



Audio File

# PUBLIC HEALTH IMPACT AND COST-EFFECTIVENESS ANALYSIS OF UNIVERSAL VARICELLA VACCINATION IN THE UNITED KINGDOM

Jamet N<sup>1</sup>, George M<sup>2</sup>, Shah H<sup>2</sup>, Oliyide A<sup>3</sup>, Zerda I<sup>4</sup>, Cristeau O<sup>5</sup>, Casabona G<sup>1</sup>

<sup>1</sup>GSK, Wavre, Belgium; <sup>2</sup>Hari Thrivikramji C/O GSK, Wavre, Belgium; <sup>3</sup>GSK, London, United Kingdom;

<sup>4</sup>Creativ-Ceutical, Cracow, Poland; <sup>5</sup>Creativ-Ceutical, Paris, France



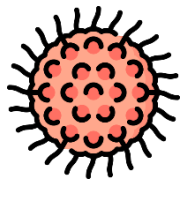
SCAN ME

EPH161

Information on QR code



## Background



Varicella is a highly infectious childhood disease caused by the varicella-zoster virus (VZV).<sup>1,2</sup>



Following primary infection, VZV remains latent in the body and can reactivate later in life as herpes zoster (HZ).<sup>3</sup> Currently, there is no universal varicella vaccination (UVV) in the United Kingdom (UK).



Varicella vaccination impact and cost-effectiveness was assessed in Akpo et al.<sup>4</sup> This model is an update including HZ immunization strategy, updating key parameters, and considering several vaccination schedules.



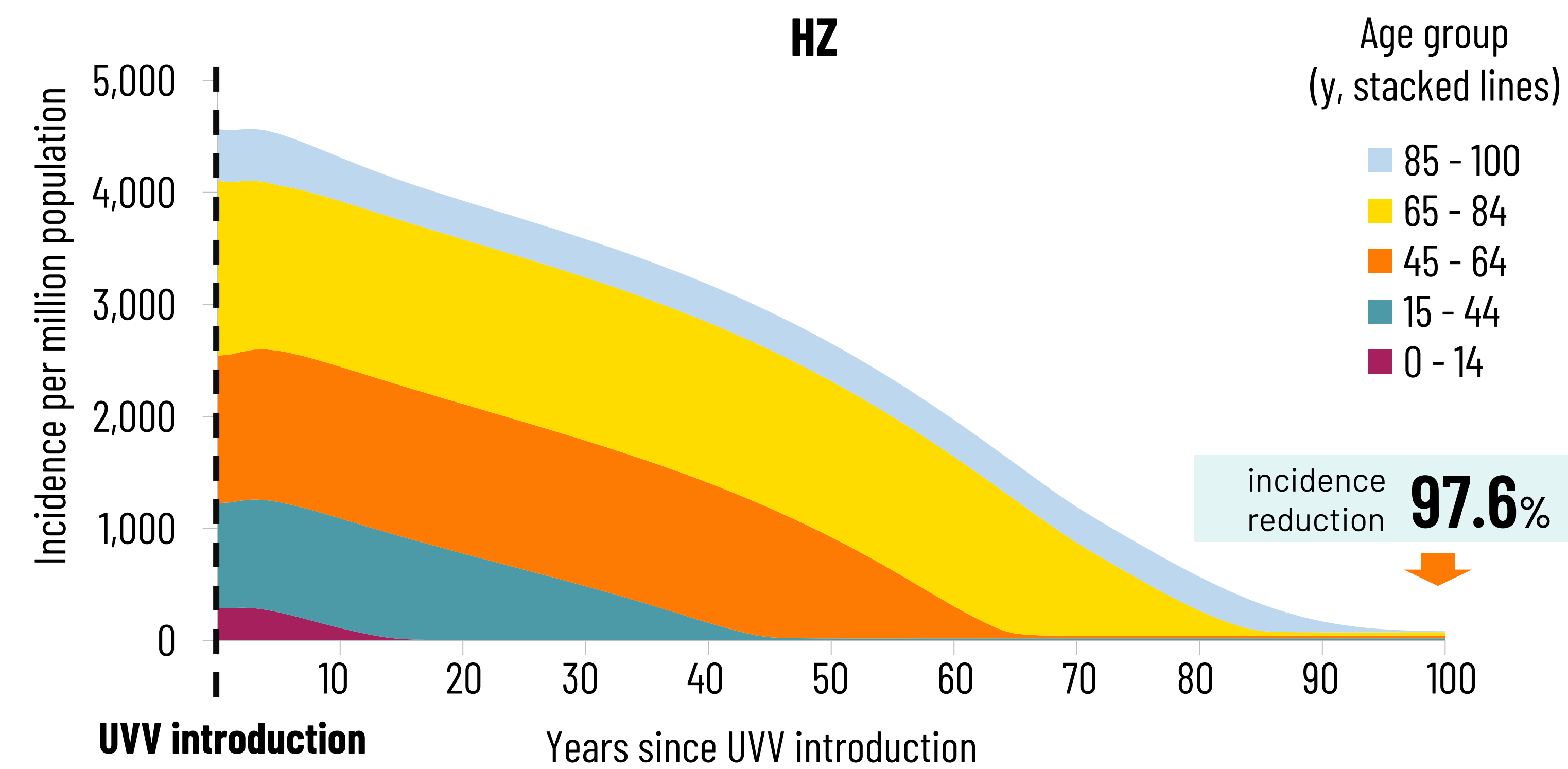
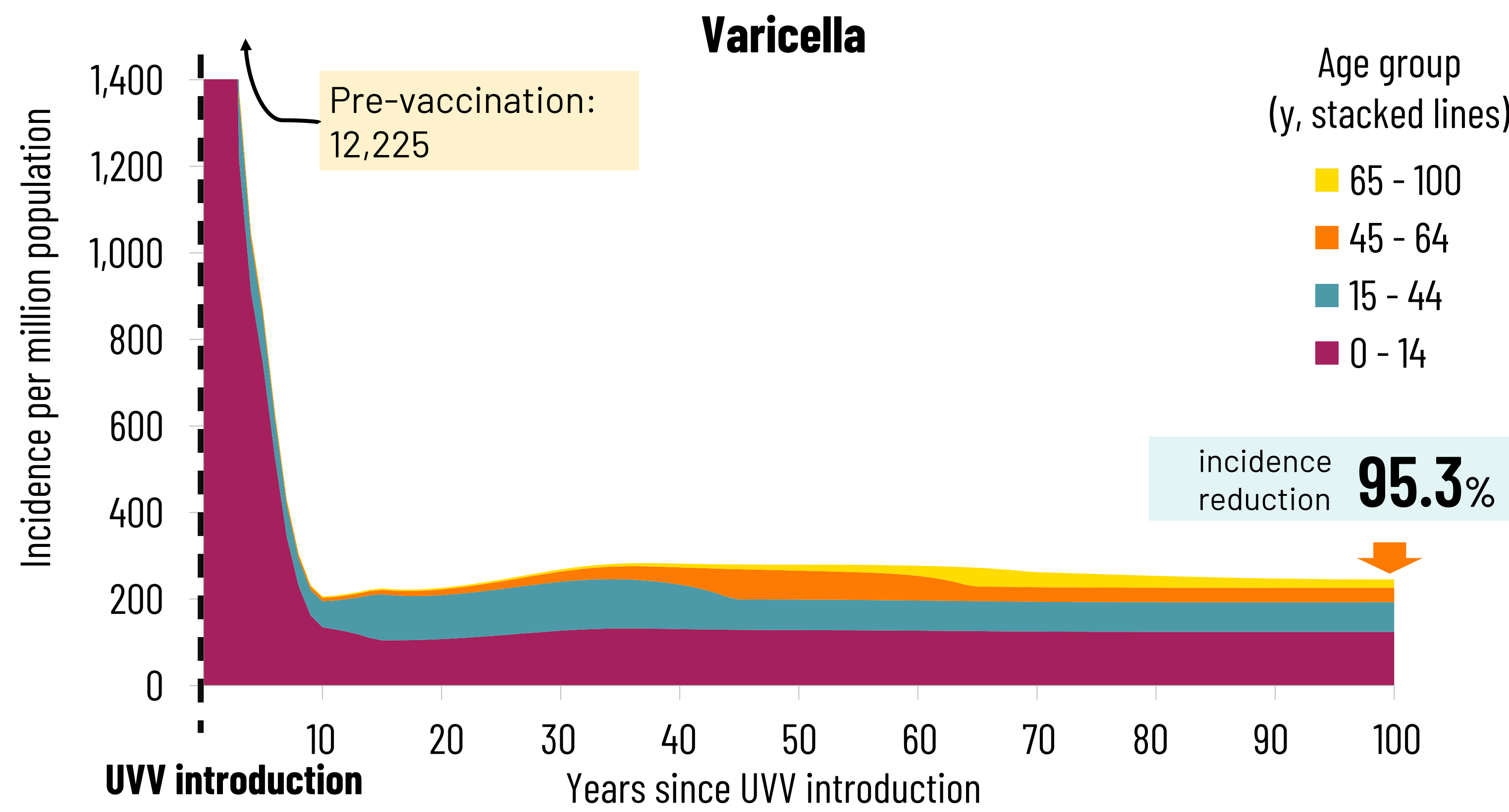
## Aims

This study aimed to **assess the public health impact** and **cost-effectiveness** of implementing **UVV** in addition to the existing HZ vaccination strategy in the **UK**.



## Results

Following UVV implementation, the **total incidence** of **varicella** and **HZ** was **reduced by 95%** (with the largest reductions occurring in the first five years) and **98%**, respectively, over 100 years in the base case.



**Abbreviations:** DTM dynamic transmission model, GP general practitioner, HZ herpes zoster, m month (of age), QALY quality adjusted life year, VZV varicella-zoster virus, UK United Kingdom, UVV universal varicella vaccination, y years (of age)



## Methods

**Model** age-structured deterministic compartmental DTM adapted from Akpo et al.<sup>4</sup>.

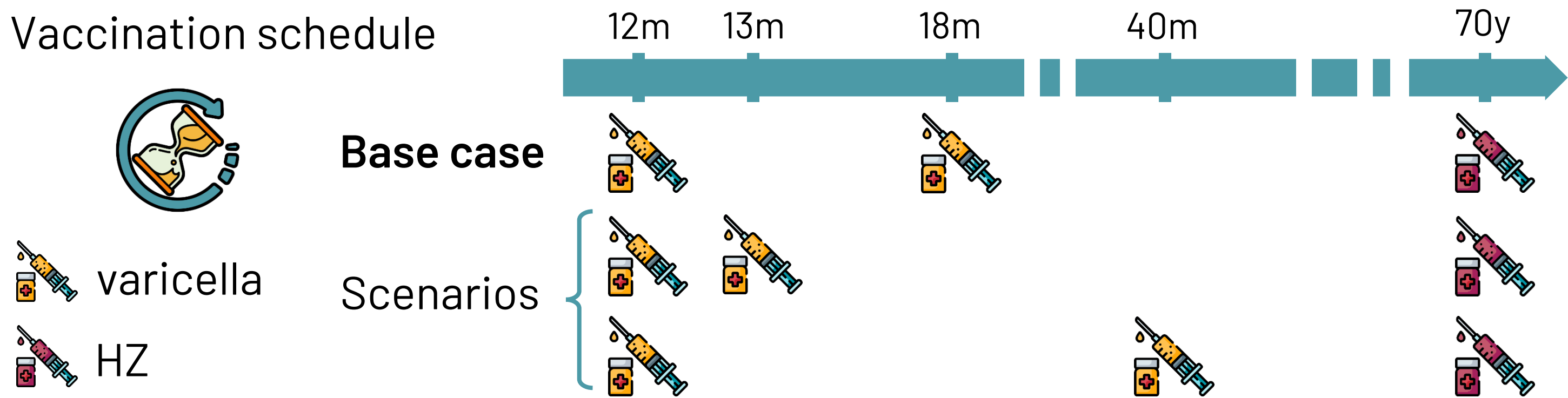
**Population** UK population (0-100 years).

**Intervention & comparator** varicella vaccination with HZ vaccination vs no varicella vaccination with HZ vaccination.

**Outcomes** direct costs, indirect costs, numbers of natural and breakthrough cases of varicella and HZ, and QALY losses.

**Perspective** 100-year time horizon with a payer and societal perspective (20-year and 50-year in scenario analyses).

**Discount rate** 3.5% applied on both costs and outcomes.



From a **payer's** perspective, the incremental cost-utility ratio was **£7,069** and **£2,984** per **QALY gained** in the base case using a 20-year and a 100-year time horizon, respectively. From a **societal** perspective, **UVV** was considered **cost saving**.

	20 years			100 years		
	UVV	No UVV	Difference	UVV	No UVV	Difference
<b>Total direct costs</b>	3,051,777	2,696,703	355,075	5,497,291	5,044,138	453,152
<b>Vaccine cost</b>	1,577,143	1,050,929	526,214	3,061,470	1,979,622	1,081,848
<b>Outpatient cost</b>	16,455	102,915	-86,460	20,663	193,858	-173,195
<b>Hospital cost</b>	1,458,180	1,542,859	-84,679	2,415,157	2,870,658	-455,501
<b>Total indirect cost</b>	2,991,345	5,841,718	-2,850,373	4,271,508	11,003,983	-6,732,475
<b>Total costs (direct+ indirect)</b>	6,043,122	8,538,421	-2,495,299	9,768,798	16,048,122	-6,279,323
<b>QALYs lost</b>	262,827	313,056	-50,229	431,643	583,504	-151,861
<b>Direct cost per QALY saved</b>	-	-	7.1	-	-	3.0
<b>Total cost per QALY saved</b>	-	-	Dominant	-	-	Dominant

**All costs are expressed in thousands £.** Direct costs include GP and specialist visit, hospitalization, vaccine acquisition and administration, indirect costs include productivity loss due to morbidity and due to premature mortality for both varicella and HZ.



Varying the timing of the second dose has **little impact on the annual incidence reduction**.



## Conclusions

- ✓ The introduction of UVV in the UK would result in reducing both varicella and HZ incidence.
- ✓ The implementation of such a vaccination program was found to be cost-effective from a payer's perspective (at £20K/QALY threshold) and cost-saving from a societal one.
- ✓ The latest evidence retrieved from real-world studies show that the introduction of UVV was not associated with an increase of HZ cases.<sup>5,6</sup> However further research is needed to apprehend the long-term evolution in HZ incidence after UVV implementation.

**Funding:** GlaxoSmithKline Biologicals SA (GSK study identifier: VEO-000191); **Acknowledgments:** Nikolaos Giannelos, Kinga Meszaros, Business & Decision Life Sciences c/o GSK (writer: Sarah Fico)

**Presenting author:** Nicolas Jamet, nicolas.x.jamet@gsk.com

ISPOR-EU | 06-09 November 2022 | Vienna, Austria and Virtual

© 2022 GSK