Clinicopathologic and Outcome Features of Superficial High-Grade and Deep Low-Grade Squamous Cell Carcinomas of the Penis

Alcides Chaux, M.D.^{1,2}

Abstract

Purpose: To report the clinicopathologic and outcome features of superficial high-grade and deep low-grade penile squamous cell carcinomas.

Methods: From a retrospectively-collected series of patients with penile cancer we identified 41 cases corresponding to 12 superficial high-grade tumors and 29 deep low-grade tumors. As outcomes we evaluated inguinal lymph node status, presence of tumor relapse, final nodal status, and cancer-specific death. Follow-up ranged from 0.8 to 386.7 months (mean, 152.5 months; median, 157.3 months).

Results: Clinicopathologic features were similar between superficial high-grade and deep low-grade tumors, except for a tendency (Fisher's exact P=0.057) of the former to include tumors with a verruciform pattern of growth. A significantly higher proportion of inguinal lymph node metastasis was found in superficial high-grade tumors compared to deep low-grade tumors [4/5 (80%) vs. 1/5 (20%) respectively, Fisher's exact P=0.02]. No significant differences were found regarding tumor relapse (Fisher's exact P=0.52), final nodal status (Mantel-Cox's P=0.42), or cancer-related death (Mantel-Cox's P=0.52).

Conclusions: Patients with superficial high-grade tumors had a significantly higher proportion of inguinal lymph node metastasis compared to patients with deep low-grade tumors. On this regard, prophylactic inguinal lymphadenectomy might be indicated in cases of superficial tumors with high-grade histology while in deeply invasive low-grade penile carcinomas a more conservative approach may be considered.

Keywords: penile cancer; squamous cell carcinoma; histological grade; prognostic factors; outcome; anatomical level.

Introduction

Pathologic features of the primary tumor affecting outcome of patients with penile cancer are multiple and include histological grade [1, 2, 3], percentage of anaplastic cells [4], anatomical level of infiltration [1], depth of invasion [5, 6], tumor stage [4, 5, 7], presence of vascular [4, 6, 7, 8] and perineural [1, 3] invasion, histological subtype [5, 9] and tumor front of invasion [7]. Risk groups systems combine histological grade with tumor extension to estimate the likelihood of inguinal nodal involvement [10, 11, 12, 13]. The combination of histological grade, anatomical level of inva-

sion and presence of perineural invasion was found to be strongly related to nodal involvement and cancerspecific survival [1]. As usually seen with tumors elsewhere, in penile cancer depth of tumor invasion and histological grade are frequently and significantly associated [9]. However, we have occasionally found superficial tumors depicting a high histological grade and deeply infiltrating malignant neoplasms showing a low-grade histology. The purpose of this study was to evaluate the clinicopathologic and outcome features of patients with such "paradoxical" tumors.

Author Affiliation: ¹Department of Scientific Research, Norte University, Paraguay; ²Centro para el Desarrollo de la Investigación Científica, Asunción, Paraguay.

Corresponding Author: Alcides Chaux, M.D. Department of Scientific Research, Norte University, Gral. Santos e/ 25 de Mayo, Asunción, Paraguay. Office: +595 (021) 203-108, ext. 142. Email: alcideschaux@uninorte.edu.py

Table 1: Clinicopathologic and Outcome Features of Superficial High-Grade and Deep Low-Grade Tumors

Table 1. Chincopathologic and Outcome Pe	Superficial High-Grade		
No. cases (%)	12 (29%)	29 (71%)	
Clinical Features	,	,	
Patients's age in years, median (IQR)	52.5 (12.5)	56 (19)	0.11
Tumor size in cm, median (IQR)	$4.5 \ (4.125)$	5(2.125)	0.95
Anatomical location (%)	, ,	, ,	0.89
Glans alone	7/26~(27%)	$19/26 \ (73\%)$	
Glans + Coronal sulcus	2/6 (33%)	4/6 (67%)	
Glans + Coronal sulcus + Forsekin	3/9~(33%)	6/9 (67%)	
Pathologic Features			
Histologic subtype (%)			0.057
Usual SCC	$11/24 \ (46\%)$	$13/24 \ (54\%)$	
Verrucous carcinoma	0/7 (0%)	7/7~(100%)	
Papillary carcinoma	$0/3 \ (0\%)$	3/3~(100%)	
Warty carcinoma	1/3~(33%)	2/3~(67%)	
Mixed carcinoma	0/4 (0%)	4/4~(100%)	
Urethral invasion (%)			0.7
Positive	3/15~(20%)	12/15~(80%)	
Negative	6/19 (32%)	13/19~(68%)	
Perineural invasion (%)			1
Positive	2/6 (33%)	4/6~(67%)	
Negative	10/34~(29%)	$24/34 \ (71\%)$	
Vascular invasion (%)			0.57
Positive	2/4~(50%)	2/4~(50%)	
Negative	10/36 (28%)	$26/36 \ (72\%)$	
Outcome Features			
Inguinal lymph node metastasis (%)			0.02
Positive	4/5~(80%)	1/5 (20%)	
Negative	8/36~(22%)	28/36~(78%)	
Tumor relapse $(\%)$			0.52
Local, regional or systemic relapse	1/2~(50%)	1/2~(50%)	
No tumor relapse	11/38~(29%)	$27/38 \ (71\%)$	
Final nodal status (%)			0.2
Positive	4/8~(50%)	4/8 (50%)	
Negative	8/33~(24%)	25/33~(76%)	
Death by disseminated cancer $(\%)$			1
Positive	0/1 (0%)	$1/1 \ (100\%)$	
Negative	12/40 (30%)	28/40 (70%)	

IQR, interquartile range; SCC, squamous cell carcinoma

P values were estimated using the Fisher's exact test for categorical data and the Kruskal-Wallis rank sum test for grouped numerical data.

Materials and Methods

Cohort of Patients

Patients were selected from a previously published series of 333 patients with invasive penile squamous cell carcinoma [9]. This dataset is publicly available at http://dx.doi.org/10.6084/m9.figshare.1290997.

Classification of Cases

The dataset was searched for tumors fulfilling the following criteria:

- Superficial High-Grade Tumors: Grade 3 tumors invading lamina propria or superficial corpus spongiosum (tumor thickness ≤ 5 mm).
- Deep Low-Grade Tumors: Grade 1 tumors invading deep corpus spongiosum (tumor thickness ≥ 10 mm) or corpus cavernosum, including tunica albuginea.

The cutoff points of 5 mm and 10 mm were selected based on previous studies [3]. Tumors not fulfilling these criteria were excluded from the dataset. From the full dataset of 333 patients, 29 patients (8.7%) were lost at follow-up, and thus excluded. The final count of selected cases for data analysis was 41, which included 12 (29%) superficial high-grade tumors and 29 (71%) deep low-grade tumors.

Follow-Up

Patients were followed-up from 0.8 to 386.7 months (mean, 152.5 months; median, 157.3 months). Two endpoints were evaluated during follow-up:

- 1. **Tumor relapse:** tumor relapse included the development of local relapse (i.e., tumor on stump), regional relapse (i.e., metastases in regional lymph nodes), or systemic relapse (i.e., metastases in systemic lymph nodes, visceral metastases) during follow-up.
- 2. **Outcome:** the possible categories of outcome included alive without disease, alive with disease, died of other causes, and died of cancer.

Final Lymph Nodes Status

The final status of inguinal lymph nodes was established as follows:

- 1. **Positive status:** the final nodal status was considered positive if pathologically-proven nodal metastases were observed (in those cases with inguinal lymphadenectomy), if regional relapse appeared during follow-up, or if the outcome was unfavorable (i.e., alive with disease or death by cancer).
- 2. **Negative status:** the final nodal status was considered negative if nodal metastases were not observed (in those cases with inguinal lymphadenectomy), if no tumor relapse (beyond local relapse) appeared during follow-up, or if the outcome was favorable (i.e., alive without disease or death by other causes).

Statistical analyses

Bivariate analyses were carried out using the Fisher's exact test for contingency tables and the Kruskal-Wallis rank sum test for grouped numerical variables. Survival curves were generated using the Kaplan-Meier method and compared with the logrank (Mantel-Cox) test. Unconditional logistic regression models were built to estimate odds ratios and 95% confidence intervals. In all cases a 2-tailed P < 0.05 was required for statistical significance. Data were analyzed and plots were generated using R version 3.2.0 "Full of Ingredients" [14]. The dataset and R scripts used for data analysis, as well as additional results (including the full analysis of the dataset), are freely available at https://github.com/alcideschaux/Penis-Paradoxical.

Results

Table 1 compares the clinicopathologic and outcome features of patients with superficial high-grade and deep low-grade tumors. None of the clinicopathologic features were significantly associated with the type of tumor, although deep low-grade tumors tended to exhibit a verruciform pattern of growth. Of the outcome features, only the presence of inguinal lymph node metastasis was significantly different between superficial high-grade and deep low-grade tumors, with a higher proportion of metastasis in the latter [4/5 (80%) cases] compared to the former [1/5 (20%) cases].

Table 2 shows the results of the logistic and Cox's regression analysis for predicting outcomes based on the type of tumor. Patients with superficial high-grade tumors had an significantly increased odds ra-

Table 2: Odds Ratios and Hazard Ratios for Superficial High-Grade vs. Deep Low-Grade Tumors by Outcomes

		Logistic Regression			Cox's Regression	
Outcome	\mathbf{OR}	95% CI	P value	$_{ m HR}$	95% CI	P value
Nodal metastasis	14	1.8, 295.7	0.026	6.13	0.68, 55.02	0.065
Tumor relapse	2.5	0.09,65.89	0.54	2.04	$0.13,\ 32.62$	0.54
Final nodal status	3.1	0.61,16.27	0.16	1.77	0.44, 7.14	0.43
Death by cancer	3.3e-08	NA, Inf	1	3.2e-09	$0.00, \mathrm{Inf}$	0.41

CI, confidence interval; HR, hazard ratios; Inf, infinite; NA, not available; OR, odds ratios

Odds ratios were estimated using unconditional binary logistic regression. Hazard ratios were estimated using Cox's proportional hazards regression. Results for death by cancer are not evaluable due to the small number of events and should be discarded

tios for inguinal lymph node metastasis compared to patients with deep low-grade tumors. Hazard ratios were also increased in patients with superficial high-grade tumors compared to patients with deep low-grade tumors, although the P value was slightly above the standard threshold. Risks were not significantly different for tumor relapse or final nodal status. Risk for cancer-related death was not evaluable due to the small number of events.

Figure 1 shows the survival curves for final nodal status and cancer-related death by type of tumor. As seen, no significant differences were observed between patients with superficial high-grade and deep low-grade tumors in regards to the aforementioned outcomes. Individuals at risk for all survival curves are included as supplementary material in the online repository at https://github.com/alcideschaux/Penis-Paradoxical.

Discussion

In this study we analyzed the clinicopathologic and outcome features of patients with superficial highgrade and deep low-grade squamous cell carcinomas of the penis. We found no significant differences in the clinicopathologic features, except for some tendency of low-grade tumors to exhibit a verruciform pattern of growth. Regarding outcome, superficial high-grade tumors showed a higher proportion of inguinal lymph node metastasis compare to deep lowgrade tumors, suggesting that histological grade is more influential on prognosis than depth of invasion in this particular setting. Nevertheless, the type of tumor had limited usefulness in predicting nodal disease (i.e., final nodal status) or cancer-related death, indicating that other factors should be taken into account for prediciting long-term outcome. Our results suggest that patients with superficial high-grade tumors may benefit from a more aggressive approach

(v.g., prophylactic inguinal lymphadenectomy), in spite of their lower pT stage. Conversely, patients with deep low-grade tumors could be suitable candidates for an active surveillance program instead of a more aggressive approach, despite their higher pT stage.

Our results are in agreement with a previous study evaluating penile tumors invading 5 to 10 mm, in which histological grade had more influence on prognosis than depth of tumor invasion [3]. Given its importance and clinical implications histological grading should be carried out using uniform and comparable criteria, moreover considering that a significant inter-observer variability has been reported for histological grading in penile carcinomas [15]. For patients included in this study, histological grading was carried out using strict (and previously validated) morphologic criteria [2]. This approach may reduce inter-observer variability, although further studies are required to evaluate the external validity and reproducibility of such criteria.

In addition, to consider only the T stage of the penile tumor to define the type and extension of primary treatment could be misleading. Some tumor variants, such as basaloid, sarcomatoid and highgrade usual carcinomas are intrinsically aggressive, regardless of the anatomical level of infiltration [16, 17, 9]. Other tumor variants, such as carcinoma cuniculatum, could invade deep erectile tissues and nevertheless be associated with good prognosis [18]. Moreover, the current TNM system for penile cancer considers invasion of corpus spongiosum or corpus cavernosum as a single pT2 category [19]. Previous studies have found that the metastatic rate of tumors invading corpus cavernosum is higher than those tumors limited to corpus spongiosum [1]. Furthermore, splitting the pT2 stage in 2 categories depending on invasion of corpus spongiosum vs. corpus cavernosum has proven useful in increasing the accuracy of

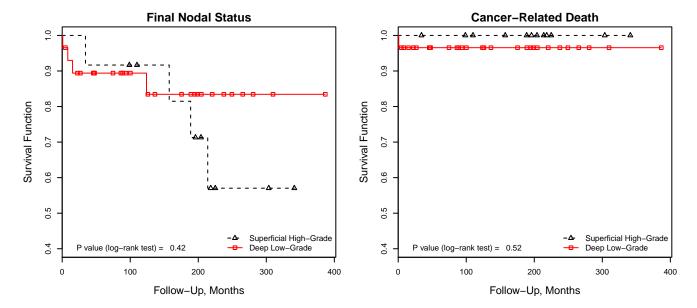


Figure 1: Kaplan-Meier survival curves for final nodal status and cancer-related death by type of tumor. No significant differences were found in the survival curves. Follow-up in months is depicted in the x-axes, while the y-axes depict survival functions. P values were estimated using the log-rank (Mantel-Cox) test.

the TNM sytem for predicting outcome [20]. It does not seem advisable to rely solely on the pT stage to plan optimal therapeutic management. An anatomical level-based approach, combined or not with measurement of tumor thickness, has been evaluated and found useful [1, 3]. Information on histological grade should be included in pathologic reports and anatomical level of infiltration weighted with this grading in order to plan optimal management of patients with penile cancer.

We are aware that the classification we used (i.e., superficial high-grade vs. deep low-grade tumors) might seen incorrect taking into account risk groups systems such as the one proposed by the European Association of Urology (EAU). In the EAU system, both superficial high-grade and deep-low grade tumors would be classified as intermediate risk tumors [11]. Thus, both types of tumor would be managed with either dynamic sentinel lymph node biopsy or inguinal modified lymphadenectomy. However, in a recent study we have found that the EAU system shows a very low accuracy for predicting inguinal lymph node metastasis [21], and thus it might not be a reliable tool for the rapeutic planning. These findings are consistent with other previous studies [22].

As mentioned before, tumors with a verruciform pattern of growth predominated in the deep low-grade category. Tumors with verrucous features includes verrucous carcinoma, papillary, and warty carcinomas [23, 24]. Verrucous carcinomas were char-

acterized by prominent acanthosis, inconspicuous fibrovascular cores, hyperkeratosis and a broad pushing base. It is noteworthy that, although verrucous carcinomas are considered as non-invasive tumors in the latest TNM classification system [19], they can affect deep erectile tissues, as shown in this study. Low-grade warty and papillary carcinomas were very similar in their pattern of growth but papillae were more irregular and architecturally complex in papillary carcinomas and koilocytosis were easily found in warty carcinomas.

The superficial high-grade category included high-grade usual and warty carcinomas. High-grade usual SCC was characterized by nuclear pleomorphism, irregular nuclear membrane, coarse chromatin, prominent nucleoli and high mitotic rate. The other subtype found in the superficial high-grade category was high-grade warty carcinoma, a tumor depicting all the typical aforementioned features of a warty carcinoma (papillae with prominent fibrovascular cores, conspicuous koilocytosis, and jagged tumor base) but showing areas of anaplastic cells, usually at the tumor front of invasion.

At least two limitations must be acknowledged. First, our study is based on a retrospectively-collected series of cases. As such, clinical features were gathered form clinical reports and pathologic features were evaluated only on available microscopic slides. This might have had an impact in the quality of the data that is hard to estimate. Second, our population sample is relatively small, thus our study

might be underpowered to detect the real impact of these tumor types on prognosis. This second limitation is bound to the relatively infrequent combination of low-grade and deep infiltration on one hand, and of high-grade and superficial infiltration on the other hand. Clearly, similar studies (preferably with prospectively-collected data) and on larger datasets of patients with penile cancer are required to determine if the differences and similarities we found are constant features of these type of tumors. Notwith-standing these limitations, our study suggests that patients with superficial high-grade and deep low-grade tumors might benefit from a different therapeutic management than the approach used in more usual clinical settings.

In conclusion, whereas in the majority of penile carcinomas higher grade and deeper tumor invasion are significantly associated, there are cases in which a high-grade tumor may invade only superficially and low-grade tumors can affect deep erectile tissues. Tumors with such features were identified in about 8% of all cases in a large dataset of patients with penile squamous cell carcinomas. We found that superficial high-grade tumors had a significantly higher proportion of inguinal lymph node metastasis compared to deep low-grade tumors. On this regard, prophylactic inguinal lymphadenectomy might be indicated in cases of superficial tumors with high-grade histology while in deeply invasive low-grade penile carcinomas a more conservative approach could be considered.

Acknowledgements

Dr. Alcides Chaux was partially supported by an award granted by the National Council of Science and Technology (CONACYT) dependent of the Presidency of the Republic of Paraguay, as a Level 2 Active Researcher of the National Incentive Program for Researchers (PRONII).

Ethical Standards

The manuscript does not contain clinical studies or patient data. Data were obtained from a publicly available dataset that included no details that might disclose the identity of the subjects.

Conflict of interest

The author declares no conflict of interest.

Authors' Contribution

A. Chaux: Data analysis, Manuscript writing

References

- [1] Alcides Chaux, Carmelo Caballero, Fernando Soares, Gustavo C Guimarães, Isabel W Cunha, Víctor Reuter, José Barreto, Ingrid Rodríguez, and Antonio L Cubilla. The prognostic index: a useful pathologic guide for prediction of nodal metastases and survival in penile squamous cell carcinoma. Am. J. Surg. Pathol., 33(7):1049–1057, July 2009.
- [2] Alcides Chaux, José Torres, Rolf Pfannl, Jose Barreto, Ingrid Rodriguez, Elsa F Velazquez, and Antonio L Cubilla. Histologic grade in penile squamous cell carcinoma: visual estimation versus digital measurement of proportions of grades, adverse prognosis with any proportion of grade 3 and correlation of a Gleason-like system with nodal metastasis. Am. J. Surg. Pathol., 33(7):1042–1048, July 2009.
- [3] Elsa F Velazquez, Gustavo Ayala, Hao Liu, Alcides Chaux, Magali Zanotti, Jose Torres, Soung I Cho, Jose E Barreto, Fernando Soares, and Antonio L Cubilla. Histologic grade and perineural invasion are more important than tumor thickness as predictor of nodal metastasis in penile squamous cell carcinoma invading 5 to 10 mm. Am. J. Surg. Pathol., 32(7):974–9, July 2008.
- [4] J W Slaton, N Morgenstern, D A Levy, M W Santos, P Tamboli, J Y Ro, A G Ayala, and C A Pettaway. Tumor stage, vascular invasion and the percentage of poorly differentiated cancer: independent prognosticators for inguinal lymph node metastasis in penile squamous cancer. J. Urol., 165(4):1138–1142, 2001.
- [5] Bo Dai, Ding Wei Ye, Yun Yi Kong, Xu Dong Yao, Hai Liang Zhang, and Yi Jun Shen. Predicting Regional Lymph Node Metastasis in Chinese Patients With Penile Squamous Cell Carcinoma: The Role of Histopathological Classification, Tumor Stage and Depth of Invasion. J. Urol., 176(4):1431–1435, 2006.
- [6] R E Emerson, T M Ulbright, J N Eble, W A Geary, G J Eckert, and L Cheng. Predicting

- cancer progression in patients with penile squamous cell carcinoma: the importance of depth of invasion and vascular invasion. *Mod. Pathol.*, 14(10):963–8, October 2001.
- [7] Gustavo Cardoso Guimarães, Ademar Lopes, Rodrigo Sousa Madeira Campos, Stenio de Cássio Zequi, Marcos Lima de Oliveira Leal, André Lopes Carvalho, Isabela Werneck da Cunha, and Fernando Augusto Soares. Front pattern of invasion in squamous cell carcinoma of the penis: new prognostic factor for predicting risk of lymph node metastases. Urology, 68(1):148–53, July 2006.
- [8] Vincenzo Ficarra, Filiberto Zattoni, Sergio Cosciani Cunico, Tommaso Prayer Galetti, Lucio Luciani, Andrea Fandella, Stefano Guazzieri, Daniele Maruzzi, Teodoro Sava, Salvatore Siracusano, Stefania Pilloni, Andrea Tasca, Guido Martignoni, Marina Gardiman, Regina Tardanico, Tiziano Zambolin, Antonio Cisternino, and Walter Artibani. Lymphatic and vascular embolizations are independent predictive variables of inguinal lymph node involvement in patients with squamous cell carcinoma of the penis: Gruppo Uro-Oncologico del Nord Est (Northeast Uro-Oncological Group) Penile Cancer data bas. Cancer, 103(12):2507–16, June 2005.
- [9] Gustavo Cardoso Guimarães, Isabela Werneck Cunha, Fernando Augusto Soares, Ademar Lopes, Jose Torres, Alcides Chaux, Elsa F Velazquez, Gustavo Ayala, and Antonio L Cubilla. Penile Squamous Cell Carcinoma Clinicopathological Features, Nodal Metastasis and Outcome in 333 Cases. J. Urol., 182(2):528–534, August 2009.
- [10] E Solsona, I Iborra, J Rubio, J L Casanova, J V Ricós, and C Calabuig. Prospective validation of the association of local tumor stage and grade as a predictive factor for occult lymph node micrometastasis in patients with penile carcinoma and clinically negative inguinal lymph nodes. J. Urol., 165(5):1506–1509, May 2001.
- [11] E Solsona, F Algaba, S Horenblas, G Pizzocaro, and T Windahl. EAU guidelines on penile cancer. *Eur. Urol.*, 46(1):1–8, 2004.
- [12] Edwin Hungerhuber, Boris Schlenker, Alexander Karl, Dominic Frimberger, Karl Heinz Rothenberger, Christian G Stief, and Peter

- Schneede. Risk stratification in penile carcinoma: 25-Year experience with surgical inguinal lymph node staging. *Urology*, 68(3):621–625, September 2006.
- [13] Antonio Augusto Ornellas, Bernardo Lindenberg Braga Nóbrega, Eduardo Wei Kin Chin, Aristóteles Wisnescky, Paulo Cesar Barbosa da Silva, and Aline Barros de Santos Schwindt. Prognostic factors in invasive squamous cell carcinoma of the penis: analysis of 196 patients treated at the Brazilian National Cancer Institute. J. Urol., 180(4):1354–1359, October 2008.
- [14] R Core Team. R: A Language and Environment for Statistical Computing.
- [15] Carsten M Naumann, Ibrahim Alkatout, Moritz F Hamann, Amr Al-Najar, Axel Hegele, Joanna B Korda, Christian Bolenz, Günter Klöppel, Klaus Peter Jünemann, and Christof Van Der Horst. Interobserver variation in grading and staging of squamous cell carcinoma of the penis in relation to the clinical outcome. BJU Int., 103(12):1660-1665, 2009.
- [16] Alcides Chaux, Victor Reuter, Cecilia Lezcano, Elsa F Velazquez, Jose Torres, and Antonio L Cubilla. Comparison of morphologic features and outcome of resected recurrent and nonrecurrent squamous cell carcinoma of the penis: a study of 81 cases. Am. J. Surg. Pathol., 33(9):1299–1306, September 2009.
- [17] Elsa F Velazquez, Jonathan Melamed, Jose E Barreto, Fatima Aguero, and Antonio L Cubilla. Sarcomatoid carcinoma of the penis: a clinicopathologic study of 15 cases. *Am. J. Surg. Pathol.*, 29(9):1152–1158, September 2005.
- [18] José E Barreto, Elsa F Velazquez, Enrique Ayala, José Torres, and Antonio L Cubilla. Carcinoma cuniculatum: a distinctive variant of penile squamous cell carcinoma: report of 7 cases. Am. J. Surg. Pathol., 31(1):71–75, January 2007.
- [19] Oliver W Hakenberg, Eva M Comperat, Suks Minhas, Andrea Necchi, Chris Protzel, and Nick Watkin. EAU guidelines on penile cancer: 2014 update. Eur. Urol., 67(1):142–150, January 2015.
- [20] Joost A P Leijte, Maarten Gallee, Ninja Antonini, and Simon Horenblas. Evaluation of Cur-

- rent TNM Classification of Penile Carcinoma. J. Urol., 180(3):933-938, 2008.
- [21] Alcides Chaux. Risk groups systems for penile cancer management: A study of 203 patients with invasive squamous cell carcinoma. *Urology*, page In press, 2015.
- [22] Giacomo Novara, Walter Artibani, Sergio Cosciani Cunico, Gioacchino De Giorgi, Marina Gardiman, Guido Martignoni, Salvatore Siracusano, Regina Tardanico, Filiberto Zattoni, and Vincenzo Ficarra. How Accurately Do Solsona and European Association of Urology Risk
- Groups Predict for Risk of Lymph Node Metastases in Patients with Squamous Cell Carcinoma of the Penis? *Urology*, 71(2):328–333, 2008.
- [23] Alcides Chaux, Elsa F Velazquez, Ferran Algaba, Gustavo Ayala, and Antonio L Cubilla. Developments in the pathology of penile squamous cell carcinomas. *Urology*, 76(2 Suppl 1):S7–S14, August 2010.
- [24] Alcides Chaux and Antonio L Cubilla. Advances in the pathology of penile carcinomas, June 2012.