

A Case of Zoster Duplex Bilateralis

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Herpes zoster involving noncontiguous dual dermatomes is very rare in both immunocompetent and immunocompromised persons. This unique presentation has been referred to as zoster duplex unilateralis or bilateralis, depending whether one or both halves of the body are involved.

A 22-year-old woman, who had been treated for acute leukemia, congestive heart failure and chronic disseminated candidiasis, was referred to our department for painful papulovesicular eruptions on the right side of the anterior chest and upper back for 2 days, and the left buttock for 1 day. Tzanck smear revealed multinucleated giant cells with intranuclear inclusion bodies. We report a rare case of zoster duplex bilateralis.

(Ann Dermatol 14(1) 59-61, 2002).

Key Words : Zoster duplex

Herpes zoster is caused by the varicella zoster virus. Following natural infection or immunization the virus remains latent in the sensory dorsal root ganglion cells and begins to replicate at some later time, traveling down the sensory nerve into the skin. The clinical manifestations are characterized by several groups of painful vesicles situated unilaterally within the distribution of the cranial or spinal sensory nerve¹. Bilateral involvement and recurrence are rare, and zoster involving two widely separated regions at one time are even rarer².

The phenomenon of zoster occurring in two noncontiguous, widely separated dermatomes has been referred to as zoster duplex unilateralis or bilateralis, depending whether one half or both halves of the body are involved³. In Korea a case of zoster duplex unilateralis has been reported³ but zoster duplex bilateralis not yet.

We report that a rare case of herpes zoster involving noncontiguous dual dermatomes on the both halves of the body that named zoster duplex bilateralis.

CASE REPORT

A 22-year-old woman was referred to our department from internal medicine because of painful vesicles for 2 days. She had been treated with chemotherapy for acute leukemia for 2 years and under complete remission state. Also she had been treated with digoxin and fluconazole for congestive heart failure and chronic disseminated candidiasis. Two days ago, several grouped erythematous papulovesicular eruptions developed with intermittent prickling pain on the right side of the anterior chest and upper back showing a band-like arrangement(right L7 dermatome, Fig. 1, 2), and the following day similar lesions with pain were also found on the left buttock(left S3 dermatome, Fig. 3).

Laboratory examinations including complete blood cell counts, routine urinalysis, liver function test, a test for VDRL and Chest PA were negative or within normal limits. Bone marrow examination showed no

Received June 4, 2001.

Accepted for publication September 15, 2001.

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blast. Hepatobiliary ultrasonography and echocardiography revealed multiple hypoechoic nodular lesions in liver and spleen, which were consistent with chronic disseminated candidiasis, and moderate left ventricular dysfunction.

Tzanck smear was done at the vesicles of left buttock (Fig. 4) and microscopic exam showed multinucleated giant cells with intranuclear inclusion bodies.

She was treated with oral administration of famcyclovir 750 mg/day for 7 days, and skin lesions and pain subsided without complications.

Herpes zoster is a relatively common disease and characterized by several groups of painful vesicles with characteristic distribution of unilateral dermatomes¹. The incidence rate is slightly different in Korean literature: Kim et al⁴ reported 0.79% in dermatologic outpatients; Kim et al⁵, 0.38%; Chun et al⁶, 1.7%; Yang et al⁷, 0.64%; Hong et al⁸, 2.84%; Kim et al⁹, 1.23%. Most of the skin lesions are in unilateral involvement



Fig. 1. The right side of the chest showed painful grouped vesicles on the erythematous base.



Fig. 2. Painful grouped erythematous papulovesicular eruptions on the right side of the upper back had a band-like arrangement.

DISCUSSION



Fig. 3. Painful grouped papules and vesicles on the erythematous base developed on the left buttock.

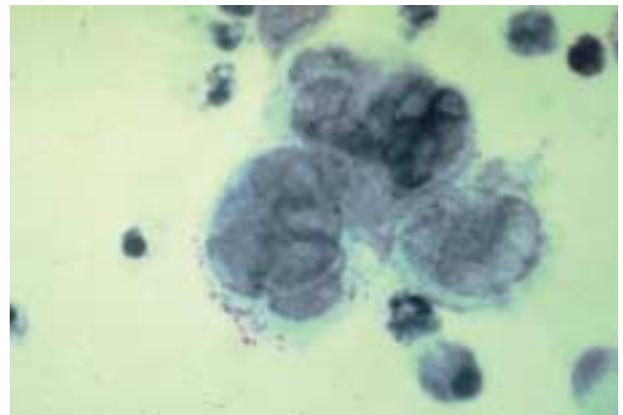


Fig. 4. Tzanck smear from the left buttock revealed multinucleated giant cells with intranuclear inclusion bodies. (Wright stain; $\times 400$)

characteristically and the dermatomes most frequently affected are thoracic. Bilateral involvement is rare and the incidence is below 0.5%¹⁰. Moreover, zoster occurring in noncontiguous dual dermatomes is very rare. On the review of the medical literature of the last 3 decades, there have only been 7 cases reported, two in an immunocompetent person^{3,11}, three in older persons on oral steroids for chronic illnesses^{12,13,14}, and two in children with cancer (1 with lymphoma and 1 with leukemia)^{14,15}. Varicella-zoster virus involving noncontiguous dermatomes is distinct from disseminated varicella-zoster virus infection that occurs in immunosuppressed renal transplant patients. The phenomenon of zoster occurring in two noncontiguous dermatomes has been referred to as zoster duplex unilateralis or bilateralis, depending whether one half or both halves of the body are involved³. Vu et al¹⁵ reported a case of herpes zoster in seven disparate dermatomes and suggested the term zoster multiplex when more than three noncontiguous dermatomes are involved.

In Korean literature, Lee et al² reported a case of bilateral zoster occurring in the dermatomes of the left T2 and the right T5, and Jung et al³ reported a case of zoster duplex unilateralis that involved the unilateral dermatomes of V1-2 and L1-2, and had no history of malignancy, immunosuppressive agents and any other immunocompromised diseases.

Patients with malignancy, especially Hodgkin's disease and leukemia, are five times more likely to develop zoster than their age-matched counterparts¹. Other patients who also have a higher incidence of zoster include patients with deficient immune systems, such as individuals who are immunosuppressed for organ transplantation, by connective tissue disease, and by the agents such as corticosteroids¹. The clinical appearance of these patients is usually identical to typical zoster, but the lesions may be more ulcerative and necrotic and may scar more severely¹.

In our patient she had the history of acute leukemia and skin lesions of character-

istic painful vesicles involved two noncontiguous dermatomes on both halves of the body. Biopsy couldn't be done because of patient's refusal, but the diagnosis of zoster duplex bilateralis was made in the basis of the characteristic clinical features and Tzanck smear. We speculate that she was under immunocompromised state that led to have defective cellular immune response and develop zoster duplex bilateralis. She was successfully treated with antiviral agent without complication.

REFERENCES

1. Odom RB, James WD, Berger TG: Viral diseases. In: Andrews' diseases of the skin. 9th ed., W.B. Saunders Co., Philadelphia, 2000, pp486-491.
2. Lee SY, Choi YS, Yu HJ, Son SJ: A case of bilateral herpes zoster. *Kor J Dermatol* 32:1119-1122, 1994.
3. Jung GD, Won JY, Choi YH, Jeon YM, Song ES: A case of zoster duplex unilateralis. *Kor J Dermatol* 38:1569-1571, 2000.
4. Kim YP, Seo JI, Kang JB: Clinical observation of herpes zoster during a 10-year-period. *Kor J Dermatol* 18:65-79, 1980.
5. Buechner SA, Ruffi T: Atrophoderma of Pasini and Pierini. Clinical and histopathological findings and antibodies to *Borrelia burgdorferi* in thirty-four patients. *J Am Acad Dermatol* 30:441-446, 1994.
6. Chun IK, Lee JJ, Won YH, Kim YP: Clinical study on relation between herpes zoster and underlying diseases. *Kor J Dermatol* 26:356-365, 1988.
7. Jablonska A, Szczepanski A: Atrophoderma Pasini-Pierini: Is it an entity? *Dermatologica* 125:226, 1962.
8. Hong JH, Kye YC, Kim SN, Lee SY: A clinical study on herpes zoster in inpatients during a 3-year-period. *Kor J Dermatol* 32:583-590, 1994.
9. Kim SY, Cho BH, Kim JH: A 5-year clinical study on herpes zoster (1990-1994). *Kor J Dermatol* 35: 266-272, 1997.
10. Burgoon CF, Burgoon JS, Baldrige GD: The natural history of herpes zoster. *JAMA* 164:265-269, 1957.
11. Cousin GC, Ferguson MM: Bilateral shingles. *Br Dent J* 160:189, 1986.
12. Hill PA, Lamey PJ: Oral herpes zoster with con-

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tralateral skin involvement. *Br Dent J* 161:217-218, 1986.

13. Lewis RJ, Mitchell JC: Systemic lupus erythematosus, miliary tuberculosis, and bilateral herpes zoster occurring in a Chinese woman. *Arch Dermatol* 104:562, 1971.
14. Takayama M, Takayama N, Hachimori K: Restriction endonuclease analysis of viral DNA from a patient with bilateral zoster lesions. *J Infect Dis* 157:392-393, 1988.
15. Vu AQ, Radonich MA, Heald PW: Herpes zoster in seven disparate dermatomes(zoster multiplex): Report of a case and review of the literature. *J Am Acad Dermatol* 40:868-869, 1999.