12 malignant tumors occurred but who did not undergo surveillance). Among those diagnosed with a cancer, survival was better in the surveillance group than the non-surveillance group (100% versus 21.4%, P < 0.001).

Lynch syndrome

Hereditary nonpolyposis colorectal cancer, also known as Lynch syndrome (LS), is an autosomal dominant disorder with germ-line mutations in the DNA mismatch repair genes MLH1, MSH2, MSH6, and PMS2. After compiling information from research articles, the hallmark characteristic of LS is noted as early-onset colorectal cancer, with significant increased risk for cancers of the stomach, small intestine, upper urinary tract, uterus, ovary, and brain. In reviewing research articles which studied eligible populations, we see that all of the articles report a reduction in life expectancy for those diagnosed with LS. The majority of previous publications have discussed the benefits of prophylactic gastrectomy rather than addressing life expectancy of individuals diagnosed with the condition.

BRCA1/2

Female BRCA1 and BRCA2 mutation carriers have reduced life expectancy mainly due to early death from breast and ovarian cancer. Life expectancy may be worse in BRCA1 carriers due to the greater risk of and poorer survival from ovarian cancer. Risk-reducing surgery is predicted to improve survival amongst female carriers of pathogenic mutations in both BRCA1 and BRCA2. Most of this benefit is likely to occur from primary prevention of breast and ovarian cancer in women currently unaffected. However, a recent decision analysis demonstrated only a minor advantage from risk reducing mastectomy over surveillance with MRI. Formal direct evidence for an improvement in survival in BRCA1/2 carriers was reported for risk-reducing bilateral salpingo-oophorectomy (RRBSO), and a survival advantage was also seen in women with RRBSO after diagnosis of breast cancer. However, no formal direct evidence of a survival advantage after bilateral risk reducing mastectomy alone has yet emerged. This may be related, in part, to better survival...
Survival in Li–Fraumeni cancers, in contrast, is often very much better in LS cancers, most notably in ovarian cancer. BRCA-related cancers appear to have better survival than expected due to and their sporadic counterparts.

Survival may also differ between the tumors in the syndromes and their sporadic counterparts. BRCA1/2-related ovarian cancers appear to have better survival than expected due to greater sensitivity to platinum-based therapy. Survival is often much better in LS cancers, most notably in ovarian cancer.

Survival in Li–Fraumeni cancers, in contrast, is often very poor. In NF1-related tumors, survival can be better than in sporadic disease, for instance, for optic pathway tumors or worse, as in MPNST.

Perhaps the greatest improvement in life expectancies has come from surgical prevention. Colectomy in FAP has increased life expectancy by 15–20 years, and extended right hemicolectomy is recommended for treatment of colorectal cancers proximal to the rectum in LS. Thyroidectomy is a vital tool for prevention in MEN2, and oophorectomy is a proven way to prolong life in BRCA1/2 mutation carriers.

Hopefully the future of personalized medicine will provide alternatives to these proven surgical options.

Conclusion
In summary, this is an attempt to fully describe the current knowledge of reduced life expectancy in autosomal dominant diseases resulting in early-onset tumors. These results are key to the understanding and development of optimal screening, prevention, and treatment programs for the management of individuals with these hereditary diseases. All the diseases examined show general reduced life expectancy among those diagnosed, though the severity of this reduction varies greatly between the conditions. MEN1 often presents with benign tumors although slightly reduced, life expectancy within these individuals has been found to be close to the general population. In contrast, VHL and Li–Fraumeni syndrome both lead to a severe reduction in life expectancy due to the high number and location of the tumors that develop. There is evidence for improvement in life expectancy in a number of conditions with well-organized systematic screening, such as in LS and FAP.

The number of studies investigating the different hereditary early-onset tumors is large and diverse. However, only a limited number of journal articles were found to examine life expectancy in these diseases, particularly in the rarer of the germ-line mutations. The high fatality rate among these uncommon autosomal dominant diseases provides strong justification for further research into life expectancy.

Disclosure
The authors report no conflicts of interest in this work.


