Dermoscopic Classification of Atypical Melanocytic Nevi (Clark Nevi)

Rainer Hofmann-Wellenhof, MD; Andreas Blum, MD; Ingrid H. Wolf, MD; Domenico Piccolo, MD; Helmut Kerl, MD; Claus Garbe, MD; H. Peter Soyer, MD

Objectives: To create a dermoscopic classification of atypical melanocytic nevi (Clark nevi) and to investigate whether individuals bear a predominant type.

Design: Digital dermoscopic images of Clark nevi were classified according to structural features, ie, reticular, globular, or homogeneous patterns or combinations of these types. The nevi were also characterized as central hypopigmented or hyperpigmented, eccentric peripheral hypopigmented or hyperpigmented, or multifocal hypopigmented or hyperpigmented.

Setting: Two pigmented skin lesion clinics.

Patients: We examined 829 Clark nevi on 23 individuals.

Main Outcome Measure: A reliable dermoscopic classification of Clark nevi and frequency of different dermoscopic types.

Results: Using the dermoscopic classification, the 829 Clark nevi were classified as follows: 221 (26.7%) as reticular, 167 (20.1%) as reticular-homogeneous, 148 (17.9%) as globular-homogeneous, 112 (13.5%) as reticular-globular, 89 (10.7%) as homogeneous, 84 (10.1%) as globular, and 8 (1.0%) as unclassified. Most individuals were prone to a predominant type of Clark nevus. Seven individuals (30%) showed a single type of Clark nevus in more than 50% of their nevi and 5 (22%) in more than 40% of their nevi.

Conclusions: The proposed dermoscopic classification of Clark nevi is easily applicable and allows a detailed characterization of the different dermoscopic types of Clark nevi. Knowledge of these dermoscopic types should reduce unnecessary surgery for benign melanocytic lesions. Exact classification of the different types of Clark nevi is a necessary prerequisite for further clinical, dermoscopic, and histopathologic studies, which will give new insights in the biology of acquired melanocytic nevi.

Arch Dermatol. 2001;137:1575-1580

CLARK NEVI (atypical melanocytic nevi) are acquired melanocytic lesions that are well-known clinically by physicians and especially by dermatologists. Clark nevi are named for Wallace H. Clark, Jr, MD, whose group first drew attention to this particular type of nevus in studies of numerous melanocytic nevi in patients with concomitant melanomas.¹ This melanocytic nevus, originally designated B-K mole,¹ has also been called dysplastic nevus²,³ or atypical mole.⁴ We prefer to use the nomenclature set forth by Ackerman and Magana-Garcia⁵ and designate these acquired melanocytic lesions as Clark nevi.

In the first published report of this special variant of acquired melanocytic nevi, Clark et al¹ described the enormous clinical variability of these melanocytic lesions, which ranged from fewer than 10 to more than 100, were prominent on the upper part of the trunk and the extremities, and varied in size (5-15 mm), shape, and color. However, in many subsequent articles about dysplastic nevi, the clinical features were not clarified,⁶,⁷ and a detailed description of the many different dermoscopic features has not yet been published.

See also pages 1583, 1590, and 1641

In the present study, we used dermoscopy to classify Clark nevi into different types. The purposes of this classification were (1) to create an easily applicable and reproducible classification of Clark nevi, and (2) to investigate whether individuals bear a predominant type of Clark nevī.
We studied 23 individuals (15 male and 8 female). The median age was 33 years (range, 7-60 years). There were 3 patients with skin type I, 15 with skin type II, and 5 with skin type III.

We examined 829 Clark nevi on all 23 patients. The number of nevi in each individual ranged from 14 to 59. Nevi were classified into the dermoscopic categories specified in Table 3. No single Clark nevus in this study showed all 3 structural components. The 8 unclassified nevi were totally hypopigmented; thus, further classification was not feasible.

The distribution of pigmentation in the different dermoscopic types of Clark nevi is given in detail in Table 3. Uniform pigmentation was found most frequently in all but 2 of the 6 types; the reticular-globular and the reticular-homogeneous types showed predominantly multifocal hyperpigmentation or hypopigmentation.

In analyzing the Clark nevus of each subject individually, we observed that the reticular pattern was the most common in 10 individuals (43%), followed by the globular-homogeneous (n = 4 [17%]), reticular-globular (n = 4 [17%]), reticular-homogeneous (n = 3 [13%]), and globular (n = 2 [9%]) patterns. Thirteen individuals (57%) had at least 1 nevus of each dermoscopic type. Eight individuals (35%) had 5 different dermoscopic types, whereas 2 individuals (9%) had only 4 different types.

Most individuals had 1 predominant type of Clark nevus. Seven individuals (30%) showed 1 type of Clark nevus in more than 50% of their nevi; 5 individuals (22%) showed 1 type in more than 40% of their nevi. In these 12 individuals, the predominant type of Clark nevus was reticular (n = 7), reticular-homogeneous (n = 2), reticular-globular (n = 1), globular (n = 1), and globular-homogeneous (n = 1). Among the 7 individuals in whom the predominant type of Clark nevus was reticular, 6 had skin type II.

Based on the present morphologic study of 829 Clark nevi (atypical melanocytic nevi) in 23 patients, we classified Clark nevi into the following 6 dermoscopic types: reticular, globular, homogeneous, reticular-globular, reticular-homogeneous, and globular-homogeneous. Using this dermoscopic classification, we were able to classify...
99.0% of the 829 Clark nevi we examined. The most common type of Clark nevus was the reticular type, followed by the reticular-homogeneous and globular-homogeneous types. In addition, the nevi were classified by pigmentation distribution into the following 3 groups of nearly equal size: 273 Clark nevi had uniform pigmentation, 236 nevi had multifocal hyperpigmentation or hypopigmentation, and the other 312 Clark nevi had central or eccentric hyperpigmentation or hypopigmentation. The most common heterogeneous distribution of pigmentation was multifocal hyperpigmentation or hypopigmentation, followed by central hypopigmentation and central hyperpigmentation. Eccentric peripheral hyperpigmentation was found in only 62 (7.5%) Clark nevi, usually in the reticular or the reticular-homogeneous type of Clark nevus. Eccentric peripheral hyperpigmentation is also often found in malignant melanoma. Thus, Clark nevi with eccentric peripheral hyperpigmentation should be regarded as the most relevant simulators of melanoma within the morphologic spectrum of Clark nevus. Therefore, this type of Clark nevus should be excised or monitored using digital dermoscopy at 3-month intervals. When the eccentric peripheral hyperpigmentation increases, excision of the lesion is necessary.

Few detailed clinical descriptions of special variants of Clark nevi have been given in the literature. Friedman et al described a dark or light target type of Clark nevus. In our classification, this description refers to Clark nevi with central hyperpigmentation or hypopigmentation. Most of these variations are seen in the reticular, reticular-homogeneous, and globular-homogeneous types.

The reticular-homogeneous type with central hyperpigmentation in our dermoscopic classification refers to the hypermelanotic nevus described recently by Cohen et al and to the new nevus of midlife described by Clark et al. Another characteristic dermoscopic pattern of Clark nevi is a patchy distribution of pigmentation, simulating, at least to a certain degree, the uneven pigmentation observed commonly in melanoma. This patchy distribution is a variation of the reticular pattern in which the presence of several small, isolated areas without pigment network leads to multifocal hypopigmentation. Another variation of the reticular type is characterized by multifocal zones of prominent, dark-brown to black pigmented network structures in a patchy distribution. Clark nevi with the reticular pattern and uneven pigmentation are especially prone to overdiagnosis as melanoma.

Remarkably, most individuals had 1 predominant type of Clark nevus. These specific features of all nevi in a given patient have to be taken into account to determine whether a pigmented lesion should be excised. A lesion that does not belong to the predominant type of Clark nevus in a given patient should be considered an atypical lesion and therefore deserving of special attention. The predominance of a single clinical type of nevus has been also documented by Grob and Bonerandi.

Figure 9. Eccentric peripheral hyperpigmented type of Clark nevus. Scale bar indicates 1 mm.

Figure 10. Central hypopigmented type of Clark nevus. Scale bar indicates 1 mm.

Figure 11. Eccentric hypopigmented type of Clark nevus. Scale bar indicates 1 mm.

Figure 12. Multifocal hyperpigmented and hypopigmented type of Clark nevus. Scale bar indicates 1 mm.
Table 2. Classification of Clark Nevi (Atypical Melanocytic Nevi) According to Pigmentation

<table>
<thead>
<tr>
<th>Pigmentation</th>
<th>Definition</th>
<th>Figure</th>
</tr>
</thead>
<tbody>
<tr>
<td>Central hyperpigmentation</td>
<td>Hyperpigmented area (significantly darker than the entire lesion) surrounded by fainter parts of the lesion</td>
<td>8</td>
</tr>
<tr>
<td>Eccentric peripheral hyperpigmentation</td>
<td>Hyperpigmented area (significantly darker than the entire lesion) reaching 1 part of the border of the lesion</td>
<td>9</td>
</tr>
<tr>
<td>Central hypopigmentation</td>
<td>Hypopigmented area (significantly fainter than the entire lesion) surrounded by darker parts of the lesion</td>
<td>10</td>
</tr>
<tr>
<td>Eccentric peripheral hypopigmentation</td>
<td>Hypopigmented area (significantly fainter than the entire lesion) reaching 1 part of the border of the lesion</td>
<td>11</td>
</tr>
<tr>
<td>Multifocal hyperpigmentation and hypopigmentation</td>
<td>Patchy distribution of hyperpigmented and hypopigmented areas</td>
<td>12</td>
</tr>
</tbody>
</table>

*Each type of nevus is depicted in the following figures.

Table 3. Major Dermoscopic Types of Clark Nevi (Atypical Melanocytic Nevi) and Their Distribution of Pigmentation*

<table>
<thead>
<tr>
<th>Type (No. [%] of Nevi)</th>
<th>Uniform Pigmentation</th>
<th>Central Hyperpigmentation</th>
<th>Eccentric Peripheral Hyperpigmentation</th>
<th>Central Hypopigmentation</th>
<th>Eccentric Peripheral Hypopigmentation</th>
<th>Multifocal Hyperpigmentation and Hypopigmentation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Reticular (n = 221 [26.9])</td>
<td>76</td>
<td>34</td>
<td>34</td>
<td>26</td>
<td>2</td>
<td>49</td>
</tr>
<tr>
<td>Reticular-globular (n = 112 [13.6])</td>
<td>23</td>
<td>19</td>
<td>2</td>
<td>15</td>
<td>5</td>
<td>48</td>
</tr>
<tr>
<td>Reticular-homogeneous</td>
<td>42</td>
<td>25</td>
<td>12</td>
<td>32</td>
<td>5</td>
<td>51</td>
</tr>
<tr>
<td>(n = 167 [20.3])</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Globular (n = 84 [10.2])</td>
<td>41</td>
<td>3</td>
<td>3</td>
<td>6</td>
<td>0</td>
<td>31</td>
</tr>
<tr>
<td>Globular-homogeneous</td>
<td>53</td>
<td>18</td>
<td>7</td>
<td>21</td>
<td>6</td>
<td>43</td>
</tr>
<tr>
<td>(n = 148 [18.0])</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Homogeneous (n = 89 [10.8])</td>
<td>38</td>
<td>9</td>
<td>4</td>
<td>13</td>
<td>11</td>
<td>14</td>
</tr>
<tr>
<td>Total (n = 821)†</td>
<td>273 (33.3)</td>
<td>108 (13.2)</td>
<td>62 (7.6)</td>
<td>113 (13.8)</td>
<td>29 (3.5)</td>
<td>236 (28.7)</td>
</tr>
</tbody>
</table>

*Data are given as number of Clark nevi classified; totals are given as number (percentage). Percentages have been rounded and may not sum 100.
†Sample for analysis was restricted to the 821 Clark nevi that could be classified from the total sample size of 829 Clark nevi.

Every new classification should fulfill at least the following 3 criteria: it should be easily applicable, it should be reproducible, and it should open new aspects. With the caveats that dermoscopic-histopathologic correlation has not yet been attempted and that the reproducibility of the classification has not yet been demonstrated, our proposed dermoscopic classification of Clark nevi fulfills all these criteria. The classification is easily applicable because only 3 main dermoscopic patterns and 2 descriptors of pigmentation have to be analyzed. The concentration on reticular, globular, and homogeneous patterns and on hypopigmentation and hyperpigmentation also minimizes the subjective variation in the assessment of dermoscopic features. Stanganelli et al. found perfect intraobserver agreement for pigment network in 100% of 150 digital dermoscopic images, for diffuse (eg, homogeneous) pigmentation in 97%, and for globules in 85%. The descriptors of chromatic characteristics are usually less reproducible, but intraobserver agreement was seen for hypopigmentation (also called depigmentation) in 91% of digital dermoscopic images. The main difficulty in our classification system was whether a nevus with unevenly distributed pigmentation should be classified as hypopigmented or hyperpigmented. However, by using the color of the main part of the nevus as a reference, this judgment also can be reproducible.

Finally, better characterization of Clark nevi offers the opportunity to study their biological features. An exact and practicable classification of Clark nevi is a prerequisite for further studies concerning the morphologic changes of these acquired melanocytic nevi by means of digital dermoscopic follow-up examinations. Furthermore, it may be important to investigate whether a particular dermoscopic type of Clark nevi is more often associated with development of cutaneous melanoma. Obviously, a meticulous dermoscopic-histopathologic correlation as outlined recently for each different dermoscopic type of Clark nevus would be suitable.

Our dermoscopic classification of Clark nevi (atypical melanocytic nevi) should be regarded not just as an academic morphologic exercise but as a classification system that will also lead to a better understanding of the biological characteristics of these common melanocytic lesions.

Accepted for publication May 23, 2001.
This study was supported by a grant from Österreichische Krebshilfe Steiermark, Graz, Austria.
We thank Barbara J. Rutledge, PhD, for editing assistance.
Corresponding author and reprints: H. Peter Soyer, MD, Department of Dermatology, University of Graz, Auernburgerplatz 8, A-8036, Graz, Austria (e-mail: peter.soyer@uni-graz.at).

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