

<u>EU Number</u>	<u>Invented name</u>	<u>Strength</u>	<u>Pharmaceutical Form</u>	<u>Route of Administration</u>	<u>Packaging</u>	<u>Content</u>	<u>Package size</u>
EU/1/10/629/001	HUMENZA	-1-	Suspension and emulsion for emulsion for injection	Intramuscular use	Suspension (H1N1): vial (glass); Emulsion (adjuvant): vial (glass)	Suspension: 1.5 ml Emulsion: 4.5 ml	10 vials (suspension) + 10 vials (emulsion)

-1-

3.8 µg HA

After mixing, 1 dose (0.5 ml) contains:

Split Influenza virus, inactivated, containing antigen\* equivalent to:

A/California/7/2009 (H1N1)-like strain (NYMC X-179A)

3.8 micrograms HA\*\*

\* propagated in eggs

\*\* haemagglutinin

AF03 adjuvant composed of squalene (12.4 milligrams), sorbitan oleate (1.9 milligrams), polyoxyethylene cetostearyl ether (2.4 milligrams) and mannitol (2.3 milligrams)