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journal of steroid biochemistry and molecular biology - d. tamae et al. / journal of steroid biochemistry & molecular biology 138 (2013) 281-283 scheme 1. the androgen metabolome is critical for male development and is therapeutically targeted in advanced prostate cancer.

journal of steroid biochemistry and molecular biology - 120 j. muncke / journal of steroid biochemistry & molecular biology 127 (2011) 118-127 ventions, and (3) exposure to substances leaching into dry foods is underestimated.

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journal of steroid biochemistry and molecular biology - 184 j.-y. moon et al. / journal of steroid biochemistry & molecular biology 139 (2014) 182-191 was confirmed in blank samples prior to running multiplexed cyp assays.

journal of steroid biochemistry and molecular biology - a. hanukoglu et al. / journal of steroid biochemistry & molecular biology 111 (2008) 268-274 269 of pha patients throughout adolescence and early adulthood.

journal of steroid biochemistry and molecular biology - s.j. ellem, g.p. risbridger / journal of steroid biochemistry & molecular biology 118 (2010) 246-251 247 emergence of pre-malignant lesions. the role of estrogen and the t:e balance in the prostate is further complicated as the specific effect of estrogen is also dictated by the differential actions of both estrogen receptors, er and er .

journal of steroid biochemistry and molecular biology - j.l. thomas et al. / journal of steroid biochemistry & molecular biology 111 (2008) 66-73 67 the structural basis of the higher affinity of 3-hsd1 for ligands relative to 3-hsd2. the aims of this study are to identify the residue that interacts with the 2-cyano group of trilostane (ser124) and to determine

journal of steroid biochemistry & molecular biology - seuter et al./journal of steroid biochemistry & molecular biology 174 (2017) 314-315. biologically plausible assumptions, such as a preponderant direction of changes in parameters as a function of 25(oh)d3 level variations. the statistical significance of the findings was evaluated using the

journal of steroid biochemistry and molecular biology - schug et al. / journal of steroid biochemistry & molecular biology 127 (2011) 204-215 fig. 1. model of the endocrine systems targeted by edcs. this figure illustrates that all major endocrine organs are vulnerable to endocrine disruption, including the hpa axis, reproductive organs, the pancreas, and the thyroid

gland.

journal of steroid biochemistry and molecular biology - x. li et al. / journal of steroid biochemistry & molecular biology 127 (2011) 9–15 as has been described for 9-cis retinoic acid, or the synthetic retinoid LG100268 in cell culture and in PPAR knockout mice [17].

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