Overseeing health product marketing

Consultez la version française

When a medicine or biological product is marketed in France, it will have undergone an assessment and been granted a marketing authorisation by ANSM or by the European Commission following a review by the European Medicines Agency.

Medical devices and in vitro diagnostic medical devices are marketed within a European framework which provides for the affixing of a CE mark certifying the compliance of the product with safety requirements. ANSM is involved in market surveillance and control in France.

Our role is to ensure that every patient treated receives products whose quality, safety, and efficacy have been demonstrated and validated.