



A guide to using

Beyfortus[®] is indicated for the prevention of respiratory syncytial virus (RSV) lower respiratory tract disease in:¹

- Neonates and infants during their first RSV season.
 - Children up to 24 months of age who remain vulnerable to severe RSV disease through their second RSV season.
- Beyfortus[®] should be used in accordance with official recommendations.¹

How to use Beyfortus[®]

First RSV season:

For neonates and infants based on **body weight at the time of dosing**¹



Weight **<5 kg**

1 x 50 mg dose



Weight **≥5 kg**

1 x 100 mg dose

Second RSV season:

For children who remain **vulnerable to severe RSV disease**, up to 24 months of age¹



Regardless of body weight

200 mg administered as
2 x 100 mg in a single appointment

Beyfortus[®] is for intramuscular injection only, preferably in the anterolateral aspect of the thigh.¹ The gluteal muscle should not be used routinely as an injection site because of the risk of damage to the sciatic nerve.¹

Consult the Summary of Product Characteristics prior to prescribing for full dosing and administration information.

Timing of administration¹

For infants during their first RSV season¹: Beyfortus[®] should be administered from birth for infants born during the RSV season. For others born outside the season, Beyfortus[®] should be administered ideally prior to the RSV season.

For children vulnerable to severe RSV disease through their second season¹: Beyfortus[®] should be administered ideally prior to the start of the second RSV season.

How to store Beyfortus[®]

Storage

- Store in a refrigerator (2°C–8°C)
- Do not freeze
- Do not shake or expose to direct heat
- Keep the pre-filled syringe in the outer carton in order to protect from light



Shelf life

- Beyfortus[®] has a shelf life of 4 years
- Beyfortus[®] may be kept at room temperature (20°C–25°C) when protected from light for a maximum of 48 hours. After this time, the syringe must be discarded



Click on the link to Sanofi's promotional website to learn more about Beyfortus[®]

<https://pro.campus.sanofi/uk/products/beyfortus-nirsevimab>

Adverse events should be reported. Reporting forms and information can be found at www.mhra.gov.uk/yellowcard or search for MHRA Yellow Card in the Google Play or Apple App Store. Adverse events should also be reported to the Sanofi drug safety department on Tel: 0800 0902 314. Alternatively, send via email to UK-drugsafety@sanofi.com.