

REASONS FOR ASSESSMENT OF TISSUE

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The contributions that tissue analysis makes to clinical management include diagnosis, staging, prediction of outcome and assistance with selection of therapy. These are often interrelated. The process of tissue assessment may make a new diagnosis or may confirm or refute a suspected or existing clinical diagnosis. There may be pointers towards a cause. Analysis may also reveal additional diagnoses that may be unsuspected. As an example, pathological assessment of an appendiceal specimen most often confirms a suspected clinical diagnosis of acute appendicitis. However, the appendix sometimes contains an incidental neuroendocrine neoplasm, mucinous contains granulomas, raising the possibility of Crohn's disease or infection. Also, a specific cause of abdominal pain other than appendicitis, e.g. endometriosis, may be apparent in the appendiceal tissue. Absence of any histological abnormality raises the possibility of an extra-appendiceal cause. Similarly, biopsies from a patient with inflammatory bowel disease may confirm the diagnosis but may sometimes reveal or suggest an alternative cause of intestinal inflammation such as tuberculosis, amoebiasis, ischaemia or mucosal prolapse. Summary box 11.1

Reasons for analysis of tissue

Tissue analysis also helps, increasingly, to determine or refine treatment and prognosis. For example, the assessment of a breast, lung, colorectal or other major cancer resection specimen helps to confirm the diagnosis but, more importantly, provides crucial information about features such as tumour stage, vascular invasion, perineural invasion and resection margin involvement, which in turn help to predict clinical outcome and determine postoperative treatment. The degree of tumour regression in a resection after neoadjuvant therapy may also have prognostic value. Additionally, pathological assessment of resections helps surgeons and radiologists to audit their accuracy and performance. Molecular pathological analysis of cancer tissue (see Diagnostic molecular pathology) increasingly contributes to management, including diagnostic categorisation, prognostic predictions and selection of drug therapy. The molecular test or group of tests that an oncologist chooses depends on patient status, tumour location, tumour morphology and stage, among other factors. The identification of a particular biomarker may provide an indication for targeted therapy. For example, detection of high microsatellite instability (MSI) in metastatic colorectal carcinoma (CRC) may predict responsiveness to immune checkpoint inhibition.

Burrill Bernard Crohn, 1884–1983, gastroenterologist, Mount Sinai Hospital, New York, NY, USA. Norman Rupert Barrett, 1903–1979, surgeon, St Thomas's Hospital, London, UK. patient. Correlation with the clinical picture and the macroscopic findings enhances the interpretation of pathological changes considerably. Therefore, absence of relevant details may cause unnecessary delays and even errors. For example, radiation therapy can have profound effects on tissue morphology, including mimicry of other inflammatory conditions or neoplasia. Accordingly, a request form with adequate information

should accompany all specimens. Examples of important details include site of biopsy/resection, clinical setting, reasons for the procedure, patient details, medications, relevant risk factors and past medical and surgical history, including previous chemotherapy and radiotherapy. A request form stating 'cancer' or 'Crohn's' is better than a form with no details but is clearly not sufficient. For small and large resection specimens, good quality macroscopic assessment and sampling is an important precursor to microscopic assessment (see Specimen processing).

Diagnosis Confirmation/rejection of a clinical diagnosis Additional diagnoses Classification of neoplasia Classification of non-neoplastic disease Staging of malignancy Prognosis Management Selection of therapy Assessment of response to treatment Cancer screening programmes and related programmes Cervical, bowel, breast, inflammatory bowel disease, Barrett's oesophagus Clinical trial support Audit

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