New National Opioids Settlement: Secondary Manufacturers
Opioids Implementation Administrator
opioidsparticipation@rubris.com

Montgomery city, TX
Reference Number: CL-1773885

TO LOCAL POLITICAL SUBDIVISIONS:

THIS PACKAGE CONTAINS DOCUMENTATION TO PARTICIPATE IN THE NEW NATIONAL OPIOIDS SECONDARY MANUFACTURERS SETTLEMENTS. YOU MUST TAKE ACTION IN ORDER TO PARTICIPATE.

Deadline: October 8, 2025

A new proposed national opioids settlement ("*Secondary Manufacturers Settlements*") has been reached with eight opioids manufacturers: Alvogen, Amneal, Apotex, Hikma, Indivior, Mylan, Sun, and Zydus ("Settling Defendants"). This *Combined Participation Package* is a follow-up communication to the *Notice of National Opioids Settlement* recently received electronically by your subdivision.

You are receiving this *Combined Participation Package* because Texas is participating in the Secondary Manufacturers Settlements.

If a state is not eligible to or does not participate in the settlement with a particular manufacturer, the subdivisions in that state are not eligible to participate in that manufacturer's settlement.

This electronic envelope contains:

- A *Combined Participation Form* for the *Secondary Manufacturers Settlements* that your subdivision is eligible to join, including a release of any claims.

The *Combined Participation Form* must be executed, without alteration, and submitted on or before October 8, 2025, in order for your subdivision to be considered for initial participation calculations and payment eligibility under the *Secondary Manufacturers Settlement*.

Based upon *Combined Participation Forms* received on or before October 8, 2025, the subdivision participation rate will be used to determine whether participation is sufficient for each settlement to move forward and whether a state earns its maximum potential payment under each settlement. If a settlement moves forward, your release will become effective. If a settlement does not move forward, that release will not become effective.

Any subdivision that does not participate cannot directly share in the settlement funds, even if the subdivision's state is settling and other participating subdivisions are sharing in settlement funds. Any subdivision that does not participate may also

reduce the amount of money for programs to remediate the opioid crisis in its state. Please note, a subdivision will not necessarily directly receive settlement funds by participating; decisions on how settlement funds will be allocated within a state are subject to intrastate agreements or state statutes.

You are encouraged to discuss the terms and benefits of the *Secondary Manufacturers Settlements* with your counsel, your Attorney General's Office, and other contacts within your state. Many states are implementing and allocating funds for this new settlement the same as they did for the prior opioids settlements but states may choose to treat this settlement differently.

Information and documents regarding the *Secondary Manufacturers Settlements*, implementation in your state, and how funds will be allocated within your state can be found on the national settlement website at https://nationalopioidsettlement.com/. This website will be supplemented as additional documents are created. You may also visit the Texas Attorney General's Office website at https://www.texasattorneygeneral.gov/globalopioidsettlement for information.

This *Participation Packet* is different than the participation packet you recently received from Rubris concerning a settlement with Purdue Pharma, L.P, and the Sackler Family. The *Secondary Manufacturers Settlements* discussed in this *Participation Packet* are different than the settlement with Purdue and the Sacklers, and you may participate in the *Secondary Manufacturers Settlements* regardless of whether you join the Purdue and Sackler settlement.

How to return signed forms:

Please note that the Texas Attorney General's Office is collecting the executed *Participation Form* differently from prior opioid settlements. There are three methods for returning the executed *Combined Participation Form* and any supporting documentation to the Implementation Administrator:

(1) *Electronic Signature via DocuSign*: Executing the *Combined Participation Form* electronically through DocuSign will return the signed form to the Implementation Administrator and associate your form with your subdivision's records. Electronic signature is the most efficient method for returning the *Combined Participation Form*, allowing for more timely participation and the potential to meet higher settlement payment thresholds, and is therefore strongly encouraged.

(2) *Manual Signature returned via DocuSign*: DocuSign allows forms to be downloaded, signed manually, then uploaded to DocuSign and returned automatically to the Implementation Administrator. Please be sure to complete all fields. As with electronic signature, returning a manually signed *Combined Participation Form* via DocuSign will associate your signed forms with your subdivision's records.

(3) *Manual Signature returned via electronic mail*: If your subdivision is unable to return an executed *Combined Participation Form* using DocuSign, the signed *Combined Participation Form* may be returned via electronic mail to opioidsparticipation@rubris.com. Please include the name, state, and reference ID of your subdivision in the body of the email and use the subject line Combined Settlement Participation Form – [Subdivision Name, Subdivision State] – [Reference ID].

Detailed instructions on how to sign and return the *Combined Participation Form*, including changing the authorized signer, can be found at https://nationalopioidsettlement.com/additional-settlements/. You may also contact opioidsparticipation@rubris.com and/or opioids@oag.texas.gov if you have any questions.

The sign-on period for subdivisions ends on October 8, 2025.

If you have any questions about executing the *Combined Participation Form*, please contact your counsel, the Implementation Administrator at opioidsparticipation@rubris.com, or the Texas Attorney General's Office at opioids@oag.texas.gov.

Thank you,

Secondary Manufacturers Settlements Implementation Administrator

The Implementation Administrator is retained to provide the settlement notice required by the Secondary Manufacturers Settlements and to manage the collection of the Combined Participation Form.

EXHIBIT K

Secondary Manufacturers' Combined Subdivision Participation and Release Form ("Combined Participation Form")

Governmental Entity: Montgomery city		State: TX
Authorized Official: Sara Countryman, Mayor		
Address 1: 101 Old Plantersville Rd		
Address 2:		
City, State, Zip: Montgomery	Texas	77316
Phone: 936-597-6434		
Email: scountryman@co.montgomery.tx.us		

The governmental entity identified above ("*Governmental Entity*"), in order to obtain and in consideration for the benefits provided to the Governmental Entity pursuant to each of the settlements which are listed in paragraph 1 below (each a "Secondary Manufacturer's Settlement" and collectively, "the Secondary Manufacturers' Settlements"), and acting through the undersigned authorized official, hereby elects to participate in each of the Secondary Manufacturers' Settlements, release all Released Claims against all Released Entities in each of the Secondary Manufacturers' Settlements, and agrees as follows.

1. The Participating Entity hereby elects to participate in each of the following Secondary Manufacturers' Settlements as a Participating Entity:

   a. Settlement Agreement for Alvogen, Inc. dated April 4, 2025.
   b. Settlement Agreement for Apotex Corp. dated April 4, 2025.
   c. Settlement Agreement for Amneal Pharmaceuticals LLC dated April 4, 2025.
   d. Settlement Agreement for Hikma Pharmaceuticals USA Inc. dated April 4, 2025.
   e. Settlement Agreement for Indivior Inc. dated April 4, 2025.
   f. Settlement Agreement for Viatris Inc. ("Mylan") dated April 4, 2025.
   g. Settlement Agreement for Sun Pharmaceutical Industries, Inc. dated April 4, 2025.
   h. Settlement Agreement for Zydus Pharmaceuticals (USA) Inc. dated April 4, 2025.

2. The Governmental Entity is aware of and has reviewed each of the Secondary Manufacturers' Settlements, understands that all capitalized terms not defined in this Combined Participation Form have the meanings defined in each of the Secondary Manufacturers' Settlements, and agrees that by executing this Combined Participation Form, the Governmental Entity elects to participate in each of the Secondary Manufacturers' Settlements and become a Participating Subdivision as provided in each of the Secondary Manufacturers' Settlements.

3. The Governmental Entity shall promptly, and in any event no later than 14 days after the Reference Date and prior to the filing of the Consent Judgment, dismiss with prejudice any Released Claims that it has filed against any Released Entity in each of the Secondary Manufacturers' Settlements. With respect to any Released Claims pending in *In re National Prescription Opiate Litigation*, MDL No. 2804, the Governmental Entity

authorizes the Plaintiffs' Executive Committee to execute and file on behalf of the Governmental Entity a Stipulation of Dismissal with Prejudice for each of the manufacturers listed in paragraph 1 above substantially in the form found at *https://nationalopioidsettlement.com/additional-settlements/.*

4. The Governmental Entity agrees to the terms of each of the Secondary Manufacturers' Settlements pertaining to Participating Subdivisions as defined therein.

5. By agreeing to the terms of each of the Secondary Manufacturers' Settlements and becoming a Releasor, the Governmental Entity is entitled to the benefits provided therein, including, if applicable, monetary payments beginning after the Effective Date.

6. The Governmental Entity agrees to use any monies it receives through each of the Secondary Manufacturers' Settlements solely for the purposes provided therein.

7. The Governmental Entity submits to the jurisdiction of the court and agrees to follow the process for resolving any disputes related to each Secondary Manufacturer's Settlement as described in each of the Secondary Manufacturers' Settlements.[1]

8. The Governmental Entity has the right to enforce each of the Secondary Manufacturers' Settlements as provided therein.

9. The Governmental Entity, as a Participating Subdivision, hereby becomes a Releasor for all purposes in each of the Secondary Manufacturers' Settlements, including without limitation all provisions related to release of any claims,[2] and along with all departments, agencies, divisions, boards, commissions, districts, instrumentalities of any kind and attorneys, and any person in his or her official capacity whether elected or appointed to serve any of the foregoing and any agency, person, or other entity claiming by or through any of the foregoing, and any other entity identified in the definition of Releasor, provides for a release to the fullest extent of its authority. As a Releasor, the Governmental Entity hereby absolutely, unconditionally, and irrevocably covenants not to bring, file, or claim, or to cause, assist or permit to be brought, filed, or claimed, or to otherwise seek to establish liability for any Released Claims against any Released Entity in each of the Secondary Manufacturers' Settlements in any forum whatsoever. The releases provided for in each of the Secondary Manufacturers' Settlements are intended by the Parties to be broad and shall be interpreted so as to give the Released Entities in each of the Secondary Manufacturers' Settlements the broadest possible bar against any liability relating in any way to Released

[1] *See* Settlement Agreement for Alvogen, Inc. Section VII.F.2; Settlement Agreement for Apotex Corp. Section VII.F.2; Settlement Agreement for Amneal Pharmaceuticals LLC Section VII.F.2; Settlement Agreement for Hikma Pharmaceuticals USA Inc. Section VII.F.2; Settlement Agreement for Indivior Section VI.F.2; Settlement Agreement for Mylan Section VI.F.2; Settlement Agreement for Sun Pharmaceutical Industries, Inc. Section VII.F.2; Settlement Agreement for Zydus Pharmaceuticals (USA) Inc. Section VII.F.2.

[2] *See* Settlement Agreement for Alvogen, Inc. Section XI; Settlement Agreement for Amneal Pharmaceuticals LLC Section X; Settlement Agreement for Apotex Corp. Section XI; Settlement Agreement for Hikma Pharmaceuticals USA Inc. Section XI; Settlement Agreement for Indivior Section X; Settlement Agreement for Mylan Section X; Settlement Agreement for Sun Pharmaceutical Industries, Inc. Section XI; Settlement Agreement for Zydus Pharmaceuticals (USA) Inc. Section XI.

Claims and extend to the full extent of the power of the Governmental Entity to release claims. Each of the Secondary Manufacturers' Settlements shall be a complete bar to any Released Claim against that manufacturer's Released Entities.

10. The Governmental Entity hereby takes on all rights and obligations of a Participating Subdivision as set forth in each of the Secondary Manufacturers' Settlements.

11. In connection with the releases provided for in each of the Secondary Manufacturers' Settlements, each Governmental Entity expressly waives, releases, and forever discharges any and all provisions, rights, and benefits conferred by any law of any state or territory of the United States or other jurisdiction, or principle of common law, which is similar, comparable, or equivalent to § 1542 of the California Civil Code, which reads:

> **General Release; extent.** A general release does not extend to claims that the creditor or releasing party does not know or suspect to exist in his or her favor at the time of executing the release that, if known by him or her would have materially affected his or her settlement with the debtor or released party.

A Releasor may hereafter discover facts other than or different from those which it knows, believes, or assumes to be true with respect to the Released Claims in each of the Secondary Manufacturers' Settlements, but each Governmental Entity hereby expressly waives and fully, finally, and forever settles, releases and discharges, upon the Effective Date, any and all Released Claims that may exist as of such date but which Releasors do not know or suspect to exist, whether through ignorance, oversight, error, negligence or through no fault whatsoever, and which, if known, would materially affect the Governmental Entities' decision to participate in each of the Secondary Manufacturers' Settlements.

12. The Governmental Entity understands and acknowledges that each of the Secondary Manufacturers' Settlements is an independent agreement with its own terms and conditions. Nothing herein is intended to modify in any way the terms of any of the Secondary Manufacturers' Settlements, to which Governmental Entity hereby agrees, aside from the exceptions in paragraph 13 below. To the extent this Combined Participation Form is interpreted differently from any of the Secondary Manufacturers' Settlements in any respect, the individual Secondary Manufacturer's Settlement controls.

13. For the avoidance of doubt, in the event that some but not all of the Secondary Manufacturers' Settlements proceed past their respective Reference Dates, all releases and other commitments or obligations shall become void ***only as to*** those Secondary Manufacturers' Settlements that fail to proceed past their Reference Dates. All releases and other commitments or obligations (including those contained in this Combined Participation Form) shall remain in full effect as to each Secondary Manufacturer's Settlement that proceeds past its Reference Date, and this Combined Participation Form need not be modified, returned, or destroyed as long as any Secondary Manufacturer's Settlement proceeds past its Reference Date.

I have all necessary power and authorization to execute this Combined Participation Form on behalf of the Governmental Entity.

Signature: _____

Name: _____

Title: _____

Date: _____