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# Challenges in developing and implementing international best practice guidance for intermediate-risk variants in cancer susceptibility genes: *APC* c.3920T>A p.(Ile1307Lys) as an exemplar

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Received 23 January 2024

Accepted 11 April 2024

Published Online First 2 May 2024

The National Health Service (NHS) in the UK is a publicly funded entity, and decisions regarding allocation of scarce resources for provision of services, including related to genetic testing and cancer surveillance, are based on clinical and cost-effectiveness of proposed interventions, and are grounded by ethical considerations related to justice—allocation of resources based on clinical need; equity—fair provision of services across the population; beneficence and non-maleficence—provision of genetic testing/surveillance for clinical benefit, and minimisation of associated harm related to overinvestigation or overdiagnosis.<sup>1</sup>

NHS-funded cancer screening in the general population is guided by Wilson and Jungner criteria.<sup>2</sup> These criteria, originally described in 1968, emphasise certain factors that should be considered in evaluating a screening test—related to the condition (natural history and severity), the test (sensitivity and specificity), the actionability of a positive finding (effective treatment), as well as acceptability and cost-effectiveness.<sup>2</sup> Priority for cancer surveillance is typically given to those individuals at highest risk of disease—where numbers needed to screen or treat to prevent disease-associated mortality are low.<sup>3</sup> NHS-funded constitutional (germline) genetic testing has traditionally been restricted, for the most part, to those individuals in whom the likelihood of identifying a causative clinically actionable constitutional pathogenic variant is at least 10%,<sup>4</sup> although expansion of testing is already increasing as associated cost of testing falls and therapeutic implications expand.<sup>5</sup> NHS-funded constitutional genetic testing is also focused, dependent on the patient phenotype, and limited to genes for which strong evidence regarding gene–disease association exists.<sup>6,7</sup> Furthermore, analysis and reporting of variants are restricted to those associated with intermediate to high penetrance and where identification of the variant has clinical utility. For situations where there is evidence suggesting low penetrance (typically defined as an OR of <2<sup>8</sup>) associated with a particular genotype or class of variant, these variants are not analysed or reported in current NHS practice. For example, analysis of *CHEK2* is largely restricted to truncating variants, which are known to be associated with higher breast cancer risks,<sup>9</sup> and testing of low-risk susceptibility alleles in genes such as *MITE*, *MC1R* or *EGFR* is not currently offered.

In other healthcare systems, thresholds at which cancer surveillance is offered may be lower than the threshold at which such surveillance is offered within the NHS. This, in part, explains international variability in the extent of genetic testing offered, eligibility criteria for genetic testing and guidelines for management of carriers of variants associated with incomplete penetrance in different cancer susceptibility genes. As a consequence, this can pose significant challenges when trying to define international best practice guidance. The International Society for Gastrointestinal Hereditary Tumours (InSiGHT) is a multidisciplinary expert group committed to enhancing care of patients with hereditary gastrointestinal cancer predisposition. This group has recently published a position statement regarding cancer risks associated with the recurrent variant NM\_000038.6(*APC*): c.3920T>A, p.(Ile1307Lys), referred to as p.I1307K hereafter for brevity.<sup>10</sup> *APC* p.I1307K has been a source of discussion internationally, given ethnicity-specific risks and variability in cancer surveillance recommendations between different jurisdictions.

The authors of the InSiGHT position statement deserve commendation for attempting to address existing variability in practice and provide a succinct and informed review of the latest data. International variability in clinical practice is almost inevitable given the specific population needs and access to intervention within each NHS or system. The UK Cancer Genetics Group (UKCGG), a constituent group of the British Society of Genomic Medicine, has issued a response to the InSiGHT position statement,<sup>11</sup> incorporating health priorities, opportunity cost and equity of access within the UK NHS, mindful that adoption of these recommendations may conflict with the aims of the authors of to standardise clinical management of carriers of this variant.

Familial adenomatous polyposis (FAP) is caused by constitutional pathogenic variants in *APC*. These variants are usually inherited from an affected parent, but may arise de novo, and mosaicism is not uncommon.<sup>12</sup> The hallmark of FAP is the development of large bowel polyps, but there is wide variation in the colonic phenotype.<sup>13</sup> Extracolonic manifestations are common and may be clinically important features. The constitutional *APC* variant p.I1307K is not associated with clinical features



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**To cite:** McVeigh TP, Laloo F, Frayling IM, et al. *J Med Genet* 2024;**61**:810–812.

of FAP but is associated with a moderately increased risk of colorectal cancer in individuals of Ashkenazi Jewish heritage.<sup>10</sup> This should be considered a separate allelic predisposition. The colorectal cancer risks associated with this variant are limited to individuals of Ashkenazi Jewish heritage and confer a low or, at most, moderate risk, with an OR of 1.5–2 in this specific group.<sup>10</sup> A high population prevalence, with 1 in 28 individuals of Ashkenazi Jewish ancestry carrying at least one copy of the variant allele, complicates things further. Homozygous carriers with wide phenotypic variability have been reported, but data are scant, such that the exact risks in such individuals are unclear.<sup>10</sup> Carrier frequency in individuals of other ancestries is much less frequent, and associated risk uncertain.<sup>10</sup> In line with this, the InSiGHT group recommends predictive testing in first-degree relatives of probands of Ashkenazi Jewish heritage as well as colorectal cancer surveillance of identified carriers (heterozygous or homozygous) every 5 years starting from 45 to 50 years if of Ashkenazi Jewish heritage, but not in individuals of other ethnicities.<sup>10</sup>

Recommendations for enhanced screening are based on family history as well as genotype. Current British Society of Gastroenterology (BSG)/Association of Coloproctology of Great Britain and Ireland/UKCGG guidelines recommend that individuals at high familial colorectal cancer risk (>6-fold) have 5-yearly colonoscopy from age 40 years, and those at moderate colorectal cancer risk (approximately 2-fold to 6-fold risk), a one-off colonoscopy at age 55 years,<sup>14</sup> before entry to the National Bowel Cancer Screening Programme (NBCSP), predicated on the impact of colonoscopy on the mitigation of cancer risk in these populations. Individuals at low population risk (<2-fold risk) are offered screening within the NBCSP, which currently includes 2-year faecal immunochemical test (FIT) from age 60 years. The role of FIT-based surveillance in carriers of the APC p.I1307K variant has not been specifically evaluated. Enhanced colonoscopic surveillance for carriers of low to intermediate risk alleles is not typically recommended in the absence of a family history unless there is a clear and proven survival benefit. Given that the estimated risks associated with APC p.I1307K are less than twofold, recommending enhanced surveillance beyond that recommended to individuals at moderate risk based on family history is inconsistent with equivalent colonoscopic surveillance strategies. In the absence of specific data detailing outcomes of endoscopic surveillance in carriers of APC p.I1307K, the age at which advanced adenomas develop is unclear, and therefore the risks and benefit of such surveillance are difficult to establish.

Identification of APC p.I1307K alone is unlikely to fully explain the phenotype of an individual fulfilling clinical testing criteria for testing of genes associated with colorectal cancer and polyposis, as outlined in the NHS Genomic Testing Directory, which broadly recommends testing in young onset or strong familial cases; thus, the clinical utility of reporting this variant as an isolated finding is uncertain. Furthermore, standard guidelines for interpretation and classification of variants in APC cannot be applied to this variant,<sup>15</sup> which has fuelled variability in reporting, with some labs classifying as ‘variant of uncertain significance’ and others as a ‘low-risk allele’.<sup>16</sup> The position statement of the InSiGHT group advocates that this variant should be reported as ‘pathogenic, low penetrance’.<sup>10</sup> However, after discussion with relevant key stakeholders, and considering that other similar low-risk alleles are not reported, the UKCGG has recommended that this variant is not considered or reported as part of NHS-funded diagnostic APC testing.

Identification of the APC p.I1307K variant in a family may arise through other channels (eg, through private testing/testing in another country). However, we do not recommend predictive testing for this variant in unaffected relatives, irrespective of the ethnicity of the proband, nor would we recommend colonoscopic surveillance unless otherwise warranted based on family history and in line with current BSG guidance. If the relative of a known carrier requires diagnostic APC testing in their own right, they should be counselled that the familial variant will not be reported by an NHS laboratory, even if present in their sample.

Given that data in an unselected population would be valuable to inform future guidance, we have also encouraged laboratories, where possible, to make data regarding variant identification available for submission to the National Disease Registration service, where possible, to facilitate review of outcomes in an unselected population in the future. We have also suggested that if screening is offered to pre-symptomatic carriers of this variant without a family history of colon cancer, that this should be within the context of a prospective research study. Ideally, prospectively collated data in an unselected patient group should be collected, to inform future evidence-based recommendations focused on improving outcomes and specific to population health needs.

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**Acknowledgements** The authors thank the clinical scientists who provided comment on current practice in their laboratories.

**Contributors** The authors (TPM, FL, IMF, AL, KS, MD, KJM, HH) conceived of the idea and wrote the manuscript. TPM is the author responsible for the overall content as the guarantor.

**Funding** HH is supported by the National Institute for Health and Care Research (NIHR) Exeter Biomedical Research Centre (NIHR203320). This study was supported by the NIHR Exeter Biomedical Research Centre.

**Disclaimer** The views expressed are those of the author(s) and not necessarily those of the NIHR or the Department of Health and Social Care.

**Competing interests** None declared.

**Patient consent for publication** Not applicable.

**Ethics approval** Not applicable.

**Provenance and peer review** Not commissioned; externally peer reviewed.

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