7. Dosage unit according to any one of the preceding claims, wherein the dosage unit contains 50-99.5 wt.% of filler selected from maltose, fructose, sucrose, lactose, glucose, galactose, trehalose, xylitol, sorbitol, erythritol, maltitol, mannitol, isomalt, microcrystalline cellulose, calcium salts and combinations thereof.

8. Dosage unit according to claim 7, wherein the dosage unit contains 50-99.5 wt.% of filler selected from lactose, xylitol, sorbitol, erythritol, mannitol, microcrystalline cellulose and combinations thereof.

9. Dosage unit according to claim 7 or 8, wherein the dosage unit contains at least 20 wt.% of sugar alcohol selected from mannitol, xylitol and combinations thereof.

10. Dosage unit according to any one of the preceding claims, wherein the dosage unit contains 0.1-20 wt.% of a disintegrating agent selected from modified starches, crosslinked polyvinyl pyrrolidone, crosslinked carmellose and combinations thereof.

11. Dosage unit according to any one of the preceding claims, wherein the dosage unit contains 0-60 wt.% of microcrystalline cellulose.

12. Dosage unit according to any one of the preceding claims, wherein the dosage unit contains 0.1-2 wt.% of lubricant selected from sodium stearyl fumarate, magnesium stearate, stearic acid, sodium lauryl sulfate, talc, polyethylene glycol, calcium stearate and mixtures thereof.

13. Use if an estetrol component for the manufacture of a medicament in the form of a solid dosage unit according to any one of claims 1-12 for medical treatment or for female hormone replacement therapy, wherein said solid dosage unit is to be administered sublingually, buccally or sublabially.

14. The use according to claim 13, wherein said solid dosage unit is to be administered once daily during a period of at least 1 week.