## SYNTHESIS AND BIOLOGICAL ACTIVITIES OF SOME NAPHTHOFURANS AS ANALOGUES OF 7.8-BENZOFLAVONES

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7.8-benzoflavone is known to be a powerful polycyclic hydrocarbons metabolism inhibitor. So, we attempted to synthesize 2-phenyl  $\alpha$ - or  $\beta$ -naphthofurans acylated in various positions according to modified Vilsmeir-Haack or Friedel-Crafts reactions.

$$\begin{array}{c}
\text{CHO} \\
\text{I} \\
\text{R = CHO} \\
\text{R = CO · CH}_3 \quad \underline{3}
\end{array}$$

$$\begin{array}{c}
\text{R = CHO} \\
\text{R = CHO} \quad \underline{4} \\
\text{R = CO · CH}_3 \quad \underline{5}
\end{array}$$

Then we tested the activity of those compounds on the metabolisation of benzo[a]pyrene incubated with rat liver microsomes and compared the results with the activity of 7.8-benzoflavone.

Compounds  $\underline{2}$  and  $\underline{3}$  show very interesting activities similar to that of 7.8-benzoflavone on the inhibition of benzo[a]pyrene metabolisation (70 % to 80 % inhibition after 40' incubation). Compounds  $\underline{1}$ ,  $\underline{4}$  and  $\underline{5}$  are less active (not over 50 % inhibition).