Dermoscopy of Abortive Hemangioma: Morphological Study of 11 Cases

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Introduction

Abortive hemangiomas (AH) or minimal or arrested growth hemangiomas constitute a distinct subgroup of infantile hemangioma (IH) with a characteristic clinical presentation and natural history that are often clinically confused with other vascular anomalies (VA) [1,2]. Dermoscopy can be a useful tool for distinguishing VAs that may be clinically similar such as IH precursor lesions, port wine stains (PWS) and, probably, AH [3]. However, dermoscopy pattern of AH has only been previously described by Toledo-Alberola et al in 2 cases [4].

Case Presentation

The aim of this study was to evaluate the dermoscopy findings of an AH single-center case series, comparing these findings with those described in other clinically similar VAs such as IH precursor lesions and PWS. We evaluated the dermoscopy patterns and structures of 11 consecutives clinically

diagnosed AH cases in our pediatric dermatology department, assessed during 2021, and we compared these findings with those described for IH precursor lesions and PWS.

Clinical and demographic findings are summarized in Table 1. Serpinginous or tortuous vessels, punctate vessels and vascular lacunae were observed in 100% of the cases clinically diagnosed as AH (Figures 1 and 2).

Conclusions

AH is a distinctive variant of IH present from birth in most cases, and, with little ability to proliferate. Although they could be clinically similar to other VAs they have different morphologic and immunostaining patterns. On histology, they show dilated thin-walled vessels within the superficial dermis, with a GLUT-1 positivity immunostaining demonstrating a clear distinction between AH and PWS or congenital hemangiomas [1,2]. Dermoscopic examination may facilitate the identification of AH for early diagnosis and treatment. In our case series, serpiginous or tortuous

Table 1. Clinical and demographic findings.

	Dermoscopy findings	Vascular	lacunae	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes
ony finding		Puntiform	vessels	SəK	Хes	Yes	Yes	Yes	Yes	Yes	SəX	Yes	Yes	Yes
Dormoc	Dermosc	Serpinginous/	tortuous vessels	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes
			Additional tests	-	-	Echography and doppler	1	Spinal echography	1	Soft tissue Echography	-	1	1	ı
			Associate lesions	1	Infantile hemangioma			1		1	ı		,	
			Treatment	Timolol	Timolol	Timolol	Timolol		Timolol	Timolol		Timolol + Propanolol		Timolol
torictics	reiistics		Location	Right leg	Left buttock	Left knee	Left thigh	Sacral fossa	Right external malleolus	Left hand	Abdomen	Right lumbar	Neck	Right thigh
) exect	Demographic characteristics		Sex	M	M	Щ	M	П	Ħ	Ħ	M	Ħ	M	H
Jomod			Age occurrence	First days	Unknown	Birth	Birth	Birth	Birth	Birth	Unknown	First days	Birth	Birth
		Age at diagnosis	(years)	9	1	2	2	0	1	1	14	9	2	1
Pa+ion+	ratiellt			1	2	3	4	5	9		8	6	10	11



Figure 1. (A) Clinically macule with telangiectasias erythematous background. (B,C) Dermoscopy image showing serpiginous or tortuous vessels, punctate vessels and vascular lacunae.

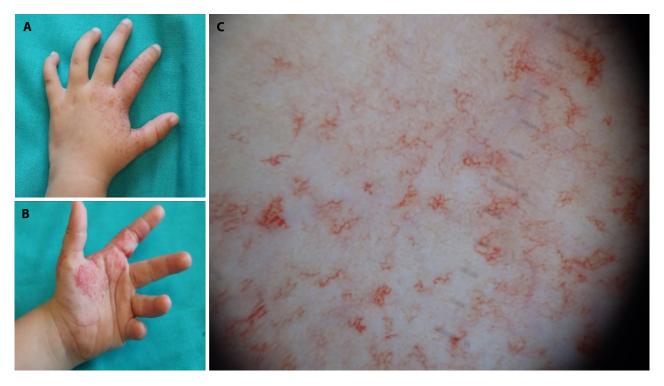


Figure 2. (A,B) Telangiectasias on an erythematous or reticulated background. (C) Dermoscopy image showing serpiginous or tortuous vessels, punctate vessels and vascular lacunae.

vessels, punctate vessels and vascular lacunae was observed in 100% of the cases clinically diagnosed as AH. IH precursor lesions dermoscopy features have been described as large linear vessel and branching vessels with erythematous background [5]. On the other hand, linear vessels, reticular vessels, sausage-like vessels, dots or globules vessels and mixed vessels, as well as white circles and whitish veil have been described in PWS [6]. When comparing our findings in AH, with those described IH precursor lesions and PWS, we found these to be differential and characteristic of AH. As limitations, no control group was included, therefore, no definitive conclusions could be made.

Dermoscopy might improve the clinical diagnosis of AH. It might help distinguish AH from IH precursor or PWS. More studies including other lesions are needed to confirm our findings.

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