**Act on handling-acceptance of standard[[1]](#footnote-2)**

Standard (s) of the following medicinal product is (are) submitting in purpose of expertise for registration, quality related changes in specifications:

|  |  |
| --- | --- |
| **Trade name, pharmaceutical form, strength, presentation form of medicinal product** |  |
| **Name and address of manufacturer (batch releaser)** |  |
| **Name and address of marketing authorization holder** |  |
| **Standard name[[2]](#footnote-3)** |  |  |  |
| **batch[[3]](#footnote-4)** |  |  |  |
| **quantity** |  |  |  |
| **shelf life** |  |  |  |
| **storage conditions[[4]](#footnote-5)** |  |  |  |
| **quality certificate** |  |  |  |
| **Who handed over** | **Who accepted** |
| **name of company** | **name of company**“Scientific Centre of Drug and Medical Technology Expertise after Academician Emil Gabrielyan” JSC |
| **name of department** | **name of department [[5]](#footnote-6)** |
| **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.*** [↑](#footnote-ref-2)
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.*** [↑](#footnote-ref-3)
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.*** [↑](#footnote-ref-4)
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.*** [↑](#footnote-ref-5)
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.*** [↑](#footnote-ref-6)