

APPROVED

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**INSTRUCTION**  
**ON SUBMITTING AN APPLICATION FOR REGISTRATION AND**  
**TRACEABILITY OF DIGITAL IDENTIFICATION DEVICES ISSUED BY THE**  
**FEDERAL STATE INFORMATION SYSTEM FOR MONITORING THE**  
**MOVEMENT OF MEDICINAL PRODUCTS FOR MEDICAL USE**  
**(IS MMMP)**  
**IN THE NATIONAL INFORMATION SYSTEM FOR THE MONITORING OF**  
**LABELING AND TRACING OF PRODUCTS**  
**"ASL BELGISI" (NIS "ASL BELGISI")**

**VERSION 1.0**

**2022**



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## 1. Introduction

According to Decree of the Cabinet of Ministers No. 149 of April 2, 2022 "On the introduction of a mandatory system of digital labeling of medicines and medical devices" (hereinafter - Decree No. 149) in the Republic of Uzbekistan introduces mandatory digital labeling of medicines and medical devices.

Digital labeling of goods, including medicines and medical devices is provided through the use of the National Information System for Monitoring Labeling and Tracking of Goods "ASL BELGISI" (hereinafter - NIS "ASL BELGISI").

According to the Regulation on the Procedure of Digital Labeling of Medicinal Products and Medical Devices, approved by Annex No. 6 to the Decree № 149 (hereinafter - the Regulation), the NIS operator "ASL BELGISI" has the right to register and subsequent traceability in the Republic of Uzbekistan of digital identification means that meet the requirements of the Regulation and issued by the operators of national information systems of digital labeling and traceability of medicines and medical devices of other countries.

14. The Operator has the right to carry out recognition and subsequent traceability on the territory of the Republic of Uzbekistan of digital identification means, complying with the requirements of this Regulation and issued by the operator of foreign information system of digital labeling and traceability of medicinal products and medical devices (hereinafter - the Foreign Operator). At the same time, a prerequisite is to provide the Operator with information on digital identification devices applied to labeled medicinal products and medical devices, information on such medicinal products and medical devices and their manufacturer.

15. Recognition of digital identification devices issued by the Foreign Operator shall be carried out:

with respect to medicines and medical devices imported (imported) into the territory of the Republic of Uzbekistan (prior to their placement under customs procedures for release for domestic consumption or re-import);

after confirmation by the Foreign Operator of information about the digital identification means and the possibility to put medicinal products and medical devices labeled with such digital identification means into circulation.

16. For the purpose of recognition of digital identification means issued by the Foreign Operator, a participant of turnover of goods which introduces labeled medicinal products and medical devices into circulation on the territory of the Republic of



Uzbekistan shall submit an application to the Operator, the form of which shall be approved by the Operator and posted on the official website of the Operator in the Internet.

17. The conditions of information interaction between the Operator and a Foreign operator, including composition and procedure for transfer of data on digital identification means, medicines and medical products and their manufacturer shall be determined by an agreement between the Operator and the Foreign Operator.

This document presents a description and condition of the procedure of registration and traceability of digital identification means issued by the operator of the Federal State Information System for Monitoring the Movement of Medicinal Products for Medical Use (hereinafter - MMMP) in NIS "ASL BELGISI".

The process of registration and traceability of digital identifiers issued by the MMMP operator in NIS "ASL BELGISI" includes the following key stages:

- Production and labeling of medicinal products (according to the procedure established by the legislation of the Russian Federation);
- Withdrawal of labeled medicinal products from circulation in the IS MMMP (in accordance with the requirements of the "ASL BELGISI" Operator);
- Registration of the participant of goods turnover in the NIS "ASL BELGISI";
- Registration of labeled medicinal products in the NIS "ASL BELGISI" subsystem "List of labeled products" (performed by the holder of registration certificate or his accredited representative on the territory of the Republic of Uzbekistan);
- Submission of application for registration and traceability of digital identification means issued by the IS MMMP in NIS "ASL BELGISI";
- Turnover of labeled medicinal products in the Republic of Uzbekistan (in accordance with the requirements of the NIS Operator "ASL BELGISI").

### 1.1 History of changes

Version	Date	Changes	Author
1.0	15.07.2022	First version of the document	Rasulev D.S.

### 1.2 Abbreviations

<b>API</b>	Application programming interface
<b>JSON</b>	JavaScript Object Notation (a text-based data exchange format based on JavaScript)
<b>CSV</b>	Comma-Separated Values is a text file format designed to provide tabular data



<b>GTIN</b>	Global Trade Item Number
<b>sGTIN</b>	Serialized Global Trade Item Number – serialized Global Trade Item Identification Number, which is a unique identifier of a secondary (consumer) package of a medicinal product (in its absence – a primary package of a medicinal product), formed by adding an individual serial number of a trade item to the Global Trade Item Identification Number
<b>SSCC</b>	Serial Shipping Container Code (logistic unit number, represented as a digital number, the uniqueness of which is guaranteed by using the GS1 company prefix, which is assigned by the GS1 National Organization)
<b>TIN</b>	Taxpayer individual number
<b>PINFL</b>	Personal identification number of an individual
<b>PC</b>	Personal Cabinet
<b>CP</b>	Consumer package identification code
<b>LC</b>	Labeling code
<b>TP</b>	Transport package identification code
<b>MPMU</b>	Medicinal products for medical use
<b>RC</b>	Registration certificate
<b>KLG</b>	Subsystem “Catalog of Labeled Goods” of the National Information System for monitoring of Labeling and tracing products “Asl Belgisi”
<b>NIS “ASL BELGISI”</b>	National Information System for monitoring labeling and tracing products (Republic of Uzbekistan)
<b>IS MMMP</b>	Federal State Information System for Monitoring the Movement of Medicinal Products for medical use (Russian Federation)
<b>PTG</b>	Participant of turnover of goods
<b>FLC</b>	Formatted-logic control
<b>EDS</b>	Electronic digital signature
<b>Authorized Body</b>	Agency for the Development of the Pharmaceutical Industry under the Ministry of Health of the Republic of Uzbekistan
<b>MMMP IS Operator</b>	Limited Liability Company “Operator CRPT”, a legal entity registered in accordance with the laws of the Russian Federation
<b>Operator NIS “ASL BELGISI”</b>	Limited Liability Company “CRPT Turon”, a legal entity registered in accordance with the laws of the Republic of Uzbekistan
<b>STC</b>	State Tax Committee of the Republic of Uzbekistan
<b>SCC</b>	State Customs Committee of the Republic of Uzbekistan

### 1.3 Terms and definitions



## **Federal State Information System for Monitoring the Movement of Medicinal Products for Medical Use (Russian Federation) -**

**Subjects of circulation of medicines** - legal entities and individual entrepreneurs engaged in production, storage, importation into the Russian Federation, release, sale, transfer, use and transfer for destruction of medicines.

**Digital Identification Issuer** - a manufacturer of medicinal products that performs the production stage of packaging of a medicinal product with digital identification means applied to the secondary (consumer) package of the medicinal product (or, in the absence thereof, to the primary package of the medicinal product), when manufacturing the medicinal product in the Russian Federation, or the holder or owner of the registration certificate of the medicinal product, when manufacturing the medicinal product.

**National Information System for Monitoring Labeling and Traceability of Products "ASL BELGISI" (the Republic of Uzbekistan)** - information system created in order to automate the processes of collection and processing of information on the circulation of goods subject to mandatory digital identification labeling, storage of such information, providing access to it, its provision and distribution, improving the efficiency of such information and ensuring traceability of the specified goods, as well as for other purposes provided by the law, as well as for other purposes.

**Catalog of Labelled Goods** is a subsystem of NIS "ASL BELGISI", designed for the formation, processing and storage of information about the goods subject to mandatory digital labeling.

**CRPT TURON LLC** is a legal entity registered on the territory of the Republic of Uzbekistan, which creates, develops, modernizes and operates NIS LTP, ensures its uninterrupted operation, as well as receives, stores and processes information.

**Participant of goods turnover** - a legal entity or individual entrepreneur, which is a tax resident or non-resident of the Republic of Uzbekistan, engaged in the production, introduction into circulation, turnover and (or) withdrawal from circulation of goods, except for legal entities and individual entrepreneurs who purchase goods for purposes not related to their subsequent realization (sale).

**Medicinal product** - dosed packaged medicinal products ready for use.

**Registration certificate** - a document confirming the fact of state registration and authorization of the Ministry of Health of the Republic of Uzbekistan on the right to use a medicinal product, medical devices, and medical equipment in medical practice.



**Goods code** (barcode) - a code assigned to a group of goods in accordance with the rules established by the standards of GS1 system, which provides accounting and storage of reliable data on the goods under the relevant commodity nomenclature.

**Individual serial number** - a sequence of characters (digits, uppercase and lowercase letters of the Latin alphabet, as well as special characters) uniquely identifying each individual item of goods based on the product code (bar code).

**Identification code** - a sequence of characters representing a unique number of product unit consisting of product code (bar code) and individual serial number.

**Verification code** - a sequence of symbols formed by the operator on the basis of identification code and allowing to detect falsification of labeling code during its verification with the use of cash register equipment and (or) other technical means of verification of labeling code.

**Labeling code** - a unique sequence of symbols formed by the operator, consisting of an identification code and a verification code.

**Digital identification means** - a labeling code in a machine-readable form presented in the form of a two-dimensional Data Matrix bar code (for consumer and group packaging) or a GS1-128 line code (for transport packaging) applied to goods, packaging of goods or other tangible medium.

**Primary (inner) packaging** - the packaging in direct contact with the dosage form.

**Secondary (outer) packaging** - the packaging in which the medicinal product in the primary packaging is placed.

**Consumer packaging** - a secondary (outer) package of the medicinal product, and in its absence the primary (closed) package of the medicinal product.

**Transport packaging** is a packaging that combines goods in consumer and (or) transport packaging, used for the storage and transportation of goods to protect them from damage during transportation, and forms an independent transport unit (including first level transport packaging and subsequent level transport packaging) in the process of aggregation.

**Aggregation** - the process of combining goods in a consumer package into a first-level transport package, as well as first-level transport packages into subsequent-level transport packages with an identification code for the transport package being created, ensuring that information about the relationship between the identification codes of each enclosed consumer package, transport package and the identification code of the transport package being created is retained.



**GS1 System of Standards** - System of Standards of the international non-profit organization GS1, designed for automatic identification, data collection and information exchange between business partners in the supply chain.

**Application Identifier (AI)** - a set of two or more characters located at the beginning of an element string. An AI identifier is a prefix that uniquely defines the meaning and format of the data field following the AI. Data following an AI identifier can contain alphabetic or numeric characters of any length, up to thirty characters. The data fields can be of fixed or variable length depending on the AI identifier.

Important:

Description of processes related to submission of information by the subjects of circulation of medicines to the IS MMMP in this document is for reference only.

Registration and traceability of means of digital identification issued by IS MMP in NIS "ASL BELGISI" does not mean recognition of results of registration of medicinal products carried out outside the Republic of Uzbekistan.

Traceability of the means of digital identification issued by the IS MMMP in NIS "ASL BELGISI" is carried out only on condition of successful processing of application for registration and traceability of the relevant means of digital identification.

<sup>1</sup> <https://gs1ca.org/gs1ca-components/documents/GS1-DataMatrix-Introduction-and-technical-overview-v.pdf>





## **2. Description of means of digital identification of consumer packages and transport packages of medicinal products used in the Republic of Uzbekistan**

According to the Regulation on the procedure of the digital labeling of medicinal products and medical devices approved by the Annex No. 6 to the Decree № 149 (hereinafter - the Regulation), the means of digital identification of the consumer packaging contains a labeling code, which is formed by the Operator and includes 4 groups of data:

- first group of data - product code consisting of 14 characters (digits) preceded by application identifier (01) and formed by the Operator based on the information submitted during the registration of the drug or medical device in the catalog of goods;
- second group of data - an individual serial number of the consumer package, consisting of 13 characters (digits, lowercase and uppercase letters of the Latin alphabet), preceded by the application identifier (21), which is generated by the operator or a participant in the circulation of goods;
- third group of data - the identifier (individual serial number) of the verification key, consisting of 4 characters (digits, lowercase and uppercase letters of the Latin alphabet), preceded by the application identifier (91), which is generated by the operator;
- fourth group of data - a verification code, consisting of 44 characters (digits, lower and upper case Latin letters, as well as special characters), preceded by the application identifier (92) and generated by the Operator.

The data groups must be arranged sequentially - from the first to the fourth.

The first and second data groups shall form the consumer package identification code, and the third and fourth data groups shall form the verification code.

The product code and the individual serial number of the consumer package shall be duplicated in readable printed text at the manufacturer's discretion.

According to the Rules for Mandatory Digital Labeling, approved by Appendix No. 1 to Cabinet of Ministers Resolution No. 833 of December 31, 2020, the means of digital identification of transport packaging contains an identification code generated by the manufacturer of the labeled goods, consists of 18 digital characters and contains the following data structure:

- application identifier (00);
- extension symbol of the transport package identification code;



- registration number of the participant of the goods turnover in the GS1 information system;
- serial number of the transport package;
- checksum (a number calculated using a special algorithm from the preceding digits and serving to ensure data integrity).

The identification code of the transport package is generated and applied to the transport package by the manufacturer of the labeled goods by printing or labeling.

Table.

Type of packaging	Template
Consumer packaging	AI01 + GTIN {14 chars} + AI21 + serial_number {13 chars} + separator {ASCII 29} + AI91 + checking_code {4 chars} + separator {ASCII 29} + AI92 + checking_key {44 chars}
Group packaging	Not provided
Transport packaging of first level	SSCC (18 numeric characters)
Transport packaging of second level	

NIS "ASL BELGISI" supports the use of two levels of product aggregation:

- Transport packaging of the first level;
- Transport packaging of the second level.

Currently, as part of the registration and traceability procedure for digital identification means issued by IS MMMP, NIS "ASL BELGISI" supports the use only in relation to consumer packaging identification codes (sGTIN).



### 3. Production, labeling and withdrawal of medicinal products

Annotation	<p>Production of medicinal products and their labeling with digital identification means shall be carried out in accordance with the procedure stipulated by the legislation of the Russian Federation: Federal Law of April 12, 2010. No. 61-FZ "On Circulation of Medicines"; Decree of the Government of the Russian Federation No. 1556 of December 14, 2018. "On Approval of the Regulations on the System of Monitoring the Movement of Medicinal Products for Medical Use"; other regulatory legal acts of the Russian Federation.</p> <p>Transmission to the IS MMMP of information on the production, storage, importation into the Russian Federation, release, sale, transfer, use and transfer for destruction of medicinal products is performed by sending files in the format and in accordance with the instruction describing the procedures for submitting information by subjects of drug circulation, posted on the official website of the IS MMMP operator on the information and telecommunications network "Internet".</p>
Participants of interaction	<ul style="list-style-type: none"><li>▪ Operator of the IS MMMP</li><li>▪ Subjects of circulation of medicinal products</li></ul>
Special conditions	<p>With regard to the means of digital identification issued by IS MMMP and applied for the purposes of registration and traceability in NIS "ASL BELGISI", information on withdrawal from circulation must be registered in IS MMMP with an indication of export to the Republic of Uzbekistan.</p> <p>The said information shall be submitted in accordance with the procedure provided by the IS MMMP Operator before applying to the NIS Operator "ASL BELGISI" with</p>



	an application for recognition and traceability of the relevant means of digital identification in the territory of the Republic of Uzbekistan.
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#### 4. Registration of the participant of goods turnover in NIS «ASL BELGISI»

Annotation	<p>Registration of the participant of the goods turnover in NIS "ASL BELGISI" is done based on an electronic application. The electronic application is signed with a digital signature key certificate.</p> <p>Data in the electronic application for registration is processed and verified by NIS "ASL BELGISI" software tools, including through interaction with external information resources.</p> <p>In accordance with the legislation on electronic digital signature, an electronic digital signature key certificate must contain:</p> <ul style="list-style-type: none"><li>▪ Information about the physical person who is the holder of the EDS key certificate;</li><li>▪ Information about the legal person if the physical person is its representative.</li></ul> <p>When submitting an application, the type of the participant of the goods turnover is determined based on TIN contained in the EDS certificate:</p> <ul style="list-style-type: none"><li>▪ TIN of legal entities that are foreign organizations begins with the number 9;</li><li>▪ In other cases, the participant of goods turnover belongs to the residents of the Republic of Uzbekistan.</li></ul> <p>According to the results of consideration of the application:</p> <ul style="list-style-type: none"><li>▪ if there are no grounds for refusal, registration of the participant of goods turnover in NIS "ASL BELGISI" is completed, a personal account is created and the participant of goods turnover is assigned a registration number of the subject of circulation in NIS "ASL BELGISI";</li></ul>
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	<ul style="list-style-type: none"> <li>▪ If there are grounds for refusal, a message of refusal to register in NIS "ASL BELGISI" and to provide a personal office is generated.</li> </ul>
Participants of interaction	<ul style="list-style-type: none"> <li>▪ State Tax Committee of the Republic of Uzbekistan</li> <li>▪ Diplomatic missions and consular offices of the Republic of Uzbekistan, located outside the territory of the Republic of Uzbekistan</li> <li>▪ Digital signature key registration center</li> <li>▪ Operator of NIS "ASL BELGISI"</li> <li>▪ Participant of goods turnover</li> </ul>
Description of the actions to be performed	
1. Obtaining a TIN	
List of transferred information and the owner of the information resource	<p>Participant of the turnover of goods, which is a resident of the Republic of Uzbekistan, receives TIN in the procedure of registration (opening) of a legal entity.</p> <p>Participant of circulation of goods, which is a foreign holder of RU, in order to obtain TIN applies to the State Tax Committee of the Republic of Uzbekistan with the presentation of the following documents:</p> <ul style="list-style-type: none"> <li>▪ The document confirming the official registration of the legal entity in the country of location, issued by the competent state body (certificate of state registration or an extract from the trade register);</li> <li>▪ Documents confirming the registration (tax) number of the legal entity under the legislation of the state of registration or its equivalent;</li> <li>▪ Copy of the identity document (civil passport) of the head of the legal entity or the authorized representative;</li> <li>▪ Power of attorney issued to the authorized representative of the legal entity, indicating full passport data and powers granted to him/her (notarized);</li> <li>▪ Registration certificate for medicines, issued by the State Center for Expertise and</li> </ul>



	Standardization of Medicines, Medical Devices and Medical Equipment of the Agency for Development of Pharmaceutical Industry under the Ministry of Health of the Republic of Uzbekistan.
Special conditions	Providing of TIN to legal entities – to non-residents of the Republic of Uzbekistan is carried out only in relation to holders of registration certificates.
List of grounds for refusal	Appeal of a person who does not have the authority to take appropriate action. Failure to submit one or more documents.
<b>2. Obtaining a digital signature key certificate</b>	
List of transferred information and the owner of the information resource	<p>Participant of goods turnover, who is a resident of the Republic of Uzbekistan, receives an EDS key certificate through public service centers or official information resources online (remotely).</p> <p>The participant of goods turnover, being a foreign RU holder of the Republic of Uzbekistan, in order to obtain an EDS key certificate, applies to a diplomatic mission or consular office of the Republic of Uzbekistan located outside the territory of the Republic of Uzbekistan, presenting the following documents:</p> <ul style="list-style-type: none"><li>▪ an application for obtaining an EDS key certificate;</li><li>▪ certificate of entering information about the legal entity in the State Register of Business Entities of the Republic of Uzbekistan;</li><li>▪ identity document (civil passport);</li><li>▪ a power of attorney to represent the interests of a legal entity, indicating the full passport details of the authorized person and the powers granted (notarized);</li><li>▪ photograph (resolution not less than 297x382 pixels);</li></ul>



	<ul style="list-style-type: none"> <li>▪ payment document confirming the payment of the fee for the issuance of the EDS key certificate (paid locally);</li> <li>▪ electronic data storage device (USB FLASH drive)</li> </ul>
List of grounds for refusal	<p>The power of attorney must contain:</p> <ul style="list-style-type: none"> <li>▪ complete and correct passport details of the authorized person;</li> <li>▪ information that the authorized person is authorized to submit applications for obtaining an EDS key certificate.</li> </ul>
Special conditions	<p>The authorized person is assigned a PINFL (personal identification number of an individual).</p> <p>The EDS key certificate and the EDS key file are issued (recorded) on an electronic data storage device (USB FLASH drive).</p> <p>The password for using the EDS key file is automatically sent to the e-mail address specified in the application for obtaining the EDS key certificate.</p>
<b>3. Applying for registration in NIS "ASL BELGISI"</b>	
List of transferred information and the owner of the information resource	<p>Registration in the NIS "ASL BELGISI" of a participant in the turnover of goods is performed using an EDS key certificate, and requires the provision of the following information:</p> <ul style="list-style-type: none"> <li>▪ tax identifier of the participant of goods turnover (TIN) - filled in automatically from the EDS key certificate;</li> <li>▪ tax identifier of the representative of the participant of the goods turnover (PINFL) - filled in automatically from the EDS key certificate;</li> <li>▪ phone number of the contact person;</li> <li>▪ email address;</li> <li>▪ actual (postal) address;</li> <li>▪ bank details;</li> <li>▪ type of participant in the turnover of goods (list of product groups and type of participation in each product group).</li> </ul>





Special conditions	An application for registration in the NIS "ASL BELGISI" is signed by an EDS key certificate issued to the head of the enterprise or to a person who has a power of attorney to perform the relevant operations.
List of grounds for refusal	<ul style="list-style-type: none"> <li>▪ Lack of information about the participant of goods turnover in the State Register of Business Entities</li> <li>▪ Error in the EDS certificate (lack of data on the TIN of the participant in the circulation of goods and (or) PINFL of the representative of the participant in the circulation of goods)</li> <li>▪ Lack of authority of the representative of the participant in the circulation of goods (inconsistency between the PINFL of the owner of the EDS key certificate and the PINFL of the representative of the enterprise in the State Register of Business Entities)</li> <li>▪ Lack of a power of attorney to represent the interests of a legal entity when interacting with NIS "ASL BELGISI".</li> </ul>
4. Registration of the participant of goods turnover in NIS "ASL BELGISI" and provision of access to the personal cabinet (if there are no grounds for refusal)	
5. Notification of the applicant on registration in the NIS "ASL BELGISI" (if there are no grounds for refusal)	
6. Generation of contracts with the NIS Operator "ASL BELGISI"	
Special conditions	To interact with the NIS "ASL BELGISI", the participant in the turnover of goods is offered: <ul style="list-style-type: none"> <li>▪ application for joining the contract and contract for connection to the NIS "ASL BELGISI"</li> <li>▪ an application for joining the agreement and an agreement for the provision of services for the issuance and traceability of labeling codes necessary for the formation of digital identification tools (only for issuers of digital identification tools)</li> </ul>
7. Signing contracts with the NIS Operator "ASL BELGISI"	
List of grounds for refusal	Contracts are signed by a person without authority.
Special conditions	Contracts are signed in one of the following ways:



	<ul style="list-style-type: none"><li>▪ in electronic form through the use of an EDS key certificate</li><li>▪ on paper in two copies, one for each of the parties to the contract.</li></ul>
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## 5. Registration of information about labeled medicinal products in the "Catalog of Labeled Products" subsystem of NIS "ASL BELGISI"

Annotation	<p>For the purposes of issuance and traceability of means of digital identification the labeled goods are subject to registration in the subsystem "Catalog of labeled Goods" of NIS "ASL BELGISI".</p> <p>Information about the labeled drugs is entered by the holder of the registration certificate (its accredited representative office in the territory of the Republic of Uzbekistan).</p> <p>Information on labeled drugs is verified (moderated) by the NIS "ASL BELGISI" Operator, including through interaction with external information systems.</p> <p>If the verification is successful, information on the labeled MPs is signed by the participant of the turnover of goods using a digital signature key certificate and published on the public resource of the "Catalog of Labeled Goods" subsystem.</p>
Participants of interaction	<ul style="list-style-type: none"> <li>▪ Operator NIS "ASL BELGISI"</li> <li>▪ Operator MMMP</li> <li>▪ GS1 Uzbekistan Association</li> <li>▪ State Center for Expertise and Standardization of Medicines, Medical Devices and Medical Equipment of Agency on Development of Pharmaceutical Industry under the Ministry of Health of the Republic of Uzbekistan</li> <li>▪ Participant of goods turnover (RU holder or its accredited representative office on the territory of the Republic of Uzbekistan)</li> </ul>
Description of the actions to be performed	
1. Submission of information about MP to NIS "ASL BELGISI"	
List of transferred information and the owner of the information resource	<p>Holder of RU or its accredited representative office on the territory of the Republic of Uzbekistan enters information about the labeled MP in the following composition:</p>

	<ul style="list-style-type: none"> <li>▪ identification data of the participants of goods turnover (owner of goods code (GTIN), manufacturer, packer, holder of RU);</li> <li>▪ classification of medicinal products (in accordance with international and national classifiers);</li> <li>▪ MP identification (product code, trade name, trademark, country of production);</li> <li>▪ classification of drugs (ATC, INN, pharmacotherapeutic group);</li> <li>▪ Registration certificate of the drug (details);</li> <li>▪ authorization documents (requisites);</li> <li>▪ product package description;</li> <li>▪ consumer characteristics of MP;</li> <li>▪ weight and size characteristics of MP;</li> <li>▪ storage conditions of MP.</li> </ul>
2. Forming a draft MP card and filling in the description in NIS "ASL BELGISI".	
3. Notification of the participant of goods turnover about the creation of the draft MP card in NIS "ASL BELGISI".	
3. Preparation of requests for MP data in external information resources	
Processing, registration and verification of information received from external information resources	
5. Verification of the correctness (moderation) of information in the draft of the MP card in NIS "ASL BELGISI"	
List of grounds for refusal	<p>Incompleteness and/or incorrectness of the submitted data on MPs</p> <p>Inconsistency between the data in the MP card in NIS "ASL BELGISI" and the data on MP received from external information resources.</p> <p>The trafficker registering the MP card is not an RU holder or its accredited representative in the territory of the Republic of Uzbekistan</p>
Special conditions	Information about MP is registered by the holder of RU or its accredited representative in the Republic of Uzbekistan
6. Publication of the MP cards in NIS "ASL BELGISI" (if there are no grounds for refusal)	



List of transferred information and the owner of the information resource	The publication of a MP card in NIS "ASL BELGISI" is carried out after the successful verification (moderation) of the MP data by means of a digital signature key certificate.
List of grounds for refusal	Expiry of EDS key certificate



## 6. Submission of application for registration and traceability of digital identification means issued by the MMMP in NIS "ASL BELGISI"

<p>Annotation</p>	<p>Import (import) into the customs territory and subsequent circulation in the territory of the Republic of Uzbekistan of LP, labeled with digital identification means, issued by IS MMMP, is allowed after registration of such digital identification means in NIS "ASL BELGISI".</p> <p>Applications for registration and traceability of digital identification means issued by IS MMMP in NIS "ASL BELGISI" are submitted by the participant of goods turnover, carrying out the procedure of importing goods into the customs territory of the Republic of Uzbekistan before placing the labeled MP under the customs procedure of release for free circulation (import).</p> <p>Registration and traceability of means of digital identification issued by IS MMMP in NIS "ASL BELGISI", subject to the following conditions:</p> <ul style="list-style-type: none"> <li>▪ the participant of the turnover of goods engaged in the importation (importation) of labeled MPs is registered in NIS "ASL BELGISI";</li> <li>▪ information on labeled MPs imported into the Republic of Uzbekistan is registered in NIS "ASL BELGISI";</li> <li>▪ confirmation of the reliability of digital identification means and the possibility of their introduction into circulation in the territory of the Republic of Uzbekistan was received from the operator of the IS MMMP.</li> </ul>
<p>Participants of interaction</p>	<ul style="list-style-type: none"> <li>▪ Operator NIS "ASL BELGISI"</li> <li>▪ MMMP Operator</li> <li>▪ Participant of goods turnover that imports labeled products into the customs territory of the Republic of Uzbekistan</li> </ul>
<p>Description of the actions to be performed</p>	



<b>1. Application for registration and traceability of digital identification means, issued by the IS MMMP, in NIS "ASL BELGISI"</b>	
List of transferred information and the owner of the information resource	<p>For the purposes of registration and traceability of digital identification means issued by the IS MMMP in NIS "ASL BELGISI", the turnover participant submits the following data:</p> <ul style="list-style-type: none"><li>▪ TIN of the participant of goods turnover - filled in automatically on the basis of NIS "ASL BELGISI" data;</li><li>▪ Country - issuer of digital identification means - filled in automatically with the value "Russian Federation";</li><li>▪ sGTIN list.</li></ul> <p>The application is signed by the digital signature key certificate of the participant of the circulation of goods, which carries out the import of labeled products into the territory of the Republic of Uzbekistan.</p>
Special conditions	<p>The application is submitted in one of the following ways:</p> <ul style="list-style-type: none"><li>▪ through the personal account of the participant of goods turnover in NIS "ASL BELGISI";</li><li>▪ by using API methods.</li></ul> <p>One application may include the means of digital identification of one or more names of labeled products.</p>
<b>2. Verification of application for registration and traceability of digital identities issued by IS MMMP in NIS "ASL BELGISI"</b>	
List of grounds for refusal	<ul style="list-style-type: none"><li>▪ the participant of the turnover of goods, who submitted the application is not registered in NIS "ASL BELGISI" and (or) has an inactive status and (or) does not have an active product group "Medicinal products";</li><li>▪ EDS key certificate error (the EDS key certificate expired and (or) the EDS key certificate does not belong to a commodity trader and (or) the EDS key certificate has been revoked);</li></ul>



	<ul style="list-style-type: none"> <li>▪ the "Catalog of Labeled Goods" subsystem of NIS "ASL BELGISI" lacks information about one or more MP indicated in the application;</li> <li>▪ the application contains means of digital identification that do not comply with the established format;</li> <li>▪ the application contains not unique (repeated) means of digital identification;</li> <li>▪ one or more means of digital identification indicated in the application have already been registered with NIS "ASL BELGISI".</li> </ul>
<p>3. Verification of Digital Identification Means in the IS MMMP</p>	
<p>List of grounds for refusal</p>	<ul style="list-style-type: none"> <li>▪ One or more means of digital identification are not registered in the IS MMMP</li> <li>▪ One or more means of digital identification are not withdrawn from circulation in the IS MMMP</li> </ul>
<p>4. Processing of application for registration and traceability of digital identification means issued by the IS MMMP in NIS "ASL BELGISI"</p>	
<p>5. Registration of digital identification means, issued in IS MMMP in NIS "ASL BELGISI"</p>	
<p>Special conditions</p>	<p>In case of successful processing of the application:</p> <ul style="list-style-type: none"> <li>▪ the status of the document "Application for registration of IS MMMP codes" changes to "Processed successfully";</li> <li>▪ means of digital identification are registered in NIS "ASL BELGISI" with the status "Applied";</li> <li>▪ registered means of digital identification are displayed in the personal office of the participant of the goods turnover.</li> </ul>





## 7. Turnover of labeled medicinal products on the territory of the Republic of Uzbekistan (in accordance with the requirements of the Operator “ASL BELGISI”)

Annotation	<p>Turnover of labeled MP on the territory of the Republic of Uzbekistan is carried out in accordance with the legislation of the Republic of Uzbekistan.</p> <p>Registration of information on the introduction into circulation, turnover and withdrawal from turnover of labeled MPs in the NIS "ASL BELGISI" is carried out in accordance with:</p> <ul style="list-style-type: none"> <li>▪ Decree of the Cabinet of Ministers of the Republic of Uzbekistan No. 833 of December 31, 2020;</li> <li>▪ Resolution No. 149 of the Cabinet of Ministers of the Republic of Uzbekistan of April 2, 2022;</li> <li>▪ Other normative legal acts of the Republic of Uzbekistan;</li> <li>▪ recommendations and technological instructions of the NIS operator “ASL BELGISI”.</li> </ul>
Participants of interaction	<ul style="list-style-type: none"> <li>▪ Participants of the turnover of goods</li> <li>▪ NIS operator “ASL BELGISI”</li> <li>▪ Operator of electronic document roaming</li> <li>▪ Operator of fiscal data</li> <li>▪ State tax committee of the Republic of Uzbekistan</li> <li>▪ State Customs Committee of the Republic of Uzbekistan</li> </ul>
Description of the actions to be performed	
1. Performing the aggregation procedure for labeled MPs (if necessary)	
2. Customs clearance of labeled MP (according to customs regulations of the Republic of Uzbekistan)	
3. Submission of a Notice of importation of labeled MP	
4. Turnover of labeled MP in the wholesale and retail level	
5. Withdrawal of labeled MP from turnover	
6. Write-off of labeled MP	
Special conditions	Registration of information about the introduction into circulation, turnover and withdrawal from turnover of



	labeled MPs in NIS "ASL BELGISI" is carried out by the participants of circulation of goods in accordance with the legislation of the Republic of Uzbekistan.
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