

TABLE 1B: EFFECTS AND EXPECTED TIME COURSE OF FEMINIZING HORMONES <sup>A</sup>

<b>Effect</b>	<b>Expected Onset<sup>B</sup></b>	<b>Expected Maximum Effect<sup>B</sup></b>
Body fat redistribution	3-6 months	2-5 years
Decreased muscle mass/ strength	3-6 months	1-2 years <sup>C</sup>
Softening of skin/decreased oiliness	3-6 months	unknown
Decreased libido	1-3 months	1-2 years
Decreased spontaneous erections	1-3 months	3-6 months
Male sexual dysfunction	variable	variable
Breast growth	3-6 months	2-3 years
Decreased testicular volume	3-6 months	2-3 years
Decreased sperm production	variable	variable
Thinning and slowed growth of body and facial hair	6-12 months	> 3 years <sup>D</sup>
Male pattern baldness	No regrowth, loss stops 1-3 months	1-2 years

<sup>A</sup> Adapted with permission from Hembree et al. (2009). Copyright 2009, The Endocrine Society.

<sup>B</sup> Estimates represent published and unpublished clinical observations.

<sup>C</sup> Significantly dependent on amount of exercise.

<sup>D</sup> Complete removal of male facial and body hair requires electrolysis, laser treatment, or both.

The degree and rate of physical effects depends in part on the dose, route of administration, and medications used, which are selected in accordance with a patient's specific medical goals (e.g., changes in gender expression, plans for sex reassignment) and medical risk profile. There is no current evidence that response to hormone therapy – with the possible exception of voice deepening in FtM persons – can be reliably predicted based on age, body habitus, ethnicity, or family appearance. All other factors being equal, there is no evidence to suggest that any medically approved type or method of administering hormones is more effective than any other in producing the desired physical changes.

TABLE 2. RISKS ASSOCIATED WITH HORMONE THERAPY. BOLDDED ITEMS ARE CLINICALLY SIGNIFICANT

Risk Level	Feminizing hormones	Masculinizing hormones
Likely increased risk	<b>Venous thromboembolic disease<sup>A</sup></b> Gallstones Elevated liver enzymes Weight gain <b>Hypertriglyceridemia</b>	Polycythemia Weight gain Acne Androgenic alopecia (balding) Sleep apnea
Likely increased risk with presence of additional risk factors <sup>B</sup>	<b>Cardiovascular disease</b>	
Possible increased risk	Hypertension Hyperprolactinemia or prolactinoma <sup>A</sup>	Elevated liver enzymes Hyperlipidemia
Possible increased risk with presence of additional risk factors <sup>B</sup>	<b>Type 2 diabetes<sup>A</sup></b>	Destabilization of certain psychiatric disorders <sup>C</sup> Cardiovascular disease Hypertension Type 2 diabetes
No increased risk or inconclusive	<b>Breast cancer</b>	Loss of bone density Breast cancer Cervical cancer Ovarian cancer Uterine cancer

<sup>A</sup> Risk is greater with oral estrogen administration than with transdermal estrogen administration.

<sup>B</sup> Additional risk factors include age.

<sup>C</sup> Includes bipolar, schizoaffective, and other disorders that may include manic or psychotic symptoms. This adverse event appears to be associated with higher doses or supraphysiologic blood levels of testosterone.