FEATURES OF USE THE ELMA-SYSTEM FOR COMPLEX ASSESMENT TO THE ACTIVITES OF PHARMACEUTICAL ENTERPRISE

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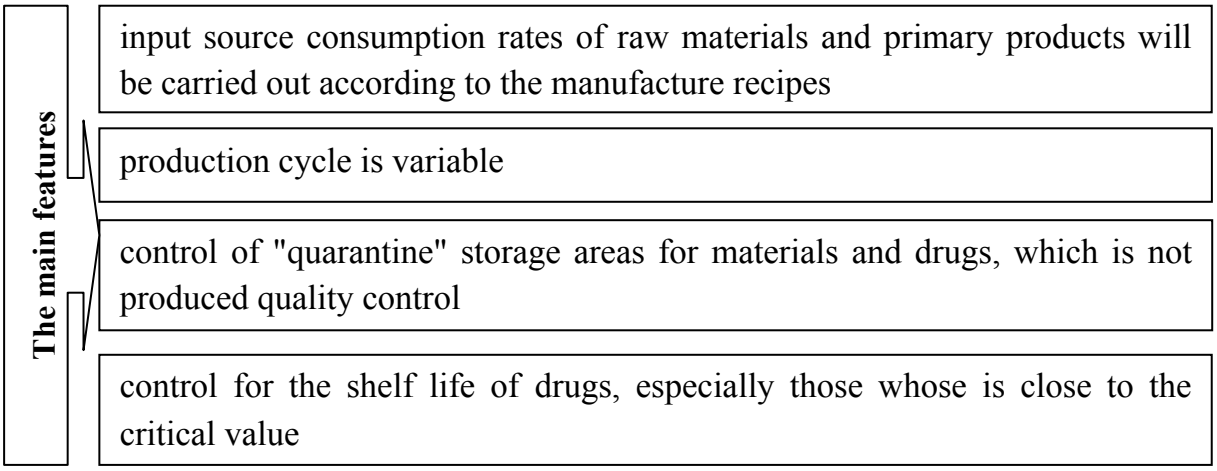
The process of save and quality drugs production is regulated by GMP standard. This standard in Ukraine implemented on all main pharmaceutical plants, because without GMP to complete with foreign manufacturers is it impossible. But GMP standard doesn't contain the requirements for pharmaceutical enterprise electronic management system.

The goal of this work is ground of opportunities of the possibility ELMA-system using in pharmaceutical enterprise.

The work with pharmaceutical enterprise documentation has main features. First of all this is documentation to primary products, raw materials and accessory substances, which have specific of pharmaceutical brunch.

The electronic document management system for pharmaceutical enterprise will be reflect main features (pic.1).



The ELMA-system allows you to create and keep the "electronic-file" with the list of documents for each drug series; reports on the shift production, quality assurance reports, drugs route maps, etc.

Thus, the considered ELMA-system can be used for the electronic documentation management system of pharmaceutical enterprise formation. This will increase the resource using and management decisions efficiency.