

## Casirivimab and Imdevimab (also known as REGEN-COV™) Co-Packaged Product Quick Reference Guide

### Information for Temporary Alternative Packaging of REGEN-COV under Emergency Use Authorization (EUA)

Scan the QR code or go to [www.REGENCOV.com](http://www.REGENCOV.com) for the FDA-authorized Fact Sheets (one for the Healthcare Provider and one for the Patient/Caregiver) for current product information for casirivimab and imdevimab.



#### It is important to note:

- Each co-packaged carton contains individual antibody solutions in separate vials as follows:
  - One (1) vial containing casirivimab; 300 mg/2.5 mL (120 mg/mL) or 1,332 mg/11.1 mL (120 mg/mL)
  - One (1) vial containing imdevimab; 300 mg/2.5 mL (120 mg/mL) or 1,332 mg/11.1 mL (120 mg/mL)
  - One (1) package leaflet which is not approved for use in the US. **This leaflet should be discarded.**  
Please refer to the authorized EUA Fact Sheets **only** (scan the QR code above).
- **The carton is labeled as “casirivimab and imdevimab 120 mg/mL concentrate for solution for infusion”.** Do not confuse this co-packaged carton with REGEN-COV (casirivimab and imdevimab) co-formulated solution.
- **The vials in the co-packaged carton may be used to prepare and administer intravenous infusions as well as subcutaneous injections** despite having the statements such as “Concentrate for solution for infusion” or “For intravenous infusion after dilution”.
- Carton and vial labels do not include an NDC. Use the NDC listed below based on the package.

Co-Packaged Carton Contents	Co-Packaged Components	Concentration	Co-Packaged Carton NDC Number
2 Vials	1 vial of casirivimab (NDC 61755-024-01)	1,332 mg/11.1 mL (120 mg/mL)	61755-042-02
	1 vial of imdevimab (NDC 61755-025-01)	1,332 mg/11.1 mL (120 mg/mL)	
2 Vials	1 vial of casirivimab (NDC 61755-026-01)	300 mg/2.5 mL (120 mg/mL)	61755-045-02
	1 vial of imdevimab (NDC 61755-027-01)	300 mg/2.5 mL (120 mg/mL)	

- **The barcode on the co-packaged carton label may not register with U.S. scanning systems.** Confirm that the barcode provides correct information when scanned and if not, consider manually inputting the product specific information into electronic systems. There is no barcode on the co-packaged vial.
- Roche manufactures the co-packaged product on behalf of Regeneron and is listed on the package.
- Refer to the “Dear Healthcare Provider Letter” for additional information on the co-packaged product.

Health care providers must submit a report on **ALL MEDICATION ERRORS and ALL SERIOUS ADVERSE EVENTS** potentially related to REGEN-COV (casirivimab and imdevimab). See the FDA Fact Sheet for Health Care Providers (Sections 8 and 9 of the Full EUA Prescribing Information) for reporting instructions.

If you have questions, please contact Regeneron at 1-844-734-6643.

**REGENERON**

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