Characterization of Adenocarcinomas of the Dorsolateral Prostate Induced in Wistar Rats by N-Methyl-N-nitrosourea, 7,12-Dimethylbenz(a)anthracene, and 3,2'-Dimethyl-4-aminobiphenyl, following Sequential Treatment with Cyproterone Acetate and Testosterone Propionate

Maarten C. Bosland, Menk K. Prinsen, Theo J. M. Dirksen, and Ben J. Spit

Department of Biological Toxicology, TNO-CIVO Toxicology and Nutrition Institute, P. O. Box 360, 3700 AJ Zeist, The Netherlands

ABSTRACT

Carcinomas of the rat prostate induced by a single injection of N-methyl-N-nitrosourea, 7,12-dimethylbenz(a)anthracene, and 3,2’-dimethyl-4-aminobiphenyl, after sequential treatment with cyproterone acetate and testosterone propionate, were evaluated as potential animal models for prostatic cancer. All ten carcinomas examined were located in the dorsolateral prostate region and did not involve the distal parts of the seminal vesicles and coagulating glands. The incidence of urinary obstruction leading to the animals’ death was 6 of 10 rats, and metastases in the lung, abdominal lymph nodes, and/or liver also occurred in 6 of 10 rats. The tumors were invasive adenocarcinomas, showing frequent peripheral invasion and a variable degree of differentiation. There were ultrastructural similarities with human prostatic carcinomas, such as intracellular lumina. Plasma acid phosphatase was increased. Enzyme histochemical analysis revealed similarities with the Dunning R3327H and -HI prostatic carcinomas but was not helpful in determining the site of origin of the tumors. The gross and microscopic appearance of the tumors and the observation of preneoplastic lesions exclusively located in the dorsolateral prostate suggest this lobe as site of origin of the carcinomas. Preneoplastic lesions (n = 9) included atypical hyperplasias (n = 5) and lesions with all histological characteristics of carcinoma except for local invasion and metastases, which were classified as carcinomas in situ (n = 4). Although androgen sensitivity could not be assessed, the observed characteristics of the tumors [their long latency time (46-80 weeks), the presence of preneoplastic lesions, and the short duration of the treatment, leaving the animals intact] all indicate that the present approach is a valid animal model for the study of prostatic carcinogenesis.

INTRODUCTION

Animal model systems of carcinogenesis can greatly contribute to our understanding of human carcinogenesis. Using such models, carcinogenic agents can be identified, factors that modify the carcinogenic process can be studied under controlled conditions, and the mechanism of action of these agents and modifying factors can be addressed. It is of utmost importance that such models of carcinogenesis be well characterized and resemble their human counterparts in such key features as morphology and histogenesis, latency time and growth rate, and biological behavior.

Desired properties of animal models for the study of biology and therapy of prostatic cancer have been discussed by Coffey and Isaacs (1). The requirements for a model of the genesis of prostatic cancer, however, have not been well defined in the literature. Based on the generalized properties of human prostatic cancer summarized by others (1-4) and on what is known about the etiology and pathogenesis of this neoplastic disease (5, 6), we have derived the following requirements for a rodent model for prostatic carcinogenesis: The adenocarcinomas should originate exclusively from the dorsolateral prostate, because that part of the rodent prostate is most likely homologous to the region of the prostate from which prostatic cancer originates in man (8-11), as pointed out elsewhere (5, 12). The great majority of human carcinomas are androgen sensitive and respond to hormonal (estrogen) therapy followed by relapse to a hormone-insensitive state. This feature should also be a hallmark of an appropriate animal model. Biochemical characteristics of the carcinomas should be comparable to those of the human tumor, e.g., the presence of androgen-metabolizing enzymes and androgen receptors. The slow growth rate and the pattern of metastatic spread to lymph nodes, lung, liver, and bone, characteristic of human prostatic cancer, should be features of such a model. Although a slow growth rate leading to long latency times is usually regarded as a disadvantage for practical animal models for carcinogenesis, a model system for rapid induction of prostate carcinomas would compare unfavorably with the human situation. Two typical characteristics of advanced human prostate cancer, i.e., elevation of serum acid phosphatase and urinary obstruction, should also occur. There is a great diversity in several of these properties of human prostatic cancer, particularly in the degree of histological differentiation, the pattern of progression and, to a lesser degree, hormone sensitivity. An appropriate animal model should mimic this diversity. Because of the central importance of androgen sensitivity for many of the human prostatic carcinomas, it is essential for a prostatic cancer model that the tumors can be induced in intact animals, i.e., animals that are not castrated or otherwise hormonally manipulated. Environmental factors affecting the progression from early noninvasive, so-called latent carcinoma to invasively growing adenocarcinomas are probably critical determinants of human prostatic cancer risk (5). It is, therefore, highly desirable that early stages of prostatic carcinogenesis be identifiable in an animal model and that induction is possible with a single or short-term exposure to a carcinogenic treatment, so that aspects of promotion and progression can be studied.

In a companion paper (12), we describe the chemical induction of carcinomas and preneoplastic lesions in the rat prostate, using an approach that may provide a useful animal model to study prostatic carcinogenesis. The purpose of this paper is to describe in detail morphological and biological characteristics of these carcinomas and preneoplastic lesions. Furthermore, to aid in determining the accessory sex gland site of origin of the induced carcinomas, their enzyme histochemical properties were compared to those of normal rat accessory sex glands and

Received 2/16/89; revised 6/9/89, 9/25/89; accepted 10/26/89.

The cost of publication of this article was defrayed in part by the payment of page charges. This article must therefore be hereby marked advertisement in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

1 Supported in part by grants CIVO 80-3 and CIVO 84-3 from the Netherlands Cancer Foundation, Koningin Wilhelmina Fonds.
2 Present address: Institute of Environmental Medicine, New York University Medical Center, 550 First Avenue, New York, NY 10016. To whom requests for reprints should be addressed. This work was submitted as part of a dissertation to the State University of Utrecht, The Netherlands, in partial fulfillment of the requirements for the Doctorate Degree.

3 Nomenclature of the male accessory sex glands is according to the system of Jesik et al. (7).
of the transplantable Dunning R3327H and R3327HI prostate carcinomas which are of proven dorsolateral prostate origin (13, 14). These two carcinoma lines were also used as positive controls for determining elevation of plasma acid phosphatase by the induced prostate carcinomas.

MATERIALS AND METHODS

Source of Tissue and Histological Procedures. The dorsolateral prostate adenocarcinomas and preneoplastic lesions that are discussed here were obtained from animals killed when moribund in the induction experiments described in a companion paper (12), as is summarized in Table 2. In these experiments, the carcinogens MNU, DMB, and DMABP were administered by a single injection during stimulation of prostatic cell proliferation by sequential treatment with cyproterone acetate and testosterone propionate. These procedures are described in detail in the companion paper (12). Necropsy procedures, fixation, tissue trimming, and histological methods used have also been detailed in this companion paper (12). For ultrastructural analysis, 3–5 mm³ pieces of tumor tissue were fixed in 3% glutaraldehyde solution in 0.1 M sodium cacodylate (pH 7.2) at 4°C for 17 h and stored in 0.1 M sodium cacodylate buffer at 4°C until further processing. Postfixation was carried out in a 4% osmium tetroxide solution in 0.1 M sodium cacodylate for 1 h. After dehydration, the tissue pieces were embedded in glycid ether-Araldite. Semithin sections were stained with methylene blue. Ultrathin sections were contrasted with uranyl acetate and lead citrate and examined using a Philips EM 201 G at 50 kV.

Plasma Acid Phosphatase. From 9 animals with macroscopically visible prostate tumors, blood was collected in heparinized tubes by cannulating the abdominal aorta while the animals were under ether anesthesia. Plasma was obtained by centrifugation (2000 × g), using Sure Sep (Organon Technika, Charlotte, NC), and stored at -20°C until assayed. Plasma acid phosphatase activity was determined according to Seiler and Nagel (15), using α-naphthyl phosphate as substrate. The tartrate-inhibitable fraction was calculated by subtracting the enzyme activity obtained in the presence of 0.0135 M L(+)-tartrate from the total activity.

Enzyme Histochemistry. From a number of dorsolateral prostatic tumors a representative, small piece (approximately 0.5 cm³) was sampled for enzyme histochemistry, frozen in isopentane over liquid nitrogen, and stored at -80°C. Pieces of the Dunning R3327H and R3327HI tumors and of normal accessory sex glands of untreated young adult male Cpb:WU rats were sampled in the same manner for comparative studies. Ten-μm-thick cryostat sections were prepared and histochemical reactions for the following enzymes were carried out, using appropriate positive control tissues and inhibitors (16): GGT according to Rutenberg et al. (17); isopropyl alcohol dehydrogenase using the procedure of Hardon (18); acid phosphatase by the lead nitrate method of Gomori with β-glycerophosphate as substrate (16); succinic dehydrogenase as described by Wachstein and Meisel (19); glucose-6-phosphatase as described by Wachstein and Meisel (19); acid phosphatase as described by Wachstein and Meisel (19); oxidative polyamine metabolism by the method of Wachstein and Meisel (19); and 5'-nucleotidase according to Nachlas et al. (22); and alkaline phosphatase using Burstone’s method with naphthol AS-BI as substrate (21); LAP according to the procedure of Nachlas et al (22); and 5'-nucleotidase by the method of Wachstein and Meisel (23). This panel of enzyme histochemical assays was selected to aid in determining the site of origin of the carcinomas, because these provide the greatest possible discrimination in histochemical (see Ref. 13) and biochemical (see Ref. 14) characteristics among the rat ventral, lateral, and dorsal prostate lobes. Staining intensity was scored blindly as absent, minimal, slight, moderate, or marked, and the distribution of the staining was recorded as focal or diffuse.

Dunning R3327H and R3327HI Prostatic Carcinomas. The Dunning R-3327H (androgen-sensitive) and R3327HI (androgen-insensitive) prostatic adenocarcinomas (see Ref. 14) were used as comparison for the chemically induced prostate carcinomas. These tumors were generously donated by Dr. N. H. Altman (Papanicolau Cancer Research Institute, Miami, FL) together with breeding pairs of the Copenhagen (COP) and Fischer (F344) rat strains. The tumors were maintained in young adult male F344 × COP F; rats, the R3327H tumor in intact animals and the R3327HI tumor in castrates. The tumors were transplanted before they reached a volume of 4 cm³, by implanting pieces of approximately 0.5 mm³ s.c. on both flanks of each animal using a trocar method (16–18-gauge needle; 1 ml L-15 medium). Histological appearance and hormone sensitivity were routinely examined to monitor tumor stability and appeared comparable to what has been reported by others (13, 14). Material from the R3327HI tumor for enzyme histochemistry and plasma for acid phosphatase determination were taken from intact animals bearing tumor transplants.

Statistical Methods. The plasma acid phosphatase data were statistically evaluated using Student’s t test and linear regression analysis. The correlations between tumor size and acid phosphatase activity and between tumor size and latency were determined by calculating the Spearman rank correlation coefficient. A one-tailed P value of 0.05 or smaller was considered significant.

RESULTS

Characteristics of ten carcinomas and nine preneoplastic epithelial lesions of the dorsolateral prostate are described in this report (Table 1). These lesions had been induced by the chemical carcinogens MNU, DMB, and DMABP as described elsewhere (12). Nine of the ten carcinomas were grossly apparent (Table 1). The latency time (time after carcinogen administration) of these 10 carcinomas was 64 ± 4 (SEM) weeks (range, 46–80 weeks).

Gross Observations

Six of the nine grossly visible tumors (67%) were symmetrically located in the region of the dorsolateral prostate, involving not only the dorsolateral prostate lobes but also the base of both seminal vesicle/coagulating gland complexes. Two of these tumors had caused bilateral urinary obstruction and hydronephrosis, whereas the other four had caused unilateral obstruction of the urinary flow (Fig. 1). Three tumors (33%) were located on one side of the dorsolateral prostate involving the base of the seminal vesicles and coagulating glands on that side. These tumors did not cause urinary obstruction, and they were smaller (diameter, 1.67 ± 0.17 cm; range, 1.5–2.0 cm) than the symmetrically located tumors (diameter, 3.40 ± 0.51 cm; range, 2–5 cm). There was no statistically significant correlation between tumor size and tumor latency. Smaller tumors were single masses, whereas some larger tumors had a multinodular ap-
characterization of chemically induced rat prostate carcinomas

Fig. 1. Tumor located in the region of the dorsolateral prostate. This tumor (T), diagnosed as adenocarcinoma, grossly involved all of the dorsolateral prostate and the base of the seminal vesicle-coagulating gland complexes (S), which were swollen. The ventral prostate (inset; V) was not involved. The urinary bladder (B) and ureters (U) were distended, because of obstruction of the urinary flow, and the kidneys (K) were mildly hydropnephrotic.

appearance. The tumors all had a firm consistency and were white-gray and homogeneous of texture on the cut surface. Distinct central necrosis was present in two large tumors, and two others showed a translucent exudate on the cut surface. The seminal vesicles and coagulating glands that were involved in the tumor processes were swollen, suggesting obstruction of drainage of secretion (Fig. 1). The ventral prostate lobes were never grossly involved in the tumors (Fig. 1).

Histopathology

Carcinomas. All ten tumors were adenocarcinomas with a glandular growth pattern. The carcinomas histologically involved the dorsal and/or lateral prostate lobes and at least one other accessory sex gland, in all cases. Most tumors histologically involved all of the accessory sex gland structures that are associated with the dorsolateral lobes: ampullary glands, seminal vesicles and coagulating glands, including the excretory ducts of these glands. Even the one microscopic carcinoma involved several of these structures (Fig. 2). (Interestingly, a microscopic carcinoma exclusively located in the dorsolateral prostate was observed in an experiment using the same approach to induce prostatic cancer as was applied here (12), with MNU as inducing agent.) All ten carcinomas grew invasively, and several showed perineural invasion (Fig. 3). The architecture of the dorsolateral prostate region was often markedly disturbed.

The degree of histological differentiation of the carcinomas varied considerably, both within a single carcinoma and between carcinomas (Table 1), ranging from well (Fig. 4) to moderately (Fig. 5) to poorly (Fig. 6) differentiated. The carcinomas were graded into these three categories according to the growth pattern that predominated (4). Carcinomas with a generally

5 M. C. Bosland, unpublished observation.
CHARACTERIZATION OF CHEMICALLY INDUCED RAT PROSTATE CARCINOMAS

well developed acinar growth pattern were considered well differentiated, whereas carcinomas with predominantly poorly developed acini were classified as moderately differentiated. Carcinomas with little evidence of acinar structures were considered poorly differentiated. With decreasing degree of differentiation there was an increase in cellular anaplasia: loss of cellular polarity; variation in cellular and nuclear size and shape and in cytoplasmic/nuclear ratio (pleomorphism); and changes in nuclear chromatin distribution and in staining properties and texture of cells and nuclei. There was no apparent correlation between degree of differentiation and size or latency time of the carcinomas.

General features of all carcinomas listed in Table 1 were the presence of abundant fibrous stroma and acini consisting of a single layer of neoplastic cells without basal cells (Figs. 4–6). A marked mixed inflammatory cell infiltrate in interstitium and acinar lumina was often but not always present (Figs. 5 and 6). In a very well differentiated microscopic carcinoma found in the earlier mentioned experiment, the mildly atypical neoplastic epithelium had somewhat retained its cellular polarity and secretory properties. The main differences with normal dorsolateral prostate were a marked focal increase in the number of acini, a much smaller size of the neoplastic acini, which were disorderly arranged, sometimes back-to-back, and a slightly increased amount of fibromuscular stroma (Fig. 4). The three well differentiated carcinomas listed in Table 1 had acini that were small and varied from round or oval to tubular and that displayed a disturbed acinar-stromal relation. The epithelium had mostly lost its normal cellular polarity and showed little or no evidence of secretory activity. The cells and nuclei were somewhat pleomorphic. The five moderately differentiated carcinomas predominantly consisted of small acini, mostly tubular or slit-like, with no evidence of secretory activity (Fig. 5). The
cells had completely lost normal polarity and showed a moderate degree of pleomorphism. They were generally cuboidal and slightly hyperbasophilic with enlarged, hypochromatic nuclei, as was also observed for most well and all poorly differentiated carcinomas. In the two poorly differentiated carcinomas most moderately to markedly anaplastic cells were arranged in small clusters or strands amid abundant stroma, and mitotic activity was higher than in well or moderately differentiated carcinomas in which very few mitotic figures occurred (Fig. 6).

Electron microscopy revealed fewer differences between well, moderately, and poorly differentiated carcinomas than light microscopy. Even the one poorly differentiated carcinoma examined showed a tendency to form lumina, but the microvilli on the luminal side of the cells were less well developed in moderately to poorly differentiated carcinomas than in well differentiated tumors (Fig. 7). Desmosomes and junctional complex-like structures were always found (Fig. 7), and sometimes there were obvious tonofilaments (Fig. 8). Structures suggestive of secretory vacuoles (Figs. 7 and 9), which normally occur in prostate secretory epithelial cells (24), were present in all tumors, even in histologically moderately to poorly differentiated carcinomas. Sometimes hyperplastic Golgi complexes, lipid droplets, and lysosomes were seen (Fig. 8). Polarization of the secretory and Golgi structures toward the apical side of the cell found in normal prostate secretory epithelium (24) was disturbed. Mitochondria were often changed, showing loss of cristae and occasional membranous inclusions (Figs. 7 and 8). Fragments of a basement membrane were present in well differentiated carcinomas but were less conspicuous or absent in moderately and poorly differentiated tumors. Distinct features

---

**Fig. 7.** Electron micrograph of a moderately-poorly differentiated adenocarcinoma of the dorsolateral prostate. There are poorly developed lumina (L) with microvilli and some secretory vacuole-like vesicles at the luminal side of the cells. Note the many mitochondria with disrupted cristae and membranous inclusions (arrow) and the junctional complex- and desmosome-like structures at the cell junctions. × 7200.

**Fig. 8.** Electron micrograph of a dorsolateral prostate adenocarcinoma cell showing tonofilaments (t), mitochondria with altered cristae (m), lipid droplets (l), lysosome-like structures (ly), and cytoplasmic inclusions in the nucleus. × 9500.

**Fig. 9.** Electron micrograph of a poorly differentiated dorsolateral prostate adenocarcinoma. One of the cells has intracellular lumina (L), one of which is filled with membranous structures. Note the highly irregular shape of the nuclei and the many secretory vacuole-like vesicles. There are stromal polymorphonuclear inflammatory cells (l), fibroblasts (f), and collagen (c). × 6000.
were marked nuclear abnormalities: irregular shape; infoldings of the nuclear membrane and cytoplasmic inclusions; and clumping and condensation of chromatin along the nuclear envelope (Figs. 7–9), regardless of the degree of histological differentiation. The presence of intracellular lumina with distinct microvilli was another striking feature of some carcinomas (Fig. 9). The absence of basal cells seen by light microscopy was confirmed. The chemically induced carcinomas were ultrastructurally very similar to the Dunning R3327H and -HI carcinomas.

**Carcinoma in Situ.** Lesions that had all characteristics of carcinoma but lacked invasive growth were classified as carcinoma in situ. These lesions were small, occupying an area of the size of one to three acini. The acinar lumen was completely replaced by microglandular proliferation of mildly atypical cells (Figs. 10 and 11), which were indistinguishable from cells observed in well differentiated carcinomas. The glands were composed of a single layer of atypical cells without obvious basal cells. There was an often marked desmoplastic reaction surrounding the lesions, sometimes suggesting early invasive growth followed by recapsulation (Fig. 11) and infiltrative growth from one acinus to an adjacent one (Fig. 10). There were occasional stromal septa within the lesions separating glandular structures, but back-to-back and cribiform-like arranged microglands were the predominant growth pattern (Fig. 11). Some inflammatory infiltrate was usually present (Fig. 11). The four such lesions seen in this study were all exclusively located in the dorsal and lateral prostate lobes and tended to be localized in the periphery of the gland (Fig. 10).

**Atypical Hyperplasia.** Lesions consisting of a hyperplasia of atypical cells that did not completely occupy one acinus were classified as atypical hyperplasia and thus coexisted with normal glandular epithelium. In all five cases observed in this study there was a mild increase in fibromuscular stroma surrounding the lesion. Two of these lesions had all cytological characteristics of carcinoma in situ, particularly the growth pattern of a single layer of cells forming microglands, and the apparent absence of basal cells. Two other lesions consisted of disorderly piled up, enlarged, hypochromatic cells that had partly lost their normal polarity and that had hypertrophic pale nuclei (Fig. 12). One lesion had characteristics of both types. The lesions were all located in the acini, not the ducts of the dorsal or lateral prostate lobes.

**Metastases and Clinical Observations**

Six of the ten carcinomas had metastasized to the lung (Fig. 13; Table 1). One had also disseminated the liver, and two had also metastasized to other structures in the abdominal cavity, such as mesenterial lymph nodes. Regional pelvic lymph node involvement was frequent; the available material did not allow determination of a precise incidence. No attempt other than a careful necropsy was made to detect bone metastases; none were grossly apparent. All six metastasizing carcinomas were moderately or poorly differentiated; well differentiated carcinomas did not metastasize. No correlation was apparent between metastasizing capacity and location or size of the tumors.

Urinary obstruction, due to blockage of the urinary flow by tumor growth through either the urethra or one or both ureters, was found in six cases (Fig. 1; Table 1). This caused unilateral or bilateral hydronephrosis and was the primary cause of morbidity in all six animals. Only larger, symmetrically located carcinomas caused urinary obstruction (see above).
CHARACTERIZATION OF CHEMICALLY INDUCED RAT PROSTATE CARCINOMAS

Plasma Acid Phosphatase

Total and tartrate-inhibitable plasma acid phosphatase activity were significantly elevated in the group of animals with grossly apparent chemically induced prostatic carcinomas, as compared with controls (Table 2). There was no significant correlation between total or tartrate-inhibitable plasma acid phosphatase levels and size of the tumors (data not shown). For comparison, acid phosphatase activity was determined in plasma from animals bearing the Dunning R3327H and R3327HI carcinomas and from appropriate controls. Total acid phosphatase activity was significantly higher in R3327HI-bearing animals, but not in animals with the R3327H tumors, than in controls (Table 2); the tartrate-inhibitable fraction was borderline significantly elevated. The average tumor burden was somewhat higher in animals with R3327HI tumors than in R3327H tumor-bearing rats (Table 2). There was no significant correlation between total or tartrate-inhibitable plasma acid phosphatase activity and the tumor burden for either the R3327H or the R3327HI tumor-bearing animals.

Enzyme Histochemistry

Table 3 presents the results of the enzyme histochemical evaluation of normal rat male accessory sex glands, the R3327H and -HI tumors, and some of the chemically induced prostatic carcinomas. Useful markers for the various accessory sex glands appeared to be GGT, LAP and, to a lesser extent, isopropyl alcohol dehydrogenase, which is an indicator of 3β-hydroxysteroid dehydrogenase activity (18). GGT was a marker for the seminal vesicles and the ampullary glands and was negative in all other structures (Fig. 14, a and b). LAP was a marker for the lateral prostate (Fig. 14c) but was minimally active in the dorsal prostate and negative in all other glands examined (Fig. 14d). Isopropyl alcohol dehydrogenase was slightly positive in the lateral prostate and seminal vesicle. No GGT activity was found in the Dunning carcinomas. One-half of the chemically induced carcinomas examined were also negative for GGT. Two were diffusely minimally-moderately positive, whereas the two remaining carcinomas were partly negative and partly minimally-slightly positive. LAP and isopropyl alcohol dehydrogenase activity were found in the Dunning tumors and all chemically induced carcinomas examined, but the staining intensity varied.

Table 2 Total and tartrate-inhibitable acid phosphatase activities in plasma from rats bearing chemically induced prostatic carcinomas or the Dunning R3327H and R3327HI transplantable prostatic carcinomas and from control animals

<table>
<thead>
<tr>
<th>Rats with prostatic carcinomas</th>
<th>Tumor burden (g)*</th>
<th>Acid phosphatase activitya</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N</td>
<td>Total</td>
</tr>
<tr>
<td>Chemically induced controls</td>
<td>9</td>
<td>12.9 ± 4.5 †</td>
</tr>
<tr>
<td>Cph:WU controls†</td>
<td>15</td>
<td>8.2 ± 0.7</td>
</tr>
<tr>
<td>R3327H tumor</td>
<td>7</td>
<td>23.5 ± 0.9</td>
</tr>
<tr>
<td>R3327HI tumor</td>
<td>6</td>
<td>31.5 ± 4.5†</td>
</tr>
<tr>
<td>F334 × COP controls</td>
<td>5</td>
<td>18.4 ± 4.2</td>
</tr>
</tbody>
</table>

* Only animals with R3327 tumors on both flanks were included in this study, and for each animal the total tumor burden (weight) was determined.
† Values are expressed in units/liter as mean ± SEM.
‡ NA, not applicable.
§ P < 0.02 for difference with control value.
¶ Untreated animals of the same strain taken from an experiment carried out in the same period and which were killed at 75-80 weeks of age.
* P < 0.05 for difference with control value.
# P < 0.067 for difference with control value.
CHARACTERIZATION OF CHEMICALLY INDUCED RAT PROSTATE CARCINOMAS

Table 3 Enzyme histochemical properties of the rat male accessory sex glands, the Dunning R3327H and -HI prostate tumors, and chemically induced prostate carcinomas

<table>
<thead>
<tr>
<th>Enzyme</th>
<th>Dorsal prostate</th>
<th>Lateral prostate</th>
<th>Ventral prostate</th>
<th>Seminal vesicle</th>
<th>Coagulating gland</th>
<th>Ampullary gland</th>
<th>R3327H tumor</th>
<th>R3327HI tumor</th>
<th>Chemically induced carcinomas</th>
</tr>
</thead>
<tbody>
<tr>
<td>γ-Glutamyl transpeptidase</td>
<td>±</td>
<td>±/−</td>
<td>±/−</td>
<td>+++</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−/++ (4)</td>
</tr>
<tr>
<td>Isopropyl alcohol dehydrogenase</td>
<td></td>
<td>±/−</td>
<td>±/−</td>
<td>NM</td>
<td>−</td>
<td>−</td>
<td>+</td>
<td>+</td>
<td>+/++ (6)</td>
</tr>
<tr>
<td>Acid phosphatase</td>
<td>++/+++</td>
<td>++</td>
<td>++</td>
<td>++</td>
<td>++/+++</td>
<td>+</td>
<td>++/+++</td>
<td>+/+++</td>
<td>+/+++ (6)</td>
</tr>
<tr>
<td>Glucose-6-phosphate dehydrogenase</td>
<td>++/+++</td>
<td>++</td>
<td>++</td>
<td>++</td>
<td>++/+++</td>
<td>+</td>
<td>++/+++</td>
<td>+/+++</td>
<td>+/+++ (6)</td>
</tr>
<tr>
<td>Alkaline phosphatase</td>
<td>−/±</td>
<td>++/+++</td>
<td>−/±</td>
<td>++</td>
<td>++/+++</td>
<td>+</td>
<td>+/+++</td>
<td>+/+++</td>
<td>+/+++ (6)</td>
</tr>
<tr>
<td>Leucine aminopeptidase</td>
<td>±/−</td>
<td>++/+++</td>
<td>±/−</td>
<td>++</td>
<td>++/+++</td>
<td>+</td>
<td>+/+++</td>
<td>+/+++</td>
<td>+/+++ (6)</td>
</tr>
<tr>
<td>Succinic dehydrogenase</td>
<td>±</td>
<td>++/+++</td>
<td>−/−</td>
<td>++/+++</td>
<td>++/+++</td>
<td>+/++</td>
<td>+/+++</td>
<td>+/+++</td>
<td>+/+++ (6)</td>
</tr>
<tr>
<td>5′-Nucleotidase</td>
<td>±</td>
<td>++/+++</td>
<td>±/−</td>
<td>++/+++</td>
<td>++/+++</td>
<td>+/++</td>
<td>+/+++</td>
<td>+/+++</td>
<td>+/+++ (6)</td>
</tr>
</tbody>
</table>

°—/−, negative; ±, marginally positive; +, slightly positive; ++, moderately positive; +++, markedly positive; /, activity varied from the score before this sign to the score after this sign; NM, not measured.

* Numbers in parentheses, number of chemically induced carcinomas that were examined.

DISCUSSION

The chemically induced carcinomas that were obtained in the experiments described in a companion paper (12) appeared to have a number of important characteristics that are desirable for an adequate animal model for the study of prostatic carcinogenesis. The tumors were adenocarcinomas. The generally long latent period and low mitotic activity indicate that these carcinomas were slow growing neoplasms. Plasma acid phosphatase activity was significantly elevated in animals with these carcinomas. The majority of these invasively growing tumors metastasized and caused urinary obstruction. There was a great diversity in characteristics such as tumor growth rate, as indicated by the latency time, and degree of histological differentiation. Probable precursor lesions were found, i.e., carcinomas in situ and atypical hyperplasias, and induction was possible in intact animals with a single carcinogen administration. The material available for this study was not suitable to determine biochemical characteristics and hormone sensitivity of the carcinomas. These important aspects are addressed in ongoing studies. Other animal models of chemical (25, 26) and hormonal induction of prostatic cancer (27, 28) have not been characterized in detail.

The carcinomas all had a glandular growth pattern, as has the majority of the human prostatic carcinomas (4, 29, 30). Cribriform growth did not occur, unlike observations in some human carcinomas (4, 29, 30) and in carcinomas that occur spontaneously in the ventral prostate of aged rats (31), particularly in the ACI/segHapBR strain (32). The presence of abundant stroma and local invasion into the perineural spaces are two features of human prostatic cancer that were also observed in the chemically induced carcinomas (4, 30, 33). The light microscopic appearance of the carcinomas was comparable to that reported for spontaneously occurring (31, 34) and chemically (25, 26) or hormonally induced carcinomas of the rat.
dorsolateral prostate (27, 28) and for transplantable carcinomas of rat dorsolateral prostate origin (13, 14, 27). Many ultrastructural features of human prostatic carcinomas (4, 29, 33) were also observed in the chemically induced tumors, e.g., loss of basal cells, loss of cellular polarization, presence of lipid droplets, and formation of intracellular lumina. These intracellular lumina, which are a characteristic finding in moderately to poorly differentiated human prostatic carcinomas, have not been described before in reports on the ultrastructure of rat prostatic carcinomas (13, 35, 36). The ultrastructural evidence for the presence of secretory structures in the carcinoma cells suggests that the tumors originated from secretory alveolar cells and not from the epithelium of the excretory ducts, which does not have secretory properties, at least in the rat ventral prostate (37).

Nonneoplastic proliferative epithelial lesions, displaying cellular atypia and confined to one or a few alveoli, were exclusively found in the dorsolateral prostate; and they occurred only in treatment groups in which carcinomas of the dorsolateral prostate were observed as detailed elsewhere (12). Therefore, these lesions are considered likely precursors of dorsolateral prostatic carcinomas. Five of these lesions were classified as atypical hyperplasias and the others as carcinomas in situ. This classification is to some extent arbitrary, inasmuch as we examined only nine lesions and have no evidence for their premalignant nature. However, the degree of cellular atypia, the disorganization of normal glandular pattern and alveolar architecture, and the loss of normal epithelial-stromal relation that characterized these lesions all support the assumption that they are preneoplastic; similar criteria have been described to distinguish premalignant changes in the human prostate (30, 33, 38, 39). The lesions classified as carcinoma in situ lacked conspicuous basal cells. Because there is consensus in the literature about the requirement of absent basal cells for the diagnosis of malignancy in the human prostate (4, 29, 33, 38, 40, 41), this classification seems justified. Furthermore, these putative preneoplastic lesions in the rat dorsolateral prostate appeared to have morphological similarity to premalignant lesions described in the human prostate gland (4, 30, 39–42). It should be noted, however, that the carcinogenic treatments applied did not result in a high incidence of preneoplastic and low-grade malignant lesions relative to the incidence of clinical prostatic cancer (12), as occurs in humans, where the incidence of latent carcinoma is many times higher than that of the clinical disease (5, 8).

Most carcinomas had metastasized, primarily to regional lymph nodes and lung. This implies that both hematogenous and lymphogenous spread occurred, as is the case for human prostatic cancer (2–4). Metastases to bone, as is very frequent in humans, specifically to pelvic and spinal bones (3, 4), were not detected. Perhaps this is due to the insensitivity of our postmortem examination. However, among spontaneous (34), hormone-induced (27, 28), and chemically induced (25, 26) rat dorsolateral prostate carcinomas and transplantable tumor cell lines that were derived from such carcinomas (13, 14, 27, 44), metastatic ability is highly variable, with lymph node and lung metastases as the most frequent sites of dissemination; bony metastases have never been described. Therefore, there may be differences between human and rat models in the venous and/or arterial blood supply of pelvic and spinal bones or in other critical factors involved in the process of metastasis that lead to this apparent species difference.

The elevation of plasma acid phosphatase in rats with chemically induced carcinomas, most of which were advanced cancers comparable to stage C and D disease in humans, resembles the situation in humans. In comparison, animals bearing the Dunning R3327HI carcinomas also had elevated plasma acid phosphatase levels, but animals with the R3327H tumors did not. Isaacs et al. (14), however, reported elevated serum acid phosphatase levels both in animals with the R3327H tumor and the R3327HI tumor. Perhaps differences in total tumor burden or in assay methodology are responsible for these discrepancies. However, it is not likely that the small difference in tumor burden between rats with the rats with the R3327H and with the R3327HI tumor accounted for the difference in acid phosphatase elevation, because there was no correlation between total tumor weight per animal and plasma acid phosphatase activity.

The results of the histochemical analysis of normal prostate and the Dunning R3327H tumors are in agreement with those reported by others (13, 45), with some minor exceptions. There are no data in the literature on the enzyme histochemical properties of the R3327HI tumor, the seminal vesicles, and the coagulating and ampullary glands reported here. The enzyme activities in the R3327H and HI tumors and in the chemically induced carcinomas were remarkably similar. However, the R3327 tumors were negative for GGT, whereas some of the chemically induced carcinomas were partly or completely positive. Because GGT appeared to be a marker for the seminal vesicle and ampullary gland, the possibility is raised that some carcinomas originated from one of these structures. The positive reaction of the carcinomas for LAP, on the other hand, contradicts such a notion; rather it points to the lateral prostate as a possible site of origin. Thus, the enzyme histochemical comparison between the chemically induced carcinomas and the various accessory sex glands does not appear to be helpful in determining the most likely site of origin of these tumors; a similar conclusion was reached by Smolev et al. (13) for the R3327H tumor.

In conclusion, the chemically induced rat prostatic carcinomas that are described here have a large number of important characteristics in common with human prostatic cancer, with the notable exception of the occurrence of bony metastases, while their androgen sensitivity remains to be investigated. Together with previously described features (12), these characteristics indicate that these chemically induced carcinomas are an appropriate model for human prostatic cancer and that the method for induction is thus a valid experimental approach for the study of human prostatic carcinogenesis.

ACKNOWLEDGMENTS

The authors wish to express their gratitude to Dr. H. E. Falke and J. Catsburg for carrying out the acid phosphatase assays; to G. Cook, F. van Welie, and B. Weir for their help with microphotography; to Dr. J. A. Joles for his editorial help, and to the histology staff of the Department of Biological Toxicology of the TNO-CIVO Toxicology and Nutrition Institute for their expert technical assistance. We thank Dr. N. H. Altman for providing animals carrying the Dunning tumors.

REFERENCES

CHARACTERIZATION OF CHEMICALLY INDUCED RAT PROSTATE CARCINOMAS


Characterization of Adenocarcinomas of the Dorsolateral Prostate Induced in Wistar Rats by \( N \)-Methyl-\( N \)-nitrosourea, 7,12-Dimethylbenz(\( a \)anthracene, and 3,2\(^{\prime}\) -Dimethyl-4-aminobiphenyl, following Sequential Treatment with Cyproterone Acetate and Testosterone Propionate
