# Stock-exchange announcement

## For media and investors only



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# GSK's *Shingrix* approved in China for prevention of shingles in adults aged 18 and over who are at increased risk due to immunodeficiency or immunosuppression

- Shingrix (GSK's Recombinant Zoster Vaccine or RZV) is the first and only vaccine approved in this population in China
- Approval adds to existing indication in adults aged 50 and over
- Approximately six million cases of shingles occur in China annually,<sup>1</sup> with immunodeficient or immunosuppressed patients at a higher risk than the general population<sup>2</sup>

GSK plc (LSE/NYSE: GSK) today announced that the China National Medical Products Administration (NMPA) has approved *Shingrix* (GSK's Recombinant Zoster Vaccine or RZV) for the prevention of shingles (herpes zoster) in adults aged 18 years and over who are at increased risk of shingles due to immunodeficiency or immunosuppression caused by known disease or therapy.

Shingles poses a significant health burden, with approximately six million cases per year in China. Besides advancing age, other factors can increase the risk of developing shingles, including immunodeficiency or immunosuppression with no other shingles vaccine approved for this population in China.

Shingles-associated pain is often described as aching, burning, stabbing or shock-like<sup>4</sup> and can have a considerable impact on patients' quality of life, e.g. affecting sleep, and ability to undertake activities of daily living including work.<sup>5,6,7</sup> This approval expands the reach of GSK's RZV, to ensure protection for those patients most vulnerable to shingles.

Sanjay Gurunathan, Senior Vice President, Vaccines and Infectious Diseases R&D, said: "This approval marks a critical milestone in expanding access to GSK's RZV for those at a higher risk of what can be a disrupting and devastating disease. Through close collaboration with regulatory bodies, we continue to drive innovation that helps protect vulnerable patient groups and shifting the focus of healthcare systems towards preventing diseases, like shingles."

The NMPA application was informed by six clinical trials in patients aged 18 years and over who had undergone recent blood-forming cells (stem cell) transplantation, kidney transplant, or have blood cancer, solid tumour, or HIV.8,9,10,11,12,13

#### About shingles

Shingles is caused by the reactivation of the varicella zoster virus (VZV), the same virus that causes chickenpox.<sup>4</sup> Shingles typically presents as a rash, with painful blisters across the chest, abdomen or face.<sup>14</sup> Following the rash, a person can also experience post-herpetic neuralgia (PHN), a long-lasting nerve pain that can last weeks or months and can occasionally persist for several years.<sup>4</sup> PHN is the most common complication of shingles, occurring in 5–30% of all shingles cases.<sup>15</sup>

#### About Shingrix

Shingrix (Recombinant Zoster Vaccine or RZV) is a non-live, recombinant subunit vaccine indicated for the prevention of shingles in adults 50 and over and in several countries and regions. RZV is also approved for adults

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aged 18 years or over at increased risk for shingles. It combines an antigen, glycoprotein E, with an adjuvant system, AS01<sub>B</sub>, and may help overcome the natural age-related decline in responses to immunisation that contributes to the challenge of protecting adults aged 50 and over from shingles. RZV is not indicated to prevent primary varicella infection (chickenpox). The use of RZV should be in accordance with official recommendations and local product label.

For product and important safety information please consult the country relevant summary of product characteristics.

#### About GSK

GSK is a global biopharma company with a purpose to unite science, technology, and talent to get ahead of disease together. Find out more at gsk.com.

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#### Cautionary statement regarding forward-looking statements

GSK cautions investors that any forward-looking statements or projections made by GSK, including those made in this announcement, are subject to risks and uncertainties that may cause actual results to differ materially from those projected. Such factors include, but are not limited to, those described in the "Risk Factors" section in GSK's Annual Report on Form 20-F for 2024, and GSK's Q2 Results for 2025.

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