REPUBLIC OF TÜRKİYE
MINISTRY OF HEALTH
MEDICINES AND MEDICAL
DEVICES AGENCY OF TÜRKİYE

Certificate No: TR/GMP/2024/151

CERTIFICATE OF GMP COMPLIANCE OF MANUFACTURER

Part 1

Issued following an inspection in accordance with current Good Manufacturing Practice Guidelines, and the Regulation on Manufacturing Plants of Medicinal Products for Human Use* and the Law No 1262 on Pharmaceutical and Medicinal Preparations. These regulations are in line with the requirements of Pharmaceutical Inspection Co-operation Scheme (PIC/s) and the Directives of the European Commission.

Turkish Medicines and Medical Devices Agency confirms the following:

Manufacturer's Name	: POLİFARMA İLAÇ SAN. VE TİC. A.Ş.
Head Office / Correspondence Address	: Vakıflar OSB Mah. Sanayi Cad. No:22/1 Ergene/TEKİRDAĞ
Site Address	: Vakıflar OSB Mahallesi Sanayi Caddesi No:22/1 Ergene/TEKİRDAĞ
Manufacturing Authorization Date	: TR/ÜY/2019/11-10
Manufacturing Authorization Number	: 01.11.2024

Has been inspected in accordance with current Good Manufacturing Practice Guidelines, the Regulation on Manufacturing Plants of Medicinal Products for Human Use, the Law No 1262 on Pharmaceutical and Medicinal Products.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 19-23.02.2024, it is considered that it complies with the requirements of Good Manufacturing Practice (GMP).

This certificate reflects the status of the manufacturing site at the time of the inspection, and Turkish Medicines and Medical Devices Agency should be consulted to verify compliance of the manufacturing site with GMP requirements if more than 3 years have elapsed since the date of inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field.

This certificate is valid only when presented with all pages and both Parts 1 and 2.

The authenticity of this certificate may be verified by Turkish Medicines and Medical Devices Agency upon request.

**This regulation is aligned with European Union Directive Directive 2003/94/EC laying down the principles and guidelines of good manufacturing practice for medicinal products for human use, and Directive 2001/83/EC on the Community code relating to medicinal products for human use.*

Eray KAPLAN
Vice President of the Agency

■ Human Medicinal Products

1 - MANUFACTURING OPERATIONS - MEDICINAL PRODUCTS	
If the company is engaged in manufacture of products with special requirements, e.g. radiopharmaceuticals or products containing penicillin, sulphanomides, cytotoxics, cephalosporins, substances with hormonal activity or other potentially hazardous active ingredients, this should be stated under the relevant product type and dosage form.	
1.1	**Sterile Products**
	1.1.2 Terminally sterilized (processing operations for the following dosage forms) 1.1.2.1 Large volume liquids - Solution for cardioplegia - Bladder irrigation - Solution for infusion - Solution for peritoneal dialysis - Irrigation solution - Emulsion for infusion 1.1.2.3 Small volume liquids - Emulsion for injection/infusion - Eye drops, emulsion
	1.1.3 Batch certification
1.2	**Non-sterile products**
	1.2.1 Non-sterile products (processing operations for the following dosage forms) 1.2.1.5 Liquids for external use - Inhalation solution
	1.2.2 Batch certification
1.3	**Biological medicinal products**
	1.3.2 Batch certification 1.3.2.2 Cell therapy products 1.3.2.6 Human or animal extract derived products
1.5	**Packaging**
	1.5.1 Primary Packaging 1.5.1.5 Liquids for external use
	1.5.2 Secondary packaging
1.6	**Quality control testing**
	1.6.1 Microbiological (sterility)
	1.6.2 Microbiological (non-sterility)
	1.6.3 Chemical/Physical
	1.6.4 Biological testing

Any restrictions or clarifying remarks related to the scope of this certificate:

1.1.2.1 Applicable to PVC bag, polypropylene bag, glass bottle and polypropylene bottle.

1.1.3 It has been found appropriate to allow parametric release activity for products named "Polifleks %0,9 İzotonik Sodyum Klorür I.V. İnfüzyon için Çözelti", "Polifleks %5 Dekstroz Sudaki I.V. İnfüzyon İçin Çözelti", "Polifleks Laktatlı Ringer I.V. İnfüzyon İçin Çözelti" (PP Torba) and "Polifleks İzolen Dengeli Elektrolit İ.V. İnfüzyon İçin Çözelti" (PP Torba).

1.3.2.2 Valid for batch release activities including batch control analyses for "CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Flakon ve CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Kullanıma Hazır Enjektör" products.

1.5.1.5: The inhalation solution is in an aluminum bottle primary packaging.

1.6.4 "CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Flakon ve CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Kullanıma Hazır Enjektör" products are valid for serial control analysis.

Eray KAPLAN
Vice President of the Agency

2 - IMPORTATION OF MEDICINAL PRODUCTS	
2.1	**Quality control testing of imported medicinal products**
	2.1.4 Biological
2.2	**Batch certification of imported medicinal products**
	2.2.3 Biological medicinal products 2.2.3.2 Immunological products

Any restrictions or clarifying remarks related to the scope of this certificate:
2.1.4 "CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Flakon ve CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Kullanıma Hazır Enjektör" products are valid for serial control analysis.
2.2.3.2 It is valid for batch release activities that include batch control analyzes for "CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Flakon ve CoronaVac 600SU/0,5ml iM Enjeksiyonluk Süspansiyon içeren Kullanıma Hazır Enjektör" products.

■ Human Investigational Medicinal Products (for Phase I, II, III Clinical trials)

1 - MANUFACTURING OPERATIONS - MEDICINAL PRODUCTS	
If the company is engaged in manufacture of products with special requirements, e.g. radiopharmaceuticals or products containing penicillin, sulphanomides, cytotoxics, cephalosporins, substances with hormonal activity or other potentially hazardous active ingredients, this should be stated under the relevant product type and dosage form.	
1.1	**Sterile Products**
	1.1.2 Terminally sterilized (processing operations for the following dosage forms) 1.1.2.1 Large volume liquids - Solution for cardioplegia - Bladder irrigation - Solution for infusion - Solution for peritoneal dialysis - Irrigation solution - Emulsion for infusion 1.1.2.3 Small volume liquids - Emulsion for injection/infusion
	1.1.3 Batch certification
1.2	**Non-sterile products**
	1.2.1 Non-sterile products (processing operations for the following dosage forms) 1.2.1.5 Liquids for external use - Inhalation solution
	1.2.2 Batch certification
1.3	**Biological medicinal products**
	1.3.2 Batch certification 1.3.2.2 Cell therapy products 1.3.2.6 Human or animal extract derived products
1.5	**Packaging**
	1.5.1 Primary Packaging 1.5.1.5 Liquids for external use
	1.5.2 Secondary packaging
1.6	**Quality control testing**
	1.6.1 Microbiological (sterility)
	1.6.2 Microbiological (non-sterility)
	1.6.3 Chemical/Physical
	1.6.4 Biological testing

Eray KAPLAN
Vice President of the Agency

Any restrictions or clarifying remarks related to the scope of this certificate:
1.1.2.1 Applicable to PVC bag, polypropylene bag, glass bottle and polypropylene bottle.
1.1.3 It has been found appropriate to allow parametric release activity for products named "Polifleks %0,9 İzotonik Sodyum Klorür I.V. İnfüzyon için Çözelti", "Polifleks %5 Dekstroz Sudaki I.V. İnfüzyon İçin Çözelti", "Polifleks Laktatlı Ringer I.V. İnfüzyon İçin Çözelti" (PP Torba) and "Polifleks İzolen Dengeli Elektrolit İ.V. İnfüzyon İçin Çözelti" (PP Torba).
1.3.2.2 Valid for batch release activities including batch control analyses for "CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Flakon ve CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Kullanıma Hazır Enjektör" products.
1.5.1.5: The inhalation solution is in an aluminum bottle primary packaging.
1.6.4 "CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Flakon ve CoronaVac 600SU/0,5ml IM Enjeksiyonluk Süspansiyon içeren Kullanıma Hazır Enjektör" products are valid for serial control analysis.

11.11.2024 TR/GMP/2024/151

Eray KAPLAN
Vice President of the Agency

Adres: Söğütözü Mahallesi 2176. Sok. No:5 06520 Çankaya/ANKARA
Tel: (0312) 218 30 00 Fax: (0312) 218 34 60

Bulgarian Drug Agency

CERTIFICATE NUMBER: ***BG/GMP/2025/308***

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER [1,2]

Part 1

Issued following an inspection in accordance with
Art. 111(5) of Directive 2001/83/EC as amended

The competent authority of Bulgaria confirms the following:
The manufacturer: ***Polifarma Ilac Sanayi Ve Ticaret A.S.***
Site address: ***Vakiflar Osb Mahallesi, Sanayi Caddesi No 22/1, Ergene, 59930, Turkey***
OMS Organisation Id. / OMS Location Id.: ***ORG-100040713*** / ***LOC-100066953***

Has been inspected in connection with marketing authorisation(s) listing manufacturers located outside of the European Economic Area in accordance with Art. 111(4) of Directive 2001/83/EC.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on ***2024-11-14***, it is considered that it complies with:

- The principles and guidelines of Good Manufacturing Practice laid down in Directive (EU) 2017/1572. [3]

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field. Updates to restrictions or clarifying remarks can be identified through the EudraGMDP website (http://eudragmdp.ema.europa.eu/). This certificate is valid only when presented with all pages and both Parts 1 and 2.

The authenticity of this certificate may be verified in EudraGMDP. If it does not appear, please contact the issuing authority.

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[1] *The certificate referred to in paragraph Art. 111(5) of Directive 2001/83/ECis also applicable to importers.*

[2] *Guidance on the interpretation of this template can be found in the Interpretation of the Union format for GMP certificate.*

[3] *These requirements fulfil the GMP recommendations of WHO.*

Part 2

Human Medicinal Products

1 MANUFACTURING OPERATIONS	
1.1	**Sterile products**
	1.1.2 Terminally Sterilised (processing operations for the following dosage forms) 1.1.2.1 Large volume liquids
	1.1.3 Batch certification
1.2	**Non-sterile products**
	1.2.1 Non-sterile products (processing operations for the following dosage forms) 1.2.1.5 Liquids for external use Special Requirements 7 Other: inhalation vapour, liquid(en)
	1.2.2 Batch certification
1.5	**Packaging**
	1.5.1 Primary Packaging 1.5.1.5 Liquids for external use Special Requirements 7 Other: inhalation vapour, liquid(en)
	1.5.2 Secondary packaging
1.6	**Quality control testing**
	1.6.1 Microbiological: sterility *1.6.2 Microbiological: non-sterility* *1.6.3 Chemical/Physical*

Clarifying remarks (for public users)

This certificate includes also manufacturing operation of bulk product – emulsion for injection/infusion.

2025-04-10

Name and signature of the authorised person of the Competent Authority of Bulgaria

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Confidential
Bulgarian Drug Agency
Tel:***Confidential***
Fax:***Confidential***

T.C. SAĞLIK BAKANLIĞI
TÜRKİYE İLAÇ VE TIBBİ CİHAZ KURUMU

RUHSATLI BEŞERİ TIBBİ ÜRÜNLER LİSTESİ

İRD-LST-15 /31.12.2019/Rev.01/17.11.2023

SIRA NO	BARKOD	ÜRÜN ADI	ETKİN MADDE	ATC KODU	RUHSAT SAHİBİ	RUHSAT TARİHİ	RUHSAT NUMARASI	DEĞİŞİKLİK	Bu hafta değişiklik yapılan ürünler ile belirtilmiştir. Yapılan Değişiklikler için DEĞİŞİKLİK kolonuna bakınız.	DEĞİŞİKLİK TARİHİ	RUHSATI ASKIDA OLMAYAN ÜRÜN: 0 MADDE-23 GEREKÇELİ ASKIDA OLAN ÜRÜN: 1 FARMAKOVİJİL/	ASKIYA ALINMA TARİHİ
19055	8681293079137	ACNEFFERİN %0,1 JEL, 1 ADET (30 G)	adapalen	D10AD03	MEGA FARMA İLAÇ VE KOZMETİK SAN. TİC. PAZ	12.07.2020	2020/149			21.08.2020	0	
19056	8699525529824	DULAMON 200 MCG/5 MCG BASINÇLI İNHALASYON, SÜSPANSİYON, 60 DOZ	mometazon furoat, formoterol fumarat	R03AK09	DEVA HOLDİNG A.Ş.	12.06.2019	2019/286			21.06.2019	0	
19057	8699525529831	DULAMON 200 MCG/5 MCG BASINÇLI İNHALASYON, SÜSPANSİYON, 120 DOZ	mometazon furoat, formoterol fumarat	R03AK09	DEVA HOLDİNG A.Ş.	12.06.2019	2019/286			21.06.2019	0	
19058	8699525529800	DULAMON 100 MCG/5 MCG BASINÇLI İNHALASYON, SÜSPANSİYON, 60 DOZ	mometazon furoat, formoterol fumarat	R03AK09	DEVA HOLDİNG A.Ş.	12.06.2019	2019/285			21.06.2019	0	
19059	8699525529817	DULAMON 100 MCG/5 MCG BASINÇLI İNHALASYON, SÜSPANSİYON, 120 DOZ	mometazon furoat, formoterol fumarat	R03AK09	DEVA HOLDİNG A.Ş.	12.06.2019	2019/285			21.06.2019	0	
19060	8680698070435	REMEMBA 5 MG/20 MG DAĞILABİLİR TABLET, 28 ADET	donepezil hcl, memantin hcl	N06DA52	ARDİ FARMA İLAÇ PAZARLAMA TİC. LTD. STİ	12.06.2019	2019/289			24.09.2021	1	07.09.2021
19061	8699580070033	REMEMBA 5 MG/10 MG DAĞILABİLİR TABLET, 28 ADET	donepezil hcl, memantin hcl	N06DA52	DROGSAN İLAÇLARI SAN. VE TİC. A.Ş.	06.04.2022	2022/208			15.04.2022	0	
19062	8699580070019	REMEMBA 5 MG/5 MG DAĞILABİLİR TABLET, 10 ADET	donezepril hcl, memantin hcl	N06DA52	DROGSAN İLAÇLARI SAN. VE TİC. A.Ş.	06.04.2022	2022/207			15.04.2022	0	
19063	8699580070026	REMEMBA 5 MG/5 MG DAĞILABİLİR TABLET, 28 ADET	donezepril hcl, memantin hcl	N06DA52	DROGSAN İLAÇLARI SAN. VE TİC. A.Ş.	06.04.2022	2022/207			15.04.2022	0	
19064	8699580070057	REMEMBA 10 MG/20 MG DAĞILABİLİR TABLET, 28 ADET	donezepril hcl, memantin hcl	N06DA52	DROGSAN İLAÇLARI SAN. VE TİC. A.Ş.	11.04.2022	2022/217			29.04.2022	0	
19065	8699580070040	REMEMBA 10 MG/10 MG DAĞILABİLİR TABLET, 28 ADET	donepezil hcl, memantin hcl	N06DA52	DROGSAN İLAÇLARI SAN. VE TİC. A.Ş.	11.04.2022	2022/218			29.04.2022	0	
19066	8699606076520	FLUSİBLE 500 MG/5 ML IV ENJEKSİYONLUK ÇÖZELTİ, 1 ADET	fluoresein sodyum	S01JA01	POLİFARMA İLAÇ SAN. VE TİC. A.Ş.	17.06.2019	2019/293			28.06.2019	0	
19067	8699565010016	LİNRA 1MG TABLET, 30 TABLET	rasajilin mesilat	N04BD02	ANGELİNİ İLAÇ SAN. VE TİC. A.Ş.	24.05.2019	2019/274			28.06.2019	0	
19068	8699527090872	OXFET 90 MG FİLM KAPLI TABLET , 30 TABLET	deferasiroks	V03AC03	SANTA FARMA İLAÇ SAN. A.Ş.	29.05.2019	2019/111			28.06.2019	0	
19069	8699527090889	OXFET 90 MG FİLM KAPLI TABLET , 90 TABLET	deferasiroks	V03AC03	SANTA FARMA İLAÇ SAN. A.Ş.	29.05.2019	2019/111			28.06.2019	0	
19070	8699527090896	OXFET 180 MG FİLM KAPLI TABLET , 30 TABLET	deferasiroks	V03AC03	SANTA FARMA İLAÇ SAN. A.Ş.	29.05.2019	2019/112			28.06.2019	0	
19071	8699527090902	OXFET 180 MG FİLM KAPLI TABLET , 90 TABLET	deferasiroks	V03AC03	SANTA FARMA İLAÇ SAN. A.Ş.	29.05.2019	2019/112			28.06.2019	0	
19072	8699527090919	OXFET 360 MG FİLM KAPLI TABLET , 30 TABLET	deferasiroks	V03AC03	SANTA FARMA İLAÇ SAN. A.Ş.	29.05.2019	2019/114			28.06.2019	0	
19073	8699527090916	OXFET 360 MG FİLM KAPLI TABLET , 90 TABLET	deferasiroks	V03AC03	SANTA FARMA İLAÇ SAN. A.Ş.	29.05.2019	2019/114			28.06.2019	0	
19074	8699514020196	RASTEL 50 MG EFERVESAN TABLET, 20 ADET	deksketoprofen trometamol	M01AE17	ABDİ İBRAHİM İLAÇ SAN. VE	17.06.2019	2019/294			28.06.2019	0	
19075	8699514020202	RASTEL 50 MG EFERVESAN TABLET, 30 ADET	deksketoprofen trometamol	M01AE17	ABDİ İBRAHİM İLAÇ SAN. VE	17.06.2019	2019/294			28.06.2019	0	
19076	8680199599190	ROSTALEPT 1 MG/ML ORAL ÇÖZELTİ, 100 ML	risperidon	N05AX08	WORLD MEDICINE İLAÇ SAN. VE TİC. A.Ş.	12.06.2019	2019/292			28.06.2019	0	
19077	8699844242145	POT-OUT 880 MG/1 G GRANÜL, 20 ADET	kalsiyum polistiren sülfonat	V03AE01	VEM İLAÇ SAN. VE TİC. A.Ş.	18.06.2019	2019/296			28.06.2019	0	
19078	8699844242169	POT-OUT 880 MG/1 G GRANÜL, 60 ADET	kalsiyum polistiren sülfonat	V03AE01	VEM İLAÇ SAN. VE TİC. A.Ş.	18.06.2019	2019/296			28.06.2019	0	
19079	8699844242176	POT-OUT 880 MG/1 G GRANÜL, 120 ADET	kalsiyum polistiren sülfonat	V03AE01	VEM İLAÇ SAN. VE TİC. A.Ş.	18.06.2019	2019/296			28.06.2019	0	
19080	8699638754540	LYSTHENON FORTE %2 ENJEKSİYONLUK ÇÖZELTİ, 25 AMPUL	suksametonyum klorür	M03AB01	TEVA İLAÇLARI SAN.VE TİC. A.Ş.	18.11.1972	113/76			12.07.2019	0	
19081	8699514570318	FENYSOL-Z 40 MG + 15 MG/5 ML ŞURUP, 100 ML	demir (ıı) glukonat, çinko	B03AE10	ABDİ İBRAHİM İLAÇ SAN. VE	24.04.2019	2019/234			09.08.2019	0	

T.C. SAĞLIK BAKANLIĞI
TÜRKİYE İLAÇ VE TIBBİ CİHAZ KURUMU

RUHSATLI BEŞERİ TIBBİ ÜRÜNLER LİSTESİ

İRD-LST-15
/31.12.2019/Rev.01/17.
11.2023

SIRA NO	BARKOD	ÜRÜN ADI	ETKİN MADDE	ATC KODU	RUHSAT SAHİBİ	RUHSAT TARİHİ	RUHSAT NUMARASI	DEĞİŞİKLİK	Bu hafta değişiklik yapılan ürünler ile belirtilmiştir. Yapılan Değişiklikler için DEĞİŞİKLİK kolonuna bakınız.	DEĞİŞİKLİK TARİHİ	RUHSATI ASKIDA OLMAYAN ÜRÜN: 0 MADDE-23 GEREKÇELİ ASKIDA OLAN ÜRÜN: 1 FARMAKOVİJİL	ASKIYA ALINMA TARİHİ
11451	8699525099310	OMETAN 40 MG FILM TABLET,FILM TABLET, 28 ADET	olmesartan medoksomil	C09CA08	DEVA HOLDİNG A.Ş.	10.03.2014	2014/207				0	
11452	8699525099327	OMETAN 40 MG FILM TABLET,FILM TABLET, 84 ADET	olmesartan medoksomil	C09CA08	DEVA HOLDİNG A.Ş.	10.03.2014	2014/207				0	
11453	8699525099341	OMETAN 5 MG FILM TABLET,FILM TABLET, 28 ADET	olmesartan medoksomil	C09CA08	DEVA HOLDİNG A.Ş.	17.04.2014	2014/320				0	
11454	8699525099358	OMETAN 5 MG FILM TABLET,FILM TABLET, 84 ADET	olmesartan medoksomil	C09CA08	DEVA HOLDİNG A.Ş.	17.04.2014	2014/320				0	
11455	8699525099402	OMETAN PLUS 20 MG/12.5 MG FILM TABLET,FILM TABLET, 28 ADET	olmesartan medoksomil + hidroklorotiyazid	C09DA08	DEVA HOLDİNG A.Ş.	01.08.2014	2014/602				0	
11456	8699525099419	OMETAN PLUS 20 MG/12.5 MG FILM TABLET,FILM TABLET, 84 ADET	olmesartan medoksomil + hidroklorotiyazid	C09DA08	DEVA HOLDİNG A.Ş.	01.08.2014	2014/602				0	
11457	8699525099426	OMETAN PLUS 20 MG/25 MG FILM TABLET,FILM TABLET, 28 ADET	olmesartan medoksomil + hidroklorotiyazid	C09DA08	DEVA HOLDİNG A.Ş.	01.08.2014	2014/603				0	
11458	8699525099433	OMETAN PLUS 20 MG/25 MG FILM TABLET,FILM TABLET, 84 ADET	olmesartan medoksomil + hidroklorotiyazid	C09DA08	DEVA HOLDİNG A.Ş.	01.08.2014	2014/603				0	
11459	8699525099440	OMETAN PLUS 40 MG/12.5 MG FILM TABLET,FILM TABLET, 28 ADET	olmesartan medoksomil + hidroklorotiyazid	C09DA08	DEVA HOLDİNG A.Ş.	01.08.2014	2014/604			25.05.2018	0	
11460	8699525099457	OMETAN PLUS 40 MG/12.5 MG FILM TABLET,FILM TABLET, 84 ADET	olmesartan medoksomil + hidroklorotiyazid	C09DA08	DEVA HOLDİNG A.Ş.	01.08.2014	2014/604			25.05.2018	0	
11461	8699525099464	OMETAN PLUS 40 MG/25 MG FILM TABLET,FILM TABLET, 28 ADET	olmesartan medoksomil + hidroklorotiyazid	C09DA08	DEVA HOLDİNG A.Ş.	01.08.2014	2014/605			25.05.2018	0	
11462	8699525099471	OMETAN PLUS 40 MG/25 MG FILM TABLET,FILM TABLET, 84 ADET	olmesartan medoksomil + hidroklorotiyazid	C09DA08	DEVA HOLDİNG A.Ş.	01.08.2014	2014/605			25.05.2018	0	
11463	8681735790163	OMEX 40 MG I.V. ENJEKSIYON IÇIN LIYOFILIZE TOZ IÇEREN FLAKO, FLAKON 1 ADET + 1 ÇÖZÜCÜ AMPUL	omeprazol sodyum	A02BC01	CENTURİON İLAÇ SAN. VE TİC. A.Ş.	21.06.2017	2017/447				0	
11464	8699688771450	OMNIPAQUE 300 MG I/ML IA IV INTRATEKAL ENJEKSİYON İÇİN ÇÖZELTİ, 100 ML	iohexol	V08AB02	OPAKİM TIBBİ ÜRÜNLER SAN. VE TİC. A.Ş.	06.07.1995	97/40				0	
11465	8699688771443	OMNIPAQUE 300 MG I/ML IA IV INTRATEKAL ENJEKSİYON İÇİN ÇÖZELTİ, 50 ML	iohexol	V08AB02	OPAKİM TIBBİ ÜRÜNLER SAN. VE TİC. A.Ş.	06.07.1995	97/40				0	
11466	8699688771559	OMNIPAQUE 350 MG I/ML IA IV INTRATEKAL ENJEKSİYON İÇİN ÇÖZELTİ, 100 ML	iohexol	V08AB02	OPAKİM TIBBİ ÜRÜNLER SAN. VE TİC. A.Ş.	06.07.1995	97/41				0	
11467	8699688771603	OMNIPAQUE 350 MG I/ML IA IV INTRATEKAL ENJEKSİYON İÇİN ÇÖZELTİ, 200 ML	iohexol	V08AB02	OPAKİM TIBBİ ÜRÜNLER SAN. VE TİC. A.Ş.	06.07.1995	97/41			07.05.2021	0	
11468	8699688771542	OMNIPAQUE 350 MG I/ML IA IV INTRATEKAL ENJEKSİYON İÇİN ÇÖZELTİ, 50 ML	iohexol	V08AB02	OPAKİM TIBBİ ÜRÜNLER SAN. VE TİC. A.Ş.	06.07.1995	97/41				0	
11469	8699606775560	OMNIPOL 300 MGI/ML IA, IV, INTRATEKAL ENJEKSIYON IÇIN ÇÖZELTI IÇEREN FLAKON, 100 ML	ioheksol	V08AB02	POLİFARMA İLAÇ SAN. VE TİC. A.Ş.	21.06.2016	2016/497				0	
11470	8699606775553	OMNIPOL 300 MGI/ML IA, IV, INTRATEKAL ENJEKSIYON IÇIN ÇÖZELTI IÇEREN FLAKON, 50 ML	ioheksol	V08AB02	POLİFARMA İLAÇ SAN. VE TİC. A.Ş.	21.06.2016	2016/497				0	
11471	8699606775584	OMNIPOL 350 MGI/ML IA, IV ENJEKSIYON IÇIN ÇÖZELTI IÇEREN FLAKON, 100 ML	ioheksol	V08AB02	POLİFARMA İLAÇ SAN. VE TİC. A.Ş.	21.06.2016	2016/496				0	
11472	8699606775591	OMNIPOL 350 MGI/ML IA, IV ENJEKSIYON IÇIN ÇÖZELTI IÇEREN FLAKON, 200 ML	ioheksol	V08AB02	POLİFARMA İLAÇ SAN. VE TİC. A.Ş.	21.06.2016	2016/496				0	
11473	8699606775577	OMNIPOL 350 MGI/ML IA, IV ENJEKSIYON IÇIN ÇÖZELTI IÇEREN FLAKON, 50 ML	ioheksol	V08AB02	POLİFARMA İLAÇ SAN. VE TİC. A.Ş.	21.06.2016	2016/496				0	
11474	8699688770507	OMNISCAN 0.5 MMOL/ML IV ENJEKSİYON İÇİN ÇÖZELTİ İÇEREN FLAKON, 10 ML	gadodiamid	V08CA03	OPAKİM TIBBİ ÜRÜNLER SAN. VE TİC. A.Ş.	29.09.1995	98/5			26.10.2018	3	01.03.2018
11475	8699688770514	OMNISCAN 0.5 MMOL/ML IV ENJEKSİYON İÇİN ÇÖZELTİ İÇEREN FLAKON, 15 ML	gadodiamid	V08CA03	OPAKİM TIBBİ ÜRÜNLER SAN. VE TİC. A.Ş.	29.09.1995	98/5			26.10.2018	3	01.03.2018
11476	8699688770521	OMNISCAN 0.5 MMOL/ML IV ENJEKSİYON İÇİN	gadodiamid	V08CA03	OPAKİM TIBBİ ÜRÜNLER	29.09.1995	98/5			26.10.2018	3	01.03.2018