Epidemiology of Skin Diseases from the Spectrum of Dermatitis and Eczema

Agnieszka Dorynska¹, *MSc*, *MPH*, Radoslaw Spiewak^{1,2}, *MD*, *PhD*, *Professor of Experimental Dermatology*

Abstract

Particular types of eczema may affect up to 29% individuals in certain populations (lifetime prevalence), thus placing the diseases among most frequent clinical problems. Nevertheless, diseases from the spectrum of dermatitis and eczema are poorly defined and frequently misdiagnosed; they also frequently overlap, making the diagnostic process even more difficult. In doubtful cases, where no further means of clinical or laboratory differentiation are available, reliable epidemiological data may provide relevant help in the diagnostic process, as the best candidate for a tentative diagnosis seems the most frequent among diseases in question, which can be verified later by the effectiveness of respective treatment regimen. However, results of epidemiological studies in the field of eczema and dermatitis may be strikingly contradictory, one of the possible reasons being definitions of various types of eczema/dermatitis that leave too much space for individual decision and thus seem hardly suitable for epidemiological research. Better studies based on unequivocal definitions of various types of eczema are necessary to achieve the quality of epidemiological data that would ensure the level of certainty expected from a diagnostic tool. The present paper collates results from available epidemiological data on various types of eczema: atopic eczema, allergic and irritant contact dermatitis, protein contact dermatitis, seborrhoeic dermatitis, asteatotic dermatitis, stasis dermatitis, nummular eczema, dyshidrotic eczema (pompholyx), hand dermatitis and occupational dermatitis. Problems and possible sources of bias in available studies are addressed and discussed along with the results from the studies.

Keywords: epidemiology; atopic eczema; allergic and irritant contact dermatitis; protein contact dermatitis; seborrhoeic dermatitis; asteatotic dermatitis; stasis dermatitis; nummular eczema; dyshidrotic eczema; pompholyx; hand dermatitis; occupational dermatitis

Correspondence

Radoslaw Spiewak ¹ Institute of Dermatology, Krakow, Poland ² Department of Experimental Dermatology and Cosmetology, Faculty of Pharmacy Jagiellonian University Medical College ul. Medyczna 9, 30-688 Krakow, Poland Tel: +48 601 22 48 13 Fax: +48 12 416 62 62 Email: spiewak.eu@gmail.com





Introduction

The knowledge of the frequency of diseases is important for policy makers, insurers, but it is also a very important diagnostic tool in the hand of a clinician. In a considerable group of patients the clinical picture does not allow for a clear-cut diagnosis, and after exhausting all available possibilities of differential diagnosis the clinician is stuck with two or more possible diagnoses.

In such cases, the knowledge of epidemiology may be resorted to as the ultimate instance of clinical decision. Knowing the prevalence rates of otherwise equally possible diseases that come in question, it seems rational to pick the more prevalent disease as the tentative diagnosis, with a possible revision if appropriate treatment turns out ineffective.

Eczema

(synonym: dermatitis) is noncontagious inflammation of the epidermis and dermis with characteristic clinical features (itch, erythema, papule, seropapule, vesicle, scale, squame, crust or lichenification that emerge simultaneously or evolve from one another) and distinct histological picture (spongiosis, acanthosis, parakeratosis, lymphocytic and granulocytic infiltrates)^{1,2}.

The debate on the differences between the terms "eczema" and "dermatitis" has been ongoing for many decades, with no definite conclusion^{3, 4}. Therefore, in the present article these terms will be considered as synonyms. The clinical spectrum of dermatitis/eczema diseases includes an array of diseases that sometimes are depicted as mutual opposites, however, their clinical features and pathomechanisms overlap to an extent making any clear-cut differentiation virtually impossible.

In epidemiological studies of the various dermatitides, most striking is the difficulty of drawing general conclusions, mainly due to imprecise definitions and incompatible outcome measures. This must be born in mind when looking at the epidemiological data discussed below.

In the analysis of diseases frequency, it is crucial to remember that different methods of collecting epidemiological data may give different outcomes. The most popular method to obtain epidemiological data on diseases is self-administered questionnaire. This method has some advantages, which are very important when conducting an epidemiological research: it is inexpensive and easy to use, so it can be applied in large populations.

Disadvantages of the questionnaire-based method are also very significant, especially the possibility of misunderstanding the questions which may lead to the probability of overestimation of the obtained results⁵.

Another method for assessing the frequency of diseases is medical examination. This method seems more objective and thus more reliable because it allows for verification of symptoms by a specialist. In comparison to the questionnaire-based method, medical examination requires much more costs and time for performing⁶. Moreover, when comparing these two methods of collecting epidemiological data, it is important to remember that questionnaire-based method is more suitable for collecting information about prevalence of diseases over a period of time (e.g., lifetime prevalence or one-year prevalence), while medical examination is more appropriate for assessing the presence of the disease at a particular point in time (point prevalence)⁷. Thus these methods should be regarded as complementary. Some estimates about the frequency of diseases come from various registers, such as hospital records, national or local statistics (e.g. occupational diseases statistics). This "ecological" (i.e. not consuming new resources) method has its advantages, for example it allows for comparison of trends at different time points. However, discrepancies may arise due to different classifications of diseases used in various data collecting systems, or in various periods of time.

A major possible disadvantage of using the "epidemiological approach" in clinical diagnosis is that of a "self-fulfilling prophecy": With poor-quality epidemiological data at the start, one may classify unclear cases of eczema for this type that is believed to be most frequent, which may not necessarily be the truth, however, by doing so the statistics are biased toward the tentative diagnosis, thus reinforcing one's beliefs into seemingly "scientific proof". It seems that this is especially true for the diseases from the spectrum of dermatitis and eczema.

Therefore, it is extremely important to be critical when looking at the frequency rates of diseases from the spectrum of dermatitis and eczema. The differences in definitions of the diseases in various studies or sometimes lack of any definitions, strongly supports this attitude. In this article, in order to be able to collate available epidemiological data, we have adopted a simplistic attitude that the diagnosis of a given disease is defined by the authors' declaration (i.e. belief) that they studied this particular disease. The following data, therefore, give us some idea about possible prevalence rates, however, due caution is recommended while using them for "epidemiological" diagnosis.

Atopic eczema

(AE, synonym: atopic dermatitis) is a chronic inflammatory skin disease that commonly begins in early infancy, runs a course of exacerbations and remissions, and is associated with a characteristic distribution and morphology of skin lesions. Furthermore, pruritus and subsequent sleeplessness are hallmarks of this disease⁸. This "minimalist" definition seems most acceptable for the time being, as it puts forward the common clinical characteristics while avoiding references to pathomechanisms, which are still subject to controversy (see below).

Prevalence of atopic dermatitis/eczema in children has been widely assessed. The most known epidemiological study on atopic eczema (AE) in children is the ISAAC Study⁹. This questionnaire-based study allows estimating one-year and lifetime prevalence rates of AE among children. Table 1 presents prevalence rates of atopic eczema according to studies based on the ISAAC questionnaire. Both indices of the disease frequency (one-year prevalence, and lifetime prevalence) showed great variability in the estimations among countries ranging from 4.5% to 20.2% (1-year prevalence) and from 2.4% to 28.7% (lifetime prevalence) ¹⁰⁻¹².

However, there has been a heated discussion on how reliable is the ISAAC questionnaire in detecting AE^{13,14}, with recent data showing that up to 50% of children with 'ISAAC eczema' may in fact be ill with allergic contact dermatitis (ACD)¹⁵. Flexural eczema - almost a "diagnostic fetish" in past epidemiological studies of AE has turned out less specific to AE than previously believed¹⁶, not least so because this clinical feature is also common in ACD¹⁷⁻²¹, and cases of ACD-related flexural eczema have been misdiagnosed as AE for decades^{22, 23}. With this respect, ISAAC studies may be looked at as an example of the "self-fulfilling prophecy" in the epidemiology of eczema in children. In order to overcome these limitations, other methods were also used when assessing the frequency of AE.

Detailed information on the prevalence of AE in children according to studies not based on the ISAAC questionnaire is shown in Table 2. Less is known on the prevalence of AE in adults - available data are collated in Table 3. The major problem with the epidemiological data of AE is that "atopic eczema" seems in fact to be a heterogeneous group of diseases with similar clinical appearance, rather than a single disease.

The spectrum of involved pathologies range from type I and IV allergy (possibly also types II and III), to barrier dysfunction, abnormalities of the innate immune response and autoimmunity, while it remains unclear, which of those are actual causes and which secondary phenomena²⁴⁻²⁷. For example, the causal role of IgE-mediated food allergy in AE seems overrated^{28, 29} and the development of food-specific IgE may, in fact, be secondary to eczema³⁰.

The name "atopic dermatitis" itself was already criticized by Rajka in 1975 as an "unfortunate choice of term"³¹, which is supported by the fact that a majority of AE patients show no evidence of atopy³². Perhaps "Hanifin-Rajka Syndrome" would be a more appropriate name for this entity, avoiding the reference to questionable aetiology and focusing instead on the common clinical picture first compiled by the authors.

Gender of Age of One-year Lifetime Country children children prevalence prevalence Austria^{33*} 6-9 (1995-97) Boys 5.0% 8.2% Girls 7.0% 10.2% 6-9 (2001-03) Boys 5.9% 10.2% Girls 7.6% 11.8% Brazil³⁴ 13-14 Boys and girls 16.2% China³⁵ 6-13 Boys and girls 5.5% -China^{36^} 14.5% 0-14 Boys and girls Croatia37 12-14 Boys and girls 5.3% 7.0% Germany³⁸ 14.3% 6-7 Boys 7.3% Girls 14.6% 6.7% 13-14 Boys 5.0% 8.2% Girls 9.4% 12.3% 6-7 13.6% Boys 6.6% Girls 9.8% 16.9% 13-14 Boys 4.5% 10.9% Girls 11.1%17.4% Ghana^{39^} 4-16 Boys 4.0% -Girls Iran⁴⁰ 13-14 Boys and girls 10.1% Italy⁴¹ 2 Boys 16.8% Girls 18.7% 3 Boys 16.2% Girls 20.2% 4 19.1% Boys _ Girls 17.2% Korea⁴² 8-11 Boys 12.7% 26.8% Girls 28.7% 14.5% Malta43 13-15 (1995) Boys and girls 12.8% 11.2% 8.5% 13-15 (2000) Boys and girls 10.1% Mexico^{44**} 6-8 10.1% 15.0% Boys and girls 11-14 10.5% 17.0% 6-8 Boys and girls 5.8% 7.3% 11-14 5.4% 7.0% Montenegro45 6-7 9.5% Boys and girls -13-14 9.1% _ Poland⁴⁶ 7 Boys and girls 9.4% _ 16 3.4% Serbia45 6-7 Boys and girls 11.2-17.2% _ 13-14 8.2-16.2% Spain47 6-7 Boys and girls 5.9% -10-11 Spain⁴⁸ Boys and girls 11.4% _ 1-2 15.0% 16.2% Sweden⁴⁹ 2-3 23.7% Boys and girls 20.2% 3-4 20.7% 25.8% 22.5% 6-7 Boys and girls 17.8% 7-8 16.6% 21.2% 8-9 20.7% 26.1% United Kingdom⁵⁰ 27.8% 6-7 Boys -Girls 27.0% _

Table 1 Prevalence rates of atopic dermatitis in children according to studies based on the ISAAC questionnaire.

* prevalence rates were estimated using questions about presence of an itchy rash in the past 12 months and lifetime symptoms of an itchy rash

** prevalence rates were estimated using questions about presence of dry itchy skin spots in the last 12 months and at any time

 $\wedge~$ calculated based on the figures provided by the authors

Country	Age of children	Method of assessment	Results
Denmark⁵¹	12-16	Q ME	<i>lifetime prevalence:</i> 21.3% (17.0% boys; 25.7% girls) <i>one-year prevalence:</i> 6.7% (5.6% boys; 7.7% girls) <i>point prevalence:</i> 3.6% (3.8% boys; 3.4% girls)
Denmark ⁵²	7	Q	lifetime prevalence: 22.9%
Gabon ³⁹	4-16	ME	point prevalence: 4.0%
Germany ⁵³	5-7	ME	point prevalence: 12.9%
Germany ⁵⁴	0-4	Q	lifetime prevalence: 21.4%
Germany ⁵²	7	Q	lifetime prevalence: 13.1%
Germany ⁵⁵	< 10	Q	<i>lifetime prevalence:</i> 13.0% (Leipzig), 13.9% (Munich)
Ghana ³⁹ ^	4-16	ME	point prevalence: 1.6%
Rwanda ³⁹	4-16	ME	point prevalence:: 0.8%
Spain ⁴⁸	10-11	ME	point prevalence: 1.9%
Sweden ⁵²	7	Q	lifetime prevalence: 15.5%
Turkey ⁵⁶	0-16	HR	lifetime prevalence: 11.8%
United Kingdom ⁵⁷	1-5	Q	<i>one-year prevalence</i> : 16.5% (22% in 1-2 y.o.; 19% in 2-3 y.o.; 13% in 3-4 y.o.; 15% in 4-5 y.o.)
United States ⁵⁸	5-9	Q	17.2% (standard scoring criteria) 6.8% (highly stringent criteria)

Table 2 Prevalence of atopic eczema in children in various studies based on different methods.

Q - questionnaire; ME - medical examination; HR - hospital record

^ own calculations based on the figures provided by the authors

Contact dermatitis

(Synonym: contact eczema) is a collective term for three dermatitides with various aetiologies, whose common feature is the development of skin inflammation in response to a direct contact with the provoking agent: 1) irritant contact dermatitis, 2) allergic contact dermatitis and 3) protein contact dermatitis⁶⁶.

Allergic contact dermatitis

(Synonym: allergic contact eczema) is inflammatory skin disease initiated by specific immune reaction to a hapten. It occurs in individuals with previously acquired contact allergy following re-exposure to the sensitizing hapten⁶⁷. In contrast to ICD, only a minority of people exposed to a particular hapten will respond with dermatitis. When looking at epidemiological data, one must remember that ACD is not the same as contact allergy (CA).

Country	Age of children	Method of assessment	Results
Australia ⁵⁹	20+	ME	<i>point prevalence:</i> 5.7% in men, 8.1% in women
Denmark ⁶⁰	18-69	Q	lifetime prevalence: 10.0%
Germany ⁶¹	0-99	Q ME	<i>lifetime prevalence: 23.5%</i> <i>point prevalence: 16.0%</i>
Japan ⁶²	20+	ME	<i>point prevalence:</i> 6.9% (participants in their 20s: 9.8%; 30s: 8.7%; 40s: 4.4%; 50/60s: 2.6%)
Norway ⁶³	18-69	Q	<i>lifetime prevalence:</i> 13.8% in men, 19.0% in women
Norway ⁶⁴	born in 1970, 1960, 1955, 1940- 1941 and 1924-1925	Q	lifetime prevalence: 8.8%
Poland ⁶⁵	18-19	Q ME	lifetime prevalence: 5.0% one-year prevalence: 3.9% point prevalence: 2.5%
Russia ⁶³	18-69	Q	<i>lifetime prevalence:</i> 10.4% in men, 12.0% in women

 Table 3 Prevalence rates of atopic eczema in adults.

Q - questionnaire; ME - medical examination

The term "contact allergy" refers to a state of altered response of the immune system to a specific substance, which is not synonymous with disease. Certain proportion of people with CA will never develop clinical symptoms. Among those symptomatic, vast majority will develop ACD, which is an inflammatory disease of the skin provoked by a hapten (a low molecular sensitizer), following the exposure to this hapten of a sensitized person⁶⁸. Confusing contact allergy with allergic contact dermatitis seems a frequent mistake of doctors and authors of clinical and epidemiological studies.

Children

A very comprehensive method of establishing the prevalence of ACD in children was used in the study conducted in Denmark. ACD in the group of 12-16 years old children was defined by the co-existence of the three criteria: 1) contact allergy diagnosed by a positive patch test 2) exposure history and 3) history or present dermatitis pattern. Lifetime prevalence of ACD was 7.2%, and point prevalence 0.7% (calculated on the basis of data provided in the article)⁵¹. A Polish study showed that among 7-year old children the lifetime prevalence of symptoms of ACD was slightly higher than among 16-year olds (7.2% versus 6.1%)⁴⁶. This is also reflected in higher contact hypersensitivity rates among children (67.0%) than adolescents (58.1%) seen in a similar cohort of Polish children⁶⁹, which may be explained by changing exposure patterns in the rapidly westernising country⁷⁰.

Adults

In the United States, in a study of university students, ACD was the cause of 3.1% of first-time visits to dermatologists, and 2.4% of total visits to dermatologists⁷¹. In Poland, prevalence of ACD was assessed among students of vocational agricultural schools. History and symptoms-based physician diagnosis estimated the frequency of ACD as: 2.0% (point prevalence), 9.3% (one-year prevalence), and 17.5% (lifetime prevalence)⁶⁵.

Irritant contact dermatitis

(ICD) is acquired inflammatory skin disease caused by chemical or physical insults leading to direct cellular injury. Most of ICD cases are associated with detergents, solvents, acids or alkali. Acute ICD (toxic dermatitis) develops rapidly (minutes to hours) after exposure to potent irritants, while chronic, cumulative variants of ICD develop gradually in response to repeated contacts with milder irritants⁷². ICD is essentially an injury, therefore, everyone will develop this disease after an individual threshold of resistance to irritants is exceeded⁷³.

The prevalence of irritant contact dermatitis (ICD) in general population is hard to determine, especially among children. Study conducted on a group of university students in the United States, showed that ICD was the cause of 2.3% of first-time visits to dermatologists, and 1.6% of total visits to dermatologists⁷¹. In Poland, estimations from the study conducted among students of a vocational school were: 0.5% (point prevalence), 4.3% (one-year prevalence), and 12.7% (lifetime prevalence)⁶⁵.

Protein contact dermatitis

(PCD) is acquired inflammatory skin disease initiated by specific immune reactions to allergens - proteins with molecular weight exceeding 10000 Daltons, usually of animal or plant origin^{74, 75}. There is lack of data on the frequency of protein contact dermatitis among children. Estimates for adults are available only for work-related settings. In Finland, protein contact dermatitis (together with contact urticaria) accounted for 11.1% of all allergic occupational diseases reported in 1991⁷⁶. Protein contact dermatitis was found in 22% of a group of 144 slaughterhouse workers in Denmark⁷⁷.

Seborrhoeic dermatitis

Seborrhoeic dermatitis is an inflammatory skin disease of the dermatitis/eczema spectrum, with a characteristic restriction to "seborrhoeic areas", i.e. areas with a high density of sebaceous glands (face, sternum, interscapular area). The aetiology remains unclear, one possibility being the excessive development of lipophilic Malassezia yeasts on the seborrheic skin with secondary development of inflammation in response to signalling molecules such as malassezin⁷⁸.

Little is known on the prevalence of seborrhoeic dermatitis. In a Turkish study of paediatric patients (0-16 years old) in a hospital registry, 4.3% children were diagnosed with seborrhoeic dermatitis⁵⁶. The prevalence of seborrhoeic dermatitis in adults was established in an Australian study based on medical examination, was 12.3% in men, and 7.3% in women⁵⁹. Among university students in the USA, seborrhoeic dermatitis was the cause of 3.1% of first-time visits, and of 2.4% of all dermatologist consultations⁷¹.

In a prospective, skin examination-based study of renal transplant recipients in the UK, seborrhoeic dermatitis was found in 9.5% of the participants⁷⁹. The prevalence of seborrhoeic dermatitis of the face and scalp diagnosed among mountain guides was 16.3%, which might hint on a role of UV irradiation in these cases⁸⁰.

Asteatotic dermatitis

Asteatotic dermatitis (dry skin dermatitis, winter itch) is an entity of unknown aetiology, characterised by the presence of dry, scaly, fissuring skin accompanied with pruritus, typically localised on the calves, with a possibility of spreading. Among exacerbating/causative factors, skin ageing with atrophy and xerosis, low humidity of ambient air, as well as frequent bathing and excessive detergent use are mentioned. Among Australian adults the prevalence of doctor-diagnosed asteatotic dermatitis was 6.6% in men, and 10.4% in women⁵⁹.

Stasis dermatitis

Stasis dermatitis is a skin manifestation of venous insufficiency and frequently is accompanied by other symptoms like the presence of varicous veins, leg oedema and ulcers, hemosiderin deposits in the skin and liposclerosis of the skin. The typical localization is calves. In the above-mentioned Australian study, the frequency of stasis dermatitis was assessed at 2.1% in men, and 1.5% in women⁵⁹.

Nummular eczema

Nummular eczema (nummular dermatitis, discoid dermatitis) is characterized by solitary or multiple, well-demarcated, round or oval-shaped itchy lesions. The typical course of the disease is chronic recurrent. The identity of this disease is built based upon the characteristic clinical appearance; however, the aetiology remains unknown. One of the more popular hypotheses considers immunological response (allergic reaction type II or IV) to circulating antigens of bacteria, fungi or parasites. On the other hand, it seems that may various types of eczema may take this clinical appearance, e.g. atopic eczema, allergic contact dermatitis (to nickel, neomycin, etc.), along with asteatotic and stasis dermatitis. In a Turkish study utilizing data of hospitalised paediatric patients, 0.4% children (0-16 years old) were diagnosed with nummular dermatitis⁵⁶.

Dyshidrotic eczema

Dyshidrotic eczema (pompholyx) is a non-infectious inflammation of the skin characterized by the appearance of pruritic vesicles on the palms and soles. The course of

Table 4 Prevalence rates of hand dermatitis/ecze
--

CHILDREN	Ago of	Mathad of	
Country	children	assessment	Results
Denmark⁵¹	12-16	Q ME	<i>lifetime prevalence:</i> 9.2% (6.3% boys, 12.2% girls) <i>one-year prevalence:</i> 7.3% (4.6% boys, 10.1% girls) <i>point prevalence:</i> 3.2% (2.2% boys, 4.2% girls)
Norway ⁸⁴	7-12	Q ME	one-year prevalence: 6.5% point prevalence: 3.5%

ADULTS

Country	Age of children	Method of assessment	Results
Denmark ^{60*}	18-69	Q	<i>lifetime prevalence:</i> 21.8% (17.0% men, 25.7% women) <i>one-year prevalence:</i> 11.7% (8.9% men, 14.0% women)
Norway ⁶⁴	born in 1970, 1960, 1955, 1940- 1941 and 1924-1925	Q	lifetime prevalence: 8.2%
Poland ⁸⁵	20-73	Q	<i>lifetime prevalence:</i> 17.3% <i>one-year prevalence:</i> 10.1% <i>point prevalence:</i> 1.9%
Sweden ⁸⁶	20-65	Q ME	one-year prevalence: 11.0% point prevalence: 5.4%
Sweden ⁸⁷	20-65	Q	<i>one-year prevalence:</i> 11.8% (1983) and 9.7% (1996)
Sweden ^{88*}	20-77	Q	<i>lifetime prevalence:</i> 11.0% (6.8% men, 14.0% women) <i>one-year prevalence:</i> 6.5% (4.5% men, 8.1% women)

Q - questionnaire; ME - medical examination

* calculated based on the figures provided by the authors

the disease may be acute, recurrent, or chronic. The skin lesions frequently are restricted to areas with high density of sweat glands and frequently accompanied by hyperhidrosis⁸¹. However, it appears that the lesions are not connected with the glands.

In the above-mentioned Turkish study, dyshidrotic eczema was diagnosed in 1.0% of paediatric hospital patients (0-16 years old)⁵⁶. In an epidemiological study of adult Dutch metalworkers, symptoms of dyshidrotic eczema were found in 7.3% of a group of metalworkers⁸².

Hand dermatitis

Hand dermatitis is a very special nosological entity that refers to the clinical picture (dermatitis localized on the hands) rather, than to the cause. Hand dermatitis/eczema may be a manifestation of ACD, ICD, atopic dermatitis, or other inflammatory diseases, which in this location are very difficult to differentiate based on the clinical picture or medical tests (including histopathology). A co-existence of more than one causes of hand dermatitis (e.g. ACD + ICD + atopic hand dermatitis) is relatively common, hence it seems practical to view hand dermatitis as a distinct clinical entity⁸³. Prevalence rates of hand dermatitis/eczema in children and adults are shown in Table 4.

Occupational dermatitis

Occupational contact dermatitis is neither clinical nor pathological entity; however, due to specific circumstances of appearance and special legal status in many countries, cases of such diseases are closely followed. OCD occurs mostly on hands (80% cases) and face (10% cases)⁸⁹. The frequency of occupational contact dermatitis (OCD) in the United Kingdom is estimated as 12.9 cases per 10 thousand full-time workers each year⁹⁰. One-year prevalence of occupational hand dermatitis, depending on the method of estimation, varies from 0.5-6.7% (medical examination) to 8.2-10.6% (questionnaire) in different populations⁹¹.

In a study based on medical examination, 4.1% Polish farmers were diagnosed with occupational hand eczema⁹². One in three of those who stated to have hand dermatitis ever, and one in five with wrist and forearm dermatitis reported on exacerbations of dermatitis due to substances present at workplace⁸⁵. Irritant contact dermatitis (ICD) and allergic contact dermatitis (ACD) contribute to most cases of OCD. Different proportions of ICD and ACD are reported in studied populations - frequency of ICD varies from 32% (USA) to 71% (Australia)⁸⁹. The differences might reflect the diagnostic routines (most importantly the use and extensiveness of patch tests).

Final remarks

The major disadvantage of available epidemiological studies of diseases from eczema and dermatitis spectrum is that they depend on clinical symptoms, which are frequently difficult to properly classify even by an experienced clinician, as clinical features and pathomechanisms of various types of eczema overlap to an extent making clearcut differentiations virtually impossible. Studies based on self-administered questionnaires, are even more susceptible to bias as conclusions are built based upon patient's own opinions and interpretations. Furthermore, various types of eczema may co-exist, while most researchers and doctors rest satisfied with a first diagnosis established. To acquire reliable data on the epidemiology of various types of dermatitides, better studies are needed in the future based on well-defined criteria that would enable accurate differentiation between analysed diseases. Specific requirements for such studies have been recently discussed elsewhere93.

Reference

- 1. Weidinger S, Ring J. Diagnosis of atopic eczema. In: Ring J, Przybilla B, Ruzicka T, editors. Handbook of Atopic Eczema. Second Edition. Berlin: Springer; 2006:84-99.
- Darsow U, Wollenberg A, Simon D, et al. ETFAD/EADV eczema task force 2009 position paper on diagnosis and treatment of atopic dermatitis. J Eur Acad Dermatol Venereol 2009;24:317-328.
- Ackerman AB, Ragaz A. A plea to expunge the word "eczema" from the lexicon of dermatology and dermatopathology. Arch Dermatol Res 1982;272:407-20.
- Altekrueger I, Ackerman AB. "Eczema" revisited. A status report based upon current textbooks of dermatology. Am J Dermatopathol 1994;16:517-22.
- 5. Dotterud LK, Falk ES. Evaluation of a self-administered questionnaire on atopic diseases. Discrepancy between self-reported symptoms and objective signs. Eur J Public Health 2000;10:105-107.

- Perreault N, Brisson C, Dionne CE, et al. Agreement between a self-administered questionnaire on musculoskeletal disorders of the neck-shoulder region and a physical examination. BMC Musculoskelet Disord 2008;17;9:34.
- Berg M. Evaluation of a questionnaire used in dermatological epidemiology. Discrepancy between self-reported symptoms and objective signs. Acta Derm Venereol Suppl (Stockh) 1991;156:13-7.
- 8. Werfel T, Breuer K. Role of food allergy in atopic dermatitis. Curr Opin Allergy Clin Immunol 2004;4:379-385.
- 9. The ISAAC Study [electronic resource, URL: http://isaac. auckland.ac.nz/about/about.php] (accessed on 5 June 2012).
- 10. Odhiambo JA, Williams HC, Clayton TO, et al.; ISAAC Phase Three Study Group. Global variations in prevalence of eczema symptoms in children from ISAAC Phase Three. J Allergy Clin Immunol 2009;124:1251-8.e23.

- Sole D, Mallol J, Wandalsen GF, Aguirre V; Latin American ISAAC Phase 3 Study Group. Prevalence of symptoms of eczema in Latin America: results of the International Study of Asthma and Allergies in Childhood (ISAAC) Phase 3. J Investig Allergol Clin Immunol 2010;20:311-23.
- 12. Flohr C. Recent perspectives on the global epidemiology of childhood eczema. Allergol Immunopathol (Madr) 2011;39:174-82.
- Flohr C, Williams HC. Childhood eczema according to the International Study of Asthma and Allergies in Childhood (ISAAC) questionnaire tool - response to Czarnobilska et al. J Eur Acad Dermatol Venereol 2011;25:993-4.
- 14. Spiewak R, Czarnobilska E. Not all that looks like eczema is atopic eczema. J Eur Acad Dermatol Venereol 2011;25:992-3.
- Czarnobilska E, Obtulowicz K, Dyga W, Spiewak R. A half of schoolchildren with 'ISAAC eczema' are ill with allergic contact dermatitis. J Eur Acad Dermatol Venereol 2011;25:1104-7.
- Flohr C, Weiland SK, Weinmayr G, et al. The role of atopic sensitization in flexural eczema: findings from the International Study of Asthma and Allergies in Childhood Phase Two. J Allergy Clin Immunol 2008;121:141-147.
- 17. Dou X, Liu LL, Zhu XJ. Nickel-elicited systemic contact dermatitis. Contact Dermatitis 2003;48:126-129.
- 18. Oiso N, Ota T, Yoshinaga E, et al. Allergic contact dermatitis mimicking atopic dermatitis due to enoxolone in a topical medicament. Contact Dermatitis 2006;54:351.
- 19. Matiz C, Jacob SE. Systemic contact dermatitis in children: how an avoidance diet can make a difference. Pediatr Dermatol 2011;28:368-374.
- 20. innicki M, Shear NH. A systematic approach to systemic contact dermatitis and symmetric drug-related intertriginous and flexural exanthema (SDRIFE): a closer look at these conditions and an approach to intertriginous eruptions. Am J Clin Dermatol 2011;12:171-180.
- 21. Tan SC, Tan JW. Symmetrical drug-related intertriginous and flexural exanthema. Curr Opin Allergy Clin Immunol 2011;11:313-318.
- 22. Shanon J. Pseudo-atopic dermatitis. Contact dermatitis due to chrome sensitivity simulating atopic dermatitis. Dermatologica 1965;131:176-190.
- 23. Herro EM, Jacob SE. Systemic contact dermatitis kids and ketchup. Pediatr Dermatol 2012 (Epub ahead of print, DOI: 10.1111/j.1525-1470.2011.01702.x).
- 24. Akdis M. The cellular orchestra in skin allergy; are differences to lung and nose relevant? Curr Opin Allergy Clin Immunol 2010;10:443-451.
- 25. Mittermann I, Aichberger KJ, Bunder R, et al. Autoimmunity and atopic dermatitis. Curr Opin Allergy Clin Immunol 2004;4:367-371.
- 26. Niebuhr M, Werfel T. Innate immunity, allergy and atopic dermatitis. Curr Opin Allergy Clin Immunol 2010;10:463-468.
- 27. Heratizadeh A, Wichmann K, Werfel T. Food allergy and atopic dermatitis: how are they connected? Curr Allergy Asthma Rep 2011;11:284-291.
- 28. Worth A, Sheikh A. Food allergy and atopic eczema. Curr Opin Allergy Clin Immunol 2010;10:226-230.
- 29. Spiewak R. Food-provoked eczema: A hypothesis on the possible role of systemic contact allergy to haptens present in both cosmetics and foods. Estetol Med Kosmetol 2011;1:35-40.
- 30. Hanifin JM. Atopic dermatitis nomenclature variants can impede harmonization. J Invest Dermatol 2012;132:472-473.
- 31. Rajka G. Atopic dermatitis. London: W. B. Saunders; 1975: 2.
- 32. Flohr C, Johansson SG, Wahlgren CF, Williams H. How atopic is atopic dermatitis? J Allergy Clin Immunol 2004;114:150-158.

- Weber AS, Haidinger G. The prevalence of atopic dermatitis in children is influenced by their parents' education: results of two cross-sectional studies conducted in Upper Austria. Pediatr Allergy Immunol 2010;21:1028-35.
- Toledo MF, Rozov T, Leone C. Prevalence of asthma and allergies in 13- to 14-year-old adolescents and the frequency of risk factors in carriers of current asthma in Taubaté, Sao Paulo, Brazil. Allergol Immunopathol (Madr) 2011;39:284-90.
- 35. Li F, Zhou Y, Li S, et al. Prevalence and risk factors of childhood allergic diseases in eight metropolitan cities in China: a multicenter study. BMC Public Health 2011;11(1):437.
- Zhao J, Bai J, Shen K, et al. Self-reported prevalence of childhood allergic diseases in three cities of China: a multicenter study. BMC Public Health 2010;13;10:551.
- Munivrana H, Vorko-Jovic A, Munivrana S, et al. The prevalence of allergic diseases among Croatian school children according to the ISAAC Phase One questionnaire. Med Sci Monit 2007;13:CR505-509.
- Maziak W, Behrens T, Brasky TM, et al. Are asthma and allergies in children and adolescents increasing? Results from ISAAC phase I and phase III surveys in Munster, Germany. Allergy 2003:58:572-579.
- 39. Hogewoning AA, Bouwes Bavinck JN, Amoah AS, et al. Point and period prevalences of eczema in rural and urban schoolchildren in Ghana, Gabon and Rwanda. J Eur Acad Dermatol Venereol 2012;26:488-94.
- Rahimi Rad MH, Hejazi ME, Behrouzian R. Asthma and other allergic diseases in 13-14-year-old schoolchildren in Urmia, Iran. East Mediterr Health J 2007;13:1005-16.
- Peroni DG, Piacentini GL, Bodini A, et al. Prevalence and risk factors for atopic dermatitis in preschool children. Br J Dermatol 2008;158:539-43.
- 42. Suh M, Kim HH, Sohn MH, et al. Prevalence of allergic diseases among Korean school-age children: a nationwide cross-sectional questionnaire study. J Korean Med Sci 2011;26:332-8.
- Montefort S, Ellul P, Montefort M, et al. A decrease in the prevalence and improved control of allergic conditions in 13- to 15-yr-old Maltese children (ISAAC). Pediatr Allergy Immunol 2011;22:e107-11.
- 44. Barraza-Villarreal A, Hernandez-Cadena L, Moreno-Macias H, et al. Trends in the prevalence of asthma and other allergic diseases in schoolchildren from Cuernavaca, Mexico. Allergy Asthma Proc 2007;28:368-74.
- Zivkovic Z, Vukasinovic Z, Cerovic S, et al. Prevalence of childhood asthma and allergies in Serbia and Montenegro. World J Pediatr 2010;6:331-6.
- Czarnobilska E, Obtulowicz K, Dyga W, et al. Contact hypersensitivity and allergic contact dermatitis among school children and teenagers with eczema. Contact Dermatitis 2009;60:264-9.
- Suarez-Varela MM, Alvarez LG, Kogan MD, et al. Diet and prevalence of atopic eczema in 6 to 7-year-old schoolchildren in Spain: ISAAC phase III. J Investig Allergol Clin Immunol 2010;20:469-75.
- 48. Batlles Garrido J, Torres-Borrego J, Bonillo Perales A, et al. Prevalence and factors linked to atopic eczema in 10- and 11year-old schoolchildren. Isaac 2 in Almeria, Spain. Allergol Immunopathol (Madr) 2010;38:174-80.
- 49. Larsson M, Hägerhed-Engman L, Sigsgaard T, et al. Incidence rates of asthma, rhinitis and eczema symptoms and influential factors in young children in Sweden. Acta Paediatr 2008;97:1210-5.
- Shamssain MH, Shamsian N. Prevalence and severity of asthma, rhinitis, and atopic eczema: the North East Study. Arch Dis Child 1999;81:313-7.

- 51. Mortz CG, Lauritsen IM, Bindlsev-Jensen C, Andersen KH. Prevalence of atopic dermatitis, asthma, allergic rhinitis, and hand and contact dermatitis in adolescents. The Odense Adolescence Cohort Study on Atopic Diseases and Dermatitis. Br J Dermatol 2001;144;523-32.
- 52. Schultz Larsen F, Diepgen T, Svensson A. The occurrence of atopic dermatitis in north Europe: an international questionnaire study. J Am Acad Dermatol 1996;34:760-4.
- 53. Schafer T, Vieluf D, Behrendt H, et al. Atopic eczema and other manifestations of atopy: results of a study in East and West Germany. Allergy 1996;51(8):532-9.
- Bockelbrink A, Heinrich J, Schäfer I, et al.; LISA Study Group. Atopic eczema in children: another harmful sequel of divorce. Allergy 2006;61:1397-402.
- 55. von Mutius E, Fritzsch C, Weiland SK, et al. Prevalence of asthma and allergic disorders among children in united Germany: a descriptive comparison. BMJ 1992;305(6866):1395-9.
- 56. Tamer E, Ilhan MN, Polat M, et al. Prevalence of skin diseases among pediatric patients in Turkey. J Dermatol 2008;35:413-8.
- 57. Emerson RM, Williams HC, Allen BR. Severity distribution of atopic dermatitis in the community and its relationship to secondary referral. Br J Dermatol 1998;139:73-6.
- 58. Laughter D, Istvan JA, Tofte SJ, Hanifin JM. The prevalence of atopic dermatitis in Oregon schoolchildren. J Am Acad Dermatol 2000;43:649-55.
- Plunkett A, Merlin K, Gill D, et al. The frequency of common nonmalignant skin conditions in adults in central Victoria, Australia. Int J Dermatol 1999;38:901-8.
- 60. Thyssen JP, Linneberg A, Menné T, et al. The effect of tobacco smoking and alcohol consumption on the prevalence of selfreported hand eczema: a cross-sectional population-based study. Br J Dermatol 2010;162:619-26.
- 61. Worm M, Forschner K, Lee HH, et al. Frequency of atopic dermatitis and relevance of food allergy in adults in Germany. Acta Derm Venereol 2006;86:119-22.
- 62. Saeki H, Tsunemi Y, Fujita H, et al. Prevalence of atopic dermatitis determined by clinical examination in Japanese adults. J Dermatol 2006;33:817-9.
- 63. Smith-Sivertsen T, Tchachtchine V, Lund E. Atopy in Norwegian and Russian adults: a population-based study from the common border area. Allergy 2003;58:357-62.
- 64. Bo K, Thoresen M, Dalgard F. Smokers report more psoriasis, but not atopic dermatitis or hand eczema: results from a Norwegian population survey among adults. Dermatology 2008;216:40-5.
- 65. Spiewak R. Occupational Dermatoses in Agriculture: Epidemiology, Etiopathogenesis, Risk Factors. Czelej Publishers, Lublin 2002. (In Polish)
- 66. Spiewak R. Contact eczema. Post Dermatol Alergol 2009;26:375-377.
- 67. Spiewak R. Immunotherapy of allergic contact dermatitis. Immunotherapy 2011;3:979-996.
- 68. Spiewak R. Patch testing for contact allergy and allergic contact dermatitis. Open Allergy J 2008;1:42-51.
- 69. Czarnobilska E, Obtulowicz K, Dyga W, Spiewak R. The most important contact sensitizers in Polish children and adolescents with atopy and chronic recurrent eczema as detected with the extended European Baseline Series. Pediatr Allergy Immunol 2011;22:252-256.
- Czarnobilska E, Dyga W, Krzystyniak D, et al. Influence of environment exposures on the frequency of contact allergies in children and adolescents. Ann Agric Environ Med 2012 23;19:11-6.

- 71. Shenefelt PD. Descriptive epidemiology of contact dermatitis in a university student population. Am J Contact Dermat 1996;7:88-93.
- 72. Akhavan A, Cohen SR. The relationship between atopic dermatitis and contact dermatitis. Clin Dermatol 2003;21:158-162.
- 73. Spiewak R. Pesticides and skin diseases in man. In: Rathore H, Nolet LM, editors. Pesticides: Evaluation of Environmental Pollution. Boca Raton: CRC Press; 2012: 525-542.
- 74. Wüthrich B. Zur Genese des Bäckerekzems. Hautarzt 1970;21:214-218.
- 75. Hjorth N, Roed-Petersen J. Occupational protein contact dermatitis in food handlers. Contact Dermatitis 1976;2:28-42.
- Kanerva L, Jolanki R, Toikkanen J. Frequencies of occupational allergic diseases and gender differences in Finland. Int Arch Occup Environ Health 1994;66:111-6.
- 77. Hansen KS, Petersen HO. Protein contact dermatitis in slaughterhouse workers. Contact Dermatitis 1989;21:221-4.
- 78. Hay RJ. Malassezia, dandruff and seborrhoeic dermatitis: an overview. Br J Dermatol 2011;165 (Suppl 2): 2-8.
- 79. Lally A, Casabonne D, Newton R, Wojnarowska F. Seborrheic dermatitis among Oxford renal transplant recipients. J Eur Acad Dermatol Venereol 2010;24:561-4.
- 80. Moehrle M, Dennenmoser B, Schlagenhauff B, et al. High prevalence of seborrhoeic dermatitis on the face and scalp in mountain guides. Dermatology 2000;201:146-7.
- Wollina U. Pompholyx: a review of clinical features, differential diagnosis, and management. Am J Clin Dermatol 2010;11:305-14.
- 82. de Boer EM, Bruynzeel DP, van Ketel WG. Dyshidrotic eczema as an occupational dermatitis in metal workers. Contact Dermatitis 1988;19:184-8.
- 83. Coenraads PJ. Hand eczema is common and multifactorial. J Invest Dermatol 2007;127:1568-70.
- 84. Dotterud LK, Falk ES. Contact allergy in relation to hand eczema and atopic diseases in north Norwegian schoolchildren. Acta Paediatr 1995;84:402-6.
- 85. Dorynska A, Pasich J, Lach L, et al. Self-reported hand, wrist and forearm dermatitis in a random sample of 691 Polish medical professionals, firefighters and students. Contact Dermatitis 2010;63(Suppl 1):96-7.
- 86. Meding B. Epidemiology of hand eczema in an industrial city. Acta Derm Venereol Suppl (Stockh) 1990;153:1-43.
- Meding B, Järvholm B. Hand eczema in Swedish adults

 changes in prevalence between 1983 and 1996. J Invest Dermatol 2002;118:719-23.
- Montnemery P, Nihlen U, Lofdahl CG, et al. Prevalence of hand eczema in an adult Swedish population and the relationship to risk occupation and smoking. Acta Derm Venereol 2005;85:429-32.
- 89. Belsito DV. Occupational contact dermatitis: etiology, prevalence, and resultant impairment/disability. J Am Acad Dermatol 2005;53:303-13.
- 90. Meyer JD, Chen Y, Holt DL, et al. Occupational contact dermatitis in the UK: a surveillance report from EPIDERM and OPRA. Occup Med (Lond) 2000;50:265-73.
- 91. Diepgen TL, Coenraads PJ. The epidemiology of occupational contact dermatitis. Int Arch Occup Environ Health 1999;72:496-506.
- 92. Spiewak R. Skin diseases related to farm work a questionnaire survey of 145 Polish farmers in Lublin Region. Post Dermatol Alergol 2001;18:194-199.
- 93. Spiewak, R. Contact dermatitis in atopic individuals. Curr Opin Allergy Clin Immunol 2012 (in press).