

we produce
for health with

58
years
experience



www.tumekip.com.tr

 **TÜM * EKİP**
İLAÇ A.Ş.

About

“ In 2000 Tüm Ekip Pharmaceuticals is established by an Equip (team) of experts on their fields under the management of Mr. Hamza Oğuz who has 58 years of experience in Pharma sector. ”

Before establishing Tum Ekip group in 2000, between 1960 –1980 he was CEO and General Manager of the biggest pharmaceutical distribution channel of whole saler group of Turkey.

Between 1981–2000 he was the CEO and shareholder of Pharmaceutical Producer Company among the first 5 of Turkish Pharma sector. As a young company, we had a secure, healthy and successful growth in a short time. Currently we have over product licenses in 100 including all the pharmaceutical presentations, and new products in the registration pipeline. Being Turkey's only Local and National specialized Injectable pharmaceutical producer; we are among leader pharmaceutical companies. We have a large expertise on development and production of all injectable forms ;including powder, liquid, lyophilized powder, SVP, IV solutions bag.

We are also unique on the production capabilities as we have 3 completely separated /dedicated sterile production area :Sterile Injectable Cephalosporin, Sterile Injectable Carbapenem, Sterile injectable Penicillin.

Additionally, with our sterile liquid ampoule, sterile liquid vial, sterile liquid bag, and sterile lyophilized injectable powder production facility, we are the only Specialized Hospital Pharmaceuticals producer who has the capacity and capability to produce 7 different sterile forms in Turkey and even among EU countries.



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OUR PRODUCTION FACILITIES



Our production facilities are located in Tuzla, Istanbul Turkey. The designs and validation procedures are compliant with EU GMP rules and conform to EMA (European Medicines Agency). Our facility is located within the territory of a PIC/S Participating Authority and have been inspected regularly to PIC/S standards. We are experienced on production of injectable products especially antibiotics, by the beginning of 2011 we are the only pharmaceutical company who have 3 completely separated/dedicated sterile production area in Turkey. On 2009 dedicated sterile injectable Cephalosporin production area started to production. The annual production capacity is 20 million units. On 2010 dedicated sterile injectable Carbapenem production area started to production in a separated building. The annual production capacity is 10 million units. On 2011 dedicated sterile injectable Penicilin production area started to production in a separated building. The annual production capacity is 20 million units. On II. Quarter 2015 sterile liquid production area started production. The annual production capacity will be 20 million units.

Mission

Our Mission

Our commitment as a team is to contribute to the general health consciousness in the region where we operate, while growing by developing professional and social collaboration with related persons and organisations, giving priority to health professionals and our business partners and by producing creative solutions.

Vision

Our Vision

To have ethical, positive and transparent relations.
To maintain a continuous organizational and personnel learning and development.
To maintain determinate, stable and systematic working.
To analyze the future by knowing the past and following the actual up closely.
To be open to changes, to be flexible.
To take fast and right decisions.

Team

Our Team

Core Personnel of Tum Ekip Pharmaceuticals are expert in Pharmaceutical Sector. These people has formed a powerful team (equip) which becomes a symbol in our company name and logo with their experiences on different expertise areas.





REGISTRATION AND R&D

The Research and Development department has a unique role in attaining both short and long term goals of Tum Ekip Pharmaceuticals as a research and development oriented generic pharmaceutical company. Regulatory Affairs and Product Development departments of the Company together develop the new registration files for the new products. Tum Ekip laboratories and all the production facilities supports these projects accordingly. The main purpose of R&D department is to develop equivalent pharmaceuticals needed in clinics and to present high quality, safe and affordable generic solutions, attaching paramount importance to quality in pharmaceuticals and to ensuring such quality during the design process. To this end, the R&D department develops injectable dosage forms in compliance with the guidelines of International Conference on Harmonization (ICH). The R&D laboratories, including the pilot manufacturing lab, are equipped with state-of the-art technology and the experiments are conducted in accordance with Good Manufacturing Practices (GMP) and Good Laboratory Practices (GLP) .

OUR LABORATORIES



Our Laboratories are conform to cGLP and is established as two departments Chemical Control Laboratory and Microbiological Control Laboratory. Our laboratories has all the modern equipment to realize all the controls and analysis of API and finished products in house: as HPLCs, GC, TOC, FTIR, Spectrophotometer, particule analyser and others in Chemical Control Laboratory and Sterile test Room, safety LAF Cabines, autoclaves, Incubators and others in Microbiological Laboratory.

R&D AND PROJECTS



Regulatory Affairs and Product Development' departments of Company has received some R&D projects support from the Scientific and Technological Research Agency Concl of Turkey and SME. Between 2018- 2023 new products planned with project application.

WHOLESALE & DISTRIBUTION



Warehouse to provide operational excellence cGMP (current Good Manufacturing Practices) cGWP (current Good Storage Practice) GDP (Good Distribution Practice) applications under the system of our company's Pharmaceutical Operations Quality Assurance (QA-PO) and in accordance with regulations, storage, packaging and shipping activities are carried out. All the our products are marked with track and trace system. Each product are labeled with a single, unique alphanumeric 2D barcode number.

Our QA SYSTEM

Our QA SYSTEM



Registration Regulations of Turkey has been prepared in line with the directive no : 2001/83/EC on Medicinal Products for Human Use, for the purpose of harmonising with the relevant legislations of the European Union pertaining for the Medicinal Products for Human Use. We have build EU standard facilities QA and QC system. All the validations, calibrations and qualifications are done regularly and under the control of independent QA management. All critical computerized systems are validated with the support of independent agreed consultants, to be complied with GAMP5 (Good Automated Manufacturing Practices 5) The essence of the system is to be complied with all cGXP rules. All procedures according the written standard operating procedures (S.O.P.) for all work carried out in the Company and Warehouse and taking all the transactions recorded are documented. All production, control, storage, packaging and shipping activities are guaranteed by the Quality Assurance System. ISO 9001 and ISO 14001 certificates are also available as a part of our quality system.

TURKISH MINISTRY OF HEALTH
Turkish Medicines and Medical Devices Agency

CERTIFICATE OF GMP (GOOD MANUFACTURING PRACTICE) COMPLIANCE OF A MANUFACTURER

Part 1

Issued following an inspection in accordance with current Good Manufacturing Practice Guidelines, and the Regulations on Manufacturing Plants of Medicinal Products for Human Use*, numbered 2830/ and dated 27.04.2013, issued based on Articles 27 and 40 of Decree Law 4963 dated 11.03.2011 on the Organization and Mandates of the Ministry of Health and its Subordinate Agencies, and Law #1262 dated 14.01.1928 on Pharmaceutical and Medicinal Products, in line with the requirements of World Health Organization.

Turkish Medicines and Medical Devices Agency certifies the following:

Manufacturer's Name: TİM EKİP İLAÇ A.Ş.
Head Office/Correspondence Address: Tuzla Köyü Sanayisizliği Organize Sanayi Bölgesi Mevkii
Ara Bulvarı Akademi Caddesi No:53
34899 Tuzla / İSTANBUL
Site Address: Tuzla Köyü Sanayisizliği Organize Sanayi Bölgesi Mevkii
Ara Bulvarı Akademi Caddesi No:53
34899 Tuzla / İSTANBUL

Manufacturing Authorization
Date and Number: 01/07/2014 2014/5

Has been inspected in accordance with current Good Manufacturing Practice Guidelines, and the Regulations on Manufacturing Plants of Medicinal Products for Human Use*, numbered 2830/ and dated 27.04.2013, issued based on Articles 27 and 40 of Decree Law 4963 dated 11.03.2011 on the Organization and Mandates of the Ministry of Health and its Subordinate Agencies, and Law #1262 dated 14.01.1928 on Pharmaceutical and Medicinal Products, in line with the requirements of World Health Organization.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted in 2019/2014 to 2019/2014, it is concluded 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. The authenticity of this certificate may be verified with Turkish Medicines and Medical Devices Agency upon request.

*This regulation is aligned with European Union Directive 91/271/EEC 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.

Human Medicinal Products

1. Manufacturing Operations of Human Medicinal Products*

If the company is engaged in manufacture of products with special requirements, e.g. radiopharmaceuticals or sterile ophthalmics, radiopharmaceuticals, ophthalmics, ophthalmics, ophthalmics, ophthalmics, ophthalmics or other potentially hazardous active ingredients, this should be noted under the relevant product type and design.

1.1 Sterile Products

1.1.1 Aseptically prepared

1.1.1.1 Filter and requires aseptic conditions, configuration, parallel

1.2 Packaging

1.2.1 Primary Packaging

1.2.2 Secondary packaging

1.3 Quality control testing

1.3.1 Microbiological (including sterility testing)

1.3.2 Microbiological (including sterility testing)

1.3.3 Chemical/physical testing

01/08/2017 - TİM/GMP/2017/125

Y. TAN

SZUTEST

SERTİFİKA

Yüzey Yıkama Sistemleri
Sistemleri - Tuzla

TİM EKİP İLAÇ A.Ş.

Tuzla Köyü Sanayisizliği Organize Sanayi Bölgesi Mevkii Ara Bulvarı Akademi Caddesi No:53 34899 Tuzla / İSTANBUL

ISO 14001:2004

Buy Tuzla, Sertifikasyon

01/08/2017

IAF

SZUTEST

SERTİFİKA

Yüzey Yıkama Sistemleri
Sistemleri - Tuzla

TİM EKİP İLAÇ A.Ş.

Tuzla Köyü Sanayisizliği Organize Sanayi Bölgesi Mevkii Ara Bulvarı Akademi Caddesi No:53 34899 Tuzla / İSTANBUL

ISO 9001:2008

Buy Tuzla, Sertifikasyon

01/08/2017

IAF

TURKISH MINISTRY OF HEALTH
Turkish Medicines and Medical Devices Agency

CERTIFICATE OF GMP (GOOD MANUFACTURING PRACTICE) COMPLIANCE OF A MANUFACTURER

Part 1

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Ara Bulvarı Akademi Caddesi No:53
34899 Tuzla / İSTANBUL

Manufacturing Authorization
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*This regulation is aligned with European Union Directive 91/271/EEC 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.

Human Medicinal Products

1. Manufacturing Operations of Human Medicinal Products*

If the company is engaged in manufacture of products with special requirements, e.g. radiopharmaceuticals or products containing penicillin, cephalosporins, ceftriaxone, cephalosporins, cephalosporins, cephalosporins or other potentially hazardous active ingredients, this should be noted under the relevant product type and design.

1.1 Sterile Products

1.1.1 Aseptically prepared

1.1.1.1 Filter and requires aseptic conditions, configuration, parallel

1.1.2 Filtered and requires aseptic conditions, configuration, parallel

1.1.3 Filtered and requires aseptic conditions, configuration, parallel

1.1.4 Filtered and requires aseptic conditions, configuration, parallel

1.1.5 Filtered and requires aseptic conditions, configuration, parallel

1.1.6 Filtered and requires aseptic conditions, configuration, parallel

1.1.7 Filtered and requires aseptic conditions, configuration, parallel

1.1.8 Filtered and requires aseptic conditions, configuration, parallel

1.1.9 Filtered and requires aseptic conditions, configuration, parallel

1.1.10 Filtered and requires aseptic conditions, configuration, parallel

1.2 Packaging

1.2.1 Primary Packaging

1.2.2 Secondary packaging

1.3 Quality control testing

1.3.1 Microbiological (including sterility testing)

1.3.2 Microbiological (including sterility testing)

1.3.3 Chemical/physical testing

01/08/2017 - TİM/GMP/2017/126

Y. TAN

OUR INNOVATIVE APPROACHES



2010 First dedicated sterile Carbapenem Production Facility	2010
2011 First Generic Injectable Meropenem product registration and launch	2011
2013 First Generic Injectable Cefoperazone + Sulbactam registration and launch	2013
2014 With the opening of Dedicated Injectable Penicillin Facility, the first and only Turkish Company who has 3 dedicated Betalactam production facility	2014
2015 The opening of Steril Liquid Productions Facility	2015
2018 First Generic Injectable Ertapenem product registration and launch	2018
2018 Tüm Ekip R&D Center approval Date	2018



8

R&D projects
Support From the
Government



100

Registered
Products

TÜM EKİP
İLAÇ A.Ş.

200

Staff
Member

58

Years
Experience

WE PRODUCE FOR HEALTH



TÜM * EKİP
İLAÇ A.Ş.





Beta - Lactam Antibacterials, Penicillins

Broad Spectrum Penicillin



- Amoxicilline

Beta-lactamase Sensitive Penicillin



- Benzathine Penicillin G

A row of Sulbaksit antibiotic packaging. From left to right: a box of Sulbaksit 1g IM (Amoxicillin 1g + Sulbactam 100 mg), a vial and ampoule of Sulbaksit 1g IM, a box of Sulbaksit 1g IM/IV (Enjeksiyonluk Çözelti), a vial and ampoule of Sulbaksit 1g IM/IV, a box of Sulbaksit 500 mg IM (Amoxicillin 500 mg + Sulbactam 100 mg), a vial and ampoule of Sulbaksit 500 mg IM, a box of Sulbaksit 500 mg IM/IV (Enjeksiyonluk Çözelti), a vial and ampoule of Sulbaksit 500 mg IM/IV, a box of Sulbaksit 250 mg IM/IV (Enjeksiyonluk Çözelti), and a vial and ampoule of Sulbaksit 250 mg IM/IV. All packaging is white with blue and green accents and includes the TUM EKİP logo.

● **Piperacillin+Tazobactam**

Other Beta-lactam Antibacterials Cephalosporins



First-generation



• Cefazolin

Second-generation



• Cefuroxime

Third-generation



• Cefotaxime



• Ceftazidime



• Cefoperazone + Sulbactam



Third-generation



• Ceftriaxone

Fourth-generation



• Cefepim

Carbapenems



First generic in TURKEY

- Ertapenem

Carbapenems



- Meropenem



- Imipenem+Cilastatin

First dedicated sterile Carbapenem Production Facility

Antibacterials for Systemic Use

Tetracyclines



• Tigecycline



Triazole Derivatives



- Fluconazole

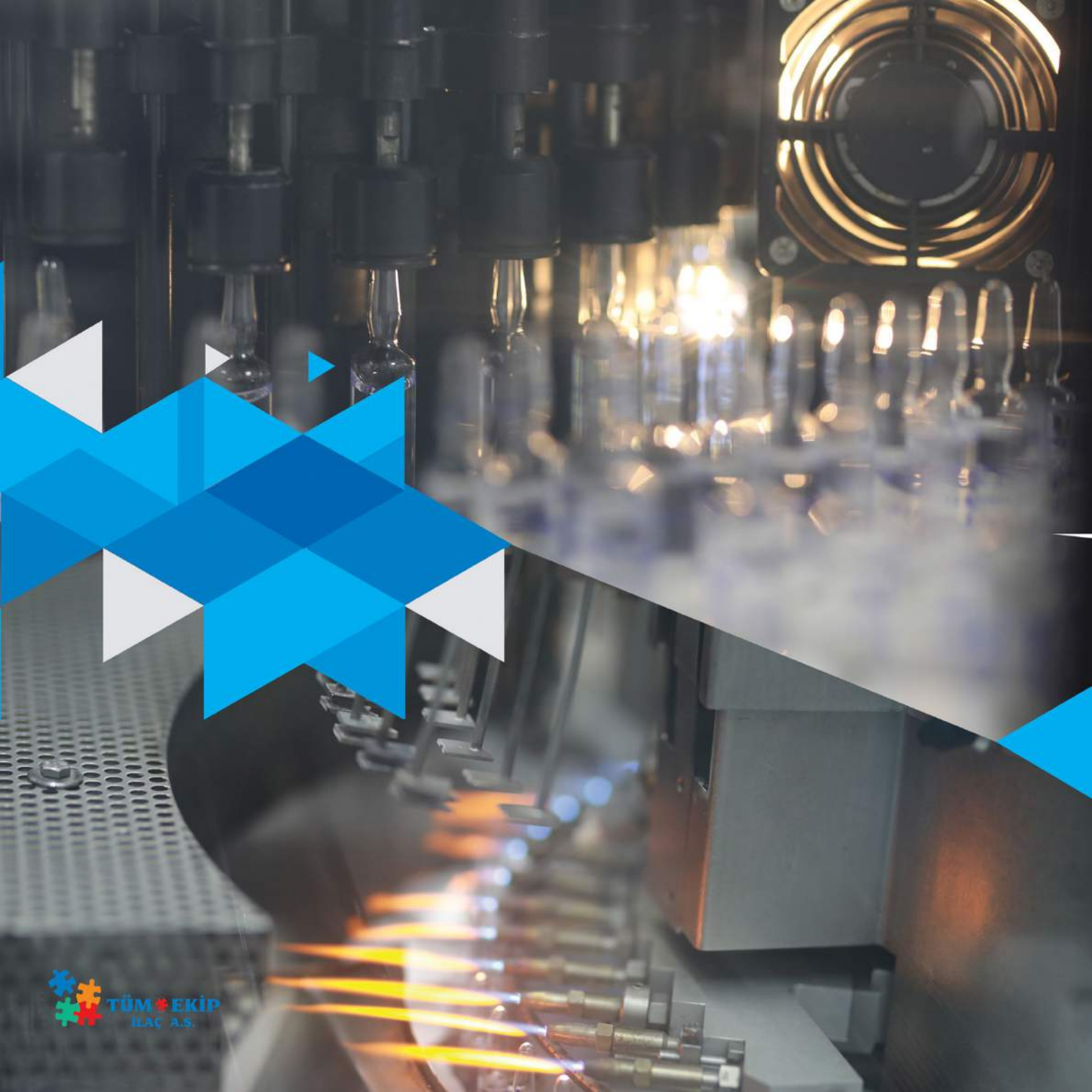


- Voriconazole

Other Antibacterials

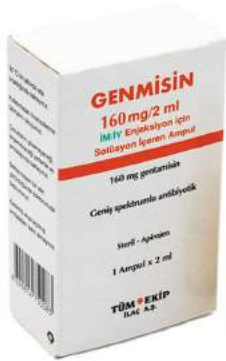


- Linezolid



Aminoglycoside Antibacterials

Other Aminoglycosides



● Gentamicin



● Amikacin

Macrolides, Lincosamides and Streptogramins



● Clindamycin



GILOWY

Keep hands clear
from moving parts

DİKKAT
MAKİNA ÇALIŞIRKEN
TEHLİKELİ VE
TEMİZLİK YAPMAYIN



TÜM EKİP
İLAÇ A.Ş.

Quinolone Antibacterials

Fluoroquinolones



- Ciprofloxacin



- Levofloxacin



- Moxifloxacin

Gastrointestinal Products

Propulsive



• Metoclopramide

Peptic Ulcer and GORD Products

H₂Receptor Antagonists



• Ranitidine

Proton Pump Inhibitors (PPI)



• Esomeprazole



• Pantoprazole



• Omeprazole

Antithrombotic Agents

Platelet Aggregation Inhibitors



• Tirofiban

Antiemetics and Antinauseants

Serotonin (5HT₃) Antagonists



- Granisetron



- Palonosetron

Xanthines



- Aminophylline

Antiarrhythmics

Vaughan Williams' Class III



- Amiodarone

PRODUCTS		ACTIVE INGREDIENT	ATC
Beta - Lactam Antibacterials, Penicillins			J01C
Broad Spectrum Penicillins Inj.			J01CA
NEOAMOX 500 MG IM/IV	1 Vial + 1 Diluent	AMOXICILLINE Na	J01CA04
NEOAMOX 1 G IM/IV	1 Vial + 1 Diluent	AMOXICILLINE Na	J01CA04
Beta-lactamase Sensitive Penicillins			J01CE
PENSILINA 1 000 000 IU IM/IV	1 Vial + 1 Diluent	PENICILLIN G K	J01CE01
BENZAPEN-LA 1 200 000 IU IM	1 Vial + 1 Diluent	BENZATHINE PENICILLIN G	J01CE08
BENZAPEN-LA 2 400 000 IU IM	1 Vial + 1 Diluent	BENZATHINE PENICILLIN G	J01CE08
BENZAPEN 6.3.3. IM	1 Vial + 1 Diluent	BENZATHINE PENICILLIN G + PROCAIN PENICILLIN G + PENICILLIN G Na	J01CE30
PROKAIN PENICILLIN 400 000 IU IM 1 Vial + 1 Diluent		PROCAIN PENICILLIN G+PENICILLIN G K	J01CE30
PROKAIN PENICILLIN 800 000 IU IM 1 Vial + 1 Diluent		PROCAIN PENICILLIN G+PENICILLIN G K	J01CE30
Combinations of Penicillins, incl., Beta-lactamase Inhibitors			J01CR
SULBAKSIT 250 MG IM	1 Vial + 1 Diluent	AMPICILLIN Na + SULBACTAM Na	J01CR01
SULBAKSIT 500 MG IM	1 Vial + 1 Diluent	AMPICILLIN Na + SULBACTAM Na	J01CR01
SULBAKSIT 1 G IM	1 Vial + 1 Diluent	AMPICILLIN Na + SULBACTAM Na	J01CR01
SULBAKSIT 250 MG IM/IV	1 Vial + 1 Diluent	AMPICILLIN Na + SULBACTAM Na	J01CR01
SULBAKSIT 500 MG IM/IV	1 Vial + 1 Diluent	AMPICILLIN Na + SULBACTAM Na	J01CR01
SULBAKSIT 1 G IM/IV	1 Vial + 1 Diluent	AMPICILLIN Na + SULBACTAM Na	J01CR01
SULBAKSIT 2 G IV	1 Vial + 1 Diluent	AMPICILLIN Na + SULBACTAM Na	J01CR01
TAZOJECT 2G/0,25G	1 Vial	PIPERACILLIN Na + TAZOBACTAM Na	J01CR05
TAZOJECT 4G/0,5G	1 Vial	PIPERACILLIN Na + TAZOBACTAM Na	J01CR05
Other Beta-lactam Antibacterials			J01D
First-generation Cephalosporins			J01DB
EQIZOLIN 250 MG IM	1 Vial + 1 Diluent	CEFAZOLIN Na	J01DB04
EQIZOLIN 500 MG IM	1 Vial + 1 Diluent	CEFAZOLIN Na	J01DB04
EQIZOLIN 1 G IM	1 Vial + 1 Diluent	CEFAZOLIN Na	J01DB04

PRODUCTS	ACTIVE INGREDIENT	ATC
Other Beta-lactam Antibacterials		J01D
First-generation Cephalosporins		J01DB
EQIZOLIN 250 MG IM/IV 1 Vial + 1 Diluent	CEFAZOLIN Na	J01DB04
EQIZOLIN 500 MG IM/IV 1 Vial + 1 Diluent	CEFAZOLIN Na	J01DB04
EQIZOLIN 1 G IM/IV 1 Vial + 1 Diluent	CEFAZOLIN Na	J01DB04
Second-generation Cephalosporins		J01DC
SEFFUR 250 MG IM/IV 1 Vial + 1 Diluent	CEFUROXIME Na	J01DC02
SEFFUR 750 MG IM/IV 1 Vial + 1 Diluent	CEFUROXIME Na	J01DC02
SEFFUR 1,5 G IV Vial 1 Vial	CEFUROXIME Na	J01DC02
SEFFUR 250 MG IM 1 Vial + 1 Diluent	CEFUROXIME Na	J01DC02
SEFFUR 750 MG IM 1 Vial + 1 Diluent	CEFUROXIME Na	J01DC02
Third-generation Cephalosporins		J01DD
EQITAX 0,5 G IM/IV 1 Vial + 1 Diluent	CEFOTAXIME Na	J01DD01
EQITAX 1 G IM/IV 1 Vial + 1 Diluent	CEFOTAXIME Na	J01DD01
EQITAX 2 G IV VIAL 1 Vial + 1 Diluent	CEFOTAXIME Na	J01DD01
ZIDIM 500 MG IM/IV 1 Vial + 1 Diluent	CEFTAZIDIME Pentahydrate	J01DD02
ZIDIM 1 G IM/IV 1 Vial + 1 Diluent	CEFTAZIDIME Pentahydrate	J01DD02
EQICEFT 1 G IM 1 Vial + 1 Diluent	CEFTRIAXONE Na	J01DD04
EQICEFT 0.5 G IM 1 Vial + 1 Diluent	CEFTRIAXONE Na	J01DD04
EQICEFT 1 G IV 1 Vial + 1 Diluent	CEFTRIAXONE Na	J01DD04
EQICEFT 0.5 G IV 1 Vial + 1 Diluent	CEFTRIAXONE Na	J01DD04
SULZON 1 G / 1G IM 1 Vial + 2 Diluents	CEFOPERAZONE Na + SULBACTAM Na	J01DD62
SULZON 1 G IM/1G IV 1 Vial + 1 Diluent	CEFOPERAZONE Na + SULBACTAM Na	J01DD62
SULZON 2 G IM/1G IV 1 Vial + 1 Diluent	CEFOPERAZONE Na + SULBACTAM Na	J01DD62
EKOBID 1 G IM/IV 1 Vial + 1 Diluent	CEFOPERAZONE Na	J01DD12

PRODUCTS	ACTIVE INGREDIENT	ATC
Other Beta-lactam Antibacterials		J01D
Fourth-generation Cephalosporins		J01DE
EKIPIM 0,5 G IM/IV 1 Vial + 1 Diluent	CEFEPIME HCl	J01DE01
EKIPIM 1 G IM/IV 1 Vial + 1 Diluent	CEFEPIME HCl	J01DE01
Other Beta-lactam Antibacterials		J01D
CARBAPENEMS		J01DH
MOPEM 500 MG IV 1 Vial	MEROPENEM Trihydrate	J01DH02
MOPEM 1 G IV 1 Vial	MEROPENEM Trihydrate	J01DH02
EKIPERTA 1 G IM/IV 1 Vial	ERTAPENEM Sodium	J01DH03
CILAPEM 500MG/500 MG IV 1 Vial	IMIPENEM Monohydrate + CILASTATIN Na	J01DH51
Antibacterials for Systemic Use		J01
Tetracyclines		J01AA
TIGEJECT 50 MG IV 10 Vials	TIGECYCLINE	J01AA12
Antimycotics for Systemic Use		J02A
Triazole Derivatives		J02AC
FLUJECT 100 MG/50 ML IV 1 Vial	FLUCONAZOLE	J02AC01
VORIJECT 200 MG IV 1 Vial	VORICONAZOLE	J02AC03
Macrolides, Lincosamides and Streptogramins		J01F
Lincosamides		J01FF
MENEKLIN 300 MG IM/IV Ampoule	CLINDAMYCIN Phosphate	J01FF01
MENEKLIN 600 MG IM/IV Ampoule	CLINDAMYCIN Phosphate	J01FF01
Aminoglycoside Antibacterials		J01G
Other Aminoglycosides		J01GB
GENMISIN 40 MG IM/IV Ampoule	GENTAMICIN Sulfate	J01GB03
GENMISIN 80 MG IM/IV Ampoule	GENTAMICIN Sulfate	J01GB03
GENMISIN 160 MG IM/IV Ampoule	GENTAMICIN Sulfate	J01GB03
AMIJEKSIN 100 MG IM/IV Ampoule	AMIKACIN Sulfate	J01GB06
AMIJEKSIN 500 MG IM/IV Ampoule	AMIKACIN Sulfate	J01GB06

PRODUCTS	ACTIVE INGREDIENT	ATC
Quinolone Antibacterials		J01M
Fluoroguinolones		J01MA
NOVAREX 200 MG/100 ML IV Vial	CIPROFLOXACIN Lactate	J01MA02
NOVAREX 400 MG/200 ML IV Vial	CIPROFLOXACIN Lactate	J01MA02
SIPROJECT 400 MG/200 ML IV Bag	CIPROFLOXACIN Lactate	J01MA02
PNEMOX 400 MG/250 ML IV PP Bag	MOXIFLOXACIN HCl	J01MA14
LEVOJECT 500 MG/100 ML IV Vial	LEVOFLOXACIN Hemihydrate	J01MA12
Other Antibacterials		J01X
LINEJECT 600 MG/300 ML IV 1 Bag	LINEZOLID	J01XX08
Peptic Ulcer and GORD Products		A02B
H2 Receptor Antagonists		A02BA
RANJECT 50 MG/2 ML IM/ IV 5 Ampoules	RANITIDINE HCl	A02BA02
Proton Pump Inhibitors (PPI)		A02BC
PROTAZ 40 MG IV 1 Vial	PANTOPRAZOLE Na sesquihydrate	A02BC02
JECTOSEM 40 MG IV 1 Vial	ESOMEPRAZOLE Na	A02BC05
LYOMEPRA 40 MG IV 1 Vial + 1 Diluent	OMEPRAZOLE Na	A02BC01
GASTROINTESTINAL PRODUCTS		A03
Propulsives		A03FA
EMOJECT 10 MG/2 ML IM/IV 5 Ampoules	METOCLOPRAMIDE HCl	A03FA01
Antiemetics and Antinauseants		A04
Serotonin (5HT3) Antagonists		A04AA
PALOJECT 250 MCG/5 ML IV Ampoule	PALONOSETRON HCl	A04AA05
SETREX 3 MG/3 ML IV 5 Ampoules	GRANISETRON HCl	A04AA02
Antithrombotic Agents		B01A
Platelet Aggregation Inhibitors excl. Heparin		B01AC
AGRABLOC 12,5 MG/50 ML 1 Vial	TIROFIBAN HCl Monohydrate	B01AC17

PRODUCTS		ACTIVE INGREDIENT	ATC
Antiarrhythmics			C01B
Vaughan Williams' Class III			C01BD
AMIDOVIN 150 MG/3 ML IV 6 Ampoules	AMIODARONE HCl		C01BD01
Xanthines			B03D
TECAR 240 MG/10 ML IV 10 Ampoules	AMINOPHYLLINE		R03DA05
Cardiac Stimulants excl. Cardiac Glycosides			C01C
Adrenergic and Dopaminergic Agents			C01CA
EPINOR 4 MG/4 ML IV 10 Ampoules	NOREPINEPHRINE Bitartrate		C01CA03
DIURETICS			C03
Sulfonamides, Plain			C03CA
FUROJECT 20MG/2 ML IM/IV 5 Ampoules	FUROSEMIDE		C03CA01
Corticosteroids for Systemic Use			H02A
Glucocorticoides			H02AB
DEXOJECT 8 MG/2 ML IM/ IV Ampoule	DEXAMETHASONE Na Phosphate		H02AB02
Antiinflammatory and Antirheumatics			M01A
Acetic Acid Derivatives and Related Substances			M01AB
DIFENJECT 75 MG/3 ML IM 10 Ampoules	DICLOFENAC Na		M01AB05
Muscle Relaxants, Peripherally Acting Agents			M03
Other Quaternary Ammonium Compounds			M03A
JECRON 50 MG/5 ML IV 10 Vials	ROCURONIUM Bromide		M03AC09
Drugs for Treatment of Bone Diseases			M05
Bisphosphonates			M05BA
BONDREX 4 MG/5 ML IV 1 Vial	ZOLEDRONIC Acid		M05BA08
Antithrombotic Agents			N02B
Pyrazolones			N02BB
NOVAMIZOL 1000 MG/2 ML IM/IV 10 Ampoules	METAMIZOLE Na		N02BB02

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