A compositions and dosages forms that allow for high drug loading for highly lipophilic drugs while maintaining excellent oral bioavailability. The pharmaceutical compositions and unit dosage forms described herein can reduce pill burden for hydrophobic drugs like (8R,9S,10R,13S,14S,17S)-10,13-dimethyl-3-oxo-1,2,6,7,8,9,11,12,14,15,16,17-dodecahydrocyclopenta[a]phenanthren-17-yl tridecanoate or (8R,9S,10R,13S,14S,17S)-10,13-dimethyl-3-oxo-1,2,6,7,8,9,11,12,14,15,16,17-dodecahydrocyclopenta[a]phenanthren-17-yl tetradecanoate, and can be formulated at advantageous drug loads (e.g., greater than 23%) while providing suitable bioavailability (e.g., capable of treating a hypogonadal male with less than 10 unit dosage forms per day) that allows for reduction in pill burden and accordingly improved patient adherence or compliance. Additionally, the composition (e.g., dosage form) has a release profile that is suitable for providing bioavailable API and the release profile is stable over time (e.g., under storage conditions).
exemple une forme galénique) présente un profil de libération qui est approprié pour fournir une API bio-disponible, et le profil de libération est stable au cours du temps (par exemple dans des conditions de stockage).

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Declarations:
Declaration made as to the identity of the inventor (PCT Rules 4.17(i) and 51bis.1(a)(i))
Declaration made as applicant's entitlement, as at the international filing date, to claim the priority of the earlier application, where the applicant is not the applicant who filed the earlier application or where the applicant's name has changed since the filing of the earlier application (Rules 4.17(iii) and 51bis.1(a)(iii))