Contents lists available at ScienceDirect



International Journal of Women's Dermatology

Original Research

## Art of prevention: The importance of bath time and avoiding extended exposure to irritating and allergenic chemicals $\stackrel{k}{\approx}$



WDS

International Iournal of

Women's Dermatology

Tracy Novosel, MD<sup>a</sup>, Chandler W. Rundle, BS<sup>b</sup>, Jiade D. Yu, MD<sup>c</sup>, Sharon E. Jacob, MD<sup>d,\*</sup>

<sup>a</sup> Department of Dermatology, Atlantic Dermatology Associates, Virginia Beach, Virginia

<sup>b</sup> School of Medicine, Loma Linda University, Loma Linda, California

<sup>c</sup> Department of Dermatology, Massachusetts General Hospital, Boston, Massachusetts

<sup>d</sup> Department of Dermatology, Loma Linda University, Loma Linda, California

#### ARTICLE INFO

Article history: Received 7 October 2018 Received in revised form 20 February 2019 Accepted 21 February 2019

*Keywords:* Irritant contact dermatitis allergic contact dermatitis skin pH skin barrier

Published by Elsevier Inc. on behalf of Women's Dermatologic Society. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

Contact dermatitis (CD) is an umbrella term for a group of inflammatory skin conditions that result from contact between a chemical substance and the skin. Irritant CD (ICD) is by far the most prevalent and can occur in anyone (no prior sensitization is necessary). ICD can be thought of as the effect observed when a sensitive area of skin is exposed to too strong a chemical, such as detergents. Allergic CD (ACD), on the other hand, represents the clinical disease state that results from exposure to an allergenic chemical in a person who is already sensitized to that chemical, such as that in poison ivy. ACD is a delayed-type hypersensitivity reaction that is diagnosed through the application of a closed (under occlusion) patch test, in which the suspected allergen is placed on intact skin (on the back or inner arm) for 24 to 48 hours (Lazzarini et al., 2013). These patches are evaluated at removal and again at a later period (72-120 hours; Lazzarini et al., 2013).

Children represent a unique group in terms of ACD, despite the condition being quite prevalent in the pediatric population (20%-25% of all childhood dermatitis; Bruckner and Weston, 2002; Militello et al., 2006; Seidenari et al., 2005). Notably, patch testing in children with recalcitrant dermatitis is often de-

Corresponding Author. *E-mail address:* sjacob@contactderm.net. (S.E. Jacob). layed (Jacob et al., 2008; McGowan et al., 2018). This is unfortunate given that there is both an increase in the reported number of children with ACD and the need for patch testing in children with dermatitis (Jacob et al., 2008). Recent studies demonstrate that allergens in personal hygiene products contribute significantly to pediatric CD (Berne et al., 1996; Goon and Goh, 2006; Jacob et al., 2005; Pratt et al., 2004)—hence the role of the Pediatric Contact Dermatitis Registry, which is a collaborative, multidisciplinary registry consisting of >250 health care providers that provide data from >1000 patch-tested children in the United States (Jacob et al., 2017).

Bathing practice–associated exposures are important sources of these allergens (Table 1). In addition, several allergens can be in one product and can range from preservatives to fragrances to emulsifiers and detergents (Smaoui and Hlima, 2012; Timmermans et al., 2007). It is also important to recognize that these chemicals may serve as a source of ICD (especially in children with eczema who demonstrate lower thresholds for irritation; Fernandez Vozmediano and Armario Hita, 2005; Lammintausta et al., 1992; Lugovic and Lipozencic, 1997; Oranje and Wolkerstorfer, 1999) and may predispose children to ACD (Marty and Cheng, 2005). Therefore, it stands to reason that if the skin has already been damaged by endogenous dermatosis (e.g., atopic dermatitis) or external trauma, skin penetration by allergens may be enhanced. Skin barrier function is often genetically predetermined at birth; however, even healthy skin is more

https://doi.org/10.1016/j.ijwd.2019.02.004

2352-6475/Published by Elsevier Inc. on behalf of Women's Dermatologic Society. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

<sup>☆</sup> Conflicts of interest: The authors have no conflicts of interest to report.

# Table 1Children producttypes used inbathing routines

Baby wipes Lotion Shampoo Baby powder Diaper cream Conditioner Bubble bath Soap-cleansers

delicate in children (compared with adults) and consequently more prone to irritant and allergic contact dermatitis (Gelmetti, 2001).

The development of CD (both irritant and allergic types) requires contact with the eliciting chemicals; thus, decreasing the time of exposure may have a clinically relevant impact (Beattie et al., 2007). Furthermore, ICD, but not the elicitation of ACD, depends on the time of exposure. ICD is dependent on the concentration and duration of exposure. Allergic sensitization is a rate-limited phenomenon of a breached threshold in relation to hapten concentration per unit area of the skin and the immune state of the skin. In addition, inflammation is thought to predispose the epidermal layer to penetration by certain allergens. Thus, decreasing the time of exposure to surfactants may decrease ICD and only indirectly ACD.

#### **Practical intervention**

We made a concerted effort to change our approach and purposefully recommended that children be allowed to have bath time first and then be washed at the end of a bath, followed by immediate rinsing to minimize exposure duration to potentially irritating and allergenic chemicals in bathing and personal hygiene products. Hot water should be avoided because excess heat can induce proprioceptors with subsequent itching. With an acidified bath with moderately warm water, a child can play up to 15 minutes. The aforementioned are hypotheses based on observations from practice.

Water can be an irritant, and parents should be sure to acidify the water to a pH of 4.5 with vinegar to reduce skin pH and improve barrier function. Three to four cups of apple cider vinegar added to a half bath (approximately 20 gallons) is sufficient to achieve these results. Alternatively, pool salts (i.e., sodium chloride formulary designed to keep pools clean) can be added as a soothing agent (2 cups per 20-gallon tub) because salts are recognized to increase tolerability, reduce skin inflammation, and decrease transepidermal water loss (Bak et al., 2012; Yoshizawa et al., 2003). Notably, pool salts added to bath water would not have the same acidifying effect as salts in a pool, largely because the electrolysis component in a pool is absent from baths. Double-blind, placebo-controlled trials for bath additives would be helpful to further support this hypothesis.

The cutaneous acid mantle is a complex and highly regulated environment that provides antimicrobial activity, epidermal structural integrity, and barrier function (Rundle et al., 2017). This milieu is maintained by several mechanisms; however, the role of the acid mantle is most highlighted when it is aberrant. Dysregulation of the acid mantle results in the activation of serine proteases, which allows for the breakdown of essential barrier proteins and enzymes, abnormal lipid organization/metabolism, and irregular desquamation of the stratum corneum. Increased desquamation further compromises the epidermis, thus perpetuating decreased barrier integrity and protective function (Rundle et al., 2017).

Preservation of the skin barrier is the most effective way to prevent the elicitation of contact dermatitis. Avoidance of irritating compounds is the first step to restore the skin barrier. Optimization of the skin pH (between 4.6 and 5.6) allows for improved activity of the enzymes (e.g., B-glucocerebrosidase and acid sphingomyelinases) responsible for ceramide and lipid production, a necessary component for the maintenance of a healthy biome (Eberting, 2014). Furthermore, studies have demonstrated that hyperacidification of the epidermal layer improves stratum corneum desquamation and barrier homeostasis while simultaneously preventing skin irritation (Berardesca et al., 1997).

In our society, bathing children and then allowing them to play in the bath water is common practice. By instituting this change in the timing of bathing and play time practice, we observed a significant improvement in clinical manageability of patients with atopic dermatitis. However, the question of whether there was enough contact with sensitizing chemicals during bathing to have a clinical impact remains. We believe that, for exquisitely sensitized individuals, even prolonged contact with very small amounts of allergens can trigger a response (Jacob and Steele, 2007; Larsen, 1989; Ringborg et al., 2016), especially in neonates and infants (Bruckner et al., 2000; Fisher, 1994a, 1994b). Notably, these interventions are based on pathophysiology rather than clinical trials.

#### Conclusions

Our observation remains largely anecdotal, but we also realize that a sensitization induction study in bathing practices on children is unlikely to be performed. Furthermore, additional studies on acidification of the skin in children and its effect on the barrier are necessary. Therefore, based on our experience and the fundamental principles behind the development of both irritant and allergic CD, we recommend changing the general bathing practices of children so that we may ultimately decrease contact sensitization rates in this underserved population.

### Acknowledgments

The authors thank the American Contact Dermatitis Society for making this study possible through their Mentoring Award.

#### References

- Bak JP, Kim YM, Son J, Kim CJ, Kim EH. Application of concentrated deep sea water inhibits the development of atopic dermatitis-like skin lesions in NC/Nga mice. BMC Complement Altern Med 2012;12:108.
- Beattie PE, Green C, Lowe G, Lewis-Jones MS. Which children should we patch test? Clin Exp Dermatol 2007;32(1):6–11.
- Berardesca E, Distante F, Vignoli GP, Oresajo C, Green B. Alpha hydroxyacids modulate stratum corneum barrier function. Br J Dermatol 1997;137(6):934–8.
- Berne B, Bostrom A, Grahnen AF, Tammela M. Adverse effects of cosmetics and toiletries reported to the Swedish Medical Products Agency 1989-1994. Contact Dermatitis 1996;34(5):359–62.
- Bruckner AL, Weston WL. Allergic contact dermatitis in children: A practical approach to management. Skin Ther Lett 2002;7(8):3–5.
- Bruckner AL, Weston WL, Morelli JG. Does sensitization to contact allergens begin in infancy? Pediatrics 2000;105(1):e3.
- Eberting CL. Irritant contact dermatitis: Mechanisms to repair. J Clin Exp Dermatol Res 2014;5(6):317–28.
- Fernandez Vozmediano JM, Armario Hita JC. Allergic contact dermatitis in children. J Eur Acad Dermatol Venereol 2005;19(1):42–6.
- Fisher A. Allergic contact dermatitis in early infancy. Cutis 1994;54:300–2.
- Fisher A. Patch testing in children including early infancy. Cutis 1994;54:387-8.

Gelmetti C. Skin cleansing in children. J Eur Acad Dermatol Venereol 2001;15(Suppl. 1):12–5.

- Goon ATJ, Goh CL. Patch testing of Singapore children and adolescents: Our experience over 18 years. Pediatr Dermatol 2006;23(2):117–20.
- Jacob SF, Steele T. Avoiding formaldehyde allergic reactions in children. Pediatr Ann 2007;36(1):55–6.
- Jacob S, Steele T, Rodriguez G. Focus on T.R.U.E. Test Allergens #21, 13 and 18: Formaldehyde and formaldehyde-releasing preservatives. Skin. Aging 2005:22–7.
- Jacob SE, Burk CJ, Connelly EA. Patch testing: Another steroid-sparing agent to consider in children. Pediatr Dermatol 2008;25(1):81–7.
- Jacob SE, McGowan M, Silverberg NB, et al. Pediatric contact dermatitis registry data on contact allergy in children with atopic dermatitis. JAMA Dermatol 2017;153 (8):765–70.

Lammintausta K, Kalimo K, Fagerlund VL. Patch test reactions in atopic patients. Contact Dermatitis 1992;26(4):234–40.

Larsen WG. How to instruct patients sensitive to fragrances. J Am Acad Dermatol 1989; 21(4):880-4.

Lazzarini R, Duarte I, Ferreira AL. Patch tests. An Bras Dermatol 2013;88(6):879–88. Lugovic L, Lipozencic J. Contact hypersensitivity in atopic dermatitis. Arh Hig Rada Toksikol 1997;48(3):287–96.

Marty CL, Cheng JF. Irritant contact dermatitis precipitating allergic contact dermatitis. Dermat Contact Atopic Occup Drug 2005;16(2):87–8 quiz 55–6.

McGowan MA, Goldenberg A, Jacob SE. Contact dermatitis in underrepresented minority. Dermat Contact Atopic Occup Drug 2018;29(2):97–9.

Militello G, Jacob SE, Crawford GH. Allergic contact dermatitis in children. Curr Opin Pediatr 2006;18(4):385–90.

Oranje AP, Wolkerstorfer A. Advances in the treatment of atopic dermatitis with special regard to children. Curr Probl Dermatol 1999;28:56–63. Pratt MD, Belsito DV, DeLeo VA, et al. North American Contact Dermatitis Group patch-test results, 2001-2002 study period. Dermat Contact Atopic Occup Drug 2004;15(4):176–83.

Ringborg E, Lidén C, Julander A. Nickel on the market: A baseline survey of articles in prolonged contact with skin. Contact Dermatitis 2016;75(2):77–81.

Rundle CW, Bergman D, Goldenberg A, Jacob SE. Contact dermatitis considerations in atopic dermatitis. Clin Dermatol 2017;35(4):367–74.

Seidenari S, Giusti F, Pepe P, Mantovani L. Contact sensitization in 1094 children undergoing patch testing over a 7-year period. Pediatr Dermatol 2005;22(1):1–5.

Smaoui S, Hlima HB. Effects of parabens and isothiazolinone on the microbiological quality of baby shampoo: The challenge test. Biocontrol Sci 2012;17(3):135–42. Timmermans A, De Hertog S, Gladys K, Vanacker H, Goossens A. Dermatologically

tested baby toilet tissues: A cause of allergic contact dermatitis in adults. Contact Dermatitis 2007;57(2):97–9.

Yoshizawa Y, Kitamura K, Kawana S, Maibach HI. Water, salts and skin barrier of normal skin. Int Soc Skin Imaging 2003;9(1):31–3.