

MD,^c Jerry D. Brewer, MD, MS,^d and Addison M. Demer, MD^d

From the Department of Dermatology, Mayo Clinic School of Graduate Medical Education, Rochester, Minnesota^a; Division of Clinical Trials and Biostatistics, Department of Quantitative Health Sciences, Mayo Clinic, Rochester, Minnesota^b; Division of Dermatopathology, Department of Dermatology, Mayo Clinic Rochester, Minnesota^c; and Division of Dermatologic Surgery, Department of Dermatology, Mayo Clinic, Rochester, Minnesota.^d

Funding sources: Mayo Clinic Department of Dermatology.

IRB approval status: Reviewed and approved by Mayo Clinic IRB (#21-000552). Reviewed and approved by Olmsted Medical Center IRB (#004-OMC-21).

This study used the resources of the Rochester Epidemiology Project (REP) medical records-linkage system, which is supported by the National Institute on Aging (NIA; AG 058738), by the Mayo Clinic Research Committee, and by fees paid annually by REP users. The content of this article is solely the responsibility of the authors and does not represent the official views of the National Institutes of Health (NIH) or the Mayo Clinic.

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

Correspondence and reprint requests to: Olivia M. Crum, MD, Department of Dermatology, Mayo Clinic School of Graduate Medical Education, 200 First St SW, Rochester, MN 55904

E-mail: Crum.Olivia@mayo.edu

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.

<https://doi.org/10.1016/j.jaad.2023.03.030>

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 I, 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.

We thank the staff of Department of Medical Research, National Taiwan University Hospital Hsin-Chu Branch for their assistance in study design, statistical analysis, and providing careful review and insightful comments regarding the articles.

I-Ling Chen, MD,^a and Hsien-Yi Chiu, MD, PhD^{b,c,d,e}

From the Departments of Family Medicine, National Taiwan University Hospital Hsin-Chu Branch, Taiwan^a; Department of Medical Research, National Taiwan University Hospital Hsin-Chu Branch, Taiwan^b; Department of

Dermatology, National Taiwan University Hospital Hsin-Chu Branch, Taiwan^c; Department of Dermatology, National Taiwan University Hospital, Taipei, Taiwan^d; and Department of Dermatology, College of Medicine, National Taiwan University, Taipei, Taiwan.^e

Funding sources: This work was funded in part by grants from National Taiwan University Hospital, Hsin-Chu branch (112-HCH092, 112-HCH065) and Taiwan Ministry of Science and Technology (MOST 111-2314-B-002-244). The funders had no role in the study design, data collection and analysis, interpretation of findings, manuscript writing, or target journal selection.

IRB approval status: Not applicable.

Key words: herpes zoster; shingles; COVID-19; COVID-19 vaccines; meta-analysis.

Reprints not available from the authors.

Correspondence to: Hsien-Yi Chiu, MD, PhD, Department of Dermatology, National Taiwan University Hospital Hsin-Chu Branch, Taiwan, NO. 25, Lane 442, Sec. 1, Jingguo Rd, Hsinchu City 300, Taiwan (R.O.C.)

E-mail: extra.ow10430@yahoo.com.tw

Conflicts of interest

H.Y.C. received speaking fees from AbbVie, Novartis Pharmaceuticals Corporation, Janssen-Cilag Pharmaceutica, Eli-Lilly, Kyowa Hakko Kirin Taiwan, and Pfizer Limited and conducted clinical trials for Eli-Lilly, AbbVie, and Sanofi Pharmaceuticals. I.L.C. has no conflicts of interest to declare.

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.

<https://doi.org/10.1016/j.jaad.2023.03.031>