

# Histopathological and patient-related characteristics of basal cell carcinomas of the head and neck influencing therapeutic management

Henrik Holtmann, Vera Fuhrmann, Karoline Sander, Christoph Sproll, Norbert R. Kübler, Daman Deep Singh, Julian Lommen

Department of Oral and Maxillofacial Surgery, Heinrich-Heine University of Düsseldorf, Düsseldorf, Germany

# Abstract

The that authors hypothesize histopathological characteristics of basal cell carcinomas of the head and neck (BCCHN) have changed over time and the correlation of BCCHN localization and histological tumour type can help improving the number and quality of necessary therapeutic interventions. Information of 222 patients with 344 BCCHN was gained. Descriptive measures were compared to prior studies to prove whether or not characteristics of basal cell carcinomas or patients have changed over time. Afterwards descriptive measures were correlated with number of conducted operations to evaluate if tumour localization, histological tumour type and number of operations depend on one another. Aggravating factors which lead to a higher number of operations were older age, greater size of BCCHN, adjacent elastosis, the localizations eye, ear and nose and histological tumour types morpheaform and nodularulcerated. In comparison to earlier studies characteristics of BCCHN and patients showed positive developments because of grown awareness of BCCHN. Furthermore, our correlations demonstrate that therapeutic results of BCCHN treatment are continuously improving. Nevertheless, treatment of aggressive morpheaform BCCHN in combination with distinctive patient characteristics still needs improvement.

# Introduction

Basal cell carcinomas (BCC) account for approximately 75% of all skin malignancies in humans.<sup>1</sup> Incidences rise on a daily basis worldwide.<sup>2</sup> The major risk factor for development of BCC is the ultra-violet radiation of the sun.<sup>3</sup> Therefore, BCC are predominantly localized in sun-exposed body regions of the head and face, *i.e.* ears, forehead, nose, lips, cheeks and neck.<sup>2</sup> Though mortality is rare, BCC abundance and therapy is costly for the regional healthcare system. Hence, they pose a popular investigational approach in many studies. BCC are slow growing, non-destructive and seldomly metastazise. There is no agreement on whether BCC derive from dysplastic stem-cells of the hairbulb or precursorcells of the interfollicular epidermis.<sup>4</sup> There are several different histopathological structures of BCC. The World Health Organization (WHO) classifies BCC in i) nodular, ii) superficial and iii) infiltrating types.<sup>5</sup> Types rarely encountered are micronodular, fibroepithelial, basosquamous, keratotic and BCC with adnexal differentiation.5

Therapy of choice is surgical excision with a recurrence rate of approximately 3%.<sup>6</sup> Alternative treatment modalities are chemotherapy, kryotherapy, photodynamic therapy and radiotherapy with the disadvantage of no available sample for subsequent microscopic examination and a recurrence rate of about 10%.<sup>7</sup>

Several studies tried to analyse the correlation of specific histopathological types and anatomical localization of BCC. With 42-90% nodular BCC represent the most common BCC in the head and neck region, whereas superficial types occur mainly on the trunk of the body.<sup>8</sup> Many studies found a male predilection for the development of BCC.<sup>9</sup> Mean age in patients with first time BCC occurance is 63-72 years.<sup>8</sup> Incomplete BCC excisions were most commonly found at the nose, ear and eyelids.<sup>10</sup> The number of necessary operations and the amount of incomplete excisions correlate with BCC size.<sup>11</sup>

To date no correlation between the histological type and the localization of BCC could be determined. Also, there is no consensus on whether BCC histology, size, age, body region or sex taken together correlate in any way with one another. In terms of early screening tests for people with BCC predisposing factors these findings could be highly relevant for prognosis and recurrence rate.

In general, correlation of BCC localization with BCC histological types is important to give patients an outline of the therapeutic approach they can expect.

# **Materials and Methods**

The present study was performed at Düsseldorf University Hospital, Germany. Comprised in this study were 222 patients with a total number of 344 basal cell carcinomas of the head and neck (BCCHN). All Correspondence: Henrik Holtmann, Department of Oral and Maxillofacial Surgery, Heinrich-Heine University of Düsseldorf, Moorenstraße 5, Building 11.52, D-40225 Düsseldorf, Germany. Tel.: +49.2066.29.1719 - +49.211.81.18877. E-mail: Henrik.Holtmann@malteser.org

Key words: Basal cell carcinoma; Histopathological characteristics; Patientrelated characteristics; Head; Neck.

Contributions: HH, VF and JL wrote the manuscript; VF collected the data presented; KS, CS, DDS and NRK reviewed the manuscript before submission.

Conflict of interest: the authors declare no potential conflict of interest.

Funding: none.

Received for publication: 18 March 2018. Accepted for publication: 26 July 2018.

This work is licensed under a Creative Commons Attribution-NonCommercial 4.0 International License (CC BY-NC 4.0).

©Copyright H. Holtmann et al., 2018 Licensee PAGEPress, Italy Dermatology Reports 2018; 10:7674 doi:10.4081/dr.2018.7674

patients were either treated at the Department of Oral and Maxillofacial Surgery or Dermatology between the years 2010 and 2014. In retrospect, the following patient and BCC data were collected from the surgerv report program Medico Release 19.00 (Cerner, Idstein, Germany): sex, age at time of diagnosis, tumour size, histological tumour type, tumour localization and the type of tumour surrounding stroma (presence or absence of adjacent elastosis or actinic keratosis). Furthermore, surgical variables, number of operations, conducted biopsies, type of anaesthesia, outpatient treatment and number of days hospitalised were listed. Additionally, operations necessary for secondary wound closure were assessed. If a patient had multiple BCCHN, variables were collected for each BCC separately. Tumour size and histological tumour type data were collected from histopathological reports. For determination of the histological tumour type classification of the WHO from 20065 was used. Mixed histological tumour types were considered separately.

The count of conducted operations included biopsies, primary resection, further resections if necessary and, if applicable, secondary wound closure. Patients who cancelled the therapy or consulted another hospital were excluded from the study. Further, patients with Gorlin-Goltz Syndrome or Xeroderma Pigmentosum were excluded from the study.

SPSS Version 22 (IBM, Armonk, USA) was used for a statistical analysis. Distributions of one variable were tested with the binominal test. Furthermore, the variables were correlated with each other. Correlations were statistically evaluated with ANOVA, Student's t-test, Chi-square or Fisher's exact-test. If Levene-test was significant Brown-Forsythe or Welch-test was used instead of ANOVA. P-value<0.05 was considered statistically significant and P-value>0.05 and <0.10 was considered marginally significant. If ANOVA or Chisquare test was significant each subcategory was compared with the rest of the category by means of Student's t-test or Fisher's exact-test. Variables with significant or marginally significant correlations were divided into improving or aggravating factors. Improving factors were defined as factors that lead to a lower number of operations, a higher amount of outpatient treatment or more operations which could be conducted under local anaesthesia. Aggravating factors resulted in a higher number of operations, a higher number of hospitalisations or a higher amount of operations conducted under general anaesthesia. Because all variables were correlated with

Table 1. Descriptive measure.

each other, possible overlaps could be shown, resulting in an improved overview.

# Results

Significantly more men than women with BCCHN were found in this study (ratio 1.4:1). The mean age of patients at first diagnosis was 72.59 years (SD 10.90; median 74 years); the youngest patient was 42 years old and the oldest 102. Average tumour size was 0.85 cm (SD 0.89 cm). Tumour sizes between 0.1-0.5 cm and 0.6-1.0 cm were represented significantly more often. The most common histological tumour type was nodular with 46.3%. Superficial BCC (15.2%) and BCCs with mixed histological tumour types (31.1%) were also often found. The amount of morpheaform BCC, which belong to the aggressive growing BCCs, was the lowest with 7.3%. The localizations could be assigned to the following eight main localizations: nose (25.2%), temple (14.5%), eyes (14.5%), ear (14.2%), forehead (12.8%), cheek (9.8%), lips (2.6%) and neck (2.6%). 45.9% of the BCCs showed healthy underlying tissue with the remaining 54.1%% showing abnormal underlying tissue like elastosis (48.8%) or actinic keratosis (2.6%) (Table 1). Mean number of operations was



2.36 (SD 1.12). In most cases patients underwent 2 operations (P<0.01). In 152 (45.51%) cases the patients had a biopsy before tumour resection. The mean number of post-resections was 0.55 (SD 0.77). In 191 (56.85%) of the cases patients had only one resection and 118 (35.12%) patients needed two resections until tumour-free margins were reached. In 170 (55.56%) BCCHN operations primary wound closure could be performed and in 123 (40.20%) cases the surgeons needed an additional operation for secondary wound closure (13 (4,24%) cases of free granulation). Most operations (72.06%) were performed under local anaesthesia. In 56.28% of cases patients had to stay in the hospital and the mean length of stay was 4 days (Table 2).

Correlation of BCC and patient characteristics with *surgical variables* lead to the following factors: improving factors (Table 3) were a lower age (mean 72.38 years) at time of first diagnosis, smaller BCCHN size, localizations cheek, neck and forehead and BCC with the histological types superficial, nodular and nodular-morpheaform. The number of operations was lower for smaller BCC, BCC on the cheek and nodular BCC. Since nodular BCC were significantly smaller than the rest, it could be assumed that both factors as a combination were responsible for the lower number of operations. The number of post-resections

Tuble It Descriptive i	icusuici								
Category					Severity				
No. of BCCs No. of patients	1 BCC 159 (71.60%)	2 BCCs 35 (15.77%		3 BCCs 15 (6.76%)	4 BCCs 7 (3.15%)	5 BCCs 3 (1.35%)	7 BCCs 1 (0.45%)	10 BCC 2 (0.90%	
Time between first and further diagnoses	50.82% of the patients Same time				Rest of the patients Ø 2.62 years after first diagnosis				
Sex	Male* 128 (57.66%)					nale (42.34%)			
Age of diagnosis	Ø 72.59 y SD=10.90 y								
Size	Ø 0.85 cm SD=0.89 cm								
Localization	Nose* 86 (25.2%)	Temple 50 (14.5%)	Eye 50 (14.5%)	Ear 49 (14.2%)	Frontal 44 (12.8%)	Cheek 33 (9.8%)	Lips 9 (2.6%)	Cervikal 9 (2.6%)	Rest 4
Histology	Nodular* 152 (46.3%)	Superficial 50 (15.2%)	Morphea 24 (7.3%)	form	Nodular, morpheaform 13 (4.0%)	Nodular, superficial 28 (8.5%)	Nodular, ulcerated 26 (7.9%)		Rest 35
Adjacent tissue			Normal ski 158 (45,9%)		Adjacent elastos 186 (54,1%)	sis or keratosis	i		
Further malignancies		5	28 pati	th further 78 ma ents had 45 SCC ents had 6 MMs	s				
Operated precancerous lesions			Patient	ts: 81 Lesions Elastosi Keratosi	s: 60				

BCC, basal cell carcinoma; SCC, Squamous cell carcinoma; MM, Malignant melanoma. \*P<0.05 in the binominal test.



was significantly lower for BCCs on the neck, nodular BCCs and superficial BCCs. Neck BCC localization significantly correlated with superficial BCC type. Therefore, it could not be distinguished to what extent these two factors contribute to the lower number of post-resections. The amount of outpatient treatments was higher for younger patients (mean age 72.38 years), for smaller BCCs (mean size 0.70 cm) and for BCCs localized on the cheek. The amount of operations that could be performed under local anaesthesia was significantly higher for BCCs with a smaller size (mean 0.79 cm), for those on the forehead, superficial and nodular-morpheaform BCCs. BCCs on the forehead were also significantly correlated with superficial BCCs.

Aggravating factors (Table 4) were older age at the time of first diagnosis, a bigger size of BCCs, localizations near the eye, on the nose, on the ear, histological types morpheaform and nodular-ulcerated as well as BCCs with adjacent abnormal tissue. The number of operations was significantly higher for patients with bigger BCCs, BCCs localized on the nose and morpheaform BCCs. The number of post-resections was significantly higher for BCCs near the eye and morpheaform BCCs. The number of hospitalisations was significantly higher for patients with older age at the time of first diagnosis (mean 75.22 years), for bigger BCCs (mean size 0.94 cm), for BCCs near the eye and the ear. For patients with nodular-ulcerated BCCs the length of hospitalization was significantly longer. General anaesthesia was performed significantly more often in patients with bigger BCCs (mean size 1.41 cm), BCCs on the ear, nodular-ulcerated BCCs and BCCs with adjacent abnormal tissue. BCCs on the ear (P=0.096) and nodular-ulcerated BCCs (P=0.059) were also correlated with a bigger size of BCCs.

#### Table 2. Improving factors.

#### Discussion

The incidence of BCCs is steadily increasing worldwide, hence, it is important to improve prevention and therapy. There are a lot of studies in which descriptive measures are pointed out. But there are no current studies that compare them with measures from earlier studies in order to overlook the latest trends in therapies. These relations may help to point out issues of BCC therapy needing improvement.

#### **Descriptive measures**

Gender distribution resulted in no difference between earlier studies and this

	Female sex	Age of diagnosis ↓	Localization nose
Occuring BCCs in general ↓	+		
Occuring mBCCs ↓	+	(+)	+ +
Occuring further malignancies ↓	+	+	+ +
		(Ø 71.26 y)	

mBCCs, multiple basal cell carcinomas. + = P<0.05 in Fisher's exact test or P<0.05 in Student's t-test in comparison to the rest.  $(+) = 0.05 \le P<0.1$  in Fisher's exact test or  $0.05 \le P<0.1$  in Student's t-test in comparison to the rest.

#### Table 3. Aggravating factors.

	Male sex	Age of diagnosis 1	Localization neck	Adjacent elastosis or keratosis	Operated precancerous lesions	mBCCs
Occuring BCCs in general 1	+					
Occuring mBCCs ↑	+	(+)	+	+	+	
Occuring further malignancies 1	+	+ (Ø 77.04 y)	+			+

mBCCs, multiple BCCs; + = P<0.05 in Fisher's exact test or P<0.05 in Student's t-test in comparison to the rest. (+) = 0.05 seP<0.1 in Fisher's exact test or 0.05 seP<0.1 in Student's t-test in comparison to the rest.

#### Table 4. Further significant correlations and trends.

Variable 1	Variable 2	P value
Male sex	Ear↑	0.007
Male sex	Nodular ulcerated 1	0.001
Male sex	Operated precancerous lesions 1	0.022
Age of diagnosis↑ (Ø 81.33 y)	Neck	0.001
Age of diagnosis↑ (Ø 82.89 y)	Lips	0.012
Age of diagnosis↓ (Ø 70.29 y)	Nose	0.001
Diameter ↑ (1.18 cm)	Ear	0.096
Diameter ↓ (0.63 cm)	Nose	0.001
Diameter ↓ (0.65 cm)	Nodular	0.001
Neck	Superficial <b>†</b>	0.021
Forehead	Superficial <b>†</b>	0.024
Neck	Nodular↓	0.073
Morpheaform	Adjacent elastosis 1	0.032
Superficial	Adjacent elastosis↓	0.077
Superficial	Operated precancerous lesions 1	0.008

The P value is the result of Fisher's exact.

study. In most studies the amount of men was higher.12 There were only a few studies with more female than male patients.<sup>13</sup> It has been suggested that one reason might be that men are more often exposed to sunlight<sup>14</sup> and that they do not protect themselves by using sunscreen regularly.15 Furthermore, women often wear makeup with sun protection factor.<sup>16</sup> The age at the time of diagnosis was similar comparing this study to former ones.17 In the studies of Heckmann et al.9 and Niederhagen et al.18 patients were 10 years younger. A reason for this might be that life expectancy was lower at that time. The most common localizations of BCC in other studies corresponded with those in this study.<sup>19</sup> The nose was the most affected localization.20 A change could be found in distribution of histological types. In earlier studies the ratio of superficial BCC was lower than in this study and the amount of morpheaform BCCs was higher.8 If the theory of a progression of BCCs from superficial via nodular to morpheaform BCCs is considered,<sup>21</sup> this points towards an earlier detection of BCCHN in this study. Furthermore, the assumption that the growth of superficial BCC is triggered by intermittent sun exposure<sup>22</sup> points towards a better sun protection nowadays. In current studies distribution of histological types equals the distribution in the present study.23 The ratio of mixed histological types was also created by an early detection of BCCs because the most frequently diagnosed histological type was nodular-superficial. This corresponds with the study of Kaur et al.12 In this study the amount of adjacent elastosis was lower than in other studies.24 This also points towards a better protection of the skin nowadays. Also of interest is that the mean size of BCC was lower than in earlier studies.<sup>25</sup> This proved a higher alertness of patients and physicians in these days. Looking at descriptive measures many characteristics could be found which showed a positive development compared to former studies. This demonstrates that the awareness for BCCs is already improving.

## **Correlations with surgical variables**

A higher age was an aggravating factor, because it led to significantly more clinical stays and longer hospitalisations. This may be due to frequent multimorbidity. Due to the significantly higher amount of outpatient treatments and treatments under local anaesthesia and the lower number of operations, patients with smaller BCCs had a better prognosis. The significantly higher number of operations for patients with bigger BCCs resulted from the fact that significantly more often an additional operation for secondary wound closure was needed. This suggests that the surgeons gave their best to keep the number of operations low because in other studies the number of postresections increased with the rise of tumour size.5,10 Considering localizations, regions like eye, ear and nose had a poorer prognosis. In contrary a higher number of postresections around the eye could not be observed in other studies.18 A reason for the high number in this study might be that the surgeons were cautious not to damage anatomic structures unnecessarily. The number of hospitalisations of patients with BCCs at the ear was high as well. The reason for this might be that BCCs on the ear were bigger than those in other regions of the head and neck. Furthermore, for BCCs at the ear the amount of operations under general anaesthesia was higher, which could also have led to a higher number of hospitalisations. The significantly higher number of operations for BCCs on the nose resulted due to the fact that the amount of additional operations for wound closure was significantly higher.

The histological types nodular, superficial and nodular-morpheaform were improving factors and the histological types morpheaform and nodular-ulcerated showed a poorer prognosis. The improving properties of nodular BCCs can be explained by the fact that nodular BCCs were significantly smaller than the rest. The significantly lower number of operations was caused by a significantly higher number of primary wound closures. The significantly lower number of post-resections in cases of nodular BCCs was confirmed by other studies.5 Superficial BCCs were significantly more often operated under local anaesthesia, which can be explained by a significantly higher amount of superficial BCCs on forehead and neck. These localizations can be easily anesthetized without general anaesthesia. Besides, superficial BCCs were more often operated without post-resections, which could not be confirmed by other studies. Aggressive morpheaform histological types counted as aggravating factors for prognosis because the number of operations was significantly higher. In other studies18 the number of incomplete excisions of morpheaform BCC was higher. Reasons for the significantly higher number of operations in this study were the significantly higher number of biopsies and additional operations for secondary wound closure. The significantly higher number of operations under general anaesthesia for nodular-ulcerated BCC in this study could be explained by the fact that nodular-ulcerated BCC were on average bigger than the rest. The amount of



operations under general anaesthesia was significantly higher for BCCs with adjacent elastosis. One reason for this might be that these BCCs seemed to be larger because of the altered adjacent tissue.

### Conclusions

Summarizing, the burden of BCC treatment on the health care system is high. The present study shows that BCC-therapy is headed in the right direction regarding prevention and therapy but there are still a few issues which could be improved to reduce costs. On the one hand the information policy, especially for men, should be improved, so that BCC can be prevented or discovered earlier. In doing so, the mean size of BCC and the number of aggressive histological types could be reduced. As a result, the number of operations, operations under general anaesthesia and the number of hospitalisations could be reduced. This in turn would decrease the economic burden on our health care system. Furthermore, the aggressive histological type morpheaform and the more complicated localizations should be operated more often with Mohs surgery or micrographic surgery with frozen section analysis, in order to decrease the number of operations and further resections.

# References

- Iftimia N, Yelamos O, Chen CJ, et al. Handheld optical coherence tomography-reflectance confocal microscopy probe for detection of basal cell carcinoma and delineation of margins. J Biomed Optics 2017;22:76006.
- Lomas A, Leonardi-Bee J, Bath-Hextall F. A systematic review of worldwide incidence of nonmelanoma skin cancer. Br J Dermatol 2012;166:1069-80.
- Gallagher RP, Hill GB, Bajdik CD, et al. Sunlight exposure, pigmentary factors, and risk of nonmelanocytic skin cancer. I. Basal cell carcinoma. Arch Dermatol 995;131:157-63.
- Youssef KK, Van Keymeulen A, Lapouge G, et al. Identification of the cell lineage at the origin of basal cell carcinoma. Nature Cell Biol 2010;12:299-305.
- Vantuchova Y, Curik R. Histological types of basal cell carcinoma. Scripta Medica (Brno) 2006;79:261-70.
- Marzuka AG, Book SE. Basal cell carcinoma: pathogenesis, epidemiology, clinical features, diagnosis, histopathology, and management. Yale J Biol Med 2015;88:167-79.
- 7. Rhodes LE, de Rie M, Enstrom Y, et al.



Photodynamic therapy using topical methyl aminolevulinate vs surgery for nodular basal cell carcinoma: results of a multicenter randomized prospective trial. Arch Dermatol 2004;140:17-23.

- Friedrich RE, Giese M, Li L, et al. Diagnosis, treatment and follow-up control in 124 patients with basal cell carcinoma of the maxillofacial region treated from 1992 to 1997. Anticancer Res 2005;25:1693-7.
- 9. Heckmann M, Zogelmeier F, Konz B. Frequency of facial basal cell carcinoma does not correlate with site-specific UV exposure. Arch Dermatol 2002;138: 1494-7.
- Husein-Elahmed H, Aneiros-Fernandez J, Gutierrez-Salmeron MT, et al. Assessment of incompletely excised basal cell carcinomas in six facial areas: influence of elastosis. Dermatology. 2012;224:177-83.
- Rivers JK, Mistry BD, Hung T, et al. A 13-year retrospective study of basal cell carcinoma in a Canadian dermatology practice: a comparison between anatomical location and histopathologic subtypes. J Cutaneous Med Surg 2016:20:233-40.
- Kaur P, Mulvaney M, Carlson JA. Basal cell carcinoma progression correlates with host immune response and stromal alterations: a histologic analysis. Am J Dermatopathol 2006;28:293-307.

- Peres LP, Fiorentin JZ, Baptista Tda S, et al. Clinical and histopathological profile of basal cell carcinoma in a population from Criciuma, Santa Catarina, Brazil. Anais Brasil Dermatol 2012;87:657-9.
- Raasch BA, Buettner PG, Garbe C. Basal cell carcinoma: histological classification and body-site distribution. Br J Dermatol 2006;155:401-7.
- de Blacam C, Dermott CM, Sugrue C, et al. Patient awareness and sun protection behaviour following excision of basal cell carcinoma. Surgeon 2017;15:12-7.
- 16. Kyrgidis A, Vahtsevanos K, Tzellos TG, et al. Clinical, histological and demographic predictors for recurrence and second primary tumours of head and neck basal cell carcinoma. A 1062 patient-cohort study from a tertiary cancer referral hospital. Eur J Dermatol 2010;20:276-82.
- 17. Richmond-Sinclair NM, Pandeya N, Williams GM, et al. Clinical signs of photodamage are associated with basal cell carcinoma multiplicity and site: a 16-year longitudinal study. Int J Cancer 2010;127:2622-9.
- Niederhagen B, von Lindern JJ, Berge S, et al. Staged operations for basal cell carcinoma of the face. Br J Oral Maxillofac Surg 2000;38:477-9.
- 19. Monballiu G. Basal cell carcinomata of

the head and neck. Br J Plastic Surg 1968;21:200-11.

- 20. Rustemeyer J, Thieme V, Gunther L, Bremerich A. [Experiences with surgical management of facial basal cell carcinoma and procedures for plastic reconstruction]. Mund-, Kiefer- und Gesichtschirurgie 2005;9:220-4.
- 21. Verkouteren JAC, Smedinga H, Steyerberg EW, et al. Predicting the risk of a second basal cell carcinoma. J Investigative Dermatol 2015;135:2649-56.
- 22. McCormack CJ, Kelly JW, Dorevitch AP. Differences in age and body site distribution of the histological subtypes of basal cell carcinoma. A possible indicator of differing causes. Arch Dermatol 1997;133:593-6.
- 23. Hallaji Z, Rahimi H, Mirshams-Shahshahani M. Comparison of risk factors of single Basal cell carcinoma with multiple Basal cell carcinomas. Indian J Dermatol 2011;56:398-402.
- 24. Epstein E. Value of follow-up after treatment of basal cell carcinoma. Arch Dermatol 1973;108:798-800.
- 25. van Iersel CA, van de Velden HV, Kusters CD, et al. Prognostic factors for a subsequent basal cell carcinoma: implications for follow-up. Br J Dermatol 2005;153:1078-80.

Article