# **ORIGINAL RESEARCH**

# Descriptive cross-sectional assessment of the comorbidities and environmental factors associated with atopic dermatitis in children and adults in dermatologyvenereology

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#### ABSTRACT

**Aim:**The objective of this work was to document the comorbidities and environmental factors associated with atopic dermatitis (AD) in dermatology Venereology.**Methods:**A descriptive cross-sectionalstudy was carried out in the Dermatology-Venereology. In the Dermatology-Venereology Departmentover the study period, we received 960 children and 2254 adults. Of these 960 children and 2254 adults, 200 children and 100 adults had AD.**Results:**Children between 0 and 5 years of age made up the majority of patients, 52% of the pediatric population, while young adults between 19 and 30 years of age made up the majority (40%) of the adult population. The median age at the time of the first episode was 4.7 years  $\pm$  4.6 in children. Among adults, the median age at the time of the first episode was 31.9 years  $\pm$  17.The main associated comorbidities were rhinitis and conjunctivitis in both children and adults but in different proportions: 50% of cases of rhinitis and 35% of cases of conjunctivitis in children against 34% and 25%, respectively, in adults. Asthma was present in 15% of children.**Conclusion:**In the Dermatology-Venereology Department, atopic dermatitis was associated with other atopic manifestations, the most frequent of which were rhinitis, conjunctivitis, and asthma. Environmental factorssuch as regular deworming, full vaccination, and living in anurban area were associated with the occurrence of AD. Heat associated with high humidity, skin irritants, and pneumallergens were reported in significant proportions.

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#### **INTRODUCTION**

Typical comorbidities in patients with atopic dermatitis (AD) include allergic asthma, allergic rhinoconjunctivitis and food allergy<sup>1</sup>. In addition, it has been shown that AD is associated with a number of other diseases, particularly autoimmune disorders such as rheumatoid arthritis, inflammatory bowel disease<sup>2</sup>, systemic lupus erythematosus<sup>3</sup>, vitiligo<sup>4</sup>,and alopecia areata<sup>5-7</sup>. As early as 1942, atopic individuals were described as being tense, nervous, depressed, introverted and anxious<sup>8</sup>.

Atopic dermatitis (AD) is a chronic and recurrent pruritic inflammatory dermatosis that preferentially

affects infants. Worldwide, its prevalence has doubled or even tripled in the last 30 years <sup>9-11</sup>. Several studies show that it is not a simple disease, but a skin condition with many different genetic and immunological mechanisms and underlying environmental factors. These factors influence the prevalence and clinical expression of the disease in different age groups, geographical regions and races<sup>12-16</sup>.

Atopic eczema (AE) is one of the most common inflammatory skin diseases<sup>17</sup>. The pathogenesis of AE is multifactorial resulting from a complex interaction between genetic and environmental factors. The onset

criteria.

of AE is usually in early childhood, that often persists into or may begin in adulthood<sup>18</sup>. The lifetime prevalence of AE ranges from 7-20% in children, and it is considered to be one of the major allergic diseases alongside asthma and allergic rhinitis <sup>19</sup>. AE is characterized by an itchy rash, with typical morphology, site and age-specific patterns, which are chronic and relapsing. This chronic condition imposes a great burden on the family and patient, regardless of age. This burden has psychosocial, educational and occupational impacts, in addition to major financial burden via direct medical costs and decreased productivity<sup>20</sup>. The Global Burden of Disease Study stated that the burden of dermatitis as assessed by Disability-Adjusted Life Years is high<sup>21</sup>.

The objective of this work was to document the comorbidities and environmental factors associated with atopic dermatitis (AD) in dermatology Venereology.

### MATERIALS AND METHODS

A descriptivecross-sectionalstudy was carried out in the Dermatology-Venereology.In the Dermatology-Venereology Departmentover the study period, we received 960 children and 2254 adults. Of these 960 children and 2254 adults, 200 children and 100 adults had AD.

The children and adults diagnosed with AD, after the free and informed consent of adult patients and regarding to children, those of their parents. The study was approved by the department head. The diagnostic criteria of the United Kingdom Working Party were used for the diagnosis of ADin children under 10 years of age<sup>9</sup>. In children over 10 years of age and adults, anamnestic (personal history of pruritic dermatosis, personal or family history of asthma, and/or allergic rhinitis or other atopic manifestations) and clinical (morphological and topographical aspects characteristic of AD in older children, adolescents, and adults, the presence of minor signs of atopy) arguments were used to establish the diagnosis of

#### RESULTS

#### Table 1: Age distribution

terns, which are childhood, before 18 years of age and late when the first onset occurred after this period<sup>22</sup>. The degree of severity was assessed with the SCORAD (scoring of atopic dermatitis). The triggering and/or aggravating environmental factors were identified on the basis of

the patients' allegations. The triggering factors sought were pneumallergens (house dustmites, pollen, mold, pet dander and smoke), infections, teething in infants, skin irritants (perfumed products, detergents and synthetic or woolen clothing), trophallergens(milk, egg white, peanuts, soy, shellfish and okra), psychological factors (stress and psychoaffective conflicts in adolescents and adults), hormonal factors in girls (premenstrual period), and physicochemical and climatic factors (sweat, heat, cold, and harmattan). Allergological tests, such as specific immunoglobulin E assays, prick tests, and patch tests, were not available to assess the relevance of the environmental factors reported by the patients. We relied on a thorough medical history (police-like questioning) to determine the notion of the cause and effect in the occurrence or recurrence of AD flare-ups in these patients. In case of a negative or doubtful answer, no factor was retained. The study was approved by local ethics committee of faculty of health sciences.

AD<sup>22,23</sup>. Sampling was exhaustive, non-probabilistic

and nonrandomized and included all patients admitted

to dermatology consultations who met the inclusion

According to the period of onset in adults, AD was

classified as persistent when the first onset occurred in

Demographic, socioeconomic, and clinical data were entered and analyzed with Epi-Data 3.1 and Epi-Info 7.0 softwares. The search for links between the occurrence of dermatosis and the other independent variables under studywas carried out by univariate analysis; Pearson's chi-squareor Fischer's test was used to search for the association between the variables. Results were significant when p<0.05.

0-5	104	52
6-10	56	28
Children 11-15	32	16
16-18	8	4
Total	200	100
19-30	40	40
31-40	25	25
Adults 41-50	15	15
51-60	12	12
>60	8	8
Total	100	100

Children between 0 and 5 years of age made up the majority of patients, 52% of the pediatric population, while young adults between 19 and 30 years of age

made up the majority (40%) of the adult population. The median age at the time of the first episode was 4.7 years  $\pm$  4.6 in children. Among adults, the median age at the time of the first episode was 31.9 years  $\pm$  17.

	ChildrenNumber(%)	<i>P</i> -value	AdultsNumber(%)	<b>P-value</b>
Rhinitis	100(50)	0.720	34(34)	0.640
Conjunctivitis	70(35)	0.430	25(25)	0.412
Asthma	30(15)	0.001	16(15)	0.240
ComorbiditiesSinusitis	8(4)	0.325	27(25.2)	0.720
Foodallergy	6(3)	—	4(3.7)	
Drugallergy	0(0)		3(2.7)	_
Prurigo strophulus	6(3)		0(0)	_
AntecedentsDeworming	120(6)	0.012	48(48)	0.732
Vaccination	110(55)	0.001		_
Urban	110(55)		60(60)	
ResidencePeriurban	80(40)	0.002	38(38)	0.036
Rural	6(3)		3(3)	
	Dry season Rainy season	90(45)	40(40)	
	Heat	84(42)	— 48(48)	
	Pneumallergens	56(28)	— 35(35)	
	Skin irritants	44(22)	— 58(58)	
Triggering and/or aggravating factors	Trophallergens	35(17.5)	— 14 (14)	
	Psychological	28(14)	— 34(34)	
	Infections	20(10)		_
	Teething	16(8)		

Table 2: Triggering or aggravating environmental factors

The main associated comorbidities were rhinitis and conjunctivitis in both children and adults but in different proportions: 50% of cases of rhinitis and 35% of cases of conjunctivitis in children against 34% and 25%, respectively, in adults. Asthma was present in 15% of children.

# DISCUSSION

Our study confirms the high prevalence of AD in children (16.71%) is pediatric prevalence is significantly higher than that reported in Abidjan (Cote d'Ivoire) in 2017<sup>24</sup>. Several studies confirm the increase in AD prevalence over the last 10 years, particularly in developing countries<sup>12-14</sup>.In Africa, the prevalence of AD varies from country to country between 4.7% and 23%<sup>25</sup>. Genetic factors alone are therefore not sufficient to explain the increase in AD around the world. There is a complex interrelation between these factors and environmental factors, which partly explains this observed disparity. The comorbidities frequently found in both children and adults were rhinitis and conjunctivitis. According to some authors, AD is considered the first manifestation of atopic gait, followed by food allergy, asthma and rhinitis or rhinoconjunctivitis<sup>9,26</sup>. These allergic manifestations are often found in high proportions in atopic patients<sup>26-28</sup>. According to some authors, the risk ofdeveloping allergic rhinitis and asthma in the presence of AD is more or less important<sup>29</sup>.

Considered as the key initiating event of atopic march, the alteration of the skin barrier accounts for the link between atopic dermatitis and subsequent atopic diseases<sup>29,30</sup>. In our study, asthma, which is

significantly more frequent in adolescents and the multiplicity of allergic diseases, associated with the onset of AD, confirm this hypothesis. However, the notion of an atopic march has recently been controversial. On the one hand, some authors believe that allergic manifestations, collected on the basis of patient claims, are overestimated in most studies <sup>31,32</sup>.On the other hand, AD is not always associated with other allergic manifestations. It has been suggested that these different manifestations, while sharing genetic and environmental risk factors, are independent conditions that can develop concomitantly or sequentially on an atopic site<sup>32</sup>. However, the concept of atopic march offers the possibility of research on the pathogenesis prospects for the prevention and treatment of atopic diseases. Correct measures to maintain or restore skin barrier function may help minimize the risk of developing allergic manifestations.

Heat was a reported contributing factor for both children and adults. The majority of flare-ups occurred during the hot season. A high temperature causes sweating, which becomes more important when humidity increases. This weather condition is the one observed in our region, hence the high frequency of this factor. Sweat can irritate the skin due to its acidic pH. This could promote Th2 inflammation, increased skin blood flow and a pruritogenic mechanism via nerve endings in the skin. There is not only a correlation between temperature and other climatic factors, namely, humidity, exposure to ultraviolet rays but also the pollen concentration in the environment, alteration of barrier function, and skin irritants. This indicates the important role of climate and even climate change, in the increase in flare-ups of AD<sup>12,16</sup>. Our study also confirms the role of skin irritants in the occurrence of flare-ups. Intrinsic barrier dysfunction can be aggravated when environmental factors such as soap and detergents cause further degradation of the epidermal barrier and irritants and allergens can interact with the immune system and promote inflammation. These irritants are thought to promote the synthesis of immunoglobulin E and sensitization to allergens<sup>33</sup>.Similarly, the use of detergent or lightening soaps was associated with specific clinical forms of AD.

# CONCLUSION

In the Dermatology-Venereology Department, atopic dermatitis was associated with other atopic manifestations, the most frequent of which were rhinitis, conjunctivitis, and asthma. Environmental factorssuch as regular deworming, full vaccination, and living in anurban area were associated with the occurrence of AD. Heat associated with high humidity, skin irritants and Pneumallergens were reported in significant proportions.

Multivariate analysis studies on a large series in the general population, supported by the demonstration of thesefactors using allergological tests and will provide a better understanding of these factors and their role in the onset or aggravation of atopic dermatitis. All these will contribute to a better knowledge of the pathophysiology of the disease and to a better therapeutic and above all preventive approach.

## REFERENCES

- AWMF
   LeitlinieNeurodermitis

   [atopischesEkzem;atopische
   Dermatitis]

   Entwicklungsstufe: S2k. AWMF online 2015.
- Schmitt J, Schwarz K, Baurecht H, Hotze M, Fölster-Holst R, Rodríguez E, Lee YA, Franke A, Degenhardt F, Lieb W, Gieger C. Atopic dermatitis is associated with an increased risk for rheumatoid arthritis and inflammatory bowel disease, and a decreased risk for type 1 diabetes. Journal of Allergy and Clinical Immunology. 2016 Jan 1;137(1):130-6.
- Wu LC, Hwang CY, Chung PI, Hua TC, Chen YD, Chu SY, Lee DD, Chang YT, Wang WJ, Liu HN, Chen CC. Autoimmune disease comorbidities in patients with atopic dermatitis: a nationwide case-control study in Taiwan. Pediatric Allergy and Immunology. 2014 Oct;25(6):586-92.
- 4. Chen YT, Chen YJ, Hwang CY, Lin MW, Chen TJ, Chen CC, Chu SY, Lee DD, Chang YT, Liu HN. Comorbidity profiles in association with vitiligo: a nationwide population-based study in Taiwan. Journal of the European Academy of Dermatology and Venereology. 2015 Jul;29(7):1362-9.
- 5. Mohan GC, Silverberg JI. Association of vitiligo and alopeciaareata with atopic dermatitis: A systematic review and meta-analysis.JAMA Dermatol 2015; 151(5): 522-8.
- 6. Magen E, Chikovani T, Waitman DA, Kahan NR. Association of alopecia areata with atopic dermatitis

and chronic spontaneousurticaria. Allergy Asthma Proc 2018; 39(2): 96–102.

- Brunner PM, Silverberg JI, Guttman-Yassky E, Paller AS, Kabashima K, Amagai M, Luger TA, Deleuran M, Werfel T, Eyerich K, Stingl G. Increasing comorbidities suggest that atopic dermatitis is a systemic disorder. Journal of Investigative Dermatology. 2017 Jan 1;137(1):18-25.
- 8. Allerhand ME, Gough HG, Grais ML. Personality factors inneurodermatitis; a preliminary study. Psychosom Med 1950;12: 386–90.
- Taieb A. Atopic dermatitis: definition, epidemiology, natural history, severity and scores. In Annals of Dermatology and Venereology 2005 Jan 1 (Vol. 132, pp. 35-43). Elsevier Masson.
- Mahé E. Dermatite atopique: épidémiologie en France, définitions, histoire naturelle, association aux autres manifestations atopiques, scores de gravité, qualité de vie. InAnnales de Dermatologie et de Venereologie 2005 Jan 1 (Vol. 132, pp. 131-150). Elsevier Masson.
- K. Ezzedine and E. Kechichian, "Epid'emiologie de la dermatite atopique," Annales de Dermatologie et de V'en'er'eologie, vol. 144, pp. VS4–VS7, 2017.
- Schmid-Grendelmeier P, Takaoka R, Ahogo KC, Belachew WA, Brown SJ, Correia JC, Correia M, Degboe B, Dorizy-Vuong V, Faye O, Fuller LC. Position statement on atopic dermatitis in sub-Saharan Africa: current status and roadmap. Journal of the European Academy of Dermatology and Venereology. 2019 Nov;33(11):2019-28.
- 13. Torrelo A. Atopic dermatitis in different skin types. What is to know?. Journal of the European Academy of Dermatology and Venereology. 2014 May;28:2-4.
- Kim Y, Blomberg M, Rifas-Shiman SL, Camargo Jr CA, Gold DR, Thyssen JP, Litonjua AA, Oken E, Asgari MM. Racial/ethnic differences in incidence and persistence of childhood atopic dermatitis. Journal of Investigative Dermatology. 2019 Apr 1;139(4):827-34.
- Zar HJ, Ehrlich RI, Workman L, Weinberg EG. The changing prevalence of asthma, allergic rhinitis and atopic eczema in African adolescents from 1995 to 2002. Pediatric allergy and immunology. 2007 Nov;18(7):560-5.
- Bonamonte D, Filoni A, Vestita M, Romita P, Foti C, Angelini G. The role of the environmental risk factors in the pathogenesis and clinical outcome of atopic dermatitis. BioMed research international. 2019 Apr 21;2019.
- Deckers IA, McLean S, Linssen S, Mommers M, Van Schayck CP, Sheikh A. Investigating international time trends in the incidence and prevalence of atopic eczema 1990-2010: a systematic review of epidemiological studies. PloS one. 2012 Jul 11;7(7):e39803.
- Margolis JS, Abuabara K, Bilker W, Hoffstad O, Margolis DJ. Persistence of mild to moderate atopic dermatitis. JAMA dermatology. 2014 Jun 1;150(6):593-600.
- Schneider L, Tilles S, Lio P, Boguniewicz M, Beck L, LeBovidge J, Novak N, Bernstein D, Blessing-Moore J, Khan D, Lang D. Atopic dermatitis: a practice parameter update 2012. Journal of Allergy and Clinical Immunology. 2013 Feb 1;131(2):295-9.
- 20. Drucker AM, Wang AR, Li WQ, Sevetson E, Block JK, Qureshi AA. The burden of atopic dermatitis: summary of a report for the National Eczema

Association. Journal of Investigative Dermatology. 2017 Jan 1;137(1):26-30.

- Karimkhani C, Dellavalle RP, Coffeng LE, Flohr C, Hay RJ, Langan SM, Nsoesie EO, Ferrari AJ, Erskine HE, Silverberg JI, Vos T. Global skin disease morbidity and mortality: an update from the global burden of disease study 2013. JAMA dermatology. 2017 May 1;153(5):406-12.
- Reguiaï Z. Atopic dermatitis in adults: clinical presentation, complications and comorbidities. In Annals of Dermatology and Venereology 2017 Dec 1 (Vol. 144, pp. VS15-VS22). Elsevier Masson.
- J. Silvestre Salvador, D. Romero-P'erez, and B. Encabo-Dur'an, "Atopic dermatitis in adults: a diagnostic challenge," Journalof Investigational Allergology and Clinical Immunology, vol. 27, no. 2, pp. 78–88, 2017.
- K. C. Ahogo, Y. I. Kouassi, I. P. Gbery, K. R. Azagoh, K. I. Yeboua, and K. A. Kouassi, "Atopic dermatitis in children: epidemiological and clinical aspects in Côte d'Ivoire," Our Dermatology Online, vol. 8, no. 1, pp. 25–27, 2017.
- Kaufman BP, Guttman-Yassky E, Alexis AF. Atopic dermatitis in diverse racial and ethnic groups-variations in epidemiology, genetics, clinical presentation and treatment. Experimental dermatology. 2018 Apr;27(4):340-57.
- 26. ChiesaFuxench ZC. Atopic dermatitis: disease background and risk factors. Management of Atopic Dermatitis: Methods and Challenges. 2017:11-9.
- A. ´Cosi´cki´c, F. Skoki´c, A. Selimovi´c, M. Muli´c, S. Suljendi´cand N. NerminaDedi´c, "Development of respiratory allergies,asthma and allergic rhinits in children with atopic dermatitis", Acta ClinicaCroatica, vol. 56, pp. 308–317, 2017.
- Chu H, Shin JU, Park CO, Lee H, Lee J, Lee KH. Clinical diversity of atopic dermatitis: a review of 5,000 patients at a single institute. Allergy, asthma & immunology research. 2017 Mar 1;9(2):158-68.
- Taniuchi S, Soejima K, Hatano Y, Takahashi M, Minami H. Dual factors may Be necessary for development of atopic march in early infancy. Journal of Nippon Medical School. 2018 Jan 15;85(1):2-10.
- van den Oord RA, Sheikh A. Filaggrin gene defects and risk of developing allergic sensitisation and allergic disorders: systematic review and metaanalysis. Bmj. 2009 Jul 9;339.
- 31. Busse WW. The atopic march: fact or folklore? Annals of Allergy, Asthma & Immunology. 2018 Feb 1;120(2):116-8.
- 32. Yang L, Fu J, Zhou Y. Research progress in atopic march. Frontiers in immunology. 2020 Aug 27;11:1907.
- 33. Kantor R, Silverberg JI. Environmental risk factors and their role in the management of atopic dermatitis. Expert review of clinical immunology. 2017 Jan 2;13(1):15-26.