

Format of the Pricelist of reimbursed medicinal products and foods for special medical purposes for reimbursement with order number starting from the level 4 or higher, SÚKL, version 21.0

No.	M/O	Label	Type	Size	Name	Description
1	O	KOD	C	7	SÚKL code	The code of the medicinal product (hereinafter referred to as "MP") allocated by SÚKL to the presentation of the MP as part of the marketing authorisation (MA) of the MP.
2	O	NAZ	C	70	Name of the MP	The name of the MP as referred to by SÚKL guideline REG-29, version 4, in compliance with Directive 2001/83/ES.
3	O	DOP	C	75	Specification of the MP	MP name supplement, which clearly defines the presentation of the MP, comprising of an integration of its pharmaceutical form, pack size, and strength. This item of the List is further specified in the items CESTA, FORMA, BALENI and SILA.
4	O	PORADI_UHRX	N	3	Reimbursement order number	Reimbursement order number starting from the level 4 or higher (designated as X).
5	O	UHRX	N	13,2	X-th Reimbursement	The amount of X-th reimbursement of the medicinal product determined by SÚKL as per Section 39d or 39db of Act No 48/1997 Coll. for the end consumer (JUHRX incremented with the maximum profit margin as per the price regulation of the Ministry of Health of the Czech Republic and VAT).
6	O	JUHRX	N	13,2	X-th Core reimbursement	The amount of X-th reimbursement of the medicinal product determined by SÚKL.
7	O	LEG_JUHRX	C	1	Legal basis for X-th core reimbursement	<p>The legal basis for the determination of the amount and conditions of X-th reimbursement of a MP/FSMP from health insurance; it may assume the following values:</p> <p>1 - The first temporary reimbursement of a highly innovative medicinal product as referred to by Section 39d of Act No 48/1997 Coll.;</p> <p>2 - The second temporary reimbursement of a highly innovative medicinal product as referred to by Section 39d of Act No 48/1997 Coll.;</p> <p>D – the amount and conditions of reimbursements for medicinal product included in the registry of orphan medicinal products determined pursuant to Section 39da of Act No 48/1997 Coll.;</p> <p>E - The decision on the amount and conditions of the first temporary reimbursement of a highly innovative medicinal product has not become final to date, and it is preliminarily enforceable;</p> <p>F - The decision on the amount and conditions of the second temporary reimbursement of a highly innovative medicinal product has not become final to date, and it is preliminarily enforceable;</p> <p>G – reimbursement of a medicinal product containing an active substance pursuant to Section 39 db of Act No. 48/1997 Coll.</p> <p>H – a decision on the reimbursement of a medicinal product containing an active substance pursuant to Section 39 db of Act No. 48/1997 Coll. has not yet become final and is provisionally enforceable.</p> <p>R – Medicinal products released from the reserve stock system according to Section 77g of the Act on Pharmaceuticals (the increase of the UHR by a special profit margin determined by the Price directive in compliance with Section 39h paragraph 1 of Act No. 48/1997 Coll.)</p> <p>T - Permanent reimbursement of a highly innovative medicinal product as per Section 39d, paragraph 4 of Act No 48/1997 Coll., in the version effective until 31.12.2021;</p> <p>X - The decision on the amount and conditions of reimbursement has not become final to date, and it is preliminarily enforceable;</p> <p>Y - The decision on the amount and conditions of permanent reimbursement of a highly innovative</p>

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						medicinal product under Section 39d, paragraph 4 of Act No 48/1997 Coll., in the version effective until 31.12.2021, has not become final to date, and it is preliminarily enforceable. Z - Other increased reimbursement determined as per Section 39b, paragraph 11 of Act No. 48/1997 Coll.
8	O	ODKAZ_JUHRX	C	20	Grounds for X-th core reimbursement amount determination	Contains the file no. of SÚKL administrative procedure.,
9	O	LIMX	C	2	X-th reimbursement reporting limit	The method of reporting MPs/FSMPs to the health insurance company. A common LIM status value implemental index is available for the LIMX fields.
10	O	OMEX	C	40	Prescribing doctor's specialisation	The specification of prescription restriction for X-th MP/FSMP reimbursement based on the specialization of the prescribing doctor. For a single MP/FSMP code it may assume several OMEX values. A common OME status value implemental index is available for the OMEX fields.
11	O	INDX	C	1	Indication restriction flag	Indication restriction (P) for the X-th reimbursement of an MP/FSMP. The DETINDX implemental index is available for the indication restriction detail (indication or clinical condition conditioning X-th reimbursement of the MP/FSMP). In respect of medicinal products reimbursed <i>ex lege</i> pursuant to Section 30 of Act No 48/1997 Coll., as amended, the particular provision of the Act is cited.
12	O	PUHRX	C	1	Full reimbursement flag	The full reimbursement flag may assume the following values: J - MPs where MFC <= UHRX. Note: Final sales under the pricing regulations of the Ministry of Health of the Czech Republic are disregarded. Where final sales for a higher price in compliance with a price regulation of the Ministry of Health of the Czech Republic is applied, the applied price for the end consumer may exceed the specified amount of reimbursement; U – MPs fully reimbursed under reimbursement agreements referred to under Section 39c, paragraph 2(d) of Act No 48/1997 Coll.
13	O	JUHRX_PLATDO	D	8	Temporary reimbursement expiry date	The temporary reimbursement is determined for the period of 24 or 12 months pursuant to Section 39d, paragraph 4 of Act No 48/1997 Coll., in the version effective until 31.12.2021 or for the period of 36 or 24 months pursuant to Section 39d, paragraph 4 of Act No 48/1997 Coll., in the version effective from 31.12.2021. The field is populated with the expiry date of the temporary reimbursement.
14	O	ODTDX	N	13,4	Usual daily therapeutic dose for X-th reimbursement	The usual daily therapeutic dose for X-th reimbursement
15	O	ODTDJX	C	5	Unit of active substance amount in ODTDJX	The usual daily therapeutic dose for reimbursement unit for ODTDJX
16	O	ODTDBALX	N	11,4	Number of ODTDJX in a MP pack	The number of usual therapeutic doses in a pack for ODTDJX.
17	O	ZU_ODTDX	N	13,4	Reimbursement for ODTDJX	X-th reimbursement of a MP for ODTDJX, determined by SÚKL as per Section 39d of Act No 48/1997 Coll.
18	O	ODKAZ_ZU_ODTDX	C	20	Grounds for EKVX determination	Contains the file no. of SÚKL administrative procedure.

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No.	M/O	Label	Type	Size	Name	Description
19	O	ZAPX	N	13,2	Eligible extra payment	An eligible extra payment for UHRX under a communication of the Ministry of Health of the Czech Republic, stipulated in compliance with Section 16b, paragraph 1 of Act No 48/1997 Coll.
20	O	NEZAPX	C	1	Limit eligibility symbol	A flag of the category of the pharmaceutical (eligibility for the limit) under a communication of the Ministry of Health of the Czech Republic, stipulated in compliance with Section 16b, paragraph 1 of Act No 48/1997 Coll. The NEZAP implemental index is available for the NEZAPX field.

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CZ Field separator "I"

The "M/O" column identifies mandatory and optional fields in the List

The "Type" column identifies the format of the fields as follows: "C" character attribute

"N" numeric attribute

"D" date in the "ddmmyyy" format

The "Size" column identifies the scope of the fields. The format of numeric fields is identified as "x,y" ("x" positions, incl. the decimal point, of which "y" are decimal)