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JOURNAL ARTICLE

Distribution and tissue expression of semenogelin I and II in man as demonstrated by in situ hybridization and immunocytochemistry

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Semenogelin I and II (SgI, SgII) are two separate gene products of chromosome 20 with extensive (80%) identity in primary structure.

They are mainly responsible for immediate gel formation of freshly ejaculated semen. Degradation of SgI and SgII is due to the

proteolytic action of prostate-specific antigen (PSA); it results

within 5-15 minutes in liquefaction of semen and release of progressively motile spermatozoa. By means of cDNA cloning and Northern blots, SgI and SgII transcripts have previously been shown to be abundant in human seminal vesicles, but SgII alone is suggested to be expressed at low levels in the epididymis. To characterize the expression and tissue distribution of SgI and SgII in greater detail, we produced monoclonal immunoglobulin Gs (IgGs for immunocytochemistry (ICC) and specific [35S]-, digoxigenin-, or alkaline phosphatase-labeled 30-mer antisense probes to SgI and SgII for in situ hybridization (ISH). Immunocytochemical staining for both SgI and SgII, and ISH detection of both SgI and SgII transcripts, were demonstrated in the cytoplasm of seminal vesicle epithelium. ISH showed SgII alone to be expressed in the epithelium of the epididymal cauda. Neither ICC nor ISH yielded any evidence of SgI or SgII expression in caput or corpus epithelium or in any stromal cells of the epididymis. Consistent with our previous findings using polyclonal IgG, monoclonal anti-SgII SgII IgGs identified epitopes on the posterior head, midpiece, and tail of ejaculated spermatozoa. Spermatozoa in the epididymal cauda were also immunoreactive, but those in the caput or corpus region of the epididymis as well as those in the testis were negative. As shown by ICC, neither SgI nor SgII were expressed in the testis, the prostate, the female genital tract, or other normal human tissue specimens. Although the significance of Sg attachment to epididymal and ejaculated spermatozoa remains to be established, monoclonal anti-Sg IgG might prove useful in establishing the origin of seminal vesicle tissue components in prostate core biopsies or other biopsy specimens.

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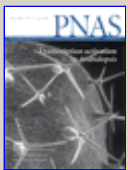
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