Colorectal cancer prognosis: is it all mutation, mutation, mutation?

A B Hassan, C Paraskeva

For the 500 000 new cases of colorectal cancer in the world each year, identification of patients with a worse prognosis and those who are more likely to respond to treatment is a challenge. There is an increasing body of evidence correlating genetic mutations with outcome in tumours derived from human colorectal cancer cohorts. K-ras, but not p53 or APC, mutations appear to be associated with poorer overall survival in colorectal cancer patients.

For the 500 000 new cases of colorectal cancer in the world this year, staging of their disease and subsequent treatment will be almost entirely based on a histopathological classification originated by Cuthbert Duke in the 1920s. While there is nothing immediately wrong with this classification and its derivatives, in particular because it has been universally adopted, it is clear that significant heterogeneity exists between patient outcomes with otherwise apparently identical pathological staging. The immediate challenge faced by the clinical community is to identify patients with a worse prognosis and those who are more likely to respond to treatments, such as surgical resection, radiotherapy, chemotherapy, and molecular therapy.

Patient selection is also now more of an important issue for a series of scientifically based reasons. These include identification of patients at high risk from colorectal cancer prior to screening and prevention programmes, those with familial predisposition (35% of total), including mutation screening of key genes (2–6% of total), and more recently in systemic treatment, as empirically applied treatments including cytotoxic chemotherapy are now being challenged by new molecular therapies. Some of the new molecules are currently administered empirically, as the co-development of molecular testing and tailored treatments has not kept pace with the enthusiasm of the immediate therapeutic testing.1,2 In time, this situation will reverse as the costs of treatments increase, and we should expect a series of predictive tests to be developed that facilitate selection based on currently available targeted therapy. In situations where targeted molecular therapy has been developed with a validated molecular diagnostic test (for example, breast (Her2), chronic myeloid leukaemia (BCR-ABL), gastrointestinal stromal tumours (C-Kit), and lung cancer (EGFR)), there have been dramatic improvements in disease management that far exceed anything that has been achieved with an empirical approach. What is critical is that the selectivity and potency of these therapies exploits mutation in the tumour cells, with amplification of Her2, translocation in BCR-ABL, and mutation of receptors C-Kit and EGFR. Although an expensive challenge, the cancer field is rightly continuing to pursue the identification and evaluation of common human mutations in cancer, as evident by the Cancer Genome project.3 This activity will no doubt drive the development of specific therapeutic interventions to specific mutated targets. Despite the success to date in other cancers, this whole activity could be an expensive mistake if a combination of mutation complexity, aneuploidy, and epigenetic alterations in tumour cells ultimately determine prognosis and response to therapy, irrespective of targeted therapy to what are presumed to be key mutations.

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So what of mutations in colorectal cancer? The paper by the Dundee group4 in this issue of Gut contributes to an increasing body of evidence correlating mutation with outcome in tumours derived from human colorectal cancer cohorts (see page 1283). Before examining their findings, it is important to recognise that mutational analysis and genetic models have already identified the key pathways critical for initiation and progression of colorectal cancer. Deregulation of the Wnt signalling pathway, by either loss of function or gain of function mutations of adenomatous polyposis coli (APC) and b-catenin, respectively, in both hereditary and sporadic colorectal adenoma, establish the principal initiation pathway.5 Although specific mutational hotspots occur that account for attenuated inherited phenotypes, most sporadic mutations of APC occur in a mutational cluster.6,7 The discriminating power and clinical utility of these mutations have proven to be less impressive, even though this pathway remains a critical therapeutic target.8 Genetic modifiers of the number, differentiation, and progression of

Abbreviations: K-ras, Kirsten-ras; APC, adenomatous polyposis coli
colorectal adenoma can either derive from the host or from coexisting and selected mutations in the tumour. Dissecting these components has been problematic, mainly because a large number of variables exist (for example, variable penetrance, gain and loss of function mutations in each gene, and synergistic effects between modifiers that coexist in both tumour and stromal cell populations). Although these limitations are being circumvented using larger patient numbers with high throughput technologies, there will have to be a practical outcome that will essentially distil down this information to a fewer number of key mutations, and potentially to where we have started from, a classification based on morphology and identification of a subpopulation of localised cells that essentially determine overall outcome (for example, those at the micro-invasive front of the tumour).

“K-ras mutations, but not p53 or APC mutations, were associated with poorer overall survival, even when correcting for Dukes’ stage, age, and sex”

The Dundee group prospectively characterised mutations in three genes implicated in colorectal cancer (APC, p53, Kirsten-ras (K-ras)) and extended a previous study.7 They observed that the mutation frequencies were variable between genes (APC = 56%, p53 = 61%, K-ras = 27%), and when corrected for expected frequencies, p53 and K-ras genes rarely coexisted (observed 2% versus expected 16%), even though the reduction in expected frequency of combined p53 and K-ras appears to have been lost when mutations of APC coexisted in addition. Conlin and colleagues4 have also now recognised that neither falls within a chromosomal nor instability also correlate with survival, with a subgroup which are best exemplified by the six hallmarks proposed by Hanahan and Weinberg,20 and to then analyse what intermediate surrogate markers may be used to subdivide and target mutation testing. This may not mean an initial screen for all the potential mutually exclusive mutations along one pathway, but a “readout” of its overall activity based on histological, proteomic, or expression profiling.26–22 The goal of the latter must be to ultimately apply specific interventions, and to test the usefulness of the mutation in the overall context of prevention, diagnosis, prognosis, and therapy. In this case, mutations, and epigenetic and expression alterations are then judged by clinical utility.

**ACKNOWLEDGEMENTS**

We acknowledge research funding support from Cancer Research UK.

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Conflict of interest: None declared.

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*Gut* 2005 54: 1209-1211
doi: 10.1136/gut.2005.070946

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