

## **Original Research Article**

# **A study on clinical presentation of Herpes zoster in a district hospital in North India**

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**Abstract:** Herpes zoster is an acute, painful, vesicular eruption distributed along a single dermatome and is associated with a prodrome of fever, malaise, headache, and pain in the dermatome. The vesicles typically crust and will heal within 2-6 weeks. The main aim is to study the clinical manifestations of patients of herpes zoster. In this the method was fifty patients suffering from herpes zoster were selected for the study. It was a randomized controlled prospective study. In results the Pain was seen in 90% patients, parasthesias were seen in 50% patients, insomnia was seen in 20% patients, was seen in 90% patients was seen in 90% patients. PHN was seen in 80% patients with age more than 50 years. HIV positivity was seen in 4 (8%) patients. Regarding the dermatomal involvement it was seen that the thoracic dermatome was the commonest dermatome involved in 40% patients followed by trigeminal nerve segment in 36% patients, lumbar segment was involved in 16% patients, cervical dermatome was involved in 8% patients, was involved in 16% patients, cervicothoracic and thoracolumbar dermatomes were involved in 4% patients each. In conclusion the Patient education and counselling is very important, so that the patients can identify early eruptions and prodromal signs and symptoms and timely antiviral therapy can be instituted.

**Keywords:** Herpes zoster; dermatome; virus; ganglion; nerve; pain

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## **INTRODUCTION**

Herpes zoster is commonly referred to as shingles. It results from reactivation of latent varicella zoster virus in sensory dorsal root or cranial nerve ganglia, and usually manifests as a painful vesicular rash along a dermatomal distribution [1, 2]. Herpes zoster usually begins with a prodrome, such as pain, itching or tingling in the area that becomes affected. This may precede the characteristic rash by days or even weeks but is rarely the only clinical manifestation of varicella zoster virus reactivation. Typically, patients experience headache, malaise and sometimes photophobia. Abnormal sensation or pain, often described as burning, throbbing or stabbing, occurs in approximately 75% of patients and may be the first noticeable feature [3, 4]. Often pruritus in the affected region is the most prominent feature. Allodynia, or pain induced by light touch, may also be described. Before the onset of the rash and depending on the location, symptoms may mimic pain caused by ischaemic heart disease, cholecystitis or renal colic. The rash is usually unilateral and may affect adjacent dermatomes, with thoracic, cervical and ophthalmic involvement being the most common. Morphologically it evolves from a maculopapular rash to one comprising clusters of vesicles that ulcerate and crust over the course of 7–10

days. Healing is usually complete by 2–4 weeks. When all lesions have crusted the rash is considered non-infectious. Residual scarring and pigmentation is common [5].

## **AIMS**

- To study the clinical manifestations of patients of herpes zoster.
- To see for any complications of herpes zoster especially post herpetic neuralgia.

## **MATERIAL AND METHODS**

Fifty patients suffering from herpes zoster were selected for the study. It was a randomized controlled prospective study. Prior permission of hospital ethical committee was taken for the study. Written informed consent was taken from all the patients before the start of the study. The study was undertaken to see the clinical profile of patients of herpes zoster. A detailed history including patient's age and prodromal signs and symptoms were noted Patient's demographic data, symptoms, location of lesions, risk factors, associated systemic disease and complications were noted in a proforma. A complete dermatological examination was done in all the patients to see the

morphology of lesions and the dermatome involved. Routine investigations along with liver and kidney function tests were done in all the patients. HIV screening was done in all the patients. Tzanck smear was done in doubtful cases.



**Fig-1: Ophthalmic herpes zoster in a 66 years old male**



**Fig 2 - Herpes zoster in a 30 years old male with thoracic dermatomal involvement**



**Fig-3: Herpes zoster in a 13 years old male child with involvement of lumbar segment**

## RESULTS

The data was collected, tabulated and the results were analysed statistically.

**Table I: showing age distribution of patients**

Sr No	Age Distribution	Number	Percentage
1	0 - 20	5	10%
2	21 - 40	15	30%
3	41 - 60	25	50%
4	> 60	5	10%
5	TOTAL	50	100%

**Table 2: Showing Dermatomal Involvement of Herpes Zoster**

Sr No	Segment Involved	Number	Percentage
1	Thoracic	20	40%
2	Lumbar	8	16%
3	Trigeminal nerve area	12	36%
4	Sacral	2	4%
5	Cervical	4	8%
6	Thoracolumbar	2	4%
7	Cervicothoracic	2	4%

**Table 3: showing clinical features and complications of herpes zoster**

Sr No	Clinical Features and Complications	Number	Percentage
1	Pain	45	90 %
2	Fever	5	10%
3	Itching	12	24%
4	Paraesthesia	25	50%
5	Ulceration/Necrosis	5	10%
6	Insomnia	10	20%
7	Secondary infection	5	10%
8	Disseminated herpes zoster	3	6%
9	Ramsay hunt syndrome	8	16%

## DISCUSSION

Regarding the age distribution of patients, it was seen that maximum (50%) patients were between 41 – 60 years of age, 30% patients were between 21 – 40 years of age, 10% patients were between 0 – 20 years of age and another 10% patients were more than 60 years of age. There were 36 males and 14 females and male: female was 2.57:1. Out of 50 patients [8]. (16%) patients had multidermatomal involvement. It was seen that post herpetic neuralgia (PHN) was seen mostly in older patients (Fig 1). PHN was seen in 80% patients with age more than 50 years. HIV positivity was seen in 4 (8%) patients. Regarding the dermatomal involvement it was seen that the thoracic dermatome (Fig 2) was the commonest dermatome involved in 40%

patients followed by trigeminal nerve segment in 36% patients, lumbar segment (Fig 3) was involved in 16% patients, cervical dermatome was involved in 8% patients, was involved in 16% patients, cervicothoracic and thoracolumbar dermatomes were involved in 4% patients each. Pain was seen in 90% patients, parasthesias were seen in 50% patients, insomnia was seen in 20% patients, was seen in 90% patients was seen in 90% patients.

Herpes zoster is a localized disease characterized by unilateral radicular pain and grouped vesicular eruption that is generally limited to the dermatome innervated by a single spinal or cranial sensory ganglion [6, 7]. It occurs as a result of reactivation of varicella zoster virus (VZV) that had persisted in latent form within sensory ganglion following an earlier attack of varicella. The rash is usually unilateral and may affect adjacent dermatomes, with thoracic, cervical and ophthalmic involvement being the most common. Morphologically it evolves from a maculopapular rash to one comprising clusters of vesicles that ulcerate and crust over the course of 7–10 days. Healing is usually complete by 2–4 weeks [8]. When all lesions have crusted the rash is considered non-infectious. Residual scarring and pigmentation is common [8]. Immune system status plays a role; patients that are treated with immunosuppressive drugs have a significantly increased risk for herpes zoster. An immunocompromised patient is more likely to have a prolonged illness, more likely to recur, and more likely to develop myelitis and vasculopathy. The risk of herpes zoster is 15 times greater in men with HIV than in men without HIV [9, 10, 11]. The virus reactivates from its dormant state in the sensory ganglion, replicates in the nerve cells, and sheds virions from the cells that are carried down the axons to the skin served by that ganglion. The local immune response results in skin blisters or ocular inflammation depending on which tissues are affected. Perineuritis causes intense pain along the nerve distribution. Aging, immunosuppression therapy, and psychological stress all could be factors resulting in reactivation of the virus.

Post herpetic neuralgia is considered the most common complication and increases with age, affecting up to 30% of people with herpes zoster over the age of 80 years [12, 13, 14]. It is generally defined as pain of at least moderate intensity persisting for three months or longer, although various definitions (and measures of pain severity) have been used in drug trials. It may occasionally last for years. Post herpetic neuralgia is characterised by constant or intermittent, usually severe, burning or lancinating pain that occurs almost daily. Allodynia is present in most cases and can make even wearing clothing an arduous task. Quality of life is invariably reduced. Features that appear to be predictive for the development of post herpetic neuralgia include

more severe initial pain, more extensive rash and age over 50 years.

Ocular involvement in herpes zoster also called as herpes zoster ophthalmic us occurs in 10–25% of cases [15, 16]. This involves the ophthalmic branch of the trigeminal nerve and results in a disproportionately high complication rate (50% in the absence of antiviral drugs) with the eye affected in several possible ways [17]. Less common manifestations of zoster include the Ramsay Hunt syndrome (involvement of the geniculate ganglion of the facial nerve) which manifests as vesicles in the external auditory canal and palate associated with loss of taste to the anterior two-thirds of the tongue and facial weakness. Most individuals with herpes zoster will have some lesions outside the primary dermatome. Disseminated zoster is defined as 20 lesions or more outside the involved dermatome [18]. It tends to occur only in immunocompromised patients and may be associated with visceral involvement (lungs, liver, gut and brain) [19].

In a study by Goh and Khoo, dermatomes most commonly involved were thoracic in 45% and cervical in 23% [20]. Ophthalmic zoster was seen only in 3% cases. Unidermatomal, but may be multidermatomal, recurrent or disseminated. In another study conducted by Dubey *et al.*; the most common prodromal symptom seen was paresthesia in 25 (23.36%) cases followed by itching in [21] (19.62%) cases [21]. Most common presenting complaint was pain in 97 (90.65%) cases followed by cervical in 17 (15.8%) cases. Uni dermatomal involvement was seen in 81 (75.7%) cases followed by multidermatomal in 18 (16.8%) cases and disseminated in 8 (7.4%) cases.

In another study conducted by Latheef *et al.*, thoracic dermatome was commonly affected and among thoracic, T4 segment was common, followed by the Trigeminal nerve [ophthalmic branch, maxillary, and mandibular] [22]. Two cases of facial nerve involvement with Ramsay hunt syndrome were present. Twenty-five patients had cervical, 16 lumbar, and 10 had sacral nerve involvement. Nine patients had more than one dermatome involvement. Twenty-seven cases had aberrant vesicles ranging from 2–16 in distant areas.

## CONCLUSIONS

Patient education and counselling is very important, so that the patients can identify early eruptions and prodromal signs and symptoms and timely antiviral therapy can be instituted. Early management with antivirals and analgesia is important and may reduce the incidence of post herpetic neuralgia.

REFERENCES

1. Pranesh N, Tandon VK, Kumar R; Herpes zoster: A clinical study. Indian J Dermatol Venereol Leprol 1972; 38:152-5.
2. Sehgal VN, Rege VL, Kharangate VN; The natural history of Herpes Zoster. Indian J Dermatol Venereol Leprol 1976; 42:86-89.
3. Peeneys N; Diseases caused by viruses. In: Elder D, editor. Lever's Histopathology of the skin. 8th ed. Philadelphia: Lippincott - Raven; 1997; 569-89.
4. Chaudhary SD, Pahwa DA; A clinico-epidemiologic profile of herpes zoster in North India. Int J Dermatol Venereol Leprol 1987; 53:213-6.
5. Ragozzino MW, Melton LJ, Kurland LT; Population based study of herpes zoster and its sequelae. Medicine (Baltimore) 1982; 6: 310 - 6.
6. Schmader K, George LK, Burchett BM, *et al.*; Racial differences in the occurrence of herpes zoster. J Infect Dis 1995; 171:701-704.
7. Thomas SL, Hall AJ; What does epidemiology tell us about risk factors for herpes zoster? Lancet Infect Dis 2004; 4:26-33.
8. Schacker T, Corey L; Herpes virus infections in HIV infected person. In: Devita VT, Hailman Samuel, Lisenberg SA, editors. Textbook of AIDS. 4th ed. Philadelphia: Lippincott - Raven; 1997; 267-80.
9. Happenjans WB, Bibler MR, Orme RL; Prolonged cutaneous herpes zoster in acquired immunodeficiency syndrome. Arch Dermatol 1988; 126:1048.
10. Bernhard P, Obel N; Chronic ulcerating acyclovir resistant varicella zoster lesions in an AIDS patient. Scand J Infect Dis 1996; 27:623-5.
11. Watson PN, Evens RJ; Post herpetic neuralgia: A review. Arch Neural 1986; 43:836 -40.
12. KostRg, Strauss SE; Post herpetic neuralgia: Pathogenesis, treatment and prevention. N Engl J Med. 1996; 335:32-41.
13. Wilson JF; Herpes zoster. Ann Intern Med 2011; 154:ITC31-15.
14. Talwar S, Shrivastava VK; Herpes zoster ophthalmic us with total ophthalmoplegia. Indian J Dermatol Venereol Leprol 1991; 56:454-5.
15. Lies gang TJ; Herpes Zoster Ophthalmicus. Ophthalmology 2008; 115:S3-S12.
16. Hutchinson J; A clinical report on herpes zoster ophthalmic us (shingles affecting the forehead and nose). Trans Am Ophthalmol oC 1942; 40:390-439.
17. Kar HK, Gautam RK, Jain RK; Disseminated cutaneous herpes zoster: A clinical predictor of human immunodeficiency virus infection. Indian J Dermatol Venereol Leprol 1995; 61:40-1.
18. Mandal BK; Herpes Zoster in the immunocompromised populations. Indian J Dermatol Venereol Leprol. 2006; 5:235-43.
19. Fueyo MA, Lookingbill DP; Herpes zoster and occult malignancy. J Am Acad Dermatol 1984; 11:480-2.
20. Goh CL, Khoo L; A retrospective study of the clinical presentation and outcome of herpes zoster in a tertiary dermatology outpatient referral clinic. Int 1997; 36:667-72.
21. Dubey AK, Jaisankar TJ, Thappa DM; Clinical and morphological characteristics of herpes zoster in south India. Indian J Dermatol 2005; 50:203-7.
22. EN Abdul Latheef, K Pavithran; Herpes zoster: A clinical study in 205 patients. Indian J Dermatol 2011; 56: 529-532.