Tumorigenesis and Neoplastic Progression

Desmosomal Plakophilins in the Prostate and Prostatic Adenocarcinomas

Implications for Diagnosis and Tumor Progression

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The plakophilins, members of the armadillo-repeat family, consist of three different proteins (PKP1-3) that are specifically recruited to desmosomal plaques in a highly cell type-specific manner. Using immunofluorescence, immunoelectron microscopy, and immunoblot, we found that all three plakophilins occurred in luminal and basal cells of the pseudostratified prostate epithelium. The analysis of 135 cases of prostatic adenocarcinomas grouped into tumors with low (Gleason score ≤ 6), intermediate (Gleason score 7), and high Gleason score ($8 \leq$ Gleason score ≤ 10) showed that the expression of PKP1 was reduced or lost in adenocarcinomas with high Gleason scores. The expression of PKP2 was unchanged in all prostatic adenocarcinomas analyzed. In contrast, PKP3 expression was increased in carcinomas with high Gleason scores in comparison with carcinomas with low Gleason scores. In DU 145 cell lines with either overexpression or knockdown of PKP3, both imbalances resulted in fewer desmosomal cell contacts. In addition, overexpression of PKP3 in DU 145 cells led to an augmentation in proliferation rate. Our data imply that both loss of PKP1 and up-regulation of PKP3 expression are biologically important events in prostate cancer and are associated with a more aggressive phenotype. (Am J Pathol 2010; 176:2509–2519; DOI: 10.2353/ajpath.2010.090737)

Cell-cell adhesion is particularly important for the formation and maintenance of normal tissue architecture. The human prostate epithelium is composed of luminal cells with secretory activity and basal cells forming a pseudostratified epithelium with dispersed neuroendocrine cells. Different types of junctions such as adherens junctions, desmosomes, and tight junctions mediate the contact between epithelial cells.^{1,2} Various transmembrane proteins are complexed with plaque proteins specific for each type of junction (eg, in desmosomes, the desmogleins and desmocollins act as Ca^{2+} -dependent transmembrane glycoproteins, which at their cytoplasmic portion, are complexed with desmoplakin, plakoglobin, and plakophilins).

Several junctional plaque proteins belong to the *armadillo* (*arm*)-repeat protein family, characterized by tandem arrays of a 42-amino acid repeat motif. This family includes β -catenin, plakoglobin, and the members of the more closely related p120 subfamily, namely a subgroup localized to adherens junctions, consisting of the name-giving protein p120, Armadillo repeat gene deleted in velocardio-facial syndrome (ARVCF), protein p0071, neurojuncin, and the desmosomal plakophilins.^{3,4} In addition to their junctional localization, most of these proteins also occur at varying degrees in the cyto- and nucleoplasm, and it has been suggested that this may reflect a dual function common to most *arm*-repeat proteins with key roles in both cell adhesion and signaling.^{5,6} For β -catenin and protein p120, cytoplasmic and nuclear binding partners have been identified and their implementation in signaling cascades (eg, the wnt signaling pathway) is well characterized.⁷

The plakophilins make up three different proteins (PKP1-3) that are specifically recruited to desmosomal

Supported by a project grant from the Deutsche Krebshilfe (107861) to I.H.
Accepted for publication January 19, 2010.

Supplemental material for this article can be found on <http://ajp.amjpathol.org>.

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Table 1. Primary Antibodies Used

Antigen	Antibody	Source
Plakophilins		
PKP1	mAb, m	Clone PP1-5C2 (Progen Biotechnik, Heidelberg, Germany) ¹¹
	As, gp	PP1 B6-4 (Progen Biotechnik) ⁶
PKP2	mAb, m	Clones PP2/62, PP2/86, PP2/150 (Progen Biotechnik) ⁵
	As, gp	Head a ⁵
	As, gp	NTb ⁵
	As, gp	HP1α ⁵
PKP3	mAb, m	Clone PKP3-270.6.2 (Progen Biotechnik)
	mAb, m	Clone 23E3/4 (Acris Antibodies, Herford, Germany)
	mAb, m	Biomol, Hamburg, Germany
	As, gp	PP3-1 (Progen Biotechnik) ¹³
Desmosomal proteins		
Desmoplakin	mAb, m	DP1&2-2.15; DP1-2.17; DP1&2-2.20 (Progen Biotechnik)
	As, gp	DP no. 495 ²²
Desmoglein 2	As, rb	rb5 (Progen Biotechnik)
Desmoglein 2	As, rb	rb8, (provided by L. Langbein, German Cancer Research Center, Heidelberg, Germany)
Desmoglein 1/2	mAb, m	DG 3.10 (Progen Biotechnik)
Plakoglobin (γ-catenin)	mAb, m	Clone 11E4 ⁵
	mAb, m	Invitrogen
Arm-repeat proteins of adherens junctions		
Protein p120	mAb, m	BD Biosciences Pharmingen, Heidelberg, Germany
	As, rb	Sigma-Aldrich, Deisenhofen, Germany
Protein p0071	mAb, m	Clone 7.7.9 (Progen Biotechnik) ^{8,9}
	mAb, m	Clone 406.3.1 (Progen Biotechnik) ^{8,9}
	As, gp	hp0071 (Progen Biotechnik) ^{8,9}
ARVCF	As, gp	2× hARVCF (Progen Biotechnik) ²³
	As, gp	3× hARVCF ³⁹
	mAb, m	Clone 1056 (Progen Biotechnik) ²³
	mAb, m	Clone 978 ²³
Neurojungin (δ-catenin)	mAb, m	Clone J19 (Progen Biotechnik)
	mAb, m	Clone 7A ²⁴
	mAb, m	BD Biosciences Pharmingen
	As, rb	KE20 (Sigma-Aldrich)
	As, rb	YV19 (Sigma-Aldrich)
β-Catenin	mAb, m	BD Biosciences Pharmingen
	As, rb	Sigma-Aldrich
Classical cadherins		
E-Cadherin	mAb, m	BD Biosciences Pharmingen
	mAb, rb	Biomol
N-Cadherin	mAb, m	A-CAM (Sigma-Aldrich)
	mAb, m	Clone 32, BD Biosciences Pharmingen

(table continues)

Table 1. Continued

Antigen	Antibody	Source
Marker proteins		
Keratin 5	As, gp	GP5.2 (Progen Biotechnik)
Keratin 8	As, gp	K8.2 (Progen Biotechnik)
Keratin 8 + 18	As, gp	K8/K18 (Progen Biotechnik)
Keratin 14	mAb, m	LL002 (Natutec, Frankfurt, Germany)
	As, gp	GP CK 14.2 (Progen Biotechnik)
Keratin 18	As, gp	GP CK18.2 (Progen Biotechnik)
Others		
α-Catenin	mAb, m	Invitrogen
	As, rb	Sigma-Aldrich
β-Actin	mAb, m	Novus Biological, Acris Antibodies
myc-tag	Rb	ab9106 (Abcam, Cambridge, UK)

mAb, monoclonal antibody; As, conventionally prepared antiserum; rb, rabbit; m, mouse; gp, guinea pig.

plaques in a highly cell type-specific manner. Protein p0071, also called PKP4, is more closely related to other members of the p120 subfamily and is localized as a nondesmosomal plaque protein in adherens junctions.^{8,9} Only PKP2 has been detected in all desmosome-producing cell types, including simple and stratified epithelia as well as nonepithelial tissues such as myocardium and lymph node follicles.^{5,10} In contrast, PKP1 is restricted to stratified and complex epithelia and urothelium,¹¹ whereas PKP3 is located in almost all desmosome-bearing cells, with the exception of desmosomes between hepatocytes and between cardiomyocytes.^{12,13}

Prostate cancer is the most frequent malignancy in men of the western civilization and early detection is essential to reduce mortality and increase survival. In recent years, evidence has been accumulated that several junctional proteins play important roles in carcinogenesis, tumor invasion and metastasis. As a prototypic example, β-catenin mutations in colorectal carcinomas result in impaired β-catenin degradation¹⁴ with profound effects on downstream signaling pathways. Loss, down-regulation, or mislocalization of another important junctional protein, protein p120, has been found to correlate with tumor progression.¹⁵ In contrast, the potential role of plakophilins in human tumorigenesis and progression is largely unexplored. A few immunohistochemical studies on squamous cell carcinoma suggested a correlation with clinical parameters^{16–18} and point to a potentially important role.

In the past, the occurrence of PKPs in normal prostate epithelium and in prostatic adenocarcinoma has not been analyzed. Therefore, we performed immunofluorescence and immunoelectron microscopy to study the distribution of PKP1-3 in human prostate. All three PKPs were detectable in normal luminal and basal cells of glandular prostatic epithelium. The analysis of adenocarcinomas of the prostate together with observations in prostatic cell lines with either overexpression or knock-down of PKPs implies that loss of PKP1 and up-regulation of PKP3 expression are biologically important events in prostate cancer.

Figure 1. Immunofluorescence results of double-labeling experiments on cryostat sections throughout the human prostate. The localization of plakophilins PKP1-3 (A, red, PKP2; D, red, PKP3; and G, red, PKP1) is compared with that of desmoplakin (B and E, green, DP) or desmoglein (H, green, DSG). The corresponding merged pictures are also shown (C, F, and I). Note the punctate staining and the pronounced localization at the apical portion of the basolateral membranes typical for desmosomes in polar epithelia. In higher magnifications (D–I), basal cells occur also positive for plakophilins. Abbreviation: L, lumen. Scale bars = 20 μ m.

Materials and Methods

Cultured Cells and Tissues

Cell cultures derived from prostatic hyperplasia and metastases used in this study were the benign cell line BPH-1¹⁹ and the malignant cell lines DU 145 and LNCaP (for further information on cell lines, see the catalog of the American Tissue Culture Collection, Manassas, VA). The BPH-1 cell line was cultivated in RPMI 1640 medium (Invitrogen, Karlsruhe, Germany) with 1% glutamine (In-

vitrogen), 10% fetal bovine serum (Biochrom, Berlin, Germany), 1 μ g/ml insulin (Sanofi-Aventis, Frankfurt, Germany), 20 ng/ml testosterone (Sigma-Aldrich, Steinheim, Germany), 5 μ g/ml transferrin (Sigma-Aldrich), and 10 ng/ml hydrocortisone (Fluka, Steinheim, Germany). The cell lines DU 145 and LNCaP were cultivated in RPMI 1640 medium with 10% fetal bovine serum and 1% glutamine. Tumor-free human prostatic tissue samples were obtained from tumor prostatectomies at the Department of Urology, Medical Faculty Heidelberg and Medical Fac-

Figure 2. Detection of PKP1 on formaldehyde-fixed, paraffin-embedded tissues of tumor-free human prostate (A, D, and G) compared with the localization of keratin 14 (B and E, K14) and keratin 8 and keratin 18 (H, K8/18). In D–F, a higher magnification of the glandular epithelium is shown. The merged pictures are given in (C, F, and I). Note PKP1 is detectable in both types of cells, luminal and basal cells. Abbreviations: L, lumen; B, blood vessel. Scale bars: 20 μ m (C and D); 10 μ m (F).

ulty Mannheim, Heidelberg University (Heidelberg, Germany), with the permission of the ethics committee of the Medical Faculty. Formaldehyde-fixed, paraffin-embedded prostatic adenocarcinomas were provided from the National Center for Tumor Diseases (Heidelberg, Germany) and the Department of Pathology, Medical Faculty Mannheim, Heidelberg University.

Patients

A total of 135 tissue samples were analyzed: 44 cases with low Gleason scores of 3 to 6, 38 cases with an intermediate Gleason score of 7, and 53 cases with high Gleason scores of 8 to 10.²⁰ The patients' age was between 52 and 77 years, and their prostate-specific antigen blood levels ranged between 0.5 and 97.6 ng/ml. A total of 71 tissue samples were investigated by paraffin multitissue arrays that consisted of five representative punches of tumor and three nonneoplastic prostate tissues per case (for technical details, see Ref. 21). With both, full-sized and microarray samples comparable results were obtained.

Antibodies

The primary antibodies used in this study are listed in Table 1.^{5,6,8,9,11,13,22–24,39} Details on references and distributors can also be found in Table 1.^{5,6,8,9,11,13,22–24,39} Secondary antibodies used for immunofluorescence microscopy were species-specific goat antibodies against Igs of guinea pig, mouse, or rabbit conjugated to Cy3 (BioTrend, Cologne, Germany) or Alexa 488 (MoBiTec, Göttingen, Germany). For immunoblot analysis, horseradish peroxidase-conjugated secondary antibodies

were applied in combination with the enhanced chemiluminescence system (AceGlow-Solution; PEQLAB Biotechnologie, Erlangen, Germany).

Immunofluorescence Microscopy and Immunoelectron Microscopy

The methods used for single- and double-label antibody immunofluorescence microscopy on sections of frozen tissues and cultured cells were described in detail elsewhere.⁸ For sectioning of paraffin-embedded tissues (3 to 5 μ m), a microtome (HM 430; Microm, Walldorf, Germany) was used. Briefly, sections were deparaffinized according to standard techniques and thereafter pretreated to retrieve masked antigens (0.05 M Tris-HCl buffer, pH 10.2, 12 minutes, 120°C for detection of PKP1, PKP2, and ARVCF and 0.1 M Tris buffer containing 10% urea, pH 11, 15 minutes, 120°C for detection of PKP3, protein p0071, and protein p120) with a multifunctional microwave processor (RHS-1; Milestone, Sorisole, Italy). Alternatively, the samples were heated in their appropriate buffer either with a standard microwave (Panasonic, Hamburg, Germany) at 600 W or a steamer (Braun, Kronberg, Germany) for 25 minutes. Sections were then treated with 0.2% Triton X-100 in PBS for 5 minutes, washed with PBS (twice 5 minutes), blocked with 5% goat serum and 2% milk powder in PBS (10 minutes), and then processed for immunofluorescence microscopy. For PKP1 and PKP2 staining sections were additionally incu-

Figure 3. Conventional electronmicroscopy of glandular epithelial cells of human prostate (A) and immunoelectron microscopic localization of plakophilins (immunogold reactions, followed by silver enhancement) to desmosomes using antibodies specific for PKP1 (B), PKP2 (C), or PKP3 (D). Note the prominent decoration of the electron-dense cytoplasmic plaques of desmosomes with all three antibodies specific for PKP1-3. Abbreviations: D, desmosome; N, nucleus. Scale bars: 500 nm (A); 200 nm (B–D).

Figure 4. Immunoblot identification of plakophilins PKP1-3 in various cultured prostatic cell lines and human prostate tissue. The reactions are shown from following total lysates: human prostate (lane 1), BPH-1 (lane 2), DU 145 (lane 3), and LNCaP (lane 4). As a loading control, an immunoblot for β -actin (actin) was performed. Molecular weight markers are as indicated in kilodaltons in the left margin.

Table 2. Immunofluorescence Results for the Plakophilins, PKP1-3 in Low, Intermediate, And High Gleason Scores of Prostatic Adenocarcinoma

	Tissue samples			
	Total	Low Gleason score	Intermediate Gleason score	High Gleason score
PKP1				
Positive	55/135 (41%)	33/44 (75%)	10/38 (26%)	12/53 (23%)
Inhomogeneous	68/135 (50%)	5/44 (11%)	26/38 (69%)	37/53 (70%)
Negative	12/135 (9%)	6/44 (14%)	2/38 (5%)	4/53 (7%)
PKP2				
Positive	106/132 (80%)	34/41 (83%)	32/38 (84%)	40/53 (75%)
Negative	26/132 (20%)	7/41 (17%)	6/38 (16%)	13/53 (25%)
PKP3				
Positive	108/132 (82%)	26/41 (63%)	35/38 (92%)	47/53 (89%)
Negative	24/132 (18%)	15/41 (37%)	3/38 (8%)	6/53 (11%)

bated before blocking with 2 μ g/ml proteinase K (Dako-Cytomation, Hamburg, Germany) for 8 minutes at 37°C, followed by 8 minutes at room temperature. The PKP1-3-specific antibodies were incubated for 4 hours at room temperature (PKP1 and PKP2) or overnight at 4°C (PKP3), the secondary antibodies for 1 hour. As control for unspecific binding, incubation with primary antibody was omitted, and the specimens were incubated with secondary antibody only. Micrographs were taken with an Axiophot microscope (Zeiss, Jena, Germany). Confocal laser scanning immunofluorescence microscopy was done on a Zeiss LSM 510 UV instrument (Zeiss) as described before.²⁵ Procedures used for conventional electron microscopy and immunoelectron microscopy were as described in detail elsewhere.²⁶

Interpretation of Immunofluorescence Signal

A representative section of each tumor sample was stained with H&E to verify tumor classification and to proof the quality of the tissue sample by light microscopy. The extent of immunofluorescence signal was subclassified as homogeneous (staining of the whole tumor) or inhomogeneous (staining only in defined tumor areas). Immunofluorescence experiments were conducted with a negative control for each tumor sample by omitting primary antibody. In addition, an internal control representing nonneoplastic prostate tissue was included. Depending on the percentage of positive cells, the immune reactions were scored into negative (<5% positive staining), inhomogeneous (5% $\leq$ positive staining $\leq$ 90%), and positive (>90% positive staining). Immunofluorescence evaluation was performed and photographs were taken with an Axiophot microscope (Zeiss).

Statistical Analyses

Statistical differences between categoric variables were determined using Yates corrected χ^2 test. Correlations between parameters in ordinal scales were analyzed by Spearman rank R test, as provided by the SPSS software (version 2000; SPSS, Chicago, IL). For all analyses, a value of $P < 0.05$ was considered significant, unless otherwise stated.

Gel Electrophoresis and Immunoblotting

Lysates were generated by the addition of Laemmli sample buffer substituted with Benzonase (Merck, Darmstadt, Germany) to cells or tissues. For immunoblot, proteins decorated with specific primary and secondary antibodies were detected by applying the enhanced chemiluminescence system to the membranes. Light emission was recorded by a photomultiplier in a gel documentation system (ChemiSmart 5000; PEQLAB Biotechnologie).

RNA Isolation and Real-Time PCR

RNA from cells was isolated using a RNA isolation kit (RNAeasy; Qiagen, Hilden, Germany). Afterward, a RT-PCR was performed to transcribe RNA into cDNA (High Capacity cDNA Reverse Transcription kit; Applied Biosystems, Darmstadt, Germany). The obtained cDNA was diluted 1/5 for real-time PCR and added to the Power SYBR Green PCR

Figure 5. Occurrence of plakophilins (PKP1, PKP2, and PKP3) in prostatic adenocarcinoma in correlation with Gleason scores (low: Gleason score 3 to 6; intermediate: Gleason score 7; high: Gleason score 8 to 10). The percentages of negative (white color), inhomogeneous (light gray color), and positive (dark gray color) cases are given.

Master Mix (Applied Biosystems). Primers against PKP1 (sequence forward: 5'-GACCAGGACAACCTCCAGTT-3', reverse: 5'-CTGCTGGTGGTCCCATAGTT-3'), PKP2 (sequence forward: 5'-GCAATGGTTTGCTCGATTT-3', reverse: 5'-GGCTGGTAATCTGCAATGGT-3'), and PKP3 (sequence forward: 5'-TGATGAGCTTCGCAAAAATG-3', reverse: 5'-CTGAGAGGCTGAGCTGAGGT-3') were used, and the occurrence of corresponding DNA fragments was analyzed via StepOnePlus Real-Time PCR System (Applied Biosystems) and controlled by electrophoresis in 1% agarose gels. Hypoxanthine-guanine phosphoribosyltransferase RNA levels were used as internal reference to evaluate RNA recovery and to exclude variations between the different cell clones.

Generation of cDNA Constructs, Transfections, and Generation of Stable Cell Lines

The PKP3-pEGFP-N1 construct²⁷ was cloned into the pLenti6.2/V5-DEST plasmid (Invitrogen) via the pENTR3C plasmid using restriction sites EcoRI and EcoRV, followed

by Clonase reaction as specified by the manufacturer. As a negative control a pLenti-IRES-EGFP plasmid containing enhanced green fluorescent protein (EGFP) only was used (provided by A. Fischer, CBTM). Additionally, a PKP3-myc tag construct was created by replacing the EGFP tag with annealed oligonucleotides coding for the myc tag (5'-CGGAGCAGAACTCATCTCAGAAGAGGATCTGTGGT-3' and 5'-CTAGATTACAGATCCTCTTCTGAGATGAGTTTCTGCTCCGGTAC-3') in the PKP3-pEGFP-N1 vector.

To generate PKP3-specific short hairpin (sh)RNAs, oligonucleotides against different PKP3-mRNA targeting sites were synthesized (shPKP3-9: 5'-GTACCTCGGGACAGGACGTGAAGATAGTTCAAGAGACTATCTTCACGTCCTGTCCCTTTTGGAAA-3' and 5'-AGCTTTTCCAAAAAGGGACAGGACGTGAAGATAGTCTCTTGAAGTATCTTCACGTCCTGTCCCGAG-3'; shPKP3-1350: 5'-GTACCTCGGATCCGTGGAACCTTTTCATTCAAGAGATGAAAGGTTCCACAGATCCTTTTGGGA-3'; and 5'-AGCTTTTCCAAAAAGGATCCTGTGGAACCTTTTCATCTCTTGAATGAAAGGTTCCACAGGATCC-3').

Figure 6. Detection of plakophilins on formaldehyde-fixed, paraffin-embedded tissue of prostatic adenocarcinoma grouped in low (A, D, G, and J), intermediate (B, E, H, and K), and high (C, F, I, and L) Gleason score by immunofluorescence. H&E is shown for a representative example of low (A, Gleason score 3 to 6), intermediate (B, Gleason score 7), and high (C, Gleason score 8 to 10) Gleason score. Immunolocalizations of PKP1 (D–F), PKP2 (G–I), and PKP3 (J–L) are presented. Note the heterogeneous detection of PKP1 in intermediate Gleason score and the loss of PKP1 in high Gleason score. Scale bars = 50 μ m.

The annealed oligonucleotides were ligated directly into the psiRNA-h7SKZeo plasmid (InvivoGen, Toulouse, France) using the unique Acc65I and HindIII restriction sites. As a negative control, a shRNA construct against Luciferase, shLuc, was used (InvivoGen).

The pLenti-PKP3-EGFP plasmid and the pLenti-EGFP plasmid were stably transduced into DU 145 cells according to the protocol of the manufacturer (Invitrogen), and cells were selected with 10 μ g/ml blasticidin (Invitrogen). For generation of stable PKP3 knockdown cell clones and of the negative control cell clone, shLuciferase, DU 145 cells were transfected with the corresponding shRNA plasmids using Eugene 6 transfection reagent (Roche, Mannheim, Germany), followed by selection with 100 μ g/ml Zeocin (Invitrogen). For selection of plasmid containing cell clones, the expression of EGFP was analyzed. A transient transfection with the PKP3-myc construct was also performed with Eugene 6 transfection reagent.

Proliferation Assay

DU 145 wild-type, PKP3 knockdown (shPKP3-1350), and PKP3 overexpressing (PKP3-EGFP) cells were seeded in a 96-well plate (6×10^2 cells/well). 5-Bromo-2'-deoxyuridine (BrdU) was added 48 hours later. Incorporation of BrdU was determined after another 24 hours, according to the protocol of the manufacturer (Cell Proliferation ELISA, BrdU (colorimetric); Roche). Absorbance at 400 nm (reference wavelength, 492 nm) was measured using an ELISA reader (Tecan, Crailsheim, Germany). Statistical differences in proliferation between cell culture lines were analyzed using Student's *t*-test for unpaired experiments.

Results

The Presence of Plakophilins in Normal Prostate

To characterize the occurrence of *arm*-repeat proteins of the p120 subfamily in normal prostate, we systematically screened human tumor-free prostate by immunofluorescence on cryosections using antibodies specific for protein p120, ARVCF, protein p0071, neurojungin, and plakophilins 1-3. Thereby, we noticed a strong staining of the glandular epithelium for all members of the p120 subfamily with the exception of neurojungin (data not shown). According to previous reports,^{2,5,11,13} the occurrence of all three plakophilins in the glandular epithelium of the prostate, especially of PKP1, was unexpected, and therefore, we followed this observation in more detail. We compared the localization PKP1-3 with that of other desmosomal plaque proteins (eg, desmoplakin) and desmosomal transmembrane proteins (eg, desmoglein) and found for all three plakophilins colocalization with desmosomal marker proteins (Figure 1). All proteins showed a punctuate staining at cell borders and a pronounced accumulation at the apical portion of basolateral membranes typical for polar epithelia. The glandular epithelium of the prostate consists of luminal cells revealing secretory activity and stretched basal cells. By careful

inspection at higher magnification, the basal cells were seen to be labeled by antibodies specific for PKP1-3. To unequivocally clarify if all three plakophilins occur in basal cells, we used antibodies specific for keratin 14, a marker for basal cells, and keratin 8 and keratin 18, markers for luminal and basal cells,²⁸ and compared the localization with that of PKP1-3. To obtain better structural preservation, we used immunofluorescence on formaldehyde-fixed and paraffin-embedded tumor-free human prostate samples. Indeed, antibodies against all three plakophilins convincingly showed a decoration of both basal and luminal cells (Figure 2 and Supplemental Figure 1, see <http://ajp.amjpathol.org>). We also applied immunoelectron microscopy to verify the observation that all three plakophilins occur in the glandular epithelium of the prostate at the ultrastructural level (Figure 3). Antibodies against all three plakophilins decorated desmosomal plaques of neighboring epithelial cells in the glandular epithelium of prostate showing the typical appearance of desmosomes with a cytoplasmic electrodense plaque where intermediate filaments are inserted.

The Detection of Plakophilins in Cultured Cells Derived from Prostatic Tissue

We compared our results obtained on human prostatic normal tissue with cultured cells derived from human prostate. Cell cultures derived from prostatic hyperplasia (eg, BHP-1 cells) and metastases (eg, LNCaP cells isolated from lymph node metastases and DU 145 cells derived from a brain metastasis) were used, and immunoblotting using antibodies specific for the plakophilins, PKP1-3, was performed. Although all three plakophilins were detectable in lysates from prostatic tissue and from the benign cell line (BPH-1), only PKP3 showed a prominent reaction in lysates from both malignant cell lines, LNCaP and DU 145 (Figure 4). In contrast, PKP2 was

Figure 7. Detection of PKP1 on formaldehyde-fixed, paraffin-embedded tissue of an adenocarcinoma of the human prostate (**A** and **C**, PKP1) compared with the localization of keratin 8 and keratin 18 (**B** and **D**, K8/K18). Note the loss of PKP1 localization in adenocarcinoma of the prostate with high Gleason pattern. Scale bars = 50 μ m.

barely traceable in LNCaP and DU 145 cells, and PKP1 was not detectable in these cell lines. We verified these observations by immunofluorescence microscopy and found that PKP3 and PKP2 were detectable in desmosomes in all cell lines, whereas PKP1 was only observed in the benign cell line BPH-1 (data not shown). Moreover, we performed RT-PCR on DU 145 cells and found mRNA expression of PKP3 (Supplemental Figure 2, see <http://ajp.amjpathol.org>) and PKP2 but not of PKP1 (data not shown).

The Plakophilins in Adenocarcinomas of the Prostate

Because not all three plakophilins were detectable in malignant prostatic cell lines, we wondered whether the loss of plakophilins correlates with different tumor characteristics. Therefore, we systematically analyzed 135 prostatic adenocarcinomas with different Gleason scores

($n = 44$ with a low Gleason score between 3 and 6; $n = 38$ with an intermediate Gleason score 7; and $n = 53$ with a high Gleason score 8 to 10) for expression of PKP1-3 by immunofluorescence. The findings are summarized in Table 2 and Figure 5. In summary, PKP1 expression was found less frequently and was generally lower in tumors with intermediate and high Gleason scores than in tumors with low Gleason scores, which was statistically highly significant ($P = 0.00006$). In contrast, PKP3 expression was significantly more frequent and generally higher in adenocarcinomas with intermediate or high Gleason scores than in tumors with low Gleason scores, of which 37% were completely negative ($P = 0.0002$). No major differences between the single groups were noted with regard to PKP2. There was no statistical correlation between PKP expression and prostate-specific antigen values (data not shown).

Representative immunofluorescence images for the presence of plakophilins in adenocarcinoma with low,

Figure 8. Influence of overexpression (D–F, PKP3-EGFP) or knockdown (G–I, shPKP3-1350) of PKP3 on DU 145-cultured cells compared with untreated cells (A–C, DU 145). The localization of endogenous PKP3 (A and G, PKP3) using monoclonal antibody clone PKP3-270.6.2 or exogenous PKP3 (D, PKP3-EGFP) is compared with the desmosomal marker desmoglein (B, E, and H, DSG), using desmoglein-specific antibody rb5. The corresponding merged pictures are shown in C, F, and I. Diffuse cytoplasmic staining in G arises from the exogenous EGFP expression introduced with the shPKP3-1350 cDNA construct. Note an imbalance of PKP3 expression in DU 145 cells results in fewer desmosomes. Scale bars = 20 μ m.

intermediate, and high Gleason score are presented in Figure 6. In tumors with a low Gleason score, PKP1 and PKP2 were localized between neighboring cells, and PKP3 was additionally noticed in the cytoplasm of glandular epithelial cells. In tumors with intermediate Gleason scores, PKP1 occurred only in larger, more regular shaped glands, whereas in smaller glands with a tendency of the epithelial cells to spread in the surrounding stroma, the PKP1 signal was lost, whereas PKP2 and PKP3 were still detectable. This observation was even more prominent in carcinomas with a high Gleason score: colocalization of PKP1 with keratin 8/18, as an epithelial cell marker, revealed large tumor areas negative for PKP1 (Figure 7).

Effects of PKP3 Overexpression and Knockdown on DU 145 Cells

To analyze the potential functional consequences of overexpression of PKP3 in more aggressive tumors, we generated stable cell lines with modified expression of PKP3. We used the malignant cell line DU 145 and introduced cDNA constructs yielding either the overexpression of the fusion protein PKP3-EGFP ("gain of function") or reduced expression ("loss of function") through a shRNA interference approach. To knock down the expression of PKP3, two different target sequences were chosen yielding the cell clones shPKP3-9 and shPKP3-1350 with different RNA and protein levels of PKP3 (Supplemental Figure 2, see <http://ajp.amjpathol.org>). In contrast, a cell clone stably expressing shRNA specific for luciferase showed RNA and protein levels comparable with untreated DU 145 cells. The cell clone, shPKP3-1350, with the higher knockdown efficiency was used for further analyses. When PKP3-EGFP was expressed in DU 145 cells, this protein was detected at cell-cell contacts in desmosomes as shown by colocalization with desmosomal proteins (eg, desmoglein) but also in a decent amount in the cytoplasm (Figure 8). In addition, a high portion of uncoupled cells was observed in stably transfected cells where the portion of cytoplasmic PKP3-EGFP was even higher. As a control, a PKP3-myc construct was introduced in DU 145 cells, and here, PKP3 fused to a smaller tag also occurred in desmosomes and in the cytoplasm (Supplemental Figure 3, see <http://ajp.amjpathol.org>). In cell lines with knockdown of PKP3, the distribution of desmosomal proteins (eg, desmoglein) was also disturbed (Figure 8). Desmoglein was only rarely detectable in desmosomes and instead mostly localized in the cytoplasm in different sized structures. Taken together, our findings clearly showed that overexpression and knockdown of PKP3 interfered with the formation of desmosomes and thus with cell-cell contacts.

To further characterize the three cell lines with different PKP3 expression levels, untreated DU 145 cells, PKP3-EGFP expressing cells, and PKP3 knockdown cells (clone shPKP3-1350) in more detail, we compared the cell proliferation rate via BrdU incorporation. For cells overexpressing PKP3 a significant higher proliferation rate was noticed (Figure 9). This observation implies that

Figure 9. Comparison of the cellular proliferation of untreated DU 145 cells (DU 145) and DU 145 cell lines with knockdown (shPKP3-1350) or overexpression (PKP3-EGFP) of PKP3 as measured by BrdU incorporation. For experimental details, see *Materials and Methods*. The data represent the mean cumulative values of three independent experiments including SD. The absorbance obtained for DU 145 cells was set as reference point 1.0. A significant difference ($*P < 0.05$) compared with DU 145 cells was observed for PKP3-overexpressing cells as assessed with Student's *t*-test for unpaired experiments.

PKP3 contributes to the regulation of cell growth and cell division.

Discussion

The present study demonstrates the presence of all three plakophilins, PKP1-3, in luminal and basal cells of the glandular epithelium of the prostate. In prostatic adenocarcinomas, PKP1 expression was decreased with increasing tumor grade, whereas the expression of PKP3 conversely was increased in tumors with high compared with low Gleason scores. PKP2 expression was similar in all tumors, irrespective of Gleason score. Interference with the level of PKP3 expression in a cell culture model influenced the formation of desmosomes and the proliferation rate.

In epithelia, the distribution of plakophilins can be considered as a marker for differentiation. Among the plakophilin family, PKP2 is the most widespread desmosomal plaque protein, occurring in simple, complex, and certain stratified epithelia.^{5,10} Another member less widely distributed is PKP3, which is found in desmosomes of most simple and almost all stratified epithelia and cell lines derived therefrom, with a remarkable exception of hepatocytes.^{12,13} In contrast, PKP1 is the least common plakophilin. It has been reported to occur in stratified epithelia, urothelium, and certain complex epithelia.^{11,29,30} Until recently, any studies on the distribution of plakophilins in the glandular epithelium of the prostate were not available. We show for the first time by immunofluorescence, immunoelectron microscopy, and immunoblot the occurrence of all three plakophilins in luminal and basal cells of pseudostratified prostate epithelium. Similarly, PKP1 has been reported to occur in other glandular epithelia such as, for example, the inner dermal ductal cells in eccrine sweat glands³⁰ or the mammary gland ducts.¹¹

We analyzed 135 cases of prostatic adenocarcinomas and noticed a generally inhomogeneous expression of PKP1 in adenocarcinomas with intermediate (69%) or

high (70%) Gleason score. In tumor areas with high Gleason pattern, PKP1 was no longer detectable, whereas the expression of the two other PKPs, PKP2 and PKP3, was either unchanged (PKP2) or even increased (PKP3). One previous study³¹ reported the presence of all three plakophilins in prostate carcinomas but did not address the association with Gleason grading. In other human tumors, the expression of plakophilins seems to differ between different entities. In oropharyngeal tumors, a reduction or an absence of PKP1 and PKP3 in desmosomes has been noticed.¹⁷ In contrast, non-small cell lung carcinomas were reported to show up-regulation of PKP3.¹⁶ An inhomogeneous expression of PKP1 in tumors has been described in several reports,^{18,30,32,33} and it has been suggested that PKP1 functions as a differentiation marker for carcinomas.³⁰ In glandular epithelium, the detection of PKP1 might be a valuable indicator of well-differentiated glandular structures. Differences in the occurrence of PKPs (eg, loss, up-regulation, or inhomogeneous expression) could reflect tumor progression, as already noticed for protein p120.¹⁵

By using DU 145 cultured cancer cells as a cell culture model, deduced from a brain metastasis of a prostatic adenocarcinoma, we interfered with the expression level of PKP3. Both the knockdown (loss of function) and the overexpression (gain of function) resulted in formation of fewer desmosomes, and the overexpression led to a higher proliferation rate. The essential importance of all three PKPs has been shown *in vitro* in cultured cells via knockdown approaches.^{18,34,35} The down-regulation of PKP1 and PKP3 resulted in a decreased formation of desmosomes and increased cell motility and invasion.^{18,35} PKPs do not only contribute to formation and maintenance of desmosomes^{22,34,36} but also function in the cyto- and nucleoplasm.^{27,37} PKP2 has been found in nuclear particles containing RNA polymerase III.³⁷ Moreover, cytoplasmic PKP1 and PKP3 are components of RNA-containing particles and can be recruited to stress granules.²⁷ The overexpression of PKP3 led to a higher portion of PKP3 in the cytoplasmic pool and thereby may regulate cell growth and cell division and effect the formation of desmosomes. A deficiency in one of the PKPs has a tremendous impact on epithelial integrity in animals and humans. Ablation of PKP1 results in skin fragility and hypohidrotic ectodermal dysplasia.³⁸ Knockout PKP2 mice exhibit lethal alterations in heart morphogenesis.³⁹ PKP3-deficient mice develop hair coat abnormalities and are prone to cutaneous inflammation.⁴⁰

Clearly, future studies will have to reveal the function of PKP1 and 3 in the cytoplasm and how exchange of the junctional and nonjunctional form is regulated in healthy and diseased states such as, for example, cancer.

Acknowledgments

We thank Prof. Dr. Werner W. Franke (Helmholtz Group for Cell Biology, German Cancer Research Center) and Prof. Dr. Hellmut G. Augustin (Joint Research Division Vascular Biology of the Medical Faculty Mannheim) for continuous and generous support.

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