Syntheses of New **b**-Bifunctionalized Derivatives and Annelated Pyrimidines Based on Natural Monoterpenes (+)-3- Carene and Limonene

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 β -Enaminones are established as versatile synthetic intermediates presumably in heterocyclic chemistry. We have investigated the synthetic routes from readily accessible enaminones **3** derived from seco-derivatives of (+)-carene (**1**) and limonene (**2**) to β -chlorovinylketones **5**, enaminones **8** and **9**, and 2-substituted pyrimidines **4** and **7**. Containing physiologically active amine moiety, enaminones **8** and **9** may be considered as prodrugs. Pharmacokinetic advantages of such compounds are supposed as follows: better stability, improved transport *via* membranes, lower toxicity. On the other hand, due to their moderate lability to hydrolisys N-acylated derivatives **10** are also of interest in case of presence of a physiologically active acid moiety. Some pyrimidine derivatives **4** are **7** are prospective as synthones for chemotherapeutical's preparations. Synthetic procedures, structure elucidation and mechanistic consideration will be presented in details.



R = fused dimethyl cyclopropane moiety, isopropenyl $R^1 = Ar$ $R^2 = H$, NH_2 $R^3 = Alk$ R^4 , $R^5 = H$, Alk, Ar