

A Comparative Study Assessing the Efficacy of Antiviral Agent Valacyclovir Compared with Famciclovir in the Treatment of Herpes Zoster

Jaya Roy¹, Md. Zamiruddin², Rohit Kumar Singh³

¹Tutor, Department of Pharmacology, Anugrah Narayan Magadh Medical College and Hospital, Gaya, Bihar, India

²Assistant Professor, Department of Pharmacology, Anugrah Narayan Magadh Medical College and Hospital, Gaya, Bihar, India

³Associate Professor and HOD, Department of Pharmacology, Anugrah Narayan Magadh Medical College and Hospital, Gaya, Bihar, India

Received: 09-07-2023 / Revised: 11-08-2023 / Accepted: 15-09-2023

Corresponding Author: Dr. Jaya Roy

Conflict of interest: Nil

Abstract:

Aim: The objective was to evaluate the efficacy of antiviral agent valacyclovir compared with famciclovir in the treatment of herpes zoster.

Methods: The present study was comprised of 200 patients who developed herpes zoster of both genders. All of the patients that were enrolled gave their consent. The study was carried out at hospital over a 2-year period. Data such as name, age, gender, etc. was recorded. All were divided into 2 groups of 50 patients each. Group I patients were prescribed valacyclovir 1000 mg thrice daily, and Group II patients were prescribed famciclovir 500 mg thrice daily. The treatment was given for 7 days.

Results: Group I had 56 males and 44 females and group II had 48 males and 52 females. Dermatome in group I and group II involved were thoracic in 60 and 52, lumbar in 12 and 12, cervical in 16 and 20 and trigeminal in 12 and 16 respectively. The difference was non-significant ($P > 0.05$). The VAS scores comparison at every follow-up visit between both the groups. The mean VAS scores on the day of presentation, i.e., the baseline scores in both the groups were almost similar, and the difference was not statistically significant. Results showed a greater number of totally pain-free patients at day 30, i.e., 82 (82%) patients in the valacyclovir group, while in the famciclovir group, 62 (62%) patients were reported to be totally pain free, although it was statistically insignificant ($P > 0.05$).

Conclusion: In comparison to famciclovir, oral valacyclovir administered for 7 days during acute zoster infection offers significant benefit by providing a well-tolerated and greater resolution of pain while maintaining a favourable safety profile, making valacyclovir more efficacious and a better drug in the management of Herpes Zoster.

Keywords: Famciclovir, herpes zoster, valacyclovir

This is an Open Access article that uses a funding model which does not charge readers or their institutions for access and distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>) and the Budapest Open Access Initiative (<http://www.budapestopenaccessinitiative.org/read>), which permit unrestricted use, distribution, and reproduction in any medium, provided original work is properly credited.

Introduction

Herpes zoster a localized disease has been known since ancient times and is often referred by different names such as varicella zoster and shingles. It is typically characterized by unilateral radicular pain along with grouped vesicular eruptions. [1] Immunosuppression and increasing age are well-established risk factors that can lead to latent virus reactivation. Pain is a common symptom that compels patients to seek medical advice. It frequently lasts even after the rash has healed, a consequence known as postherpetic neuralgia. This distinctive presentation of signs and symptoms is usually sufficient enough to reach a clinical diagnosis. [2]

Among the antiviral agents, acyclovir is the most commonly used, but its prodrug, valacyclovir, has been observed to be better than acyclovir. [3] Valacyclovir is known to accelerate the resolution of acute pain associated with herpes zoster and also decrease the number of patients complaining of persistent pain. [4] Famciclovir is another antiviral agent that is a prodrug of penciclovir and has the advantage of a longer intracellular half-life and better bioavailability. Some studies claim famciclovir to be a better drug when compared to valacyclovir for the relief of pain. [5,6]

The pharmacotherapy for its management comprises antiviral agents, analgesics, and corticosteroids. The ideal time to start the treatment is within 3 days from when the rash first appears. Opioids and tricyclic antidepressants are often used for the treatment of significant persisting pain. Among the antiviral agents, acyclovir is the most commonly used agent, but its prodrug valacyclovir has been observed to be better than acyclovir. [6] Valacyclovir is known to accelerate the resolution of acute pain associated with herpes zoster and also decreases the number of patients complaining of persistent pain. [6] Famciclovir is another antiviral agent, which is a prodrug of penciclovir available with the advantage of a longer intracellular half-life and a better bioavailability. Some studies claim famciclovir to be a better drug when compared to valacyclovir for relief of pain, [7] while others state valacyclovir to be a better drug when compared to famciclovir. [8]

The objective was to evaluate the efficacy of antiviral agent valacyclovir compared with famciclovir in the treatment of herpes zoster.

Materials and Methods

The present study was comprised of 200 patients who developed herpes zoster of both genders. All of the patients that were enrolled gave their consent. The study was carried out at hospital, Anugrah Narayan Magadh Medical College and Hospital, Gaya Bihar, India over a 2 year period. Data such as name, age, gender, etc. was recorded. All were divided into 2 groups of 50 patients each. Group I patients were prescribed valacyclovir 1000 mg thrice daily, and Group II patients were prescribed famciclovir 500 mg thrice daily. The treatment was given for 7 days. During the next 10 months of the trial, the same alternating medication method was used during the study period. One dose of 40 milligrams of methylprednisone was given once daily in the morning for one week following the presentation, followed by tapering over the next 2 weeks, and was added to all the patients. For the first week, acetaminophen 500 mg TDS was also added. Follow-up was done on days 4, 9, 16, 22, and 30. The data thus obtained was subjected to statistical analysis. A P value of 0.05 was considered significant.

Results

Table 1: Distribution of patients

Groups	Group I	Group II
Drug	Valacyclovir 1000 mg	Famciclovir 500 mg
M:F	56/44	48/52

Group I had 56 males and 44 females and group II had 48 males and 52 females.

Table 2: Dermatome distribution

Dermatome	Group I	Group II	P value
Thoracic	60	52	0.20
Lumbar	12	12	
Cervical	16	20	
Trigeminal	12	16	

Dermatome in group I and group II involved were thoracic in 60 and 52, lumbar in 12 and 12, cervical in 16 and 20 and trigeminal in 12 and 16 respectively. The difference was non-significant ($P > 0.05$).

Table 3: Mean visual analog scale scores comparison at every follow-up visit between both the study groups

Days	Mean±SD		P
	Valacyclovir	Famciclovir	
0	6.53±1.51	6.27±1.67	0.5290
4	4.13±0.99	4.33±1.29	0.5030
9	2.0±1.36	2.33±1.59	0.3910
16	1.07±1.28	1.33±1.50	0.4730
22	0.60±0.83	1.2±1.42	0.0500
30	0.2±0.41	0.87±1.19	0.0050

The VAS scores comparison at every follow-up visit between both the groups. The mean VAS scores on the day of presentation, i.e., the baseline scores in both the groups were almost similar, and the difference was not statistically significant.

Table 4: Comparing the number of patients free of zoster pain in both the study groups at each follow-up

Days	Mean±SD		P
	Valacyclovir	Famciclovir	
4	0	0	-
9	22 (22)	22 (22)	1.000
16	52 (52)	48 (48)	0.7963
22	62 (62)	58 (58)	0.7945
30	82 (82)	62 (62)	0.1590

Results showed a greater number of totally pain-free patients at day 30, i. e., 82 (82%) patients in the valacyclovir group, while in the famciclovir group, 62 (62%) patients were reported to be totally pain free, although it was statistically insignificant ($P > 0.05$).

Discussion

Viral herpes zoster (HZ) is characterised by a unilateral dermatome rash and pain due to the reactivation and amplification of endogenous varicella zoster virus (VZV) dormant in sensory ganglia due to basic varicella infection (VZI). [9,10] Varicella zoster virus reactivation is the cause of herpes zoster, also known as shingles. People over the age of 50 are more likely to acquire herpes zoster because of immunosenescence, although it can affect anyone, especially those with a reduced cell-mediated immunity owing to any condition or medication. [11] Herpes zoster has been linked to complications involving the cerebral, splanchnic, motor nerves and ophthalmic. However, post-herpetic neuralgia is the most frequent complication. [12,13] Ophthalmic zoster infection can cause permanent damage to the eyes, including blindness. Approximately 4 to 6 cases per 1000 individuals of general population have been observed every year. [14]

Group I had 56 males and 44 females and group II had 48 males and 52 females. Dermatome in group I and group II involved were thoracic in 60 and 52, lumbar in 12 and 12, cervical in 16 and 20 and trigeminal in 12 and 16 respectively. The difference was non-significant ($P > 0.05$). Valacyclovir is a prodrug of acyclovir. [15] Famciclovir is also a prodrug with an active metabolite penciclovir. They both inhibit viral DNA polymerase enzyme, thus preventing the viral replication. [16] When given within 3 days of the appearance of the first lesion, both the drugs provide a reduction in duration not only loss of acute pain but also of full crusting and healing of the zoster associated lesions. [17] Both valacyclovir and famciclovir were well tolerated. No serious adverse effects were observed in either of the study groups to warrant withdrawal of any patient. A male preponderance was observed as in other studies conducted previously. [18] On observing the dermatome distribution, both the groups showed

thoracic dermatomes to be the most commonly involved dermatome segments.

The mean time to full crusting of herpes zoster lesions was 15.033 days in the acyclovir group and 14.840 days in the famciclovir group (log-rank p -value = 0.820). Headache, diarrhoea, nausea, back pain, cold, and drowsiness were the most common side effects in the pooled groups, but none of them were deemed clinically significant. Dermatome in group I and group II involved were thoracic in 60 and 52, lumbar in 12 and 12, cervical in 16 and 20 and trigeminal in 12 and 16 respectively. The difference was non-significant ($P > 0.05$). Bist et al [19] evaluated the efficacy of the antiviral agent valacyclovir compared with famciclovir in the treatment of herpes zoster. The first group of patients received valacyclovir tablets of 1000 mg thrice daily, whereas those in the second group were given famciclovir tablets of 500 mg thrice daily. Both medications were administered for seven days. The effects of the administered medications were assessed on a regular basis till the 29th day. On day 29, when pain scores were compared between the two groups using the visual analogue scale, the valacyclovir group scored significantly lower than the famciclovir group. Furthermore, when compared to famciclovir, valacyclovir treatment accelerated the cure of zoster-associated pain in a greater number of patients. Basicckes V et al [18] found that in comparison to famciclovir, oral valacyclovir administered for 7 days during acute zoster infection offers significant benefit by providing a well-tolerated and greater resolution of pain while maintaining a favourable safety profile, making valacyclovir more efficacious and a better drug in the management of Herpes Zoster. Results showed a greater number of totally pain-free patients at day 30, i. e., 82 (82%) patients in the valacyclovir group, while in the famciclovir group, 62 (62%) patients were reported to be totally pain free, although it was statistically insignificant ($P > 0.05$). The most common symptom of shingles is pain, which affects about 75% of patients in the form of altered sensitivity or pain circumscribed to the affected dermatome, where the rash will appear later. In the course of viral reactivation, acute hyperalgesia is usually the first symptom and occurs in approximately 70–80% of patients. [20] However,

the type and degree of pain can change over time, and it can occur at all stages of the disease.

Conclusion

In comparison to famciclovir, oral valacyclovir administered for 7 days during acute zoster infection offers significant benefit by providing a well-tolerated and greater resolution of pain while maintaining a favourable safety profile, making valacyclovir more efficacious and a better drug in the management of Herpes Zoster. In the present study, we found that oral valacyclovir in acute zoster infection was found to be better as compared to famciclovir.

References

1. Strauss SE, Oxman MN, Schmader KE. Varicella and herpes zoster. In: Wolff K, Goldsmith LA, Katz SI, Gilchrist BA, Paller AS, Leffell DJ, editors. Fitzpatrick's Dermatology in General Medicine. 7th ed. New York: Mc Graw Hill Publishing Division; 2008. pp. 1885–98.
2. Thomas SL, Hall AJ. What does epidemiology tell us about risk factors for herpes zoster?. The Lancet infectious diseases. 2004 Jan 1;4(1):26-33.
3. Wareham DW, Breuer J. Herpes zoster. BMJ. 2007;334:1211–5.
4. Jeon YH. Herpes zoster and postherpetic neuralgia: practical consideration for prevention and treatment. The Korean journal of pain. 2015 Jul 1;28(3):177-84.
5. Glaser RB. Clinical aspects of herpes zoster. Western Journal of Medicine. 1983 Nov;139(5):718.
6. Beutner KR, Friedman DJ, Forszpaniak C, Andersen PL, Wood MJ. Valaciclovir compared with acyclovir for improved therapy for herpes zoster in immunocompetent adults. Antimicrobial Agents and Chemotherapy. 1995 Jul;39(7):1546-53.
7. Ono F, Yasumoto S, Furumura M, Hamada T, Ishii N, Gyotoku T, Higuchi M, Inokuchi K, Jyo K, Koga H, Komai A, Maruta K, Mashiko T, Mihara T, Miyahara H, Miyasato M, Muto K, Nagase K, Nagata M, Sakihama H, Tanahashi T, Ueda A, Yamakawa K, Ohata C, Dainichi T, Tsuruta D, Hashimoto T. Comparison between famciclovir and valacyclovir for acute pain in adult Japanese immunocompetent patients with herpes zoster. J Dermatol. 2012 Nov;39(11):902-8.
8. Wald A, Selke S, Warren T, Aoki FY, Sacks S, Diaz-Mitoma F, Corey L. Comparative efficacy of famciclovir and valacyclovir for suppression of recurrent genital herpes and viral shedding. Sex Transm Dis. 2006 Sep;33(9):529-33
9. Schmader KE, Oxman MN. Varicella & Herpes zoster. In: Fitzpatrick's Dermatology in General Medicine. Wolff K, Goldsmith LA, Katz SI, Gilchrist BA, Paller AS, Leffell DJ. 8th ed, New York: McGraw- Hill; 2012:2383-2401.
10. Suaya JA, Chen SY, Li Q, Burstin SJ, Levin MJ. Incidence of herpes zoster and persistent post-zoster pain in adults with or without diabetes in the United States. In Open forum infectious diseases 2014 Sep 1 (Vol. 1, No. 2, p. ofu049). Oxford University Press.
11. Lal H, Cunningham AL, Godeaux O, Chlibek R, Diez-Domingo J, Hwang SJ, Levin MJ, McElhaney JE, Poder A, Puig-Barberà J, Vesikari T. Efficacy of an adjuvanted herpes zoster subunit vaccine in older adults. New England Journal of Medicine. 2015 May 28;372(22):2087-96.
12. Gershon AA, Gershon MD, Breuer J, Levin MJ, Oaklander AL, Griffiths PD. Advances in the understanding of the pathogenesis and epidemiology of herpes zoster. Journal of clinical virology. 2010 May 1;48:S2-7.
13. Ansaldi F, Trucchi C, Alicino C, Paganino C, Orsi A, Icardi G. Real-world effectiveness and safety of a live-attenuated herpes zoster vaccine: a comprehensive review. Advances in Therapy. 2016 Jul;33:1094-104.
14. Shaikh S, Ta CN. Evaluation and management of herpes zoster ophthalmicus. Am Fam Physician. 2002; 66(9):1723-30.
15. Strauss SE, Oxman MN, Schmader KE. Varicella and herpes zoster. In: Wolff K, Goldsmith LA, Katz SI, Gilchrist BA, Paller AS, Leffell DJ, editors. Fitzpatrick's Dermatology in General Medicine. 7th ed. New York: Mc Graw Hill Publishing Division; 2008. pp. 1885–98.
16. Araújo LQ, Macintyre CR, Vujacich C. Epidemiology and burden of herpes zoster and postherpetic neuralgia in Australia, Asia and South America. Herpes. 2007 Sep;14 Suppl 2:40-4.
17. Heininger U, Seward JF. Varicella. Lancet. 2006 Oct 14;368(9544):1365-76.
18. Basic-Kes V, Demarin V. Postherpetic neuralgia. Acta Clinica Croatica. 2007 Sep 1;46(3):279-82.
19. Bist A, Savitha A, Gumma KM. Efficacy of valacyclovir and famciclovir in herpes zoster: A comparative study. Indian Journal of Pharmacology. 2020 Nov;52(6):472.
20. Indradevi R, Manoharan K, Oudeacoumar P, Karthikraja S, Jaffar NA. A comparative study to evaluate the efficacy and safety of valacyclovir and famciclovir in the management of herpes zoster in a tertiary care hospital in Puducherry. Journal of Evolution of Medical and Dental Sciences. 2014 May 26;3(21) :5804-13.