## **CLINICAL REPORT**



# 1,2-Ethanedithiol-induced Erythema Multiforme-like Contact Dermatitis

Jeng-Wei TJIU, Chia-Yu CHU and Chee-Ching SUN

Department of Dermatology, National Taiwan University Hospital and National Taiwan University College of Medicine, Taipei 100, Taiwan

Contact dermatitis simulating erythema multiforme can be caused by many allergens. The chemical agent 1,2ethanedithiol, which serves as a protective group in chemical synthesis, has hitherto only been implicated as an irritant. We report on a 22-year-old female chemistry student who developed widespread erythema multiformelike lesions after local contact with 1,2-ethanedithiol. Many target lesions were observed bilaterally on her hands, forearms, arms, and on her forehead. One such lesion was histologically compatible with erythema multiforme. The patient had a positive patch test to 1,2-ethanedithiol, whereas none of 30 healthy subjects showed a positive reaction. However, eight of the 30 controls (26.7%) developed irritant reactions to 1,2ethanedithiol. Cautious handling of the compound is a prudent precaution. Key words: allergic contact dermatitis; erythema multiforme; 1,2-ethanedithiol.

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Chia-Yu Chu, Department of Dermatology, National Taiwan University Hospital, 7, Chung-Shan South Road, Taipei, Taiwan. E-mail: chiayuju@seed.net.tw

Erythema multiforme (EM)-like contact dermatitis is a non-eczematous allergic contact dermatitis (1), which is histopathologically indistinguishable from EM (2). The EM-like contact dermatitis can be caused by potent contact allergens including topical medicaments, chemicals and plants (1). True EM, on the other hand, is usually a consequence of infection with herpes simplex virus (HSV) (3).

Compounds that are chemically very reactive and widely used represent candidate contact allergens. One such compound is 1,2-ethanedithiol (CAS540-63-6; HS-CH<sub>2</sub>-CH<sub>2</sub>-SH). The chemical agent is widely used as a protective group in many chemical experiments. However, to the best of our knowledge, EM-like contact dermatitis caused by 1,2-ethanedithiol has not been hitherto reported in the literature. We report a case of systemic EM-like contact dermatitis induced by 1,2-ethanedithiol. To elucidate the potential of skin irritancy or contact sensitivity after contact with 1,2-ethanedithiol, a prospective patch testing study was undertaken on healthy volunteers.

## CASE REPORT

### History

The patient was a 22-year-old Taiwanese female, who was a university graduate student in chemistry. She had no past medical history of mycoplasma or HSV infection. In the year prior to her admittance to our hospital, she had periodic and minimal contact with 1,2-ethanedithiol during the course of laboratory experiments. In July 2001, an accident resulted in the spillage of a large amount of the compound over her right forearm. Within 24 h, many variously sized, erythematous, oedematous, target lesions appeared on her right forearm (Fig. 1). This was followed by the bilateral spread of EM-like lesions to her hands, forearms, and arms, as well as to her forehead on the third day following the spill. By the fifth day after the spill, the skin lesions had worsened and the patient had developed fever, malaise, and dizziness. Laboratory examinations revealed an elevated white blood cell count  $(10.6 \times 10^{9}/l)$ , as compared to the normal range of  $4.0 - 10.0 \times 10^{9}$ /l).

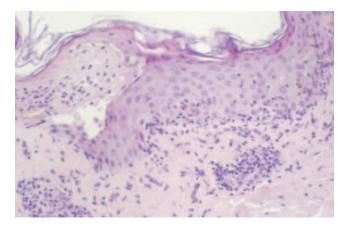
Upon a diagnosis of EM-like contact dermatitis, the patient was hospitalized and treated intravenously with methylprednisolone (160 mg/day) for three consecutive days. Prednisolone was then administered orally until the skin lesions had cleared.

## Histopathology

A skin biopsy specimen was taken from one of the EMlike lesions. Microscopic examination of the biopsy sections showed frequent dyskeratosis, focal epidermal



*Fig. 1.* Several circular, wheal-like erythematous plaques and target-like lesions with confluence located on the right forearm.



*Fig.* 2. Histopathological examinations showed frequent dyskeratosis, focal epidermal keratinocyte necrosis, basal vacuolization, and cleft formation in the dermal-epidermal junction (haematoxylin & eosin, original magnification  $\times$  200).

keratinocyte necrosis, basal vacuolization, and cleft formation in the dermal-epidermal junction (Fig. 2). These microscopic features were compatible with EM.

#### Patch testing

Patch testing was performed according to international standards (4). Briefly, test substances were applied to the upper back with Finn chambers, where they remained in contact with the skin for 48 h. Observations were made at 72 and 120 h.

Reactions were scored according to the scale recommended by the International Contact Dermatitis Research Group (ICDRG). A positive result was defined as an erythema with infiltration (1+), erythema with infiltration and papules (2+), or erythema with infiltration, papules and vesicles (3+) (4). The patient was patch-tested with the European standard series and 1,2-ethanedithiol concentrations of 1% and 5% prepared in petrolatum.

After 72 h the patient produced negative results to the European standard series but had positive reactions (1+) to both concentrations of 1,2-ethanedithiol. After 120 h, the 5% concentration of 1,2-ethanedithiol produced a 3+ positive reaction (Table I).

#### CONTROL STUDY

Thirty healthy volunteers (12 men and 18 women, aged 22-64 years, mean age  $27.7\pm8.2$  years) were enrolled

Table I. Results of patch testing in the patient

	Day 3	Day 5
European standard series	Negative	Negative
1,2-Ethanedithiol (1% pet.)	1+	1+
1,2-Ethanedithiol (5% pet.)	1+	3+

in the study. None of them had received systemic medications or topical steroids, nor had they a previous contact history of exposure to 1,2-ethanedithiol. The study was approved by the ethical committee of the National Taiwan University Hospital and informed consent was obtained from each participant.

Patch testing was performed as summarized above. All of the control subjects were patch-tested with 1% and 5% 1,2-ethanedithiol in petrolatum. The effect of contact exposure was assessed 72 h after application of the patch.

All of the control subjects registered a negative reaction to both concentrations of 1,2-ethanedithiol. However, the 5% concentration produced an irritant reaction in 8 of the 30 control subjects (26.7%) within 5-10 min after applying the Finn chambers to the back.

#### DISCUSSION

As summarized in Table II, many allergens have been reported to cause EM-like contact dermatitis (1, 5-26). To the best of our knowledge, until the present report, 1,2-ethanedithiol has not been implicated in allergic contact dermatitis.

Clinically, EM-like contact dermatitis manifests as erythematous, oedematous plaques with target lesions (3). The skin lesions can be localized in the contact area or can occur more generally on the trunk and extremities (3). Pathologically, EM-like contact dermatitis shows the same features as EM (3).

EM is currently considered to represent a cellmediated immune reaction usually targeting keratinocytes that express HSV antigens (27). On the other hand, EM-like contact dermatitis appears to be principally elicited by type IV hypersensitivity (3), although widespread, cutaneous lesions or systemic symptoms, such as were evident in the present case may instead result from circulating immune complex (type III hypersensitivity) (15, 23). The positive patch testing result in our patient suggests that type IV hypersensitivity is involved in the pathogenesis of EM-like contact dermatitis.

In addition to the cutaneous manifestations, the patient had systemic toxic signs including fever, malaise, and dizziness. It has been reported that one patient died of EM-like contact dermatitis and toxic epidermal necrolysis after exposure to spray cologne (28). Thus, toxic complications and severe systemic complications can occur in EM-like contact dermatitis, and may be life-threatening.

1,2-Ethanedithiol is a chemical agent used as a protecting group in chemical synthesis (29). It is readily volatile and vaporizes easily. The fetid odor associated with its presence can be detected in the air even at a very low concentration (1 ppm) (30). The compound can cause eye and skin irritation. Indeed, in the present control study more than one-quarter of the healthy

Causative agent (Ref.)	Age/sex	Involved site	Form/phase of the contactants
Topical medicament			
Bufexamac (5)	34/M	Local, then systemic	Ointment
Budesonide (6)	19/F	Hand, face, forearms	Ointment
Bufexamac (7)	52/M	Face, trunk	Ointment
Povidone-iodine (8)	30/M	Limbs, trunk	Topical
Phenylbutazone (9)	65/F	Thorax, right axilla	Ointment
Proflavine (10)	24/M	Legs and knee	Powder
Lincomycin (11)	37/F	Ear, face, neck, trunk	Ointment
Mephenesin (12)	30/M	Arms, legs	Ointment
Chemicals			
Ethyl ethoxymethylene cyanoacetate (13)	36/M	Face, extremity	Crystal
Beta-cyclocostunolide (14)	26/M	Upper limbs	Oil
Rubber gloves (1)	48/F	Bilateral forearms	Solid
Natural rubber latex (15)	26/F	Thigh, face	Solid
Oxybenzone (16)	44/F	Forearm, leg	Sunscreen
Dimethoate (17)	41/F	Trunk, back, hand	Insecticide
Plants			
Rhus (lacquer) (18)	43.8 (mean)	Body	Ingestion
	M18, F13		
Rhus (lacquer) (19)	54/M	Hand, arm, trunk	Contact
Sesquiterpene (20)	31/F	Presternal, trunk	Solid
Rosewood (21)	36/M	Neck, thigh, penis	Solid
Primula (22)	30/F	Back, hands, forearms	Plants
Others			
Hair dyes (23)	70/F	Hands, forearms, lip, thigh	Liquid
Capsicum (24)	65/F	Right knee, body	Tincture
Epoxy sealant (25)	46/F	Upper extremity, arm, hand	Paints
Cutting oil (26)	48/M	Forearm, hand, trunk	Liquid

Table II. Review of EM-like contact dermatitis with a positive patch test

volunteers displayed an irritant reaction, including erythema, itching, and a burning sensation almost immediately after the patch application of the compound.

Given this irritant potency, 1,2-ethanedithiol should be used in a ventilated fume hood and handled with protective gloves, goggles and clothing (31). Finally, as examplified by the present case, because 1,2ethanedithiol can be inhaled or transcutaneously absorbed, the possibility of systemic contact dermatitis cannot be excluded. Cautious handling of 1,2-ethanedithiol is recommended.

#### REFERENCES

- 1. Lu CY, Sun CC. Localized erythema-multiforme-like contact dermatitis from rubber gloves. Contact Dermatitis 2001; 45: 311–312.
- 2. Puig L, Fernandez-Figueras MT, Montero MA, Ferrandiz C, Alomar A. Erythema-multiforme-like eruption due to topical contactants: expression of adhesion molecules and their ligands and characterization of the infiltrate. Contact Dermatitis 1995; 33: 329–332.
- 3. Huff JC. Erythema multiforme and latent herpes simplex infection. Semin Dermatol 1992; 11: 207–210.
- Rietschel RL, Fowler JF Jr. Practical aspects of patch testing. In: Rietschel RL, Fowler JF Jr, eds. Fisher's contact dermatitis, 5th edn. Philadelphia, PA: Lippincott Williams & Wilkins, 2001: 9–26.
- 5. Koch P, Bahmer FA. Erythema-multiforme-like, urticarial papular and plaque eruptions from bufexamac: report of 4 cases. Contact Dermatitis 1994; 31: 97–101.

- Stingeni L, Caraffini S, Assalve D, Lapomarda V, Lisi P. Erythema-multiforme-like contact dermatitis from budesonide. Contact Dermatitis 1996; 34: 154–155.
- 7. Kurumaji Y. Photo koebner phenomen in erythemamultiforme-like eruption induced by contact dermatitis due to bufexamac. Dermatology 1998; 197: 183–186.
- Torinuki W. Generalized erythema-multiforme-like eruption following allergic contact dermatitis. Contact Dermatitits 1990; 23: 202–203.
- 9. Kerre S, Busschots A, Dooms-Goossens A. Erythemamultiforme-like contact dermatitis due to phenylbutazone. Contact Dermatitis 1995; 33: 213–214.
- Goh CL. Erythema multiforme-like and purpuric eruption due to contact allergy to proflavine. Contact Dermatitis 1987; 17: 53-54.
- Conde-Salazar L, Guimaraens D, Romero L, Gonzalez M, Yus S. Erythema multiforme-like contact dermatitis from lincomycin. Contact Dermatitis 1985; 12: 59–61.
- Degreef H, Bonamie A, van Derheyden D, Dooms-Goossens A. Mephenesin contact dermatitis with erythema multiforme features. Contact Dermatitis 1984; 10: 220-223.
- Hsu CK, Sun CC, Su MS, Kuo EF, Wu YC. Systemic contact allergy from occupational contact with ethyl ethoxymethylene cyanoacetate. Contact Dermatitis 1992; 27: 58-59.
- 14. Le Coz CJ, Lepoittevin JP. Occupational erythemamultiforme-like dermatitis from sensitization to costus resinoid, followed by flare-up and systemic contact dermatitis from beta-cyclocostunolide in a chemistry student. Contact Dermatitis 2001; 44: 310–311.
- 15. Bourrain JL, Woodward C, Dumas V, Caperan D, Beani JC, Amblard P. Natural rubber latex contact dermatitis

with features of erythema multiforme. Contact Dermatitis 1996; 35: 55–56.

- Zhang XM, Nakagawa M, Kawai K, Kawai K. Erythema-multiforme-like eruption following photoallergic contact dermatitis from oxybenzone. Contact Dermatitis 1998; 38: 43–44.
- Schena D, Barba A. Erythema-multiforme-like contact dermatitis from dimethoate. Contact Dermatitis 1992; 27: 116–117.
- Park SD, Lee SW, Chun JH, Cha SH. Clinical features of 31 patients with systemic contact dermatitis due to the ingestion of Rhus (lacquer). Br J Dermatol 2000; 142: 937-942.
- Schwartz RS, Downham TF 2nd. Erythema multiforme associated with Rhus contact dermatitis. Cutis 1981; 27: 85-86.
- Mateo MP, Velasco M, Miquel FJ, de la Cuadra J. Erythema multiforme-like eruption following allergic contact dermatitis from sesquiterpene lactones in herbal medicine. Contact Dermatitis 1995; 33: 449–450.
- Irvine C, Reynolds A, Finlay AY. Erythema multiformelike reaction to "rosewood". Contact Dermatitis 1988; 19: 224-225.
- 22. Lengrand F, Tellart AS, Segard M, Dejobert Y, Thomas P. Erythema multiforme-like eruption: an unusual presentation of primula contact allergy. Contact Dermatitis 2001; 44: 35.
- 23. Tosti A, Bardazzi F, Valeri F, Toni F. Erythema

multiforme with contact dermatitis to hair dyes. Contact Dermatitis 1987; 17: 321–322.

- Raccagni AA, Bardazzi F, Baldari U, Righini MG. Erythema multiforme-like contact dermatitis due to capsicum. Contact Dermatitis 1995; 33: 353-354.
- Whitfeld MJ, Rivers JK. Erythema multiforme after contact dermatitis in response to an epoxy sealant. J Am Acad Dermatol 1991; 25: 386–388.
- 26. Hata M, Tokura Y, Takigawa M. Erythema multiformelike eruption associated with contact dermatitis to cutting oil. Eur J Dermatol 2001; 11: 247–248.
- 27. Brice SL, Krzemien D, Weston WL, Huff JC. Detection of herpes simplex virus DNA in cutaneous lesions of erythema multiforme. J Invest Dermatol 1989; 93: 183–187.
- Thompson JA Jr, Wansker BA. A case of contact dermatitis, erythema multiforme, and toxic epidermal necrolysis. J Am Acad Dermatol 1981; 5: 666–669.
- 29. Greene TW, Wuts PGM. Protection for the carbonyl group. In: Greene TW, Wuts PGM, eds. Protective groups in organic synthesis, 2nd edn. New York: John Wiley & Sons, 1991: 175–223.
- Kocienski PJ. Carbonyl protecting groups. In: Kocienski PJ, ed. Protecting groups, 1st edn, New York: Thieme, 1994: 155-184.
- 1,2-Ethanedithiol. Material Safety Data Sheets (MSDS) [database online] 2003 [cited 2003 Dec 21]; [11screens]. Available from: URL: http://msds.ehs.cornell.edu/msds/ msdsdod/a432/m215696.htm.