**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 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 61755-024-00) | 1,332 mg/11.1 mL (120 mg/mL) | 61755-042-02 |
| | **1 vial of imdevimab** (NDC 61755-025-00) | 1,332 mg/11.1 mL (120 mg/mL) | |
| **2 Vials** | **1 vial of casirivimab** (NDC 61755-026-00) | 300 mg/2.5 mL (120 mg/mL) | 61755-045-02 |
| | **1 vial of imdevimab** (NDC 61755-027-00) | 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

Manufactured for:
Regeneron Pharmaceuticals, Inc.
777 Old Saw Mill River Road
Tarrytown, NY 10591-6707