

Vaccines

> Consultez la version française

Vaccines fall within the remit of ANSM's scope of action. They are medicinal products. The Agency is involved from the <u>clinical trial</u> phase. It then grants a national marketing <u>authorisation</u> (MA) according to the benefit-risk assessment. MAs may also be granted for the entire European Union by the European Commission, following recommendations from European Medicines Agency (EMA) ANSM continues to oversee <u>batch release procedures</u>, <u>surveillance</u> of safety of use, and control over <u>advertising</u> aimed at the public and healthcare professionals.