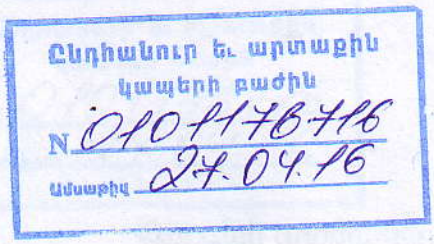
«ԱԿԱԴԵՄԻԿՈՍ ԷՄԻԼ ԳԱԲՐԻԵԼՅԱՆԻ ԱՆՎԱՆ ԴԵՂԵՐԻ ԵՎ ԲԺՇԿԱԿԱՆ ՏԵԽՆՈԼՈԳԻԱՆԵՐԻ ՓՈՐՁԱԳԻՏԱԿԱՆ ԿԵՆՏՐՈՆ» ՓԲԸ
«НАУЧНЫЙ ЦЕНТР ЭКСПЕРТИЗЫ ЛЕКАРСТВ И МЕДИЦИНСКИХ ТЕХНОЛОГИЙ ИМЕНИ АКАДЕМИКА ЭМИЛЯ ГАБРИЕЛЯНА» АОЗТ
«SCIENTIFIC CENTRE OF DRUG AND MEDICAL TECHNOLOGY EXPERTISE AFTER ACADEMICIAN EMIL GABRIELYAN» CJSC

Կոմիտաս պող. 49/4, Երևան 0051, Հայաստան
пр. Комитаса 49/4, Ереван 0051, Армения
49/4 Komitas aven., Yerevan 0051, Armenia

Հեռ. (+374 10) 23-16-82, 23-08-96, 23-47-32, 23-20-91
Tel.: (+374 10) 23-16-82, 23-08-96, 23-47-32, 23-20-91
Fax: (+374 10) 23-29-42, 23-21-18
E-mail: admin@pharm.am ; URL: http://www.pharm.am



Dear Sir/Madam,

Considering the fact that according to the new law "On Medicines", discussing in the National Assembly, the registration dossier of a medicinal product should be submitted in the common technical document format (CTD), hereinafter it is recommended to follow attached list (Appendix 1) and guideline (Appendix 2) while compiling new registration dossier. The use of the logically structured format not only improves the effectiveness and transparency of expertise, but also contributes to registration of safe, good quality and effective medicinal products providing in separate sections detailed data on finished product, as well as active substance and excipients. Similar structure is also planned for documents submitting for registration of medicinal products in the framework of the Eurasian Economic Union.

Best regards,

Director Hakob Topchyan