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Key words: epidemiology; female; incidence; males; melanoma; mortality; public health; skin cancer; survival; young adults.

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Conflicts of interest

None disclosed.

REFERENCES

1. Reed KB, Brewer JD, Lohse CM, Bringe KE, Pruitt CN, Gibson LE. Increasing incidence of melanoma among young adults: an epidemiological study in Olmsted County, Minnesota. *Mayo Clin Proc.* 2012;87(4):328-334.
2. Guy GP Jr, Watson M, Seidenberg AB, Hartman AM, Holman DM, Perna F. Trends in indoor tanning and its association with sunburn among US adults. *J Am Acad Dermatol.* 2017;76(6):1191-1193.
3. Indoor Tanning Services; Cosmetic Services; Excise Taxes. Federal Register Volume 75, Issue 114 (June 15, 2010). Accessed April 16, 2023. <https://www.govinfo.gov/app/details/FR-2010-06-15/2010-14398>
4. Use By Minors Prohibited. Minnesota Legislators. Accessed April 16, 2023. <https://www.revisor.mn.gov/statutes/cite/325H.085> <https://www.revisor.mn.gov/laws/2014/0/291/>
5. St Sauver JL, Grossardt BR, Leibson CL, Yawn BP, Melton LJ III, Rocca WA. Generalizability of epidemiological findings and public health decisions: an illustration from the Rochester Epidemiology Project. *Mayo Clin Proc.* 2012;87(2):151-160.

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Association of herpes zoster with COVID-19 vaccination: A systematic review and meta-analysis



To the Editor: Herpes zoster (HZ), which is reactivation of the varicella zoster virus (VZV), is associated with older age, use of immunomodulatory drugs, trauma, family history, and other comorbidities.¹ Moreover, case studies have reported development of HZ following administration of vaccinations against influenza, Japanese encephalitis, hepatitis A, and rabies.² Recently, several cases of HZ after COVID-19 vaccinations have been reported¹; however, the association remains a matter of debate due to the small number of cases and lack of control groups.

We performed a systematic search within PubMed, EMBASE, and Web of Science for relevant publications from inception to November 2022 following the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines. The protocol was registered in PROSPERO (CRD42023381589). Cohort studies, case-control cross-sectional studies, and randomized controlled trials reporting HZ outcomes in patients receiving COVID-19 vaccinations and control subjects were included. Studies with overlapping cases, lack of HZ outcomes, incomplete data, and case studies without controls were excluded (Supplementary Fig 1, available via Mendeley at <https://data.mendeley.com/datasets/6jg7jkn65r/1>). This meta-analysis (MA) used a random-effects model to calculate the pooled odds ratios (ORs) and 95% CIs to determine the risk of HZ in the COVID-19 vaccination group versus control groups. Subgroup analyses comparing the risk of HZ in the mRNA versus adenovirus vaccination groups and Moderna versus BioNTech vaccination groups were performed. Sixteen studies were included from initially identified 465 articles (Supplementary Table 1, available via Mendeley at <https://data.mendeley.com/datasets/6jg7jkn65r/1>). COVID-19 vaccination was associated with a significantly increased risk of HZ (OR, 1.32; 95% CI, 1.09-1.62, $P = .006$) compared with controls (Supplementary Fig 2, available via Mendeley at <https://data.mendeley.com/datasets/6jg7jkn65r/1>). In subgroup analysis, the mRNA vaccination was associated with a higher

risk of HZ compared with the adenovirus vaccination (OR, 1.67; 95% CI, 1.19-2.35, $P = .003$) (Supplementary Fig 3, available via Mendeley at <https://data.mendeley.com/datasets/6jg7jkn65r/1>). Further MA of studies comparing different brands of COVID-19 vaccinations showed no significant difference of HZ risks between Moderna and BioNTech (OR, 0.64; 95% CI, 0.18-2.21) (Supplementary Fig 4, available via Mendeley at <https://data.mendeley.com/datasets/6jg7jkn65r/1>).

The mechanism underlying the link between COVID-19 vaccination and HZ remains elusive; however, vaccination-induced immunomodulation has been proposed. Vaccine-induced massive shift of CD8⁺ T cells and CD4⁺ helper T cells may cause temporary inability to suppress latent VZV, allowing for its reactivation.³ Previous studies indicate that immunocompromised status and older age are associated with a higher risk of VZV reactivations after vaccination.⁴ The reported median time to onset of HZ after COVID-19 vaccination was 7 to 10 days (range, 2-51 days).^{1,4,5} Although most cases of VZV reactivations were dermatome-limited, 2 cases of HZ infection after COVID-19 mRNA vaccination were disseminated.⁵ Our research has limitations. First, because most randomized controlled trials of COVID-19 vaccination did not report HZ as a separate individual adverse effect, the number of randomized controlled trials included in this MA is quite limited. Second, heterogeneity exists among the included studies, and the quality of the included studies is not very high (Supplementary Fig 5, available via Mendeley at <https://data.mendeley.com/datasets/6jg7jkn65r/1>). Nevertheless, our MA suggests an increased risk of HZ in patients receiving COVID-19 vaccination, and the mRNA vaccination is associated with a higher risk of HZ than the adenovirus vaccination. This highlights the awareness of possible reactivation of HZ following COVID-19 vaccination, particularly for high-risk individuals.

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Conflicts of interest

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REFERENCES

1. Rodríguez-Jiménez P, Chicharro P, Cabrera LM, et al. Varicella-zoster virus reactivation after SARS-CoV-2 BNT162b2 mRNA vaccination: report of 5 cases. *JAAD Case Rep.* 2021;12:58-59.
2. Walter R, Hartmann K, Fleisch F, Reinhart WH, Kuhn M. Reactivation of herpesvirus infections after vaccinations? *Lancet.* 1999;353(9155):810.
3. Psichogiou M, Samarkos M, Mikos N, Hatzakis A. Reactivation of varicella zoster virus after vaccination for SARS-CoV-2. *Vaccines (Basel).* 2021;9(6):572.
4. Préta LH, Contejean A, Salvo F, Treluyer JM, Charlier C, Chouchana L. Association study between herpes zoster reporting and mRNA COVID-19 vaccines (BNT162b2 and mRNA-1273). *Br J Clin Pharmacol.* 2022;88(7):3529-3534.
5. Said JT, Virgen CA, Lian CG, Cutler CS, Merola JF, LeBoeuf NR. Disseminated varicella-zoster virus infections following messenger RNA-based COVID-19 vaccination. *JAAD Case Rep.* 2021;17:126-129.