CLINICAL GUIDELINES PROGRAM

NEW YORK STATE DEPARTMENT OF HEALTH AIDS INSTITUTE | HIV · HCV · SUBSTANCE USE · LGBT HEALTH

Resource: ART Drug-Drug Interactions

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Table 47: Gender-Affirming	g Hormones [Hembree, et al. 2017; Irving and Lehault 2017] (also see dru	g package inserts)
→ Cyproterone acetate, estradiol, finasteride, goserelin, leuprolide acetate, spironolactone, testosterone		
Class or Drug	Mechanism of Action	Clinical Comments
ARV medications, all	 Cyproterone acetate: Interaction with ARVs has not been studied. Estradiol: Interaction between ARVs and estradiol in transgender women has not been studied. Finasteride: Interaction with ARVs has not been studied. Finasteride is metabolized by CYP3A4; levels may increase when taken concomitantly with COBI-boosted ARVs, but clinical significance is expected to be minimal. Goserelin: Interaction with ARVs has not been studied. Based on what is known about metabolism of goserelin, no clinically significant interactions are expected. Leuprolide acetate: Interaction with ARVs has not been studied. Based on what is known about metabolism of leuprolide acetate, no clinically significant interactions are expected. Testosterone: Interaction between ARVs and testosterone in transgender men has not been studied. Testosterone has been used in androgen-deficient cisgender men with HIV without clinical drug interactions. Spironolactone: No interactions expected. 	 Estradiol: When prescribing ARVs, consider use of medications not expected to interact with estradiol. Finasteride: No dose adjustments recommended.
Cobicistat (COBI)	 Estradiol: Based on known mechanisms of metabolism, COBI- boosted PIs or other ARVs may have mixed effects on estradiol levels. COBI does not induce CYP1A2, and as such may increase estradiol levels by inhibition of CYP3A. Finasteride: When taken concomitantly, finasteride levels may be increased, but with minimal clinical significance. Testosterone: Based on known mechanisms of metabolism, there is limited potential that COBI-boosted PIs or other ARVs may increase testosterone levels. Relevance of this interaction is expected to be low in transgender men. 	 Estradiol: When taken concomitantly with COBI-boosted ARVs monitor for signs of estrogen deficiency or excess. Finasteride, testosterone: No dose adjustments are recommended.

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→ Cyproterone acetate, estradiol, finasteride, goserelin, leuprolide acetate, spironolactone, testosterone			
Class or Drug	Mechanism of Action	Clinical Comments	
Doravirine (DOR)	Estradiol, testosterone: No interactions expected.	N/A	
Efavirenz (EFV)	 Estradiol: EFV could induce CYP3A and could decrease estradiol levels. Finasteride, testosterone: Levels may decrease when taken concomitantly with EFV. 	 Estradiol: No dose adjustments are recommended, but when taken concomitantly with EFV, monitor for signs of estrogen deficiency or excess. Finasteride, testosterone: No dose adjustments recommended. 	
Etravirine (ETR)	 Estradiol: ETR could induce CYP3A and could decrease estradiol levels. Finasteride, testosterone: Levels may decrease when taken concomitantly with ETR. 	 Estradiol: No dose adjustments are recommended, but when taken concomitantly with ETR, monitor for signs of estrogen deficiency or excess. Finasteride, testosterone: No dose adjustments are recommended. 	
Etravirine (ETR)	 Estradiol: ETR could induce CYP3A and could decrease estradiol levels. Finasteride, testosterone: Levels may decrease when taken concomitantly with ETR. 	 Estradiol: No dose adjustments are recommended, but when taken concomitantly with ETR, monitor for signs of estrogen deficiency or excess. Finasteride, testosterone: No dose adjustments are recommended. 	
 Rilpivirine (RPV) INSTIs, non-boosted NRTIs, non-boosted 	Estradiol, finasteride, testosterone: No interactions are expected.	N/A	
Ritonavir (RTV)	 Estradiol: RTV may induce CYP1A2, which could decrease estradiol levels. This outweighs RTV inhibition of CYP3A. Testosterone: No interactions are expected. 	N/A	

References

Hembree WC, Cohen-Kettenis PT, Gooren L, et al. Endocrine treatment of gender-dysphoric/gender-incongruent persons: an Endocrine Society clinical practice guideline. Endocr Pract 2017;23(12):1437. [PMID: 29320642] https://pubmed.ncbi.nlm.nih.gov/29320642

Irving A, Lehault WB. Clinical pearls of gender-affirming hormone therapy in transgender patients. *Ment Health Clin* 2017;7(4):164-67. [PMID: 29955517] https://pubmed.ncbi.nlm.nih.gov/29955517