Arborizing Vessels on Dermoscopy in Various Skin Diseases Other Than Basal Cell Carcinoma

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Background: Arborizing vessels (AVs) are dermoscopically defined as telangiectasias with distinct treelike branching, and are a characteristic feature of basal cell carcinoma (BCC). However, AVs are observed in various conditions other than BCC. Objective: The aim of this study was to investigate skin diseases showing AV and investigates dermoscopic differences between BCC and non-BCC. Methods: Dermoscopic images showing AV were prospectively collected and classified into BCC/non-BCC. Non-BCC was further classified into tumors (benign cystic, benign non-cystic, premalignant, and malignant) and non-tumors. We compared AV focusing, widest diameter of stem vessels, widest diameter ratio of stem vessel to first branch, and number of ramifications between groups. Results: Among 124 images, 54.0% were BCC and 46.0% were non-BCC. Non-BCC included epidermal cysts, hypertrophic scars/keloids, intradermal nevi, actinic keratoses, etc. The proportion of focused AV in BCC was significantly higher and the proportion of unfocused AV in BCC was lower than that of premalignant and malignant non-BCC. The widest diameter ratio of stem vessel to first branch was higher in non-BCC. Number of ramifications was significantly less in benign cystic non-BCC than BCC.

Conclusion: Various skin diseases showed AV, so that diagnoses other than BCC should be considered. The findings in this study could help discriminate BCC from non-BCC. (Ann Dermatol 29(3) 288~294, 2017)

Keywords: Arborizing telangiectasia, Arborizing vessel, Basal cell carcinoma, Dermoscopy, Vascular structure

INTRODUCTION

An arborizing vessel (AV) is composed of a stem vessel and irregular treelike branching vessels. Among a variety of vascular structures on dermoscopy, this appearance of AV is very characteristic, and is regarded as an important dermoscopic feature of basal cell carcinoma (BCC). Thus, AV is included in the dermoscopic diagnostic criteria of pigmented BCC proposed by Menzies et al. As a result, most dermatologists who perform dermoscopic exam instinctively consider BCC if they find AV on dermoscopy. Although there have been some reports that AV is found in various skin disorders other than BCC, we experienced AV in more skin conditions than in previous reports. We performed this study to clarify what skin conditions other than BCC can show AV, and what dermoscopic differences of AV differentiate BCC from non-BCC.

MATERIALS AND METHODS

Patients

Among patients who visited the dermatologic clinics in Pusan National University Hospital and Pusan National University Yangsan Hospital from May 2012 to March
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2015, patients whose lesions showed AV on dermoscopy were prospectively enrolled in this study. After the final diagnosis was made with skin biopsy, we classified the lesions into BCC and non-BCC. Non-BCCs were classified into tumors and non-tumors. Tumors were further divided into 3 subgroups of benign cystic, benign non-cystic, pre-malignant, and malignant tumors. This study was approved by Institutional Review Board of Pusan National University Hospital Clinical Trials Center (IRB no. E-2015106).

Methods

1) Photographs
Clinical photographs were taken with a Canon EOS 50D digital single lens reflex camera (Canon, Tokyo, Japan). For dermoscopic images, Dermlite II Pro HR equipment (3Gen; LLC, Dana Point, CA, USA) was used, and dermoscopic photographs were taken with a DSC-W290 (Sony, Tokyo, Japan).

2) Systematic literature search
To identify previous reports demonstrating AV in dermatoses other than BCC, we used PubMed with the following combinations of keywords: “arborizing vessel,” “arborizing vessel AND dermoscopy,” and “vascular AND dermoscopy.” We limited the search to articles published in English before March 31, 2015.

3) Characteristics of AV on the dermoscopy
We examined whether AV was focused or unfocused (Fig. 1). We also compared the widest diameter of stem vessel, the widest diameter ratio of stem vessel to the first branch, and the number of ramifications in both groups. Focused AV was defined as bright red AV, which was sharply focused and passed over the central parts of the lesion. Unfocused AV was blurred in color and distributed at the periphery of the lesion. The widest diameter of stem and branching vessels was measured by using graduation marks of the dermoscope.

4) Statistical analysis
The proportions of focused or unfocused AVs in BCC and non-BCC groups were compared with the chi-square test. The independent t-test was used to evaluate differences between BCC and non-BCC groups (total and individual) in regard to the diameter of the stem vessel, the widest diameter ratio of stem vessel to the first branch and number of ramifications. Statistical significance was defined as a p-value of < 0.05 and statistical analysis was performed with IBM SPSS Statistics ver. 21.0 (IBM Co., Armonk, NY, USA).

RESULTS

Demographics
A total of 124 dermoscopic images showing AV were included in this study. Among these, 67 (54.0%) were BCC and 57 (46.0%) were non-BCC. The ratio of male to female subjects was 1 to 1 (BCC, 1:1.09 and non-BCC, 1:0.90) and the average age was 60.15 years (BCC, 69.12 and non-BCC, 49.60).

Non-BCC with AV

1) Present study
Various skin diseases showed AV (Fig. 2). There were 53 tumors and 4 non-tumors. Among tumors, the benign non-cystic subgroup was most common, followed by benign cystic and premalignant and malignant subgroups (Table 1). The most common dermatosis was epidermal

![Fig. 1. Basal cell carcinoma: (A) with focused arborizing vessel (AV), (B) unfocused AV, and (C) mixed AV. *: focused AV, ^: unfocused AV.](image)
Fig. 2. Clinical (inset) and dermoscopic appearance of arborizing vessels observed in various skin diseases other than basal cell carcinoma: (A) epidermal cyst, (B) digital mucous cyst, (C) intradermal nevus, (D) xanthogranuloma, (E) scar, (F) venous hemangioma, (G) nipple adenoma, (H) chondroid syringoma, (I) lipoma, (J) trichoblastoma, (K) dermatofibrosarcoma protuberans, and (L) squamous cell carcinoma.
Table 1. Various non-BCC dermatoses with AV in the present study (n=57)

| Diagnosis                              | n (%)   |
|----------------------------------------|---------|
| Tumors                                 |         |
| Benign cystic (n=14)                   |         |
| Epidermal cyst                         | 12 (21.1) |
| Digital mucous cyst                    | 1 (1.8)  |
| Milia                                  | 1 (1.8)  |
| Benign non-cystic (n=28)               |         |
| Hypertrophic scar and keloid           | 7 (12.3) |
| Intradermal nevus                      | 6 (10.5) |
| Xanthogranuloma                        | 3 (5.3)  |
| Nipple adenoma                         | 2 (3.5)  |
| Angioleiomyoma                         | 1 (1.8)  |
| Dermatofibroma                         | 1 (1.8)  |
| Apocrine mixed tumor                   | 1 (1.8)  |
| Chondroid syringoma                    | 1 (1.8)  |
| Hidrocystoma                           | 1 (1.8)  |
| Lentigo                                | 1 (1.8)  |
| Lipoma                                 | 1 (1.8)  |
| Pilomatrixicoma                        | 1 (1.8)  |
| Trichoblastoma                         | 1 (1.8)  |
| Venous hemangioma                      | 1 (1.8)  |
| Premalignant & malignant (n=11)        |         |
| Actinic keratosis                      | 5 (8.8)  |
| Bowen’s disease                        | 2 (3.5)  |
| Dermatofibrosarcoma protuberance       | 2 (3.5)  |
| Sebaceous carcinoma                    | 1 (1.8)  |
| Squamous cell carcinoma                | 1 (1.8)  |
| Non-tumors (n=4)                       |         |
| Necrobiosis lipoidica                  | 2 (3.5)  |
| Morphea                                | 1 (1.8)  |
| Porokeratosis                          | 1 (1.8)  |

BCC: basal cell carcinoma, AV: arborizing vessel.

cyst (n=12, 21.1%), followed by hypertrophic scar and keloid (n=7, 12.3%), intradermal nevus (n=6, 10.5%), and actinic keratosis (n=5, 8.8%). Non-tumors included necrobiosis lipoidica, morphea, and porokeratosis.

2) Systematic literature review

There were some reports of non-BCC with AV (Table 2)3-14. According to the keywords, 9 of 81 articles found with “arborizing vessel,” 8 of 21 articles with “arborizing vessel AND dermoscopy,” and 10 of 250 articles with “vascular AND dermoscopy” were included. Some articles were duplicated, and 12 of 352 were finally selected. Among these, 5 articles were case reports and 7 were original articles.

Characteristics of AV on dermoscopy

1) Comparison between BCC and non-BCC

Compared to non-BCC, focused AV was more prevalent in the BCC group, but unfocused and mixed AV was less prevalent in the BCC group (Table 3). The widest diameter of stem vessel and the widest diameter ratio of stem vessel to the first branch were less in BCC than in non-BCC. The number of ramifications was greater in BCC than in non-BCC. Of these, only the widest diameter ratio of stem vessel to the first branch was statistically significant at 1.52 in BCC and 1.78 in non-BCC (p=0.03).

2) Comparison between BCC and each subgroup of non-BCC

Statistical significance was only found for whether or not AV were focused in BCC and premalignant and malignant non-BCC, and for ramification numbers in BCC and benign cystic non-BCC (Table 3).

DISCUSSION

Dermoscopy is a noninvasive and very useful diagnostic tool not only for pigmented skin tumors but also non-pigmented dermatoses, including nonpigmented skin tumors, and inflammatory and infectious skin diseases. In dermoscopic analysis of nonpigmented dermatoses, vascular structures can be the main focus. Various types of vascular structures, such as comma-like, dotted, linear-irregular, hairpin, glomerular, and AVs can be observed on dermoscopy, and most of these were described as characteristic for specific entities1,15. For example, linear-irregular vessels were detected in melanoma, glomerular and dotted vessels arranged in small clusters were described as a specific feature of Bowen’s disease, and hairpin, dotted, and/or linear-irregular vessels were commonly observed in squamous cell carcinoma15,16. Among these, AV was reported as a dermoscopic hallmark of nodular, sclerodermiform, or cystic BCC, with high diagnostic accuracy2,17.

The morphology of AV on dermoscopy is very distinctive, and is easily recognizable by dermatologists. However, this could be a pitfall, because AV can be observed in many other skin conditions, including tumors or inflam-mations. In the present study, AV was observed in a variety of diseases, and many have not been previously reported to show AV. Among the various diseases in this study, only digital mucous cysts, hidrocystomas, keloids, and dermatofibrosarcoma protuberans were reported previously8,10,12. Therefore, dermatologists should not simply
confirm BCC when AV is observed on dermoscopy, and should consider possibilities other than BCC.

Accordingly, we tried to identify morphologic differences in AV between BCC and non-BCC; to our knowledge, these have not previously been reported. Recently, criteria for morphologic discrimination between focused and unfocused AV have been adopted. By definition, focused AVs show classical arborizing telangiectasias, which resemble branches of a tree. These are composed of large-diameter stem vessels, with branches tapering irregularly into finer capillaries. They are bright red in color and sharply focused on dermoscopy. This is a hallmark of nodular BCC, and passes over the central part of the lesion. On the other hand, unfocused AVs are branched vessels with blurring located at the periphery of the lesion. Unfocused AV were described as striking vascular features

### Table 2. Various non-BCC dermatoses with AV

| Report                  | Diagnosis                                                                 | n/N* | Other dermoscopic features except AV                                                                                                                                 |
|-------------------------|------------------------------------------------------------------------------|------|--------------------------------------------------------------------------------------------------------------------------------|
| Llambrich et al. 4 (2009) | Cutaneous leishmaniasis                                                       | 3/26 | Erythema, hyperkeratosis, ulceration, yellow tears, white starburst-like pattern, vascular structures                                                           |
| Fujimura et al. 5 (2012) | Pseudolymphomatous folliculitis                                               | 1/1  | Follicular yellowish spots                                                                                                                                       |
| Dalle et al. 6 (2012)   | Merkel cell carcinoma                                                        | 6/10 | Milky-red globules, linear-irregular vessels, dotted vessels, glomerular vessels, structureless areas, blue-gray veil, scale |
| Moscarella et al. 7 (2013) | Sebaceous adenoma & sebaceoma in Muir-Torre syndrome                        | 7/20 | Yellow globules, elongated crown vessels, opaque structureless white/yellow areas, ulceration, yellowish keratotic plug |
| Zaballos, et al. 8 (2014) | Apocrine hidrocystomas                                                        | 15/22| Blush homogeneous pigmentation, whitish structures, linear-irregular vessels                                                                                   |
| Giacomel et al. 3 (2013) | Basosquamous carcinoma                                                       | 19/22| Ulceration, blue-grey blotches, pigmentation structures, keratin mass, white structureless areas, surface scale, white structures, dotted, linear irregular, hairpin vessels |
| Khelifa et al. 9 (2013)  | Desmoplastic trichoepithelioma                                                | 1/1  | Well-defined borders, ivory-white color                                                                                                                          |
| Bernard et al. 10 (2013) | Dermatofibrosarcoma protuberance                                             | 11/15| Pigmented network, structureless light brown areas, shiny white streaks, pink background coloration, structureless hypo- or depigmented areas |
| Aydingoz et al. 11 (2013) | Neurothekeoma                                                               | 1/1  | None                                                                                                                                                             |
| Saleni et al. 12 (2014) | Digital mucous cyst                                                          | 3/3  | Translucent aspect with presence of bright white structures upon compression                                                                                     |
| Yoo and Kim 13 (2014)   | Keloid                                                                       | 17/30| Linear-irregular vessels, and comma-shaped vessels                                                                                                                |
| Collgros and Guitera 14 (2015) | Eccrine spiradenoma                                                          | 1/1  | Blue-gray ovoid nests and a yellowish clod                                                                                                                       |

Articles from PubMed search with keywords “arborizing vessel,” “arborizing vessel AND dermoscopy,” and “vascular AND dermoscopy,” which were published in English before March 31, 2015. BCC: basal cell carcinoma, AV: arborizing vessel. *Number of cases showing AV/number of cases examined.

### Table 3. Comparison of AV characteristics in BCC and non-BCC

|                  | BCCs (n = 67) | Non-BCCs (n = 57) | p   | Benign cystic (n = 14) | p    | Benign non-cystic (n = 28) | p    | Premalignant & malignant (n = 11) | p    | Non-tumor (n = 4) | p    |
|------------------|---------------|-------------------|-----|-----------------------|------|----------------------------|------|----------------------------------|------|------------------|------|
| Focused AV       | 43 (64.2)     | 30 (52.6)         | 0.19| 11 (78.6)             | 0.30 | 15 (53.6)                  | 0.33 | 3 (27.3)*                       | 0.02 | 1 (25.0)         | 0.12 |
| Unfocused AV     | 34 (50.7)     | 36 (63.2)         | 0.17| 6 (42.9)              | 0.59 | 17 (60.7)                  | 0.37 | 10 (90.9)*                      | 0.01 | 3 (75.0)         | 0.35 |
| Mixed AV         | 10 (14.9)     | 9 (15.8)          | 0.89| 3 (21.4)              | 0.55 | 4 (14.3)                   | 0.94 | 2 (18.2)                        | 0.68 | 0 (0.0)          | -    |
| Widest diameter  | 0.14          | 0.15              | 0.54| 0.12                  | 0.58 | 0.17                       | 0.28 | 0.15                            | 0.68 | 0.15             | 0.56 |
| Widest diameter ratio of stem vessel to the first branch   | 1.52*         | 1.78*             | 0.03| 1.84                  | 0.25 | 1.75                       | 0.10 | 1.79                            | 0.14 | 1.76             | 0.68 |
| Number of ramification | 3.63          | 3.40              | 0.09| 3.21*                 | 0.03 | 3.50                       | 0.49 | 3.36                            | 0.25 | 3.32             | 0.70 |

Values are presented as number (%) or number only. AV: arborizing vessel, BCC: basal cell carcinoma. *Significant findings (p<0.05).
of basosquamous cell carcinoma, observed either alone or in combination with dotted or focused AV. Unfocused AV was also reported as a major vascular structure in dermatofibrosarcoma protuberans, observed alone or mixed with focused AV. In the present study, focused AV were more frequently observed in BCC, but the frequency showed no statistical significance compared to non-BCC. In particular, in benign cystic non-BCC, focused AV was more prevalent than in BCC. Therefore, focused AV as a diagnostic feature of BCC needs to be reconsidered. However, we think that classification of AV by whether or not they are focused can help discriminate BCC from premalignant and malignant non-BCC. Before this study, we surmised that the stem vessel in BCC might be thicker than that of non-BCC. On the contrary, the diameter of stem vessels in non-BCC was greater than in BCC. In particular, the diameter of stem vessels is greatest in benign cystic lesions. The widest diameter ratio of stem vessel to the first branch was less in BCC than in non-BCC. This means that the diameter of the branch vessel remains larger. This is supposedly related to cancer angiogenesis, but further investigation is needed to determine the relationship between the morphology of AV and clinical characteristics, such as aggressive progression. The number of ramifications was greater in BCC than that of non-BCC, and this difference was statistically significant compared to benign cystic non-BCC. This could also be related to cancer angiogenesis in BCC. The morphologic differences of AV found in this study could be helpful in discriminating BCC from non-BCC.

In the present study, we investigated what skin diseases other than BCC can display AV on dermoscopy. Besides BCC, AV was observed in various dermatoses, including benign cystic, non-cystic, premalignant, and malignant tumors, and even non-tumors, including necrobiosis lipoidica, morphea, and porokeratosis. Even though AV is a very distinct and remarkable dermoscopic vascular structure, and is reportedly very specific in the diagnosis of BCC using dermoscopy, dermatologists should be careful in interpreting the significance of AV. Furthermore, we tried to identify dermoscopic differences of AV between BCC and non-BCC. Focused AV were observed more frequently in BCC, and unfocused AV were dominant in premalignant and malignant non-BCC. In non-BCC, the number of ramifications was less than in BCC, and the diameter of vessels decreased more abruptly from the stem vessel to the first branch. In ambiguous cases with AV, these dermoscopic differences could help differentiate lesions.

CONFLICTS OF INTEREST

The authors have nothing to disclose.

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