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NEW SEMISYNTHETIC CEPHALOSPORINS WITH ANTIBACTERIAL, ANTIVIRAL AND ANTIFUNGAL ACTIVITY

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NEW SEMISYNTHETIC CEPHALOSPORINS WITH ANTIBACTERIAL, ANTIVIRAL AND ANTIFUNGAL ACTIVITY

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The synthesis of new derivatives of thiazolidine 2-alkylidene-4-on-5-aceta-midocephalosporanic acids or thiazolidine 2-arylidenazino-4-on-5-acetamido-cephalosporanic acids and their salts of general formula (I) has been achieved.

$$\begin{array}{c} \text{CH}_{3} \\ \text{R}_{1} \\ \text{R}_{2} \\ \text{N} \end{array} \begin{array}{c} \text{CH}_{2} \\ \text{CONH} \\ \text{COOR}_{3} \\ \end{array}$$

 $R_1 = CH_3, C_2H_5, C_4H_9, C_6H_5$

 $R_2 = H, C_6H_5, CH_3C_6H_4$

 $R_3 = H$, Na, K

Synthesis of new semisynthetic cephalosporins was performed using the mixed anhydride, acid chloride and carbodiimide methods. The mixed anhydride method has given the best results regarding yields and purity. The structure of the new compounds has been established by elementary analysis, NMR spectrometry and TLC.

Same of these compounds showed broad antibacterial, antiviral and antifungal activity.