

Abstracts from the 1998 Annual Meeting of the Hong Kong Division of the International Academy of Pathology

MEMBRANOUS GLOMERULONEPHRITIS WITH MESANGIAL IGA DEPOSITION: AN UNUSUAL CASE OF NEPHROTIC SYNDROME

IK, W. Chan, T. M. Chan

Queen Mary Hospital, Hong Kong

A 30-year-old man who had been in good past health presented with the nephrotic syndrome, a skin rash, and deranged liver function. Renal biopsy showed diffuse granular deposition of IgG, C3, and C1q along the capillary basement membrane. Mesangial IgA and C3 deposition was also present. Subepithelial and mesangial electron dense deposits were readily identified. Serological markers for autoimmune diseases were absent. The biopsy findings were those of stage II membranous glomerulonephritis and mesangial IgA nephropathy. Skin biopsy showed dermal lymphocytic infiltration which raised the suspicion of cutaneous malignant lymphoma. The nephrotic syndrome spontaneously remitted without specific treatment. The underlying cause of the symptoms was revealed by a positive VDRL test, a history of recent venereal exposure, and self-healing genital ulcer. Coexistent membranous glomerulonephritis and primary mesangial IgA nephropathy are rarely encountered. Syphilis is an uncommon but well documented cause of membranous glomerulonephritis associated with mesangial IgA deposition. Given these renal biopsy findings, syphilis must be considered in the clinical diagnosis so that specific therapy to this curable disease would not be delayed.

GLOMERULAR DISEASES IN HONG KONG: A BIOPSY STUDY

IK, W. Chan

Queen Mary Hospital, Hong Kong

A prospective analysis of 1,629 consecutive percutaneous renal biopsies performed on native kidneys and examined in the Queen Mary Hospital, Hong Kong, during 1993-1997, showed that IgA nephropathy (n=317) was the most frequently biopsied glomerular lesion. This was closely followed by lupus nephritis (n=289). A highly unusual case of truncated IgA heavy chain deposition nephropathy was encountered. Minimal change nephrotic syndrome (n=125) with its variant IgM nephropathy (n=11) was the commonest glomerular disease responsible for the nephrotic syndrome in the overall patient population and in children under the age of 12. Male patients outnumbered the female by a ratio of 1.8:1 in minimal change nephrotic syndrome, IgA nephropathy and membranous glomerulonephritis affected about equal numbers of male and female patients. Female patients biopsied for lupus nephritis were more than 14 times as many as the male. This basement membrane nephropathy in females tripled that in the males. The commonest reason for renal biopsy was for the investigation of asymptomatic proteinuria (n=717). This was followed in frequency by the nephrotic syndrome (n=386), microscopic haematuria (n=155), chronic renal failure (n=73), gross haematuria (n=46), and acute renal failure (n=16).

VAGINAL METASTASIS AS THE FIRST PRESENTATION OF RENAL CELL CARCINOMA

T. Hlaing, S. L. Wong, L. W. Chan, P. C. L. Choi, K. F. To, M. Y. Yu
The Chinese University of Hong Kong, Prince of Wales Hospital, Shatin, Hong Kong

Vaginal metastasis as the first presentation of occult renal cell carcinoma has been sparsely reported in the literature. Histologically, this may be mistaken as müllerian clear cell carcinoma.

We report a case of metastatic renal cell carcinoma, firstly presented in a 77-year-old Chinese lady with postmenopausal bleeding. Per vaginal examination revealed a vaginal polyp. Biopsy showed clear cell carcinoma forming solid sheets and lobules of polygonal clear cells without distinct mucicoid and is separated by thin fibrovascular septae. Hobnail cells or

pavement-like cells, typical of müllerian type clear cell carcinoma, are absent. Since primary vaginal clear cell carcinoma is extremely rare in postmenopausal women, metastasis has to be excluded including kidney and other parts of female genital tract. Histochemical and immunohistochemical results are as follows: the tumor cells contain abundant cytoplasmic lipid (oil-red-o positive), glycogen (PAS -positive, PASD -negative), and are diffusely positive for epithelial markers (AE1/AE3, Cam 5.2). However, no neutral mucin is present. (PASD -negative, Mucicarmine -negative). The presence of cytoplasmic lipid is in favor of metastatic clear cell carcinoma of renal cell origin. Subsequent CT SCAN examination confirmed presence of a 7 cm tumor mass at right kidney.

This case demonstrates that renal cell carcinoma could be presented as peculiar secondary including vaginal metastasis.

AGGRESSIVE ANGIOMYXOMA—A CYTOGENETIC AND IMMUNOHISTOCHEMICAL STUDY

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Department of Pathology, PLA Chinese General Hospital, Beijing and *Department of Pathology, The University of Hong Kong, Queen Mary Hospital

Aggressive angiomyxoma is a rare, slowly growing gynecologic soft tissue neoplasm most commonly found in premenopausal women. This disease usually occurs in the vulva, perineal and pelvic region with a propensity for local infiltration and multiple recurrences. Cytogenetic information about aggressive angiomyxoma are few. Besides the rare occurrence of the tumor, technical difficulties associated with conventional karyotype studies may also be responsible. In order to investigate into possible numerical chromosomal changes in aggressive angiomyxoma, the technique of chromosome *in situ* hybridization (CISH) was performed using DNA probes specific for the pericentromeric regions of chromosome 11, 12, 16, and X. Immunohistochemical analysis of estrogen (ER) and progesterone (PR) receptors, MIB1 (Ki67), bcl-2, p53, and c-erbB2 was also performed to characterize the proliferative capacity of the tumor and the potential role of hormone receptors and oncogenes in its development. Archival paraffin-embedded tissue sections of five aggressive angiomyxoma and two vaginal polyps were used. Normal copy numbers of chromosome X, 12, 16, 11 were found in all five cases of aggressive angiomyxoma. High level of ER and PR expression was detected in these tumors. This, together with the relatively site-specific location of aggressive angiomyxoma and the strong female predominance, tend to suggest that hormonal factors are important in the evolution of this neoplasm. Low Ki67 index was detected and this correlates well with the low mitotic figures found in this tumor and its slow-growing nature. All the aggressive angiomyxomas we studied were negative for c-erbB2 and p53 (wild or mutant type) gene products. Such negative findings imply that these oncogenes may not be important in its pathogenesis.

HUMAN PAPILOMA VIRUS (HPV) SUBTYPES IN UTERINE CERVICAL NEOPLASMS IN HONG KONG

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Human Papilloma Virus (HPV) has been associated with uterine cervical neoplasms and also squamous carcinomas in many other organs. The aim of this study is to use sensitive polymerase chain reaction (PCR) to evaluate the prevalence of different HPV subtypes in uterine cervical neoplasms in a district general hospital in Hong Kong.

A total of 120 patients were retrieved from the surgical pathology archives from 1996-1998. This included 21 condylomas, 22 CIN I, 24 CIN 2, 29 CIN 3, and 24 carcinomas. The mean and median age of each group

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impairment in spatial construction but great strength in face processing and sociability By capitalizing on these dissociations
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