

**Comparison of Claims Between U.S. Patent Nos. 6,774,122; 7,456,160; 8,329,680; 8,466,139**

	<b>6,774,122</b> <i>Independent Claims 1 and 5</i>	<b>7,456,160</b> <i>Independent Claims 1 and 2</i>	<b>8,329,680</b> <i>Independent Claims 1 and 9</i>	<b>8,466,139</b> <i>Independent Claim 1 and 11</i>
Treatment	A method of treating a hormonal dependent benign or malignant disease of the breast or reproductive tract			
Injection Type	Intramuscularly to a human			
Fulvestrant	>45 mgml <sup>-1</sup> of fulvestrant ( <i>claim 5</i> )	>45 mgml <sup>-1</sup> of fulvestrant ( <i>claim 2</i> )	about 50 mgml <sup>-1</sup> fulvestrant about 50 mgml <sup>-1</sup> fulvestrant	about 50 mgml <sup>-1</sup> fulvestrant
Ethanol	10% w/v	10–30% w/v combined	about 10% w/v	17–23% w/v combined
Benzyl Alcohol	10% w/v	10–25% w/v	about 10% w/v	12–18% w/v
Benzyl Benzoate	15% w/v		about 15% w/v	12–18% w/v
Castor Oil		a sufficient amount <sup>1</sup>		
PK Result	whereby a therapeutically significant blood plasma fulvestrant concentration of at least 2.5 ngml <sup>-1</sup> is attained for at least 2 weeks after injection ( <i>claim 1 only</i> )	whereby a therapeutically significant blood plasma fulvestrant concentration of at least 2.5 ngml <sup>-1</sup> is attained for at least 2 weeks after injection ( <i>claim 1 only</i> )	wherein the method achieves a therapeutically significant blood plasma fulvestrant concentration of at least 2.5 ngml <sup>-1</sup> for at least four weeks	wherein the method achieves a blood plasma fulvestrant concentration of at least 2.5 ngml <sup>-1</sup> for at least two weeks

<sup>1</sup> Omitted from U.S. Patent No. 8,329,680, claim 9.

		Dependent Claims			
		Independent Claims 1 and 5		7,456,160 Independent Claims 1 and 2	8,329,680 Independent Claims 1 and 9
Breast Cancer	6,774,122 Independent Claims 1 and 5	2,9	12	3,6,11,14	5,10,15,20
PK Result (Raising Blood Plasma Conc. Levels or Extending Time Periods)	3-4 <i>Claim 3: ≥2.5 ngml<sup>-1</sup> for ≥4 weeks Claim 4: ≥2.5 ngml<sup>-1</sup> for ≥2-5 wks.</i>	5-9 <i>Claim 5: ≥2.5 ngml<sup>-1</sup> for ≥3 weeks Claim 6: ≥2.5 ngml<sup>-1</sup> for ≥4 weeks Claim 7: ≥3 ngml<sup>-1</sup> for ≥2 weeks Claim 8: ≥8.5 ngml<sup>-1</sup> for ≥2 weeks Claim 9: ≥8.5 ngml<sup>-1</sup> for ≥4 weeks</i>	2,10 <i>Claim 2: ≥8.5 ngml<sup>-1</sup> for ≥4 wks. Claim 10: ≥2.5 ngml<sup>-1</sup> for ≥4 wks.</i>	4,6,10,14,16,20 <i>Claims 4, 14: ≥8.5 ngml<sup>-1</sup> for ≥2 weeks Claims 6, 10, 16, 20: ≥2.5 ngml<sup>-1</sup> for ≥4 weeks</i>	
Divided Dose			17-20		9,19
Once Monthly				5,8,13,16	8,18
Limiting Injection Volume	6-8 <i>Claims 6, 7: ≤6 ml Claim 8: 5-5.25 ml</i>	10-11 <i>Claims 10, 11: ≤6 ml</i>	4,7,12,15 <i>Claims 4, 7,12, 15: 5 ml</i>	7,17 <i>Claims 7, 17: 5 ml</i>	
Limiting Conc. & Dose	6-8 <i>Claims 5,6: ≥45 mgml<sup>-1</sup> Claim 7: ≥250 mg Claim 8: 250mg</i>	10-11 <i>Claim 2,10: ≥45 mgml<sup>-1</sup> Claim 11: ≥250 mg</i>	3-4 <i>Claim 3: 15-25% w/v ethanol &amp; benzyl alcohol, 12-20% w/v benzyl benzoate</i>	2-3,12-13 <i>Claims 2, 12: 19-21% w/v ethanol &amp; benzyl alcohol, 14-16% w/v benzyl benzoate</i>	
Further Narrowing Excipient %				Claims 3, 13: <i>about 10% w/v ethanol, about 10% w/v benzyl alcohol, about 15% w/v benzyl benzoate</i>	