SEC. 1. IMPROVING FDA AUTHORITIES FOR OPIOIDS.

Section 505–1(f)(3) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355–1(f)(3)) is amended—

(1) in subparagraph (E), by striking "or" at the end;

(2) in subparagraph (F), by striking the period and inserting a semicolon; and

(3) by adding at the end the following:

"(G) the drug be made available for dispensing to patients in unit dose packaging or another packaging system that the Secretary determines appropriate; or

"(H) the drug be dispensed to patients with a safe disposal packaging or safe disposal system that the Secretary determines appropriate for purposes of disposing of any unused dose of the dispensed drug.".