

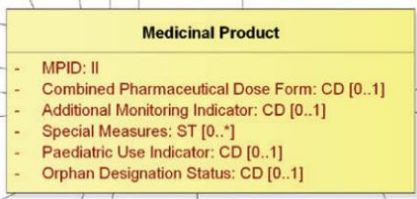




CUPRIOR Filmtabl 150 mg 72 Stk


Cuprior, Filmtabletten		
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Quelle		
Compendium	CUPRIOR Filmtabl 150 mg	
SAI	CUPRIOR Filmtabl 150 mg 72 Stk	

Medicinal Product		
MPID	CH-7640109110007-6771901	
Approved Dose Form (EXT)	FILM	Filmtabletten
		 <ul style="list-style-type: none"> - MPID: II - Combined Pharmaceutical Dose Form: CD [0..1] - Additional Monitoring Indicator: CD [0..1] - Special Measures: ST [0..*] - Paediatric Use Indicator: CD [0..1] - Orphan Designation Status: CD [0..1]

Product Classification		
ATC	A16AX12	
Heilmittelcode	S – Synthetika	
		 <ul style="list-style-type: none"> - Code System: CD.codeSystem - Value: CD.code

Medicinal Product Name		
Type (EXT)	Original	
Full Name	Cuprior 150 mg, Filmtabletten	
Invented Name Part	Cuprior	
Pharmaceutical Dose Form Part	Filmtabletten	
Strength Part	150 mg	
Delimiter	,	
		 <ul style="list-style-type: none"> - Full Name: ST - Invented Name Part: ENXP [0..*] - Scientific Name Part: ENXP [0..*] - Strength Part: ENXP [0..*] - Pharmaceutical Dose Form Part: ENXP [0..*] - Formulation Part: ENXP [0..*] - Intended Use Part: ENXP [0..*] - Target Population Part: ENXP [0..*] - Container or Pack Part: ENXP [0..*] - Device Part: ENXP [0..*] - Trademark or Company Name Part: ENXP [0..*] - Time / Period Part: ENXP [0..*] - Flavour Part: ENXP [0..*] - Delimiter Part: ENXP [0..*]

Country / Language		
Language	de	Deutsch
		 <ul style="list-style-type: none"> - Country: CD - Jurisdiction: CD [0..1] - Language: CD

Medicinal Product Name		
Type (EXT)	BAG	
Full Name	CUPRIOR Filmtabl 150 mg	
Invented Name Part	CUPRIOR	
Pharmaceutical Dose Form Part	Filmtabl	
Strength Part	150 mg	

Medicinal Product Name

- Full Name: ST
- Invented Name Part: ENXP [0..*]
- Scientific Name Part: ENXP [0..*]
- Strength Part: ENXP [0..*]
- Pharmaceutical Dose Form Part: ENXP [0..*]
- Formulation Part: ENXP [0..*]
- Intended Use Part: ENXP [0..*]
- Target Population Part: ENXP [0..*]
- Container or Pack Part: ENXP [0..*]
- Device Part: ENXP [0..*]
- Trademark or Company Name Part: ENXP [0..*]
- Time / Period Part: ENXP [0..*]
- Flavour Part: ENXP [0..*]
- Delimiter Part: ENXP [0..*]

Country / Language		
Language	de	Deutsch

Country / Language

- Country: CD
- Jurisdiction: CD [0..1]
- Language: CD

Marketing Authorisation (Medicinal Product)		
Marketing Authorisation Type (EXT)	Marktzulassung	
Marketing Authorisation Number	6771901	
Legal Status of Supply	B	Abgabe auf ärztliche oder tierärztliche Verschreibung (B)
Authorisation Status	Z	Zugelassen
Authorisation Status Date	21.01.2021	
Validity Period End	20.01.2026	
Date of First Authorisation	21.01.2021	

Marketing Authorisation

- Marketing Authorisation Number: II
- Country: CD [1..*]
- Legal Status of Supply: CD
- Authorisation Status: CD
- Authorisation Status Date: TS
- Validity Period Start: TS
- Validity Period End: TS
- Data Exclusivity Start Date: TS [0..1]
- Data Exclusivity End Date: TS [0..1]
- Date of First Authorisation: TS [0..1]
- International Birth Date: TS

Medicinal Product

- MPID: II
- Combined Pharmaceutical Dose Form: CD [0..1]
- Additional Monitoring Indicator: CD [0..1]
- Special Measures: ST [0..*]
- Paediatric Use Indicator: CD [0..1]
- Orphan Designation Status: CD [0..1]

Marketing Authorisation Holder (Organisation)		
Name	Curatis AG	
GLN	7640109110007	

Marketing Authorisation Holder (Organisation)

Hinweis: Pack Size, 72 Filmtabletten		
Packaged Medicinal Product		
PCID	CH-7640109110007-6771901-001	
Package Description	CUPRIOR Filmtabl 150 mg 72 Stk	
Pack Size (EXT)	72 Tablette(n)	

Packaged Medicinal Product

- PCID: II
- Package Description: ST

Data Carrier Identifier		
GTIN (Code System)	7680677190018	<div> Data Carrier Identifier <ul style="list-style-type: none"> - Code System: CD.codeSystem - Value: CD.code </div>

Packaged Item (Container)		
Package Item (Container) Type	30009000	Box
Package Item (Container) Quantity	1	
Material	Karton	

Package Item (Container)

- Package Item (Container) Type: CD
- Package Item (Container) Quantity: PQ
- Material: CD [1..*]
- Alternate Material: CD [0..*]

Packaged Item (Container)		
Package Item (Container) Type	30007000	Blister
Package Item (Container) Quantity	Unknown	
Material	Aluminium, Aluminium	

Package Item (Container)

- Package Item (Container) Type: CD
- Package Item (Container) Quantity: PQ
- Material: CD [1..*]
- Alternate Material: CD [0..*]

Packaged Item (Manufactured Item)		
Manufactured Item Quantity	72	Tabletten (per box)

Manufactured Item

- Manufactured Dose Form: CD
- Unit of Presentation: CD [0..1]
- Manufactured Item Quantity: PQ

Marketing Authorisation (Packaged Medicinal Product)		
Marketing Authorisation Type	tbd	
Marketing Authorisation Number	67719001	
Legal Status of Supply	B	Abgabe auf ärztliche oder tierärztliche Verschreibung (B)
Authorisation Status	Z	Zugelassen
Authorisation Status Date	21.01.2021	
Validity Period End	20.01.2026	

Marketing Authorisation

- Marketing Authorisation Number: II
- Country: CD [1..*]
- Legal Status of Supply: CD
- Authorisation Status: CD
- Authorisation Status Date: TS
- Validity Period Start: TS
- Validity Period End: TS
- Data Exclusivity Start Date: TS [0..1]
- Data Exclusivity End Date: TS [0..1]
- Date of First Authorisation: TS [0..1]
- International Birth Date: TS

Packaged Medicinal Product

- PCID: II
- Package Description: ST

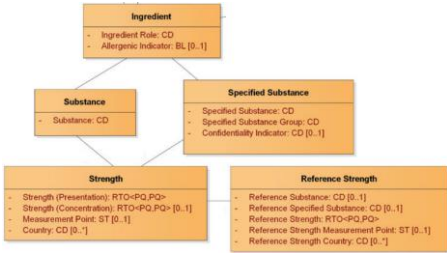
0..1

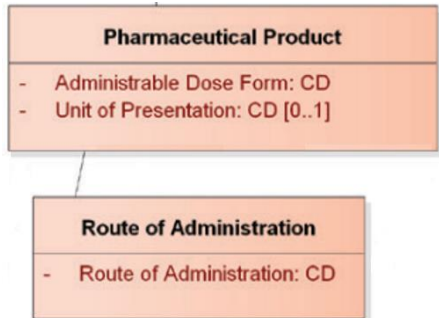
Shelf Life / Storage			
Shelf Life Type	100000073403	Closed Package	<div>Shelf Life / Storage</div> <ul style="list-style-type: none"> - Shelf Life Type: CD - Shelf Life Time Period: PQ - Special Precautions for Storage: CD [0..*]
Shelf Life Time Period	24 mo		
Special Precautions of Storage	NO30	Nicht über 30°C lagern	


Manufactured Item			
Manufactured Dose Form	10221000	Filmtablette	<div>Manufactured Item</div> <ul style="list-style-type: none"> - Manufactured Dose Form: CD - Unit of Presentation: CD [0..1] - Manufactured Item Quantity: PQ
Unit of Presentation	15054000	Tablette	

Ingredient (Manufactured Item) / Substance / Strength / Reference Strength			
Ingredient Role	WIRKS	Wirkstoff	<pre> graph TD Ingredient[Ingredient] --> Substance[Substance] Ingredient --> SpecifiedSubstance[Specified Substance] Substance --> Strength[Strength] SpecifiedSubstance --> ReferenceStrength[Reference Strength] </pre>
Substance	SJ76Y07H5F	Trientinum	
Strength (presentation)	150 mg (per UoP)		

Ingredient (Manufactured Item) / Substance / Strength / Reference Strength			
Ingredient Role	WIRKS	Wirkstoff	<pre> graph TD Ingredient[Ingredient] --> Substance[Substance] Ingredient --> SpecifiedSubstance[Specified Substance] Substance --> Strength[Strength] SpecifiedSubstance --> ReferenceStrength[Reference Strength] </pre>
Substance		Trientini tetrahydrochloridum	

Ingredient (Manufactured Item) / Substance / Strength / Reference Strength			
Ingredient Role	HNIDK	Hilfsstoff	 <pre> graph TD Ingredient[Ingredient] --> Substance[Substance] Ingredient --> SpecifiedSubstance[Specified Substance] Substance --> Strength[Strength] SpecifiedSubstance --> ReferenceStrength[Reference Strength] </pre> <p>Ingredient</p> <ul style="list-style-type: none"> - Ingredient Role: CD - Allergenic Indicator: BL [0..1] <p>Substance</p> <ul style="list-style-type: none"> - Substance: CD <p>Specified Substance</p> <ul style="list-style-type: none"> - Specified Substance: CD - Specified Substance Group: CD - Confidentiality Indicator: CD [0..1] <p>Strength</p> <ul style="list-style-type: none"> - Strength (Presentation): RTO=PQ,PQ> - Strength (Concentration): RTO=PQ,PQ> [0..1] - Measurement Point: ST [0..1] - Country: CD [0..1] <p>Reference Strength</p> <ul style="list-style-type: none"> - Reference Substance: CD [0..1] - Reference Specified Substance: CD [0..1] - Reference Strength: RTO=PQ,PQ> - Reference Strength Measurement Point: ST [0..1] - Reference Strength Country: CD [0..1]
Substance	3OWL53L36A	Mannitolum	
	ETJ7Z6XBU4	Silica colloidalis anhydrica	
	R8WTH25YS2	Glyceroli dibehenas	
	532B59J990	Poly (alcohol vinylicus)	
	7SEV7J4R1U	Talc Talcum	
	15FIX9V2JP	TITANII DIOXIDUM CL 77891 E 171	
	HYE7O27HAO	Glyceroli monocaprylocapras	
	EX438O2MRT	E 172 (flavum) YELLOW LAKE CLF 3076 (E104 AND E172)	
	368GB5141J	Natrii laurilsulfas	
	9NEZ333N27	0.057 mg Natrium	

Pharmaceutical Product / Route of Administration			
Administrable Dose Form	10221000	Filtablette	 <pre> graph TD PharmaceuticalProduct[Pharmaceutical Product] --> RouteOfAdministration[Route of Administration] </pre> <p>Pharmaceutical Product</p> <ul style="list-style-type: none"> - Administrable Dose Form: CD - Unit of Presentation: CD [0..1] <p>Route of Administration</p> <ul style="list-style-type: none"> - Route of Administration: CD
Unit of Presentation	15054000	Tablet	
Route of Administration	20053000	Oral use	
Methode of Administration (Swissmedic)	200000002052	Swallowing	

Therapeutic Indication			
Indication as	10061091	Copper metabolism disorder (Morbus Wilson)	 <pre> graph TD TherapeuticIndication[Therapeutic Indication] </pre> <p>Therapeutic Indication</p> <ul style="list-style-type: none"> - Indication Text: ST - Indication as "Disease / Symptom / Procedure": CD [0..1] - Disease Status: CD [0..1] - Comorbidity: CD [0..1] - Intended Effect: CD [0..1] - Timing / Duration: PQ [0..1]
Intended Effect	200000003194	Therapie	