

**UNITED STATES DISTRICT COURT
FOR THE WESTERN DISTRICT OF LOUISIANA
LAFAYETTE DIVISION**

THE STATE OF LOUISIANA, by and
through its Attorney General, LIZ
MURRILL, and ROSALIE MARKEZICH,

Plaintiffs,

v.

U.S. FOOD AND DRUG
ADMINISTRATION, et al.,

Defendants.

Case No. 6:25-cv-01491

Judge David C. Joseph

Magistrate Judge David J. Ayo

**BRIEF FOR FORMER U.S. DEPARTMENT OF JUSTICE OFFICIALS AS
AMICI CURIAE IN SUPPORT OF DEFENDANTS**

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INTEREST OF AMICI CURIAE¹

Amici are 15 former high-ranking U.S. Department of Justice officials who served as Deputy Attorney General, Solicitor General, Assistant Attorney General, and/or U.S. Attorney. Amici were responsible for enforcing federal criminal laws, including the Comstock laws, 18 U.S.C. §§ 1461-1462, and represented the United States in criminal matters in all levels of the judiciary around the country. A full list of amici appears in the Appendix.

Amici hold diverse views regarding the moral and jurisprudential questions surrounding abortion but agree that Plaintiffs' argument that the 2023 Risk Evaluation and Mitigation Strategy ("REMS") for mifepristone violates the Comstock laws is erroneous. The Food & Drug Administration ("FDA") was not authorized to consider, interpret, or apply federal criminal laws as part of its drug-approval process, including its 2023 REMS decision. Further, Plaintiffs gravely misinterpret the Comstock laws, expanding their scope beyond Congress's intent. Plaintiffs' misreading of the Comstock laws fails to account for the holdings of all four courts of appeals to have addressed the question and the construction embraced by the U.S. Justice Department, the sole agency responsible for prosecuting violations of the Comstock laws. The Court need not address Plaintiffs' misinterpretation of the Comstock laws at all, but if it does so, amici urge the Court to reject Plaintiffs' strained interpretation.

INTRODUCTION AND SUMMARY OF ARGUMENT

Plaintiffs' argument that FDA's 2023 actions violated the Comstock laws is erroneous. First, the Comstock laws provide no basis to enjoin or invalidate¹ the challenged FDA actions. Regardless, Plaintiffs' interpretation of the Comstock laws ignores the decades-long

¹ No counsel for a party authored this brief in whole or in part, and no entity or person other than amici and their counsel made a monetary contribution to fund the preparation or submission of this brief.

understanding of the Comstock laws shared by Congress, the Judiciary, and the Executive Branch.

As a threshold matter, the Comstock laws are irrelevant to the validity of the challenged FDA actions. Congress charged FDA with determining whether a drug is safe and effective and set forth specific factors the agency is required to consider when imposing a REMS. Those factors do not include consideration, application, or interpretation of any criminal laws. Congress did not authorize FDA to limit access to safe and effective medication based on its interpretation and application of the Comstock laws. Accordingly, FDA's determinations did not purport to address whether distributions of the drug were or were not lawful under the Comstock laws (just as they did not purport to declare whether such distributions were or were not lawful under other criminal statutes). The Comstock laws are therefore irrelevant to the validity of the challenged FDA actions.

But even if the Comstock laws were relevant to FDA's actions, those actions would be valid because they accord with those statutes. Plaintiffs misinterpret the Comstock laws to reach items intended to produce both lawful and unlawful abortions. As four circuits concluded in decisions issued between 1915 and 1944—decisions resting on lengthy statutory analyses—the Comstock laws reach the distribution of items only if intended to produce unlawful abortions. This unanimous interpretation is the only one that both makes sense of all the Comstock laws—not just 18 U.S.C. §§ 1461-1462 but also 19 U.S.C. § 1305—and also avoids absurd and likely unconstitutional implications. And critically, this interpretation was acknowledged by Congress in 1948 and (if more were needed) further ratified repeatedly since then by Congress's reenactments and amendments of the Comstock laws without relevant alteration.

Adopting Plaintiffs’ erroneous interpretation of the Comstock laws would risk a profound destabilization of medical care. Plaintiffs’ construction of the Comstock laws cannot be limited to telemedicine abortion or mifepristone; it necessarily would affect in-person abortion and a broad range of other kinds of healthcare. That is because mifepristone and other drugs and items used for in-person abortion are likewise delivered through the mails. And many abortion-inducing drugs and devices are used for non-abortion-related care. For instance, mifepristone is regularly used to treat miscarriages, and misoprostol is frequently used during labor and delivery. This Court should reject Plaintiffs’ broad-ranging interpretation.

ARGUMENT

I. The Comstock Laws Are Irrelevant To The Validity Of The Challenged FDA Actions

The Comstock laws, however interpreted, are irrelevant here. FDA’s actions were within the bounds of its legal authority to assess, pursuant to specific statutorily enumerated factors, whether mifepristone would be safe and effective under specified conditions. Indeed, FDA would not be authorized to consider the Comstock laws in deciding whether to approve mifepristone and what restrictions to apply to it.

A. FDA Had No Power Or Duty To Consider The Comstock Laws In Deciding Whether Or With What Use Restrictions To Approve Mifepristone

As is true with any agency, FDA’s “power to regulate . . . must always be grounded in a valid grant of authority from Congress.” *FDA v. Brown & Williamson Tobacco Corporation*, 529 U.S. 120, 126, 161 (2000). In terms of approving a drug for use in the United States, Congress specified that FDA’s role is to assess whether the drug is safe and effective for the indicated use. If FDA determines the drug *is* safe and effective, it must approve the drug. This approval serves only to remove one particular legal barrier to the drug’s distribution, namely the bar on distribution of drugs not approved by FDA as safe and effective. Whether any *other* laws

restrict or prohibit distribution of a drug is a “factor[] which Congress has not intended [the agency] to consider.” *Motor Vehicle Manufacturers Association v. State Farm Mutual Automobile Insurance Company*, 463 U.S. 29, 43 (1983).

The Food, Drug, and Cosmetic Act (“FDCA”) requires that FDA, in deciding whether to approve a drug application, consider whether the drug will be safe and effective under the conditions of use described in the proposed label. *See* 21 U.S.C. § 355(b)(1)(A)(i), (d); *Merck Sharp & Dohme Corporation v. Albrecht*, 587 U.S. 299, 302 (2019). The statute specifies seven “grounds for refusing [a drug] application,” five relating to safety and efficacy, one requiring the filing of patent information, and one relating to the label’s accuracy. *See* 21 U.S.C. § 355(d). The FDCA gives FDA no authority to deny a drug application for any other reason. To the contrary, the FDCA commands that, if none of the patent-filing or safety-and-efficacy grounds for denial is present, FDA “shall” approve the application. *Id.*; *see also* 21 C.F.R. §§ 314.105, 314.125. Therefore, FDA cannot deny a drug application based on any potential restrictions on distribution imposed by the Comstock laws (or any other law FDA does not administer).

FDA’s framework for REMS is similarly focused on safety and efficacy as well as assuring patient access. The FDCA requires an applicant to propose a REMS if FDA determines one “is necessary to ensure that the benefits of the drug outweigh the risks.” 21 U.S.C. § 355-1(a)(1); *see also id.* § 355-1(a)(2), (b)(1), (4)(5). Accordingly, a REMS must identify means to mitigate risks to patients’ health. *See id.* § 355-1(c), (e)-(f). Nothing authorizes FDA to consider the Comstock laws (or, again, any other law FDA does not administer) in the REMS process. Nor has Congress expected FDA to have done so. Indeed, although members of Congress have engaged in exacting oversight of FDA’s approval of mifepristone, including via initiating U.S. Government Accountability Office investigations and subcommittee hearings, they have not

questioned FDA approval on the grounds that the agency failed to consider the Comstock laws. See GAO-08-751, *FDA: Approval and Oversight of the Drug Mifepristone* at 1 (August 2008) (identifying Congressional requesters); GAO-18-292, *FDA: Information on Mifepristone Labeling Changes and Ongoing Monitoring Efforts* at 28 (March 2018) (same).

That FDA did not consider the Comstock laws in approving mifepristone or in the REMS process is therefore entirely unsurprising. Indeed, it would be not only unlawful but also entirely impractical for FDA to catalog and evaluate the countless laws it does *not* administer or have expertise on but that nonetheless might apply to the drugs it reviews—including continually monitoring changes in such laws and reevaluating its prior decisions in light of those changes.

FDA’s approval, moreover, is merely a necessary condition for introducing a drug into interstate commerce; the FDCA provides that “[n]o person shall introduce . . . into interstate commerce any new drug, unless” FDA has approved the drug. 21 U.S.C. § 355(a); *see also, e.g., Mutual Pharmaceutical Company v. Bartlett*, 570 U.S. 472, 476 (2013) (“drug manufacturers must gain approval from the [FDA] before marketing any drug in interstate commerce”). But FDA approval means nothing with respect to the applicability of federal laws outside FDA’s purview; FDA’s approval (and REMS decisions) do not purport to override such laws. In fact, FDA routinely approves drugs that are restricted by laws FDA does not administer. One such law is the Controlled Substances Act, which is enforced by the U.S. Department of Justice and criminalizes the distribution, dispensing, and possession of many FDA-approved substances, such as fentanyl and methadone. *See* 21 U.S.C. §§ 811(a), 812, 823, 841(a)(1), 844(a).

Plaintiffs argue that the “Administrative Procedure Act requires federal courts to set aside federal agency action that is ‘not in accordance with law’—which means, of course, *any* law, and not merely those laws that the agency itself is charged with administering.” Pls.’ Mem. Supp.

Mot. Prelim. Inj. (ECF No. 20-26) at 14 (quoting *FCC v. NextWave Personal Communications, Inc.*, 537 U.S. 293, 300 (2003) (quoting 5 U.S.C. § 706(2)(A))). Plaintiffs’ reliance on *NextWave* is misplaced. In *NextWave*, the Court held that the Federal Communications Commission violated a law directly applicable to it when it revoked a broadband license in violation of a Bankruptcy Code provision specifically prohibiting such revocations by any “governmental unit.” 537 U.S. 300-301. By contrast, Plaintiffs do not and cannot suggest that FDA itself somehow violated the Comstock laws’ provisions governing illegal mailing when it approved the REMS. Again, the Comstock laws do not govern FDA’s drug-approval and REMS decision making; they govern the distribution of some abortion-producing items.

B. FDA’s Actions Do Not Purport To Declare Any Distributions Of Mifepristone Lawful Under The Comstock Laws (However Interpreted)

FDA’s 2023 REMS conform to FDA’s limited statutory authority. In adopting the REMS, FDA explained that “[u]nder the Mifepristone REMS Program, mifepristone may be dispensed in person or by mail.” *See* Information about Mifepristone for Medical Termination of Pregnancy Through Ten Weeks Gestation, Food & Drug Admin. (Jan. 30, 2025); *see also* Approved Risk Evaluation and Mitigation Strategies (REMS), Mifepristone, Food & Drug Admin. (Sept. 30, 2025). In doing so, FDA expressed nothing more than its determination that such distribution would not undermine mifepristone’s safety or efficacy. That is, as explained, the only consideration FDA may assess. The agency certainly did not purport to opine on whether any particular distributions of mifepristone are lawful under the Comstock laws.

II. FDA’s Actions Accord With The Comstock Laws Because The Comstock Laws Reach Only Distributions Intended For Unlawful Abortion

Because FDA’s authority in approving medications does not encompass interpreting or applying federal criminal statutes like the Comstock laws, the proper interpretation of those laws has no bearing on the validity of the agency’s challenged actions. But even if the Comstock laws

were relevant here, Plaintiffs’ interpretation—that 18 U.S.C. §§ 1461-1462 prohibit distribution of items intended to produce not only *unlawful* abortions but also *lawful* ones—is incorrect. For more than a century, the Comstock laws have been understood—by Congress, the Judiciary, and the Executive Branch—not to prohibit sending or receiving abortion-related items where an individual lacks the intent that the items be used for unlawful abortion. Indeed, as every circuit court to address the question has recognized, Plaintiffs’ reading is absurd and raises serious constitutional concerns, especially given the interaction between §§ 1461-1462 and § 1305. Critically, Congress’s 1948 reenactment of the Comstock laws explicitly took note of those judicial decisions and specifically adopted the courts’ interpretation. Congress then repeatedly ratified that interpretation. And for nearly a century, the Executive Branch has recognized this settled meaning too, including in a 2022 opinion issued by the Department of Justice’s Office of Legal Counsel reaffirming this construction.

A. Congress Enacted §§ 1461-1462 Specifically Intending That They Be Interpreted To Reach Items Only If Intended For Unlawful Abortion

The Comstock laws were enacted in the late 1800s. *See* Act of Mar. 3, 1873, ch. 258, § 2, 17 Stat. 598, 599; Act of Feb. 8, 1897, ch. 172, 29 Stat. 512. In 1909, Congress revised the Comstock laws to substantially the language found today at 18 U.S.C. §§ 1461-1462. One section—what is now § 1461—prohibited “knowingly deposit[ing]” in the mails “every article or thing designed, adapted, or intended for preventing conception or producing abortion, or for any indecent or immoral use.” Pub. L. No. 60-350, § 211, 35 Stat. 1088, 1129 (1909). Another provision—what is now § 1462—prohibited “bring[ing] . . . into the United States” and “knowingly deposit[ing] . . . with any express company or other common carrier for [interstate] carriage . . . any drug, medicine, article, or thing designed, adapted, or intended for preventing conception, or producing abortion, or for any indecent or immoral use[.]” *Id.* § 245, 35 Stat. at

1138. While those provisions speak plainly to reach mailings for “*any* indecent or immoral purpose,” § 1461 (emphasis added); *accord* § 1462, they contain no similar language suggesting they reach items related to *any* abortion or *any* contraception. Indeed, the Act referred to “producing abortion” or “procuring abortion,” which then had a specialized criminal-law meaning and referred not to all abortions, but specifically to the particular crime of producing or procuring an abortion that violated the law, either because it occurred after detectable fetal movement, known at the time as “quickening,” or because it was not therapeutic.²

Between 1915 and 1944, four federal circuit courts issued six decisions interpreting this statutory language. Each one rejected the proposition that these provisions reached all items for preventing conception and producing abortion regardless of the intended circumstances of their use. The Seventh Circuit, for example, explained that it was not “reasonable” to suppose Congress intended “the statute [to] cover all acts of abortion.” *Bours v. United States*, 229 F. 960, 964-965 (7th Cir. 1915). The Second Circuit likewise observed that “[i]t would seem reasonable” to interpret the statute “as requiring an intent on the part of the sender that the article mailed or shipped by common carrier be used for *illegal* contraception or abortion or for indecent or immoral purposes.” *Youngs Rubber Corporation v. C.I. Lee & Company*, 45 F.2d 103, 107-108 (2d Cir. 1930) (emphasis added); *accord United States v. Nicholas*, 97 F.2d 510, 512 (2d Cir. 1938); *United States v. One Package*, 86 F.2d 737, 738-739 (2d Cir. 1936). And the Sixth Circuit adopted the Second Circuit’s reading based on the “soundness of its reasoning” and because “the statute must be given a reasonable construction.” *Davis v. United States*, 62 F.2d 473, 474-475 (6th Cir. 1933). Finally, the D.C. Circuit chose “to follow the interpretation which

² See Siegel & Ziegler, *Comstockery: How Government Censorship Gave Birth to the Law of Sexual and Reproductive Freedom, and May Again Threaten It*, 134 Yale L.J. 1068, 1096-1098 & n.139 (Feb. 2025).

has been adopted in other circuits.” *Consumers Union of United States v. Walker*, 145 F.2d 33, 33, 35 (D.C. Cir. 1944). No court of appeals has ever adopted a contrary construction.³

In 1948, against the backdrop of this uniform judicial precedent, Congress reenacted these provisions at 18 U.S.C. §§ 1461-1462, without change to the relevant language. *See* Pub. L. No. 80-772, 62 Stat. 683, 768-769 (1948). This action—without more—“is convincing support for the conclusion that Congress accepted and ratified the unanimous holdings of the Courts of Appeals” regarding the proper interpretation of the provisions. *Texas Department of Housing & Community Affairs v. Inclusive Communities Project, Inc.*, 576 U.S. 519, 536 (2015). That is because “[i]f a word or phrase has been . . . given a uniform interpretation by inferior courts . . . , a later version of that act perpetuating the wording is presumed to carry forward that interpretation.” Scalia & Garner, *Reading Law* 322 (2012); *accord Texas Department of Housing*, 576 U.S. at 537 (citing cases to the same effect).

But here there is more: Congress’s attention was specifically drawn to most of the circuit decisions discussed above. In particular, a note included in the House Judiciary Committee’s 1947 report accompanying the bill stated: “The attention of Congress is invited to the following decisions of the Federal courts construing [proposed § 1461] and section 1462.” H.R. Rep. No. 80-304, at A104-A105 (1947). These “decisions” to which Congress’s “attention” was “invited” were four of the relevant circuit cases. First, the report explained that *Youngs Rubber* concluded that “the more reasonable interpretation” of the language “as used in [proposed § 1461] and section 1462” was “to construe the whole phrase ‘designed, adapted or intended’ as requiring ‘an

³ Contrary to amici’s suggestion, the fact that some of these cases involved the mailing of contraception items rather than abortion-producing items is irrelevant. *See* Br. of Ethics and Public Policy Center (ECF No. 101) at 10; Br. of Members of Congress (ECF No. 141) at 17. The statutory language defining the requisite mens rea applies to all the unmailable items listed in the Comstock laws, see 18 U.S.C. §§ 1461, 1462, and so the construction given to that language applies equally to abortion-producing items as to contraception items.

intent on the part of the sender that the article mailed or shipped by common carrier be used for *illegal* contraception or abortion.” *Id.* at A105 (emphasis added). Next, the report stated that *Nicholas* “held that the importation or sending through the mails of contraceptive [or abortion] articles is not forbidden absolutely, but only when such articles or publications are *unlawfully* employed.” *Id.* (emphasis added). Finally, the report added that “[t]he same rule was followed” by *Davis* and *One Package*. *Id.*

Notes issued in connection with the 1948 codification project have been described by the Supreme Court as “obviously authoritative.” *United States v. National City Lines, Inc.*, 337 U.S. 78, 81 (1949); *see also Ex parte Collett*, 337 U.S. 55, 71 (1949) (“flatly reject[ing]” the notion that “Congress did not appreciate what it was enacting” in light of a revisers’ note); *United States v. Thompson*, 319 F.2d 665, 669 (2d Cir. 1963) (“[T]he revisers’ notes to the 1948 codification of the Judicial Code . . . are authoritative guides to congressional intent.”); *Acron Investments, Inc. v. Federal Savings & Loan Insurance Corporation*, 363 F.2d 236, 240 (9th Cir. 1966) (similar). Congress thus understood when it enacted §§ 1461-1462 that the language “as used in” those sections had been consistently interpreted to reach items for producing abortion only if intended to produce unlawful abortion. The fact that Congress enacted that language with that understanding and without expressing any reservation about that interpretation shows conclusively that Congress intended by its enactment of §§ 1461-1462 to ratify that judicial interpretation.⁴

⁴ Amici Members of Congress (ECF No. 141 at 18) suggest there was no pre-1948 consensus by pointing to three Court of Appeals decisions involving prosecutions under the Comstock laws that did not expressly address whether the statute reached mailing for lawful abortions. Those decisions do not suggest that the cases in fact concerned *lawful* abortions. *See, e.g., Weathers v. U.S.*, 126 F.2d 118, 119 (5th Cir. 1942) (referring to the abortions at issue as “a State crime”). And in any event, the revisers’ note—which pointed to the consensus of courts to squarely address the issue—is the “authoritative” word on Congress’s intent in reenacting the Comstock laws in 1948.

Indeed, having been alerted to this uniform precedent, Congress not only left the statutory text unchanged, but titled the new 18 U.S.C. § 1461 “Mailing obscene or *crime-inciting* matter” (emphasis added). Congress’s framing of § 1461 as a statute concerned with conduct inciting crime reflects its understanding that the Act does not apply in connection with lawful abortions. *See Almendarez-Torres v. United States*, 523 U.S. 224, 234 (1998) (“‘[T]he title of a statute and the heading of a section’ are ‘tools available for the resolution of a doubt’ about the meaning of a statute.”) (citation omitted).

B. Congress Repeatedly Ratified The Circuits’ Unanimous Interpretation Of The Comstock Laws

While no more is needed to establish that Congress intended §§ 1461-1462 to reach abortion items only if intended for unlawful abortion, the decades-long post-1948 dialogue between Congress, the courts, and the executive agencies charged with enforcing the Comstock laws confirms that intent.

For example, in 1950 and again in 1955, Congress revised §§ 1461-1462 while preserving the key language. Pub. L. No. 81-531, § 1, 64 Stat. 194, 194 (1950); Pub. L. No. 84-95, §§ 1-2, 69 Stat. 183, 183 (1955). As explained, that shows Congress’s intent to ratify the circuits’ unanimous interpretation of the laws. In 1957, a district court recognized that “[t]he cases” interpreting §§ 1461-1462 hold “that only contraceptives [and abortion items] intended for ‘unlawful’ use were banned.” *United States v. 31 Photographs*, 156 F. Supp. 350, 357 (S.D.N.Y. 1957) (citing *Bours*, *One Package*, *Nicholas*, *Youngs Rubber*, *Davis*, and *Consumers Union*). The next year, Congress again revised §§ 1461-1462 while preserving the abortion-related language—once again ratifying the interpretation that had just been recognized in *31 Photographs*. Pub. L. No. 85-796, § 2, 72 Stat. 962, 962 (1958).

Not long after that, another district court deemed it “well established that the defendants should not be convicted [under §§ 1461-1462] unless it is established beyond a reasonable doubt that at the time they mailed the sample packages of prophylactics that they intended them to ‘be used for *illegal* contraception.’” *United States v. H.L. Blake Company*, 189 F. Supp. 930, 934-935 (W.D. Ark. 1960) (emphasis added) (citing *Bours, Nicholas, One Package, Youngs Rubber*, and *Davis*). And in 1961, Justice Harlan similarly noted the “judicial interpretation . . . that the absolute prohibitions of the [Comstock] law . . . exclude professional medical use.” *Poe v. Ullman*, 367 U.S. 497, 546 n.12 (1961) (Harlan, J., dissenting) (citing *Youngs Rubber, Davis*, and *One Package*). The following year, still another district court explained that it was “clear under the authorities that in order to make out an offense under [§§ 1461-1462], the Government should be required to allege and prove that . . . devices are shipped and received with intent that they be used for illegal contraception or abortion.” *United States v. Gentile*, 211 F. Supp. 383, 385 n.5 (D. Md. 1962) (citing *Youngs Rubber, Davis*, and *Nicholas*).

Against this backdrop, Congress took up §§ 1461-1462 in the early 1970s. And it again left the language of §§ 1461-1462 intact with respect to abortion (while removing references to contraception in response to *Griswold v. Connecticut*, 381 U.S. 479 (1965)). Pub. L. No. 91-662, §§ 3-4, 84 Stat. 1973, 1973 (1971). Notably, in the leadup to the 1971 amendment, the United States Postal Service (“USPS”) informed Congress that both USPS and the federal courts had adopted the long-standing, repeatedly ratified, narrow interpretation of the Act described above, as described in a House Report that accompanied the amendment. *See* H.R. Rep. No. 91-1105, at 3-4 (1970). Specifically, the Postmaster General informed the House Committee on Ways and Means that the Act had been construed not to extend to mailing items “for lawful purposes,” and advocated for the removal of prohibitions related to contraception (which had become broadly

lawful, and against which the Act had become “as presently written[,] . . . unenforceable”). *Id.* at 4. Amicus Ethics and Public Policy Center emphasizes that the same House Report states that existing statutes “completely prohibit” the importation and mailing of contraceptive materials. Br. (ECF No. 101) at 15 (citing H.R. Rep. No. 91-1105 (1970)). But amicus takes this language entirely out of context. The statement was clearly an introductory reference to the literal text of the provisions, as opposed to their settled meaning, as the Report then discussed the Postmaster General’s description of the settled narrow construction of the statute. H.R. Rep. No. 91-1105, at 3-4 (1970).

In 1994 and 1996, Congress again amended §§ 1461-1462 without material alteration. *See* Violent Crime Control and Law Enforcement Act, Pub. L. No. 103-322, 108 Stat. 1796 (1994); Communications Decency Act, Pub. L. No. 104-104, tit. V, § 507(a), 110 Stat. 133, 137 (1996). These repeated post-1948 ratifications are still further evidence of Congress’s understanding and approval of the circuits’ unanimous interpretation of §§ 1461-1462 as excluding distributions intended for legal abortions. Plaintiffs and some amici reference a failed amendment to § 1461 that would have confirmed that the statute applied only to “illegal abortion[s].” Pls.’ Reply Br. at 18 (ECF No. 111); Br. of Ethics and Public Policy Center at 15 (ECF No. 101); Br. of Members of Congress at 19 (ECF No. 141); Br. of Advancing American Freedom (ECF No. 123) at 18. But as the Supreme Court has consistently warned, “failed legislative proposals are ‘a particularly dangerous ground on which to rest an interpretation of a prior statute.’” *Central Bank of Denver, N.A. v. First Interstate Bank of Denver, N.A.*, 511 U.S. 164, 187 (1994).

Moreover, Congress re-enacted the Comstock laws while abortion was being openly, publicly, and legally provided—even before the Supreme Court recognized a federal

constitutional right to abortion. In 1970, prior to the three most recent reenactments of the Comstock laws and before *Roe*, “a total of 16 states had liberalized their abortion laws.” CDC, *Abortion Surveillance Annual Summary 1972* at 2 (April 1974). Those States “reported to the CDC” “over 193,000 legal abortions,” *id.* at 1-2, tbl.1. In 1972, more than 500,000 abortions were reported by just 17 States, *id.* at 2, tbl. 1. If Congress understood the Comstock laws to prohibit sending items intended for lawful abortions, such widespread provision of legal abortion—formally reported to an arm of the Executive Branch—certainly put Congress on notice that neither the public nor the coordinate branches of government shared that understanding. But Congress took no responsive action, which one would have expected if there were such a stark discrepancy between Congress’ intent and public understanding. Rather, Congress “perpetuat[ed] the wording” of the Comstock laws consistently interpreted by courts to prohibit only conduct intended to result in unlawful abortions, and Congress is accordingly “presumed to carry forward that interpretation.” *Texas Department of Housing*, 576 U.S. at 536.

Although some of the congressional ratification actions postdate *Roe v. Wade*, 410 U.S. 113 (1973), and *Planned Parenthood of Southeastern Pennsylvania v. Casey*, 505 U.S. 833 (1992), they are still meaningful because *Roe* and *Casey* did not prohibit all restrictions on abortions and some States permitted abortions that were not constitutionally protected. *See, e.g.*, Ore. Rev. Stat. § 659.880; D.C. Code § 2-1401.06 (repealed Feb. 23, 2023); N.J. Stat. § 10:7-2; Colo. Rev. Stat. § 25-6-403. Applying the Comstock laws to the distribution of items for producing abortion that was lawful in those States, therefore, would not necessarily have infringed the constitutional right to abortion and thus would have been an option for Congress every time it amended the Comstock laws. But, as explained, Congress simply reenacted the same language long understood *not* to reach items related to those lawful abortions. Thus,

Congress's actions—both before *Roe* and after it—made clear that Congress intended for the Comstock laws to reach only unlawful abortions.

Furthermore, amici are aware of no prosecutions brought under the Comstock laws for sending or receiving materials related to the provision of lawful abortion care since Congress's 1948 codification of the Act. Even before the Supreme Court recognized a federal constitutional right to abortion in *Roe v. Wade*, 410 U.S. 113 (1973), the Comstock laws were not used to bring prosecutions for sending or receiving supplies for lawful abortion care, despite the fact, as noted above, that more than half a million abortions were openly provided—and indeed reported to the federal government—the year before *Roe* was decided. Nor were there such prosecutions after the Supreme Court's decision in *Dobbs*. And following that decision, the Department of Justice's Office of Legal Counsel ("OLC") issued an opinion in response to a request from USPS, reiterating that 18 U.S.C. §§ 1461-1462 do not prohibit sending or receiving mifepristone or misoprostol when the sender or receiver does not intend for the recipient of the drugs to use them unlawfully. *See Application of the Comstock Act to the Mailing of Prescription Drugs That Can Be Used for Abortions*, 46 Op. O.L.C. ___, 1-2, 2 n.3, n.27 (Dec. 23, 2022). The OLC Opinion joined the longstanding and uniform recognition by legislative, judicial, and executive actors that the Comstock laws do not apply to sending or receiving items absent the intent they be used unlawfully.

C. The Comstock Laws' Text And Structure Show Congress Intended That They Reach Only Items Intended For Unlawful Abortions

Even without Congress's actions in 1948 and thereafter, the Comstock laws' text and structure would require that §§ 1461-1462 be interpreted to reach items only if intended for unlawful abortion.

As the Supreme Court has explained, a court’s “duty . . . is to construe statutes, not isolated provisions.” *King v. Burwell*, 576 U.S. 473, 486 (2015) (quotation marks omitted). Sections 1461-1462 must therefore be read in harmony with 19 U.S.C. §1305(a), which prohibits the “import[ation]” of “any drug or medicine or any article whatever for causing *unlawful* abortion” (emphasis added) and which also derives from the originally enacted Comstock Act, Act of Mar. 3, 1873, ch. 258, §3, 17 Stat. 598, 599. Moreover, statutory “interpretations . . . which would produce absurd results are to be avoided if alternative interpretations consistent with the legislative purpose are available.” *Griffin v. Oceanic Contractors, Inc.*, 458 U.S. 564, 575 (1982).

Plaintiffs’ interpretation of §§ 1461-1462 is not faithful to these canons. Indeed, it creates two absurdities in light of § 1305(a). First, it would mean that items intended for lawful abortion could be imported under § 1305(a) but not then distributed under §§ 1461-1462, or at least not distributed through the primary modes of interstate distribution for imported items. Such a regime makes no sense. Second, it would mean that items intended for lawful abortion could be imported under § 1305(a), but the importer could be prosecuted for doing so under § 1462, which prohibits importing abortion-producing items. Creating such a trap—where a person could be convicted of a crime for an act that another federal law expressly permits—is not only senseless but also would raise serious due-process concerns, contrary to the admonition that “statutes should be read where possible to avoid unconstitutionality.” *Dobbs v. Jackson Women’s Health Organization*, 597 U.S. 215, 287 (2022).

The court of appeals decisions discussed earlier recognized these problems. For example, in *One Package* the Second Circuit found it “hard to suppose” that Congress intended that “articles intended for use in procuring abortions were prohibited in all cases” under §§ 1461-

1462 but “only prohibited when intended for use in an ‘unlawful abortion’” under § 1305. 86 F.2d at 739. Concurring, Judge Learned Hand amplified the point, observing that “it is of considerable importance that the law as to importations should be the same as that as to the mails; we ought not impute differences of intention upon slight distinctions in expression.” *Id.* at 740.

And even apart from § 1305, the Seventh Circuit recognized, it is not “reasonable” to suppose Congress intended “the statute [to] cover all acts of abortion.” *Bours*, 229 F. at 964. As the Second Circuit elaborated, “[t]he intention to prevent a proper medical use of drugs or other articles merely because they are *capable* of illegal uses is not lightly to be ascribed to Congress.” *Youngs Rubber*, 45 F.2d at 108 (emphasis added). Therefore, the court said, it would not be “reasonable” to read the statute “to forbid the transportation by mail or common carriage of anything ‘adapted’ . . . for preventing conception . . . even though the article might also be capable of legitimate uses and the sender in good faith supposed that it would be used only legitimately.” *Id.*; *see also Davis*, 62 F.2d at 474-475. And the Second Circuit affirmed the reasoning in *One Package*, rejecting the notion that Congress intended to “bar [distribution of] articles for preventing conception though employed by a physician in the practice of his profession in order to protect the health of his patients or to save them from infection,” 86 F.2d at 739; *see also id.* at 740 (“It seems unreasonable to suppose that the national scheme of legislation involves such inconsistencies and requires the complete suppression of articles, the use of which in many cases is advocated by such a weight of authority in the medical world.”). Likewise, the D.C. Circuit concluded that, consistent with its “duty to avoid absurdity or injustice,” the statutory language should not be “tak[en] out of context” but rather should be construed to make exception for legitimate medical use. *Consumers Union*, 145 F.2d at 34-35.

Reading the Comstock laws contrary to the narrow construction Congress ratified also would disrupt and displace the States’ authority to regulate abortion within the bounds of the federal Constitution and any federal statutory protections. “[I]f Congress intends to alter the ‘usual constitutional balance between the States and the Federal Government,’ it must make its intention to do so ‘unmistakably clear in the language of the statute.’” *Will v. Michigan Department of State Police*, 491 U.S. 58, 65 (1989) (citation omitted). Plaintiffs’ interpretation of the Comstock laws as essentially an indirect ban on abortion care in States that choose to keep abortion legal runs counter to this “clear statement” requirement. Indeed, when overturning *Roe* and *Casey*, the Supreme Court underscored that States would be free to permit abortion. *See Dobbs*, 597 U.S. at 217-218 (“[T]he people of the various States may evaluate those interests differently. The Nation’s historical understanding of ordered liberty does not prevent the people’s elected representatives from deciding how abortion should be regulated.”); *see also id.* at 339 (Kavanaugh, J., concurring) (“[T]he ‘States may, if they wish, permit abortion on demand.’” (quoting *Casey*, 505 U.S. at 979 (1992) (Scalia, J., concurring in judgment in part and dissenting in part))); *id.* (“Today’s decision therefore does not prevent the numerous States that readily allow abortion from continuing to readily allow abortion.”). Plaintiffs’ interpretation would flatly contradict this authority, effectively nullifying state laws that permit abortion by blocking the supply of the materials necessary to engage in conduct deemed lawful—and indeed, in many instances, constitutionally protected—by individual States.

III. Plaintiffs’ Incorrect Interpretation Of The Comstock Laws Threatens To Affect All Abortions—Not Just Telemedicine Abortions—And Even Non-Abortion Healthcare

The Court should reject Plaintiffs’ interpretation of the Comstock laws because it is legally unsound for all the reasons discussed above. In addition, the Court should reject that interpretation because it could have devastating and far-reaching impacts on the provision of in-

person abortion care and other essential healthcare—effects that underscore the implausibility of Plaintiffs’ interpretation.

According to Plaintiffs, allowing for the mailing of mifepristone to patients is unlawful because the Comstock laws make it criminal to mail abortion-producing items. But nothing in the nature of that construction is limited to telemedicine abortion or to mifepristone. Mifepristone is also regularly used for *in-person* abortion care. National Abortion Federation, *Abortion in Clinic: What to Expect*. And the hospitals and clinics that provide that in-person care receive their supply of mifepristone through the mail or common carrier. *See, e.g.*, GenBioPro, *Ordering for Dispensing*. Plaintiffs’ interpretation would reach such mailing, threatening to block a common method of lawful, in-person abortion care. Plaintiffs’ interpretation could also impact the availability of a wide range of other drugs, devices, and items that may be used to produce an abortion and that hospitals and clinics receive through the mail or common carrier—not just mifepristone. *See Felix et al., The Comstock Act: Implications for Abortion Care Nationwide*, Kaiser Family Foundation (Apr. 15, 2024). Amici Women Injured By Abortion et al. suggest that “no woman in any state where abortion is legal will be denied an abortion because surgical abortion is still widely available and legal in all those states.” Br. (ECF No. 92) at 17. But amici fail to explain why Plaintiffs’ erroneous interpretation of the Comstock laws would not also reach items necessary for surgical abortions, such as manual vacuum aspirators, as the Comstock laws are not limited to drugs.

Critically, blocking the supply of many of these drugs and items could impact non-abortion healthcare as well. Mifepristone itself is central to miscarriage management. Gordon & McCammon, *A Drug That Eases Miscarriages Is Difficult For Women To Get*, NPR (Jan. 10, 2019). In a world in which Plaintiffs’ interpretation of the Comstock laws blocked the

availability of this medicine, healthcare providers would be unable to treat patients experiencing miscarriage with the standard-of-care treatments. And misoprostol, the other medication used in the two-drug medication abortion regimen, is also used in various settings outside the abortion context. Healthcare providers regularly use misoprostol for induction of labor to deliver a baby at term, for a range of common gynecological procedures, and to prevent and treat postpartum hemorrhage. Li et al., *Classifying Misoprostol and Mifepristone as Controlled Substances: Implications for the Management of Non-Abortion Related Conditions*, Kaiser Family Foundation (Apr. 3, 2025).

In sum, the Court should reject Plaintiffs' incorrect interpretation of the Comstock laws, which would significantly destabilize both abortion and non-abortion healthcare across the United States.

CONCLUSION

The Court should deny Plaintiffs' motion for a preliminary injunction.

Dated: February 20, 2026

**WILMER CUTLER PICKERING
HALE & DORR LLP**

Alan Schoenfeld*
M. L. Io Jones*
7 World Trade Center
250 Greenwich Street
New York, NY 10007
Telephone: (212) 295-8000
Fax: (212) 230-8888
alan.schoenfeld@wilmerhale.com
io.jones@wilmerhale.com

Kimberly A. Parker*
2100 Pennsylvania Avenue NW
Washington, DC 20037
Telephone: (202) 663-6000
Fax: (202) 663-6363
kimberly.parker@wilmerhale.com

Kyle Edwards Haugh*
50 California Street, Suite 3600
San Francisco, CA 94111
Telephone: (628) 235-1000
Fax: (628) 235-1001
kyle.haugh@wilmerhale.com

Arielle Herzberg*
1225 17th Street, Suite 2600
Denver, CO 80202
Telephone: (720) 274-3135
Fax: (720) 274-3133
arielle.herzberg@wilmerhale.com

**Pro hac vice application forthcoming*

Respectfully submitted,

/s/ Richard G. Perque_____

LAW OFFICE OF RICHARD G. PERQUE

Richard G. Perque
LA State Bar No.: 30669
700 Camp Street
New Orleans, LA 70130
Telephone: (504) 681-2003
Fax: (504) 681-2004
richard@perquelaw.com

Counsel for Amici Curiae

APPENDIX

LIST OF AMICI CURIAE

- Leslie R. Caldwell, Assistant Attorney General for the Criminal Division, U.S. Department of Justice (2014-2017)
- Michael W. Cotter, U.S. Attorney, District of Montana (2009-2017)
- Edward L. Dowd Jr., U.S. Attorney, Eastern District of Missouri (1993-1999)
- Jamie S. Gorelick, Deputy Attorney General, U.S. Department of Justice (1994-1997)
- Dwight C. Holton, U.S. Attorney, District of Oregon (2010-2011)
- William C. Killian, U.S. Attorney, Eastern District of Tennessee (2010-2015)
- James A. Lewis, U.S. Attorney, Central District of Illinois (2010-2016)
- Kenneth Magidson, U.S. Attorney, Southern District of Texas (2011-2017)
- Denise E. O'Donnell, U.S. Attorney, Western District of New York (1997-2001)
- David W. Ogden, Deputy Attorney General of the U.S., U.S. Department of Justice, (2009-2010); Assistant Attorney General for Civil Division, U.S. Department of Justice (1999-2001)
- Sarah R. Saldaña, U.S. Attorney, Northern District of Texas (2011-2014)
- Anne Tompkins, U.S. Attorney, Western District of North Carolina (2010-2015)
- John W. Vaudreuil, U.S. Attorney, Western District of Wisconsin (2010-2017)
- John F. Walsh, U.S. Attorney, District of Colorado (2010-2016)
- Seth P. Waxman, Solicitor General of the United States, U.S. Department of Justice (1997-2001)

CERTIFICATE OF SERVICE

I hereby certify that on February 20, 2026, I presented the foregoing Amici Curiae Brief of Former U.S. Department of Justice Officials to the Clerk of Court for filing and uploading to the CM/ECF system which will send notification of such filing to all registered CM/ECF users associated with this case.

/s/ Richard G. Perque

Richard G. Perque
LA State Bar No.: 30669
LAW OFFICE OF RICHARD G. PERQUE
700 Camp Street
New Orleans, LA 70130
Telephone: (504) 681-2003
Fax: (504) 681-2004
richard@perquelaw.com

Counsel for Amici Curiae

**UNITED STATES DISTRICT COURT
FOR THE WESTERN DISTRICT OF LOUISIANA
LAFAYETTE DIVISION**

THE STATE OF LOUISIANA, by and
through its Attorney General, LIZ
MURRILL, and ROSALIE MARKEZICH,

Plaintiffs,

v.

U.S. FOOD AND DRUG
ADMINISTRATION, *et al.*,

Defendants.

Case No. 6:25-cv-01491

Judge David C. Joseph

Magistrate Judge David J. Ayo

**MOTION OF FORMER U.S. DEPARTMENT OF JUSTICE OFFICIALS FOR LEAVE
TO FILE BRIEF OF AMICI CURIAE IN SUPPORT OF DEFENDANTS**

Proposed amici are 15 former high-ranking United States Department of Justice officials who served as Deputy Attorney General, Solicitor General, Assistant Attorney General, and/or U.S. Attorney. Amici were responsible for enforcing federal criminal laws, including the Comstock laws, 18 U.S.C. §§ 1461-1462, and represented the United States in criminal matters in all levels of the judiciary around the country. While amici hold diverse views regarding the moral and jurisprudential questions surrounding abortion, amici have broad experience in enforcing federal criminal laws that informs the analysis in the proposed brief and offers an important perspective to the Court. Proposed amici respectfully move for leave to file the

attached amicus brief in support of Defendants’ Opposition to Plaintiffs’ Motion for Preliminary Injunction.¹

The proposed brief explains why Plaintiffs’ argument that the 2023 Risk Evaluation and Mitigation Strategy (“REMS”) for mifepristone violates the Comstock laws is erroneous and provides additional analysis on issues not fully explored by the parties. First, the Food & Drug Administration (“FDA”) was not authorized to consider, interpret, or apply federal criminal laws as part of its drug-approval process, including its 2023 REMS decision. Furthermore, Plaintiffs gravely misinterpret the Comstock laws, expanding their scope beyond Congress’s intent. Plaintiffs’ misreading of the Comstock laws fails to consider the holdings of all four courts of appeals to have addressed the question, a construction that was ratified by Congress and embraced by the Justice Department, the sole agency responsible for prosecuting violations of the Comstock laws. Finally, amici explain how adopting Plaintiffs’ erroneous interpretation of the Comstock laws would risk a profound destabilization of medical care—not just abortion healthcare but other kinds of healthcare as well.

Courts enjoy broad discretion to allow the participation of amici curiae. *Lefebure v. D’Aquila*, 15 F.4th 670, 673 (5th Cir. 2021). The parties have already consented to the filing of amicus briefs. Granting leave to file the attached brief will not prejudice any party. Additionally, this brief is timely filed by the February 20, 2026 deadline for amicus briefs filed in support of Defendants.

For these reasons, amici respectfully request the Court grant the motion and file the attached brief.

¹ Amici are making this motion without conferring with the parties pursuant to this Court’s Order Regarding Joint Motion and Notice of Stipulation Regarding Amicus Curiae Briefs, Answer Deadline, and Briefing Schedule, Dkt. 39 (Jan. 20, 2026).

Dated: February 20, 2026

Respectfully submitted,

/s/ Richard G. Perque

**WILMER CUTLER PICKERING
HALE & DORR LLP**

Alan Schoenfeld*
M. L. Io Jones*
7 World Trade Center
250 Greenwich Street
New York, NY 10007
Telephone: (212) 295-8000
Fax: (212) 230-8888
alan.schoenfeld@wilmerhale.com
io.jones@wilmerhale.com

Kimberly A. Parker*
2100 Pennsylvania Avenue NW
Washington, DC 20037
Telephone: (202) 663-6000
Fax: (202) 663-6363
kimberly.parker@wilmerhale.com

Kyle Edwards Haugh*
50 California Street, Suite 3600
San Francisco, CA 94111
Telephone: (628) 235-1000
Fax: (628) 235-1001
kyle.haugh@wilmerhale.com

Arielle K. Herzberg*
1225 17th Street, Suite 2600
Denver, CO 80202
Telephone: (720) 274-3135
Fax: (720) 274-3133
arielle.herzberg@wilmerhale.com

**Pro hac vice application forthcoming*

LAW OFFICE OF RICHARD G. PERQUE

Richard G. Perque
LA State Bar No.: 30669
700 Camp Street
New Orleans, LA 70130
Telephone: (504) 681-2003
Fax: (504) 681-2004
richard@perquelaw.com

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/s/ Richard G. Perque

Richard G. Perque
LA State Bar No.: 30669
LAW OFFICE OF RICHARD G. PERQUE
700 Camp Street
New Orleans, LA 70130
Telephone: (504) 681-2003
Fax: (504) 681-2004
richard@perquelaw.com

Counsel for Amici Curiae

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Judge David C. Joseph

Magistrate Judge David J. Ayo

**ORDER ON MOTION OF FORMER U.S. DEPARTMENT OF JUSTICE OFFICIALS
FOR LEAVE TO FILE BRIEF OF AMICI CURIAE IN SUPPORT OF DEFENDANTS**

Considering the motion of former U.S. Department of Justice officials (“Proposed Amici”), and considering this Court’s Order Regarding Joint Motion and Notice of Stipulation Regarding Amicus Curiae Briefs, Answer Deadline, and Briefing Schedule, it is therefore ordered that the motion is GRANTED.

The proposed amicus brief shall be filed into the record.

SO ORDERED on this the ____ day of _____, 2026.

JUDGE
UNITED STATES DISTRICT COURT