Psammomatosus carcinoid of the rectum

Psammoma bodies have been documented in carcinoid tumours of the duodenum, but not in carcinoids at other sites in the gastrointestinal tract. We describe a psammomatosus carcinoid tumour in the rectum.

A 50 year old woman presented with a two month history of abdominal pain and bleeding from the rectum. On examination, she had palpor, in addition to fullness and tenderness in the left iliac fossa. There was no evidence of mass or free fluid in the abdomen on abdominal examination. Rectal examination revealed a growth approximately 5 cm above the anal verge. Her erythrocyte sedimentation rate was 80 mm at the end of the first hour and she had a neutrophilic leucocytosis. Her liver function tests were largely unremarkable, except for a mild increase in transaminases (less than two times upper limit of normal). Flexible sigmoidoscopy showed a large ulcerated friable growth involving more than 50% of the bowel circumference.

Biopsies of the tumour showed a neoplasm composed of glandular structures with individual cells having round, uniform nuclei with a “salt and pepper” appearance. Abundant psammoma bodies were seen throughout the biopsy (fig 1). Immunohistochemistry with antibodies to synaptophysin (Dako, Glostrup, Denmark; 1/100 dilution) and chromogranin (Dako; 1/100 dilution) showed cytoplasmic staining. The features were those of a glandular carcinoid, with extensive psammomatous calcification. An abdomino-perineal resection was planned. At surgery, the patient was found to have extensive inoperable disease. A sigmoid proctectomy with colostomy (Hartmann’s procedure) was performed to relieve her obstruction. The sections through the resected colon showed similar histological features—that is, a glandular carcinoid with psammoma bodies. The tumour infiltrated the full thickness of the bowel wall. The regional lymph nodes contained metastatic carcinoid tumour.

Carcinoid tumours of the gut are classified, based on site of origin, into foregut (stomach and duodenum), midgut (small intestine and proximal colon), and hindgut (terminal intestine and rectum) tumours. Psammoma bodies have been described in about 15–20% of carcinoid tumours of the duodenum and are found almost exclusively at the ampulla of Vater. Most of these are histologically characterised by glandular structures and psammoma bodies. These neoplasms often express somatostatin by immunohistochemistry and are termed somatostatinomas. They rarely result in the somatostatinoma syndrome of diabetes mellitus, cholelithiasis, and steatorrhoea, which is commonly seen with pancreatic somatostatin containing neuroendocrine tumours. An association has also been reported between psammomatosus carcinoid of the duodenum and neurofibromatosis. The rectum is not an uncommon site for carcinoids. However, they are conventional carcinoids and display solid sheets, ribbons, and trabeculae or gland-like structures composed of round cells with monotonous nuclei and stippled chromatin. Psammoma bodies have not been documented in rectal carcinoids previously. The reason for this predilection of psammoma bodies for foregut carcinoids is not known.

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References

Figure 1. Psammoma bodies seen in the glandular carcinoid.
Psammomatous carcinoid of the rectum

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Establishing a contract for bench training of specialist registrars in medical microbiology

In common with other medical microbiology laboratories in the UK, the biomedical scientist staff (BMS) at Cambridge have faced an increasing burden of specialist registrar (SpR) training from our own, and other specialties, in the face of falling BMS numbers as a result of efficiency savings and an increasing diagnostic workload. This has led to periods of several months in each year in which SpR training by the BMS has had to be abandoned locally in favour of services with formal laboratory contracts.

Developing a long-term strategy to deal with this problem necessitated the preparation of a business case for SpR training by the BMS, which required an estimation of the local BMS SpR training load. In the first two years, SpRs receive the same basic training in diagnostic methodologies provided for trainee BMS. SpRs spend four months annually in full time bench training, apprenticed to qualified BMS, who spend approximately half their working time on direct supervision of the trainee, equivalent to four months BMS time for each SpR over the first two years. In our own laboratory, we have nine trainees in bacteriology and virology, representing 36 months of BMS time over a notional five year period of SpR training. This calculation assumes that all trainees achieve their annual milestones without delay. In practice, half the trainees sitting the practical component of the examination for membership of the Royal College of Pathologists have required a further six or 12 months of bench training in preparation for a re-sit of the examination. Assuming that half of all the trainees require a nine-month extension, this represents an increase in training load at this stage of 27 months, equivalent to an additional 13.5 months of dedicated BMS time, resulting in a total BMS training time of 42.75 months. From their third year of training, SpRs receive three months of advanced training annually, in focused preparation for the RCPath practical examination. The BMS provide and supervise the processing of simulated clinical samples, to be processed independently by the trainee, on a programme that runs continuously throughout the year. The BMS contract required for this programme amounts to one day each week. This is equivalent to 2.4 months of BMS time annually, in years 3, 4, and 5—that is, 7.2 months over the five year period, in both bacteriology and virology, amounting to 14.4 months in total. Therefore, the total of BMS time required for SpR training is 57.15 months—11.43 months of working time each year. Allowing for six weeks of leave entitlement, this is equivalent to one full time BMS. The experience and skills are of those at grade 2 of the BMS pay spine, for which the salary, with on costs, is £26 663 to £33 737 per annum.

Having established a business case for a full time training contract, we sought a source of funding from those with an interest in training. None was prepared to fund this comparatively large recurrent sum, but three local stakeholders were prepared to fund one third each: the deanery with its responsibility for the delivery of SpR training; the workforce development directorate (the training and development arm of the Strategic Health Authority), with its overarching responsibilities to training laboratory staff; and the local primary care trust, which has an interest in preserving and developing the laboratory’s services.

The core duty of the training BMS (grade 2) is to preserve SpR training. Whenever BMS numbers are inadequate to allow SpR training, this function is assumed in full by the training BMS. At other times, the training of SpRs continues to be shared between all the BMS, and the training BMS engages in related educational activities, which have been tailored to the sources of funding. These duties are, first, supporting and developing the education, training, and research of SpRs in medical microbiology, as directed by the deanery’s programme director in medical microbiology. Second, duties supporting the laboratory training manager in the provision of training of SpRs, BMS, medical laboratory assistants, and visitors to the laboratory. Third, maintaining the training BMS’s own professional status and microbiological skills by participating in the rota for provision of the laboratory’s routine diagnostic service, under the direction of the laboratory manager. Because the job description contains three major components with different line managers, a clear division of the working week is necessary to ensure realistic expectations and harmonious working relations. The time allocated to the first, second, and third duties are three, one, and one day each week, respectively.

We resolved our local crisis in the provision of SpR technical training by negotiating a training contract, analogous to the laboratory’s service contract, with local parties with a training interest. We recommend this model, which we believe to be unique in the UK, to laboratories experiencing difficulties similar to our own.
Audit of the histological definition of cervical transformation zone

We have noticed in our routine practice that there is some variation in the histological definition of the transformation zone. In another study, the incidence of myelodysplastic syndrome (MDS) was found to be 10.9% in an analysis of the incidence of myelodysplastic syndrome in long-term outcome after allogeneic transplantation. The diagnosis of AML was confirmed by the presence of squamous and columnar epithelium in continuity and/or the presence of squamous epithelium with underlying glands.

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References

BOOK REVIEWS

WHO Classification of Tumours: Pathology and Genetics of Tumours of the Breast and Female Genital Organs


Pathology and Genetics of Tumours of the Breast and Female Genital Organs is the fifth volume of the series by the World Health Organisation on the Classification of Tumours. This book fills a gap in the availability of a ready reference manual for tumours of the breast and the female genital tract. It provides a well written, concise, and comprehensive reference book for pathologists and oncologists involved in these disciplines.

The authors use a very well researched approach to the topics included in the manual and the reader friendly layout of the material provides clarity to complex issues in diagnostic pathology. I found that the contents of the “blue boxes” were most confirm sampling of the transformation zone because of the absence of glandular epithelium. The fact that some of the respondents in this audit chose multiple options probably reflects the difficulty of defining this dynamic zone. We would cautiously recommend that the transformation zone should be defined by the presence of squamous and columnar epithelium in continuity and/or the presence of squamous epithelium with underlying glands.

Table 1 Questionnaire results

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Table 2 Combinations

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<td>A+B+C+E</td>
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<td>Total</td>
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</table>
informative and provided helpful diagnostic hints. One of the chapters on breast pathology shows an individual author’s bias on the subject, with the inclusion of different terminologies in the already confused subject of intraepithelial proliferative lesions of the breast. Fortunately, this trend does not persist in the remaining chapters of the book.

The immunological and molecular profiles of different tumours is exhaustive, well presented, and provides a valuable diagnostic aid to pathologists in practice, as well as a ready reckoner to research oriented clinicians.

The inclusion of the final section on inherited tumour syndromes is a valuable source of information. A final word of praise on the excellent images of tumours, which will be a delight to the picture matching eclectic. I highly recommend this book and consider it an indispensable reference volume for both pathologists and oncologists.

R Vaipeyi

Colour Atlas of Anatomical Pathology, 3rd Edition


This atlas has formed an essential part of the training of both new anatomical pathology registrars and pathology undergraduate students. It is probably more essential now given the declining rate of necropsies and the slight ambivalence and aversion of new trainees towards necropsies in general. Knowledge of gross pathology is a fundamental component of anatomical pathology and this atlas is the next best thing to seeing fresh, wet macroscopic pathology.

The illustrations in the book are of uniform high quality and portray the pertinent pathological features. I have no real criticisms of this compendium of gross photos and would recommend that it be on the shelves of training departments and medical school libraries.

R Chetty


Edited by J L Callum, P H Pinkerton. 2003 (paperback). Published by Sunnybrook and Women’s College Health Sciences Centre, pp 115. ISBN 0 96813 4424

This is a useful book directed at medical staff who are responsible for the transfusion of blood, blood components, and blood products to patients. It is compact and would fit easily into a pocket. The approach is very practical and will provide clinical staff with the information necessary to answer patients’ questions. The style is didactic but important statements are supported by references provided in small print in an appendix. This is a sensible approach because the uncluttered pages are easy to read and can be referred to quickly, whereas the reader who wants to know more is not denied the necessary evidence base. The book links to a useful website, which is designed for those seeking a greater depth of information.

The detailed information given is specifically applicable to Canada. Much, but not all, is readily transferable to other countries. Anyone wishing to provide this book for the use of clinical staff might wish to have an insert giving details of local dictation systems. Two blank pages at the back of the book would permit this to be pasted in.

On a personal note, I do not like the title—it would make me less likely to recommend this book.

B J Bain

Forensic Pathology Reviews: Volume 1

Edited by M Tsokos. Published by Humana Press, 2004, £52.23 (hardback), pp 384. ISBN 1 58829 4145

As a rule, new forensic pathology books do not sprout original or unique information that is not already present in the print medium. It was with a sense of excitement that I agreed to review this book, which according to the cover spoke of an impressive list of international collaborators. The selection of contributors is excellent; an example would be the chapters on neonaticide written by the prolific pen of Roger Byard, who was also responsible for coauthoring the chapter on sudden infant death syndrome (SIDS) with Harry Krous.

The book is not presented in a standard “A to Z” form but instead has 15 chapters dealing with disparate issues, varying from common topics such as SIDS to unusual subjects such as ileopsoas haemorrhage. These chapters are placed into the subheadings of: death from environmental conditions, trauma, neurotraumatology, forensic neuropathology (separating these last two topics is not warranted in my opinion), sudden death from natural causes, child abuse, neglect and infanticide, SIDS, infectious diseases, death scene investigation, maternal death in pregnancy, iatrogenic injury, toxicology, and forensic differential diagnosis. The major advantage of such a format with short precise individual chapters is that one can take random “dips” into the book looking at topics that may catch your eye at a specific moment in time.

The text is current and contains a useful sprinkling of hints and pearls that would be of use in death investigation. A good example is the practical approach to sudden cardiac death in chapter 9.

A shortcoming, in my opinion, is the lack of colour illustrations. I know that this would increase the costs but perhaps the editor should consider a separate companion CD that could have additional text and illustrations.

It is a great compliment to the editor that the book flows seamlessly from one chapter to another, despite the diversity of the contributors. This is a well compiled book, which is a refreshing addition to the forensic pathology genre. I would recommend this book to anyone in the medico–legal arena who has an interest in forensic pathology. The book has something to offer to both the novice and the expert. It would make a useful reference in any departmental or institutional library. I look forward to volume 2.

M A Dada

RETRACTION


At the time of writing, the authors were not aware that the patient had undergone a panhysterectomy for a possible ovarian tumour six years previously. Details of surgery and pathology are still not available. In view of this, and the negative cytokeratin 7 and positive cytokeratin 20 results (kindly performed by Dr J Aidan Carney, Mayo Clinic, Rochester, Minnesota, USA), we realise that the neoplasm is not a psammomatous carcinoid.

CALENDAR OF EVENTS

Full details of events to be included should be sent to Maggie Butler, Technical Editor JCP, The Cedars, 36 Queen Street, Castle Hedingham, Essex CO9 3HA, UK; e-mail: maggie.butler2@btopenworld.com

Practical Pulmonary Pathology
26–29 July 2005, Royal Brompton Hospital, London, UK
Further details: Professor B Corrin, Brompton Hospital, London SW3 6NP, UK. (Fax +44 (0)20 7 351 8293; e-mail b.corrin@ic.ac.uk)

Association of Clinical Pathologists’ National Scientific Meeting
16–17 June 2005, Royal College of Physicians, Londondon, UK
Further details: ACP Central Office, 189 Dyke Road, Hove BN3 1TL, UK. (Tel +44 (0)1273 775700; e-mail info@pathologists.org.uk)

Breast Diagnostic Histopathology Update
22–23 September 2005, Hammersmith Hospital and Imperial College, London, UK
Further details: Wollson Conference Centre, Hammersmith Hospital, Du Cane Road, London W12 ONN, UK. (Tel +44 (0)20 8383 3117/3227/3245; Fax +44 (0)20 8383 2428; e-mail wcc@ic.ac.uk)