SCABIES MIMICING CHILD ABUSE – A CASE REPORT

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INTRODUCTION

Scabies is an infestation of the skin by the mite Sarcoptes scabiei var. hominis that results in an intensely pruritic eruption with a characteristic distribution pattern. The estimated prevalence ranges from 0.2 to 71 percent, with as many as 100 million people affected worldwide (Romani 2015). Transmission of scabies is usually from person to person by direct contact, by wearing or handling heavily contaminated clothing, or by sleeping in an unchanged bed recently occupied by an infested individual. Transmission from parents to children, and especially from mother to infant, is routine. Schools do not ordinarily provide the level of contact necessary for transmission. In young adults, the mode of transmission is usually sexual contact (Fuller 2013, Chosidow 2006, Heukelbach 2006). The essential lesion is a small, erythematous, nondescript papule, often excoriated and tipped with hemorrhagic crusts (Pomares 2014, Eshagh 2014). In other cases scabies can present with unusual forms (e.g. vesicular, bullous, nodular, crusted, urticarial) which can make diagnosis difficult even for experienced dermatologist (Orkin 1985). The diagnosis of scabies is generally made from the history and the distribution of lesions, as well as the skin scraping, dermoscopy, and the adhesive tape test (Mahé 2005, Heukelbach 2005, Walter 2011, Dupuy 2007). Topical permethrin 5% cream and oral ivermectin are the first-line therapies (Chosidow 2006, Heukelbach 2006, Ly 2009).

In rare instances there are unusual complications of scabies such as cutaneous vasculitis and glomerulonephritis, which can sometimes overshadow the primary disease (Mazzatenta 1996). Vascular purpura has rarely been reported (Valks 1996, Estève 2001).

CASE REPORT

A 6-year old male patient was admitted to University Department of Pediatrics with a 10-day history of rash resembling bruises. The dermatosis started with ecchymotic and a few petechial skin changes on the back of the lower legsand progressed proximally to thighs, buttocks, trunk, upper extremities andface. He was referred to our Department for evaluation of skin changes resembling child abuse-like bruises (Figure 1 and 2). Parents have denied any kind of child abuse.At the follow up we noticed the progression of skin changes with generalization of nonblanching ecchymotic and petechial lesions, accompanied with intensive pruritic sensation and mild edema and arthralgia of talocrural joints, without any gastrointestinal involvement (Figure 3 and 4). According to the worsening of clinical picture during the hospitalization, any possibility of child abuse was excluded.

Laboratory tests showed slightly increased levels of CRP: 6.8 mg/dL; tIgE: 80.8 klU/L and LDH: 329 U/L. Antistreptolysin O titer was 600 IU/L. The full blood count, serum urea, creatinine, electrolytes, protein electrophoresis, immunoelectrophoresis, coagulogram, C3, C4, ANA, ENA profil, c-ANCA, p-ANCA, EBV-VCA IgM, Anti-HAV IgM, HbsAg, Anti-HBc IgG were within the reference range. Urinalysis showed no hematuria or proteinuria. Coproculture, stool ova and parasites tests were negative, as well as the nasal and throat swabs. Microscopic examination of material scraped from few purpuric skin changes demonstrated numerous living mites and eggs leading to the diagnosis of scabies. Histological examination revealed hyperkeratosis, focal parakeratosis, acantosis, spongiosis and a dense infiltrate of lymphocytes, histiocytes and eosinophils through the epidermis and dermis. Infiltrating cells were arranged around small dermal vessels that showed signs of a lymphocytic vasculitis. Scabies associated leucocytoclastic vasculitis was diagnosed. Treatment consisted of two applications of topical 5% permethrin cream followed by a systemic steroid therapy which resulted in a complete recovery.

DISCUSSION

Generalized cutaneous vasculitis is a rare complication of scabies, and only few cases have been reported (Menné 1984). The exact mechanism by which infestation of the skin with Sarcoptes mite causes vasculitis is unknown. The possible immunological reaction to scabies probably occurs trough a veriety of mechanisms including immediate, cell-mediated and possibly immune complex pathways (Jarret 1998). Scabietic leucocytoclastic vasculitis with focal glomerulonephritis has been reported in two patients (Menné 1984, Jarret 1998).



Figure 1. Skin changes resembling child abuse-like bruises on the patient's face

Vasculitis has also been reported in a patient with Norwegian scabies and an human immunodeficiency virus positive man (Skinner 1992).

Physical abuse of children is a common occurrence, and it carries a significant morbidity and mortality rate (Mikulic 2013). The presence of various age bruising is



Figure 3. Progression of skin changes with generalization of nonblanching ecchymotic and petechial lesions on buttocks and lower extremities



Figure 2. Skin changes resembling child abuse-like bruises on the patient's trunk

suspicious (Ricci 1991). Bruises on relatively protected places such as the upper arms, medial and rear thighs, arms, torso, cheeks, ears, neck, genitals and buttocks, should raise suspicion of abuse, especially if they are extensive and ifbruises of different ages coexist (Kos 2006). According to the presence of various age bruises-like skin



Figure 4. Progression of skin changes with generalization of nonblanching ecchymotic and petechial lesions on lower extremities

lesions in presented case, it was reasonable to suspect child abuse. Due to the worsening of clinical picture during the hospitalization, the possibility of child abuse in our case was excluded.

It can be presumed that the persistence of mites and their products caused by a delay in the diagnosis, inducing sensitization, could have resulted in the development of the leucocytoclastic vasculitis (Stinco 2008).

CONCLUSION

Regarding to the increased number of atypical clinical manifestations of scabies, it would be wise to exclude scabies in any pruritic skin lesion, regardless of its clinical manifestation.

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Anita Gunarić: Design of the study, literature searches, literature and patient data analyses;

- Kristina Jurišić : Design of the study, literature searches, patient's data analyses;
- Dubravka Šimić: Design of the study, literature searches, patient's data analyses;

Jasna Zeljko-Penavić: Design of the study, literature searches, patient's data analyses;

Sandra Jozić: Patient's data analyses;

Ivana Goluža: Patient's data analyses.

References

- 1. Chosidow O: Clinical practices. Scabies. N Engl J Med 2006; 354:1718.
- 2. Dupuy A, Dehen L, Bourrat E, Lacroix C, Benderdouche M, Dubertret L et al.: Accuracy of standard dermoscopy for diagnosing scabies. J Am Acad Dermatol 2007; 56:53.
- 3. Eshagh K, DeKlotz CM, Friedlander SF: Infant with a papular eruption localized to the back. JAMA Pediatr 2014; 168:379-80.
- Estève E, Maitre F, Legac E: Purpura vasculaire au cours d'une gale sévère. Ann Dermatol Venereol 2001; 128:911-14.
- 5. Fuller LC: Epidemiology of scabies. Curr Opin Infect Dis 2013; 26:123-6.

- 6. Heukelbach J, Feldmeier H: Scabies. Lancet 2006; 367:1767.
- 7. Heukelbach J, Wilcke T, Winter B, Feldmeier H: Epidemiology and morbidity of scabies and pediculosis capitis in resource-poor communities in Brazil. Br J Dermatol 2005; 153:150.
- 8. Jarrett P, Snow J: Scabies presenting as a necrotizing vasculitis in the presence of lupus anticoagulant. Br J Dermatol 1998; 139:701-3.
- 9. Kos L, Shwayder T: Cutaneous manifestations of child abuse. Pediatr Dermatol 2006; 23:311-20.
- 10. Ly F, Caumes E, Ndaw CA, Ndiaye B, Mahé A: Ivermectin versus benzyl benzoate applied once or twice to treat human scabies in Dakar, Senegal: a randomized controlled trial. Bull World Health Organ 2009; 87:424-30.
- MahéA, Faye O, N'Diaye HT, Ly F, KonaréH, Kéita S et al.: Definition of an algorithm for the management of common skin diseases at primary health care level in sub-Saharan Africa. Trans R Soc Trop Med Hyg 2005; 99:39.
- 12. Mazzatenta C, Fimiani M, Flori ML, Andreassi L: Leucocytoclastic vasculitis following nodular scabies. J Eur Acad Dermatol Venereol 1996; 7:179-81.
- 13. Menné T, Christophersen J, Gram N, Bjerrehus T: Scabietic leucocytoclastic vasculitis with focal glomerulonephritis. Acta Derm Venereol 1984; 64:445-7.
- 14. Mikulic M, Jurisic K: The physicians role in recognizing physical abuse of children. Pediatrics Today 2013; 9:64-71.
- 15. Orkin M: Special forms of scabies. En/In: Orkin M, Maibach HI. Cutaneous infestations and insect bites. New York. Marcel Dekker Inc 1985: 25-30.
- 16. Pomares C, Marty P, Delaunay P: Isolated itching of the genitals. Am J Trop Med Hyg 2014; 90:589-90.
- 17. Ricci LR: Photographing the physically abused child. Principles and practice. Am J Dis Child 1991; 145:275-81.
- Romani L, Steer AC, Whitfeld MJ, Kaldor JM: Prevalence of scabies and impetigo worldwide: a systematic review. Lancet Infect Dis 2015; 15:960-7.
- 19. Skinner SM, DeVillez RL: Sepsis associated with Norwegian scabies in patients with acquired immunodeficiency syndrome. Cutis 1992; 50:213-6.
- Stinco G, Governatori G, Intersimone D, Frattasio A, Patrone P: Scabetic leukocytoclastic vasculitis. Eur J Dermatol 2008; 18:479-81.
- 21. Valks R, Buezo GF, Dauden E: Scabies and leukocytoclastic vasculitis in an HIV-seropositive man. Int J Dermatol 1996; 35:605-6.
- 22. Walter B, Heukelbach J, Fengler G, Worth C, Hengge U, Feldmeier H: Comparison of dermoscopy, skin scraping, and the adhesive tape test for the diagnosis of scabies in a resource-poor setting. Arch Dermatol 2011; 147:468.

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