

All influenza vaccines marketed in the UK for the 2025 to 2026 season

| Supplier | Product | Vaccine type | Age indications | Ovalbumin content micrograms/dose | Contact details |
|-----------------------|--------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------|------------------------------------------------------|-------------------------------------------------------------|--------------------|
| AstraZeneca UK Ltd | Fluenz® | Trivalent LAIV (live attenuated influenza vaccine) supplied as nasal spray suspension | From 24 months to less than 18 years of age | Less than 0.024 micrograms per 0.2 ml dose | 0845 139 0000 |
| Sanofi | Vaxigrip ¹ | TIVe (standard egg- grown trivalent influenza vaccine), split virion, inactivated | From 6 months | Equal to or less than 0.05 micrograms per 0.5 ml dose | 0800 854 430 |
| Viatris | Influvac [®] sub- unit TIV ▼ | TIVe (standard egg- grown trivalent influenza vaccine) surface antigen, inactivated | From 6 months | Equal to or less than 0.1 micrograms per 0.5 ml dose | 0800 358 7468 |
| CSL Seqirus UK | Cell-based Trivalent Influenza Vaccine Seqirus ▼ | TIVc (cell-based Trivalent influenza vaccine) surface antigen, inactivated | From 6 months | Egg-free | 0345 0093 804 |
| Sanofi | Supemtek TIVr ▼ ¹ | Supemtek TIV (trivalent influenza vaccine (recombinant, prepared in cell culture)) | From 18 years | None | 0800 854 430 |
| Sanofi | Efluelda TIV- HD ▼ ¹ | TIV-HD (High-dose egg- grown trivalent influenza vaccine), split virion, inactivated 60 micrograms HA/strain | From 60 years | Equal to or less than 1 microgram per 0.5ml dose | 0800 854 430 |
| CSL Seqirus UK | Adjuvanted Trivalent Influenza Vaccine Seqirus ▼ | aTIV (adjuvanted egg- grown Trivalent influenza vaccine) surface antigen, inactivated, adjuvanted with MF59C.1 | | Equal to or less than 1 microgram per 0.5ml dose | 0345 0093 804 |

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¹ Sanofi intends to supply trivalent formulations for the 2025 to 2026 season. Relevant license applications are under assessment and Medicines and Healthcare products Regulatory Agency (MHRA) approval is pending. Other potential formulations (quadrivalent) would have the same antigen content per strain, and ovalbumin content threshold (where relevant) as presented in the table.