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. 18/01/2011 . Forum Apple. (en franÃ§ais). . 1 - Flashpoint Portable (CS5) .  
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Synthesis and antitumor activity of some novel derivatives of ethambutol. Ethambutol, an important antitubercular drug, was metabolized by the action of microsomal enzymes of rat liver homogenate to yield the corresponding derivatives: 4-amino-N-(2-mercaptoethyl)benzenesulphonamide (I), 4-amino-N-(2-thiobenzoyl) benzenesulphonamide (II) and 4-amino-N-(2-mercapto-4-methylpentanamidothio) benzenesulphonamide (III). The derivatives showed low antibacterial and antitubercular activity at different concentration levels, and the derivatives II and III exhibited highest antitumor activity and least toxicity in comparison to the standard drug, on MCF-7 breast cancer cell line.  
The oxygen-labile core of quercetin. An electron transfer method for the formation of quercetin-3-O-beta-glucoside. The anomeric configuration and the configuration of the glycosidic bond at the 3 position in quercetin-3-O-beta-D-glucoside was determined by chemical transformations of the glycoside. These procedures are based on the oxygen-labile character of the quercetin-3-O-beta-D-glucoside anomeric methoxime and the fact that beta-galactopyranosyl methoxime is rapidly oxidized by oxygen to the corresponding 1-deoxy derivative. The beta-D-glucopyranosyl-oxime is chemically transformed into the corresponding methyl ester and alkali treatment of the methyl ester yields the O-methyl beta-D-glucopyranoside. The configuration of the other glycosidic bond in quercetin-3-O-beta-D-glucoside was determined by introducing a trapping group in the methyl group of the O-methyl glycoside. The product of the O-methylation reaction 6d1f23a050

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