«ԴԵՂԵՐԻ ԵՎ ԲԺՇԿԱԿԱՆ ՏԵԽՆՈԼՈԳԻԱՆԵՐԻ ՓՈՐՁԱԳԻՏԱԿԱՆ ԿԵՆՏՐՈՆ» ՓԲԸ
«НАУЧНЫЙ ЦЕНТР ЭКСПЕРТИЗЫ ЛЕКАРСТВ И МЕДИЦИНСКИХ ТЕХНОЛОГИЙ» АОЗТ
«SCIENTIFIC CENTRE OF DRUG AND MEDICAL TECHNOLOGY EXPERTISE» JSC

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Dear Sir/Madam,

Please be informed that **Monitoring of the adverse drug reactions in the RA** is carried out by Scientific Center of Drug and Medical Technology Expertise of Minister of Health of the Republic of Armenia.

According to the requirement of Article 17 Law "On Medicines" of Republic of Armenia the health facilitates, pharmacies and establishments and organizations engaged in drug realization and use should immediately notify the body authorized by the Government about all cases of unknown side effects of medicines.

Thus, based on requirements of Article 17 of Law "On Medicines" of Republic of Armenia and point 2.1.4 of Contract " On rendering expertise services", please be informed that it is necessary to submit to the Centre the information about all cases of serious and/or unexpected adverse drug reactions occurred in post-registration period within 15 days of its occurrence. All other cases of adverse drug reactions must be submitted once in a 3 month.

Director H. Topchyan