

<b><u>MA (EU) number</u></b>	<b><u>(Invented) name</u></b>	<b><u>Strength</u></b>	<b><u>Pharmaceutical Form</u></b>	<b><u>Route of Administration</u></b>	<b><u>Immediate Packaging</u></b>	<b><u>Content (concentration)</u></b>	<b><u>Pack size</u></b>
EU/1/18/1272/001	Shingrix	-- <sup>1</sup>	Powder and suspension for injection	Intramuscular use	powder: vial (glass) suspension: vial (glass)	Powder: 50 µg Suspension: 0.5 ml	1 vial (powder) + 1 vial (suspension)
EU/1/18/1272/002	Shingrix	-- <sup>1</sup>	Powder and suspension for injection	Intramuscular use	powder: vial (glass) suspension: vial (glass)	Powder: 50 µg Suspension: 0.5 ml	10 vials (powder) + 10 vials (suspension)

--<sup>1</sup> After reconstitution, 1 dose (0.5 ml) contains 50 micrograms of gE antigen<sup>1</sup> adjuvanted with AS01<sub>B</sub><sup>2</sup>.

<sup>1</sup> Varicella Zoster Virus (VZV) glycoprotein E (gE) produced by recombinant DNA technology in Chinese Hamster Ovarian (CHO) cells

<sup>2</sup> AS01<sub>B</sub> Adjuvant System is composed of the plant extract *Quillaja saponaria* Molina, fraction 21 (QS-21) (50 micrograms) and 3-O-desacyl-4'-monophosphoryl lipid A (MPL) from *Salmonella minnesota* (50 micrograms)