Inhibition of Human Placenta Aldose Reductase by Tannic Acid.

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Tannic acid was found to be a highly potent inhibitor of human placenta aldose reductase. The most potent inhibitory component of the tannic acid was isolated and identified as penta-\(O\)-galloyl-\(\beta\)-D-glucose, which showed an IC\(_{50}\) value of 70 nM. The inhibition by the gallotannin was reversible and of mixed type with respect to DL-glyceraldehyde as the varied substrate.

Purification and Properties of Multiple Forms of Dihydriodiol Dehydrogenase from Monkey Liver.

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Two major and two minor forms of dihydriodiol dehydrogenase with similar molecular weights of around 36,000 were purified from monkey liver cytosol. All the forms oxidized trans-dihydriodiols of benzene and naphthalene and reduced aromatic aldehydes, but showed differences in specificity for other substrates and inhibitor sensitivity. The results indicate that one major (pI 8.7) and one minor (pI 7.9) form of the enzyme are indanol dehydrogenases, the other major form (pI 6.2) is 3α-hydroxysteroid dehydrogenase and the other minor form (pI 5.8) cross-reacted with human aldehyde reductase is aldehyde reductase.

Identification of Two Dihydriodiol Dehydrogenases Associated with 3(17)α-Hydroxysteroid Dehydrogenase Activity in Mouse Kidney.

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Three minor and one major forms of dihydriodiol dehydrogenase were in the kidney cytosol. Two minor forms are immunologically identical to hepatic 3α-hydroxysteroid dehydrogenase and aldehyde reductase, respectively. The other minor and the major forms Mr 39,000 distinct from the hepatic enzymes oxidized trans-dihydriodiols of benzene and naphthalene, cis-benzene dihydriodiol and alicyclic alcohols in the presence of NAD(P)\(^+\), and reduced several xenobiotic aldehydes and ketones with NAD(P)H as a cofactor. The enzymes also oxidized 3α-hydroxysteroids and epitestosterone, and reduced 3- and 17-ketosteroids. They may be identical to 3(17)α-hydroxysteroid dehydrogenases.