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

Sample(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** |  | | |
| **Sample batch [[2]](#footnote-3)** |  | |  |
| **quantity** |  | |  |
| **shelf life** |  | |  |
| **marking language** |  | |  |
| **storage conditions[[3]](#footnote-4)** |  | |  |
| **instruction for use**  **(leaflet insert)** |  | |  |
| **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 [[4]](#footnote-5)** | |
| **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 and 3rd columns of table in case you submit different batches of samples of the same medicinal product.*** [↑](#footnote-ref-3)
3. ***It is necessary to mark “not complied” in the line “storage conditions” of the table in case required special storage conditions for submitted samples are not kept.//*** [↑](#footnote-ref-4)
4. ***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-5)