# Act on handling-acceptance of standard[1]

Standard(s) of the following medicinal product is(are) submitting in purpose of registration expertise:

| | | | |
|---|---|---|---|
| **Trade name, dosage form and strength of medicinal product** | | | |
| **Name and address of manufacturer** | | | |
| **Name and address of marketing authorization holder** | | | |
| **Standard name[2]** | | | |
| **batch[3]** | | | |
| **quantity** | | | |
| **shelf life** | | | |
| **storage conditions[4]** | | | |
| **quality certificate** | ☐ | ☐ | ☐ |

| **Who handed over** | **Who accepted** |
|---|---|
| **name of company** | **name of company**<br>"Scientific Centre of Drug and Medical Technology Expertise after Academician Emil Gabrielyan" JSC |
| **name of department** | **name of department [5]**<br>General and external affairs department<br>Narcotics and other controlled substances management department |
| **name, last name** | **name, last name** |
| **date, signature, seal/stamp** | **date, signature, stamp** |

[1] *The act shall be submitted either by marketing authorization holder or by its authorized representative in 2 copies filled, printed, signed and sealed/stamped in advance.*

[2] *It is necessary to fill in the 2nd, 3rd and 4th columns of table in case you submit different standards of the same medicinal product.*

[3] *It is necessary to fill in the 2nd, 3rd and 4th columns of table in case you submit different batches of the same standard for the same medicinal product.*

[4] ***It is necessary to mark "not complied" in the line "storage conditions" of the table in case required special*** *storage conditions for submitted standards are not kept.*

[5] *Choose appropriate department. NOTE. Samples of narcotics or other controlled substances should be handed over to the Head of Narcotics and other controlled substances department of the Center.*