COUNCIL OF EUROPE
COMMITTEE OF MINISTERS

(PARTIAL AGREEMENT IN THE SOCIAL AND PUBLIC HEALTH FIELD)

RESOLUTION AP (77) 1

ON THE CLASSIFICATION OF MEDICINES WHICH ARE OBTAINABLE ONLY ON MEDICAL PRESCRIPTION[1]

(Adopted by the Committee of Ministers on 22 March 1977 at the 267th meeting of the Ministers' Deputies)

The Representatives on the Committee of Ministers of Belgium, France, the Federal Republic of Germany, Italy, Luxembourg, the Netherlands, the United Kingdom of Great Britain and Northern Ireland, these states being parties to the Partial Agreement in the social and public health field, and the Representatives of Austria, Denmark, Ireland and Switzerland, states which have participated in the public health activities carried out within the above-mentioned Partial Agreement since 1 October 1974, 2 April 1968, 23 September 1969 and 5 May 1964, respectively,

1. Having regard to the recommendation on the classification of medicines which are obtainable only on medical prescription, adopted by the Partial Agreement Public Health Committee on 8 October 1976 ;

2. Considering that, under the terms of its Statute, the aim of the Council of Europe is to achieve a greater unity between its Members for the purpose of safeguarding and realising the ideals and principles which are their common heritage and facilitating their economic and social progress ;

3. Having regard to the provisions of the Brussels Treaty signed on 17 March 1948, by virtue of which Belgium, France, Luxembourg, the Netherlands and the United Kingdom of Great Britain and Northern Ireland declared themselves resolved to strengthen the social ties by which they were already united ;

4. Having regard to the Protocol modifying and completing the Brussels Treaty, signed on 23 October 1954 by the signatory states of the Brussels Treaty, on the one hand, and the Federal Republic of Germany and Italy, on the other hand ;

5. Observing that the seven states parties to the Partial Agreement, which have resumed, within the Council of Europe, the social work hitherto undertaken by the Brussels Treaty Organisation and then by Western European Union, which derived from the Brussels Treaty as modified by the Protocol mentioned at paragraph 4 above, as well as Austria, Denmark, Ireland and Switzerland, who participate in Partial Agreement activities in the field of public health, have always endeavoured to be in the forefront of progress in social matters and also in the associated field of public health, and have for many years undertaken action towards harmonisation of their legislation ;

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1. This resolution supersedes Resolution AP (69) 2 of 25 January 1969. When it was adopted, the delegation of Austria, in accordance with Article 10.2.c of the Rules of Procedure for the meetings of the Ministers' Deputies, reserved the right of their government to comply with it or not.

6. Considering further that the lack of uniform laws on the supply of medicines restricted to medical prescription has created problems in the field of public health and, at the present time, raises difficulties at international level ;

7. Considering in particular that the tendency towards auto-medication and over-medication is increasing ;

8. Believing that it is therefore increasingly necessary to achieve harmonisation of national laws in this field,

I. Recommend to the governments of the seven states parties to the Partial Agreement as well as to those of Austria, Denmark, Ireland and Switzerland, that they restrict to medical prescription the supply of the medicines mentioned in the General Provisions set out hereafter ;

II. Invite the same governments to accept amendments which may be made later to the list set out in the appendix attached hereto by the Public Health Committee, which shall carry out an annual revision of this list in the light of the General Provisions set out hereafter ;

III. Invite the said governments to keep the Secretary General of the Council of Europe fully informed every five years of the action taken by them with regard to the present resolution.

GENERAL PROVISIONS

governing the drawing up and periodical revision of the lists of medicines which are obtainable on medical prescription

1. The only substances affected by the recommended provisions are those used for medical purposes.

2. Narcotic drugs are not referred to when they are already covered by special common provisions concerning the rules governing their supply.

3. This resolution does not apply to homeopathic preparations or to other similar non-allopathic minute dose preparations on the market in the member states. Sale and supply of these preparations are governed by legal provisions in force in each member state.

4. The lists of medicines which are obtainable only on medical prescription are drawn up with reference not only to their toxicity, but also to all the risks, direct or indirect, which they may represent to human health and, in particular, according to :

a. their acute and chronic toxicity ;

b. results of clinical trials and experience in use (adverse reactions, tolerances etc.) ;

c. their intended actions and uses.

Salts, esters and salts of ester are subject to the same classifications as the substances themselves, unless otherwise specified in the lists.

In cases where several drugs are present in a preparation, the classification should take account of the phenomenon of synergy—whether potentiating or not—antagonism or changes in the effects of the components.

5. A medicine may not be supplied except on a medical prescription when one or more of the following conditions apply :

a. it contains an active principle not previously used for medical purposes, and, in particular, a new chemical molecule. In such cases a final decision concerning any restrictions to be permanently applied shall be taken within three years of introduction on to the market, and the participating states will be notified accordingly ;

b. it is used parenterally ; [1,2]

1. By "parenteral" are intended the sub-cutaneous, intracutaneous, intramuscular, intravenous, intraspinal, intracisternal, epidural, intraperitoneal, intra-articular, intracardiac and other similar routes.

2. Reservation of the Federal Republic of Germany on this point.

c. it contains one or more substances in Lists I or II annexed hereto, to which the following criteria apply :

List I

The supply of a medicine containing one of the substances in this list may only be repeated if the prescriber so specifies on the prescription.

List II

The supply of a medicine containing one of the substances in this list may be repeated without the prescriber having so specified, provided that he did not explicitly forbid such repetition :

— during a period of three months following the date of issue of the prescription, and

— not more than five times during the same period provided that such a supply does not permit an increase in the posology indicated by the prescriber.

Exemptions from Lists I and II

For certain substances, exemptions from the "prescription only" requirement may appear in Lists I and II :

— in respect of low dosage or concentration ;

— according to the route of administration and the composition of the drug.

Preamble to the list of medicines which are obtainable only on medical prescription

1. *Minimum requirements*

The classifications shown are to be considered as minimum requirements ; consequently, governments are free to apply stricter rules in any given case.

2. *Provisional classifications*

Certain medicines are provisionally classified, especially in cases where further information or consultation with interested parties might be useful. Provisionally classified medicines are suitably marked, i.e. with an asterisk.

3. *Nomenclature*

Wherever possible, the nomenclature used is that of the International Non-Proprietary Names (INN) of the World Health Organisation.

4. *Revisions*

Annual revisions will deal with :

— final classification of medicines provisionally classified ;

— proposals for adding or deleting medicines ;

— proposals for removing or diminishing the number of exemptions.

Proposals for revision will be submitted by 1 January of each year in order to be examined by the competent bodies before 1 July. In cases of urgency, proposals for revision may be submitted at any time.

5. *Date of adoption*

The published list will indicate in each case the date on which it was adopted by the Public Health Committee and transmitted to the governments concerned.

LIST OF MEDICINES ADOPTED BY THE PUBLIC HEALTH COMMITTEE (PARTIAL AGREEMENT) UP TO 1 JANUARY 1977

Explanations

* a. *Single active principle*
List I, without exemptions.

b. *In association with another hypnotic sedative or with any potentialiser*
List I, without exemptions.

c. *In association with active principles other than those mentioned under paragraph* b *above*
List II, provided the dose of barbituric acid derivatives does not exceed 50 mg per unit dose.
Without prescription, provided the dose of barbituric acid derivatives does not exceed 15 mg per unit dose.

** *Exemptions in respect of nasal and ophthalmic drops*

*** *Prescribed as an ovulatory suppressor* : List II.

Abbreviations

W.P. : without prescription
U.D. : unit dose. For the purposes of this resolution, "unit dose" is understood to be the maximum dose for which exemption from prescription is provided.
E.U. : external use
I.U. : internal use
w/v : weight/volume
w/w : weight/weight
S.W. No. 1: special warning for Phenacetine derivatives
S.W. No. 2: special warning for pyrazole derivatives
S.W. No. 3: special warning for glaucoma

N.B. Special warnings apply solely to preparations which are not subject to medical prescription.

A

ACECARBROMAL	II	W.P. : tablets and suppositories containing not more than 500 mg of an active ingredient or of a combination of open-chain ureides
ACENOCOUMAROL	II	
ACEPROMAZINE	II	
ACETANILIDE	II	
ACETARSOL	II	W.P. : preparations for dental application
ACETAZOLAMIDE	II	
ACETOHEXAMIDE	II	
ACETOPHENAZINE	II	
ACETYLCHOLINE CHLORIDE	II	W.P. : — E.U. — collyria
ACETYLDIHYDROCODEINE	II	
ACETYLPHENYLHYDANTOIN	II	
ACETYLSULFAMETHOXYPYRIDAZINE**	I	
ACETYLSULFISOXAZOLE**	I	
ACONITINE	I	

ACONITE, galenical preparations	II	W.P. : — E.U. — preparations for I.U. containing not more than 25 mcg of aconitine per U.D.
ACRISORCIN	II	W.P. : E.U.
ACTINOMYCIN C	I	
ADICILLINE	I	
ADIPHENINE	II	W.P. : — U.D. not exceeding 30 mg — ophthalmic preparations *S.W. No. 3*
AJMALINE	II	
ALCLOFENAC	II	
ALCURONIUM CHLORIDE	I	
ALDOSTERONE	I	
ALIMEMAZINE	II	W.P. : — U.D. not exceeding 10 mg — syrups containing not more than 0.05%
ALLENOESTROL	I	E.U. : II
ALLOBARBITAL*		
ALLOMETHADIONE	II	
ALLOPURINOL	II	
ALLYL-5 (β-HYDROXYPROPYL) 5 BARBITURIC ACID*		
ALLYLESTRENOL	I	
ALPRENOLOL	II	
ALTIZIDE	II	
AMADINONE	I	
AMANOZINE	II	
AMANTADINE	II	
AMBENONIUM CHLORIDE	I	
AMBOMYCIN	I	
AMFECLORAL	I	
AMFEPENTOREX	I	
AMFEPRAMONE	I	
AMFOMYCIN	I	E.U. : II
AMILORIDE	II	
AMINOANTIPYRINE	II	W.P. : U.D. not exceeding 500 mg *S.W. No. 2*
AMINOCAPROIC ACID	II	
AMINOGLUTETHIMIDE	II	
AMINOMETRADINE	II	
AMINOPHENAZONE	II	W.P. : — E.U. — U.D. not exceeding 500 mg *S.W. No. 2*
AMINOPHENAZONE CARBETHYL SALICYLATE	II	W.P. : U.D. not exceeding 1.5 g *S.W. No. 2*
AMINOPROMAZINE	II	
AMINOPTERIN SODIUM	I	
AMINOREX	I	
AMIPHENAZOLE	II	
AMISOMETRADINE	II	
AMITRIPTYLINE	I	
AMOBARBITAL*		
AMOXICILLIN	I	
AMPHENONE B	II	
AMPHETAMINE	I	
AMPHOTERICIN B	I	
AMPICILLIN	I	
AMPYRIMINE	II	
ANADORE	I	
ANAGESTONE	I	
ANDROGENOL	I	
ANDROISOXAZOLE	II	

ANDROSTANOLONE	II	
ANDROSTENEDIOL	I	
ANDROSTERONE	I	
ANGIOTENSINAMIDE	II	
ANISINDIONE	II	
ANISOMYCIN	I	E.U. : II
ANTAPENITE	II	
ANTELMYCIN	I	E.U. : II
ANTHIOLIMINE	II	
ANTRAMYCIN	I	
APOMORPHINE	II	W.P. : — preparations containing not more than 0.2% w/v — U.D. less than 2 mg
APROBARBITAL*		
APRONALIDE	II	
ARECOLINE	II	
ARGIPRESSIN	I	
ARSPHENAMINE	II	
ARSTHINOL	II	
ATROPINE	II	W.P. : — U.D. not exceeding 300 mcg — dermatological preparations *S.W. No. 3*
	I	ophthalmic preparations
ATROPINE-OXIDE	II	W.P. : U.D. not exceeding 500 mcg *S.W. No. 3*
AZACYCLONOL	II	
8-AZAGUANINE	I	
AZAPERONE	I	
AZARIBINE	I	
AZASERINE	I	
AZATHIOPRINE	I	
AZAURIDINE	I	
AZIDAMFENICOL	I	
AZIDOCILLIN	I	
AZOMYCIN	II	
AZOTOMYCIN	I	

B

BACITRACIN	I	W.P. : — E.U. — lozenges
BACLOFEN	II	
BARBEXACLONE	II	
BARBITAL*		
BAROTAL*		
BECLAMIDE	II	
BECLOMETASONE	I	
BELLADONNA, galenical preparations	II	W.P. : — preparations containing not more than 300 mcg per U.D. calculated as hyoscyamine — dermatological preparations *S.W. No. 3*
BEMEGRIDE	II	
BENACTYZINE	II	
BENAPRYZINE	II	
BENDROFLUMETHIAZIDE	II	
BENPERIDOL	I	
BENZATROPINE	II	
BENZESTROL	I	E.U. : II
BENZILONIUM BROMIDE	II	W.P. : U.D. not exceeding 10 mg *S.W. No. 3*
BENZONATATE	II	W.P. : U.D. not exceeding 100 mg
BENZPHETAMINE	I	

BENZPYRINIUM CHLORIDE	I	
BENZQUINAMIDE	II	
BENZQUINONIUM	I	
BENZTHIAZIDE	II	
BENZYDAMINE	II	
BENZYLPENICILLIN	I	
BENZYLSULFAMIDE**	I	
BETAHISTINE	II	
BETAMETHASONE	I	
BETANIDINE	II	
BETHANECHOL	II	
BIALAMICOL	II	
BIETAMIVERINE	II	W.P. : U.D. not exceeding 50 mg
BIPERIDEN	II	
BLEOMYCIN	I	
BOLANDIOL	I	
BOLASTERONE	I	
BOLDENONE	I	
BOLENOL	I	
BOLMANTALATE	I	
BRALLOBARBITAL*		
BRETYLIUM TOSILATE	II	
BROMACRYLIDE	I	
BROMALLYLAMYLBARBITURIC ACID*		
BROMINDIONE	II	
BROMISOVAL	II	W.P. (cf. ACECARBROMAL)
BROMISOVALERYL HYDANTOIN	II	
BROMO-5-URACIL	I	
BROPARESTROL	I	E.U. : II
BROPHEBARBITAL*		
BUCLIZINE	II	W.P. : — tablets containing not more than 25 mg — syrups containing not more than 0.1% — ointments containing not more than 2%
BUFORMIN	II	
BUMETANIDE	II	
BUPRANOLOL	II	
BUSULFAN	I	
BUTALBITAL*		
BUTALLYLONAL*		
BUTAPERAZINE	II	
BUTAZOLAMIDE	II	
BUTHALITAL SODIUM*		
BUTIZIDE	II	
BUTOBARBITAL*		
BUTOPIPRINE	II	

C

CALCITONIN	II	
CALCIUM-5-ALLYL-5-BUTYL-BARBITURATE*		
CALCIUM CARBIMIDE	II	
CANDICIDIN	I	
CAPREOMYCIN	I	
CAPTODIAME	II	
CARBACHOL	II	
CARBAMAZEPINE	II	
CARBARSONE	II	
CARBENICILLIN	I	
CARBIMAZOLE	II	
CARBOMYCIN	I	
CARBROMAL	II	W.P. (cf. ACECARBROMAL)
CARBUTAMIDE	II	

CARFECILLIN	I	
CARFENAZINE	II	
CARINDACILLIN	I	
CARISOPRODOL	II	
CARPERIDINE	I	
CARSALAM	II	
CARZENIDE	II	
CEFACETRILE	I	
CEFALEXIN	I	
CEFALORIDINE	I	
CEFALOTIN	I	
CEFAPIRIN	I	
CEFAZOLIN	I	
CEFRADINE	I	
CHLORAL HYDRATE	II	W.P. : — E.U. — U.D. not exceeding 1 g
CHLORALOSE	II	
CHLORAMBUCIL	I	
CHLORAMINOPHENAMIDE	II	
CHLORAMPHENICOL	I	
CHLORCYCLIZINE	II	W.P. : — E.U. — U.D. not exceeding 50 mg
CHLORDIAZEPOXIDE	II	
CHLORDRONOLONE	I	
CHLORETHYLAMINOURACIL	I	
CHLORMADINONE	I	
CHLORMERODRIN	II	
CHLORMETHINE	I	
CHLORMETHINE OXIDE	I	
CHLORMEZANONE	II	
CHLORMIDAZOLE	II	W.P. : E.U.
(CHLORO-2-ACETYL) AJMALIN	II	
6-CHLOROPURINE	I	
CHLOROQUINE	II	W.P. : U.D. not exceeding 300 mg
CHLOROTHIAZIDE	II	
CHLOROTRIANISENE	I	E.U. : II
CHLORPHENOXAMINE	II	
CHLORPHENTERMINE	I	
CHLORPROETHAZINE	II	
CHLORPROMAZINE	II	
CHLORPROPAMIDE	II	
CHLORPROTHIXENE	II	
CHLORTALIDONE	II	
CHLORTETRACYCLINE	I	
CHORIONIC GONADOTROPHIN	I	
CICLACILLIN	I	
CINCHOPHEN	II	
CINGESTOL	I	
CISMADINONE	I	
CLIDINIUM BROMIDE	II	
CLINDAMYCIN	I	
CLOBETASOL	I	
CLOBETASONE	I	
CLOCORTOLONE	I	
CLODANTOIN	II	
CLOFAZIMINE	II	
CLOFENAMIDE	II	
CLOFEZONE	II	
CLOFIBRATE	II	
CLOFIBRIC ACID	II	
CLOFOREX	I	
CLOGESTONE	I	

CLOMEGESTONE	I	
CLOMETERONE	I	
CLOMETHIAZOLE	II	
CLOMETOCILLIN	I	
CLOMIFENE	I	E.U. : II
CLOMINOREX	I	
CLOMIPRAMINE	I	
CLOMOCYCLINE	I	
CLONIDINE	II	
CLOPAMIDE	II	
CLOPENTHIXOL	II	
CLOQUINATE	II	
CLOREXOLONE	II	
CLORINDIONE	II	
CLOSTEBOL	I	
CLOTIAPINE	II	
CLOTRIMAZOLE	II	
CLOXACILLIN	I	
CLOXESTRADIOL	I	E.U. : II
CLOXOTESTOSTERONE	I	
COBAMAMIDE	I	
COCAINE	I	W.P. : preparations containing not more than 0.1%
CODEINE	II	W.P. : U.D. not exceeding 20 mg
CODEINAMINOXYDE	II	
CODEINE-5, 5-DIPROPYL BARBITURATE*		
COLASPASE	I	
COLCHICUM, alkaloids and glycosides	I	
COLISTIN	I	
COLPORMONE	I	E.U. : II
CONVALLATOXINE	II	
CORBADRINE	II	
CORTICOTROPHIN	I	
CORTISONE	I	
CORTIVAZOL	I	
COUMACHLOR	II	
COUMETAROL	II	
CROMOGLYCATE SODIUM	II	W.P. : nasal application
CROPROPAMIDE	II	W.P. : preparations containing not more than 100 mg
CROTETHAMIDE	II	W.P. : preparations containing not more than 100 mg
CROTARBITAL*		
N_1-CROTONYLSULFANILYL-N_2-n-BUTYL CARBAMIDE	II	
CURARE, and preparations of	I	
CYAMEMAZINE	II	
CYCLAZOCINE	I	
CYCLIZINE	II	W.P. : — tablets : U.D. not exceeding 50 mg — suppositories : U.D. not exceeding 100 mg
CYCLOBARBITAL*		
CYCLOCOUMAROL	II	
CYCLOFENIL	I	
CYCLOPENTHIAZIDE	II	
CYCLOPENTOBARBITAL*		
CYCLOPENTOLATE	II	
CYCLOPHOSPHAMIDE	I	
CYCLOPREGNOL	I	
CYCLOSERINE	I	
CYCLOTHIAZIDE	II	
CYCRIMINE	II	
CYPROTERONE	I	
CYTARABINE	I	

D

DACTINOMYCIN	I	
DAPSONE	II	
DAUNORUBICIN	I	
DEBRISOQUINE	II	
DECAMETHONIUM BROMIDE	I	
DEFEROXAMINE	II	
DEFOSFAMIDE	I	
DEHYDROANDROSTANOLONE	I	
DEHYDROANDROSTERONE	II	
DELMADINONE	I	
DEMECARIUM BROMIDE	I	
DEMECLOCYCLINE	I	
DEMECOLCINE	I	
DEMOXYTOCIN	I	
DEPRODONE	I	
DEPTROPINE	II	
DESASPIDIN	II	
DESCINOLONE	I	
DESERPIDINE	II	
DESIPRAMINE	I	
DESLANOSIDE	II	
DESONIDE	I	
DESOXIMETASONE	I	
DESOXYCORTONE	I	
DEXAMETHASONE	I	
DEXAMPHETAMINE	I	
DEXTROMETHORPHAN	II	W.P. : U.D. not exceeding 15 mg and in combination with substances other than centrally-acting antitussives and narcotic analgesics
DEXTROPROPOXYPHENE	II	
DEXTROTHYROXINE SODIUM	II	
DIAMINOPURINE	I	
DIAZEPAM	II	
6-DIAZO-5-OXO-L-NORLEUCINE	I	
DIAZOXIDE	II	
DIBENAMINE	I	
DIBENZEPIN	I	
DICHLORISONE	I	
DICLOFENAMIDE	II	
DICLOXACILLIN	I	
DICOUMAROL	II	
DICYCLOVERINE	II	W.P. : U.D. not exceeding 20 mg *S.W. No. 3*
DIENESTROL	I	E.U. : II
DIETHAZINE	II	
DIETHYLAMINOETHYLPHENOBARBITAL*		
DIETHYLSTILBESTROL	I	E.U. : II
DIFEBARBAMATE*		
DIFENCLOXAZINE	II	
DIFETARSONE	II	
DIFLUCORTOLONE	I	
DIFLUOROPHATE	II	
DIGLUCOMETHOXANE	II	
DIGOXIN	II	
DIHEXYVERINE	II	W.P. : U.D. not exceeding 20 mg *S.W. No. 3*
DIHYDRALAZINE	II	
DIHYDROCODEINE	II	W.P. : U.D. not exceeding 20 mg
DIHYDROERGOCORNINE	II	
DIHYDROERGOCRISTINE	II	

DIHYDROERGOKRYPTINE	II	
DIHYDROERGOTAMINE	II	
DIHYDROERGOTOXINE	II	
DIHYDRONOVOBIOCINE	I	
DIHYDROSTREPTOMYCIN	I	
DIHYDROXYPROGESTERONE ACETOPHENIDE	I	
DIGITALIS : plants, galenical preparations, heterosides and their derivatives	II	
DIHYPRYLONE	II	W.P. : syrups containing not more than 1.5% w/w
DIIODOTYROSINE	II	
DILOXANIDE	II	
DIMANTINE	II	
DIMEPHENOPANE	I	
DIMEPREGNEN	I	
DIMETACRINE	I	
DIMETHISTERONE	I	
DIMETHOXANATE	II	W.P. : — U.D. not exceeding 10 mg — syrups containing not more than 2.5 mg/ml
DIMETHOXYDIETHYLSTILBENE	I	E.U. : II
DIMETHYLTUBOCURARINE	I	
DIMETOTIAZINE	II	
DINOPROST	I	
DINOPROSTONE	I	
DIPHEMANIL	II	
DIPHEMETHOXIDINE	I	
DIPHENADIONE	II	
DIPHENOXINE + ATROPINE	I	
DIPHENOXYLATE + ATROPINE	I	
DIPHENYLMETHANE-4,4-DISULFONAMIDE	II	
DIPHENYL TETRAHYDROCIMIDAZOL	II	
DIPIPROVERINE	II	
DIPOTASSIUM CLORAZEPATE	II	
DIPROPYL BARBITURIC ACID*		
DIPROQUALONE	II	
DIPYRONE	II	W.P. : — E.U. — U.D. not exceeding 1g *S.W. No. 2*
DISOPYRAMIDE	II	
DISTIGMINE	II	
DISULFIRAM	II	
DITHIAZANINE IODIDE	I	
DITOLAMIDE	II	
DIXYRAZINE	II	
DODECAMETHYLENEDIGUANIDE	II	
DOISYNOESTROL	I	E.U. : II
DOPAMINE	I	
DOXAPRAM	I	
DOXEPIN	I	
DOXORUBICIN	I	
DOXYCYCLINE	I	
DROPERIDOL	I	
DROSTANOLONE	I	
DUAZOMYCIN	I	
DYDROGESTERONE	I	

E

ECOTHIOPATE IODIDE	I	
ECTYLUREA	II	
EDOGESTRONE	I	
EDROPHONIUM CHLORIDE	I	

EMYLCAMATE	II	
ENALLYLPROPYMAL*		
ENRAMYCIN	I	
EPHEDRINE	II	W.P. : — U.D. not exceeding 30 mg — nasal application : not exceeding 2% — dermatological preparations
EPICILLIN	I	
EPIESTRIOL	I	E.U. : II
EPINEPHRINE	II	W.P. : topical use, including application to the skin
EPITIOSTANOL	I	
EPITIZIDE	II	
ERGOCORNINE	II	
ERGOCRISTINE	II	
ERGOKRYPTINE	II	
ERGOMETRINE	II	
ERGOTAMINE	II	W.P. : U.D. not exceeding 1 mg in a solution containing not more than 0.1%
ERGOTOXINE	II	
ERYTHROMYCIN	I	
ESERINE	II	
ESTRADIOL	I	E.U. : II
ETAMOCYCLINE	I	
ETEBENECID	II	
ETHACRYNIC ACID	II	
ETHADIONE	II	
ETHALLOBARBITAL*		
ETHCHLORVYNOL	II	
ETHIAZIDE	II	
ETHINAMATE	II	
ETHINYLESTRADIOL***	I	E.U. : II
ETHIONAMIDE	II	
ETHISTERONE	I	
ETHOHEPTAZINE	I	
ETHOSUXIMIDE	II	
ETHOTOIN	II	
ETHOXYZOLAMIDE	II	
ETHYLAMPHETAMINE	I	
ETHYL-5(BICYCLO 3,2,1)-5 OCTENYLBARBITURIC ACID*		
ETHYL-5(1-METHYLPROPYL)-5 THIOBARBITURIC ACID*		
ETHYL BUTYLETHYLMALONAMATE	II	
ETHYL BISCOUMACETATE	II	
ETHYLESTRENOL	II	
ETHYLIDENEDICOUMAROL	II	
ETHYLMORPHINE	II	W.P. : U.D. not exceeding 20 mg
ETHYLNORADRENALINE	II	
ETHYNERONE	I	
ETILEFRINE	II	W.P. : U.D. not exceeding 5 mg
ETODROXIZINE	II	
ETOGLUCID	I	
ETOZOLIN	II	
ETYBENZATROPINE	II	
ETYMEMAZINE	II	
ETYNODIOL***	I	
EUCATROPINE	II	

F

FEBARBAMATE*		
FELYPRESSIN	I	
FENADIAZOLE	II	

FENBENICILLIN	I	
FENBUTRAZATE	I	
FENCAMFAMIN	I	W.P. : U.D. not exceeding 10 mg
FENETHAZINE	II	
FENETYLLINE	I	
FENFLURAMINE	II	
FENMETRAMIDE	I	
FENOMISAL	II	
FENOTEROL	II	
FENPROPOREX	I	
FENYRAMIDOL	II	
FIBRINOLYSIN (human)	II	
FLOPROPIONE	II	
FLOXURIDINE	I	
FLUANISONE	I	
FLUCLOROLONE ACETONIDE	I	
FLUCLOXACILLIN	I	
FLUDROCORTISONE	I	
FLUDROXYCORTIDE	I	
FLUFENAMIC ACID	II	
FLUGESTONE	I	
FLUMEDROXONE	I	
FLUMETASONE	I	
FLUMETHIAZIDE	II	
FLUMINOREX	I	
FLUNISOLIDE	I	
FLUOCINOLONE	I	
FLUOCINOLONE ACETONIDE	I	
FLUOCINONIDE	I	
FLUOCORTOLONE	I	
FLUOROMETHOLONE	I	
FLUOROURACIL	I	
FLUOROURIDINE	I	
FLUOXYMESTERONE	I	
FLUPENTIXOL	II	
FLUPEROLONE	I	
FLUPHENAZINE	II	
FLUPREDNIDENE	I	
FLUPREDNISOLONE	I	
FLURANDRENOLONE	I	
FLUSPIRILENE	I	
FOLLOTROPINE	I	
FORMINITRAZOLE	II	
FORMOCORTAL	I	
FORMYLGITOXINE	II	
FRAMYCETIN	I	W.P. : — E.U. — lozenges
FUMAGILLIN	I	
FURALTADONE	II	W.P. : E.U.
FURAZABOL	I	
FURAZOLIDONE	II	W.P. : E.U.
FURFENOREX	I	
FURFURYLISOPROPYLBARBITURIC ACID*		
FURMETHOXADONE	II	W.P. : E.U.
FUROSEMIDE	II	
FUROSTILBESTROL	I	E.U. : II
FURTERENE	II	
FUSAFUNGINE	I	
FUSIDIC ACID	I	

G

GABOB (γ-AMINO-β-HYDROXY-BUTYRIC ACID)	II	
GALLANTAMINE	I	
GALLAMINE TRIETHIODIDE	I	
GELSEMINE	II	W.P. : U.D. not exceeding 0.5 mg
GENESERINE	II	W.P. : U.D. not exceeding 0.5 mg
GENTAMICIN	I	
GESTACLONE	I	
GESTONORONE CAPROATE	I	
GLAFENINE	II	W.P. : U.D. not exceeding 500 mg
GLIBENCLAMIDE	II	
GLIBORNURIDE	II	
GLICLAZIDE	II	
GLIDANILE	II	
GLISOXEPIDE	II	
GLUCOSULFAMIDE**	I	
GLUTETHIMIDE	II	
GLYBUTHIAZOL	II	
GLYBUZOLE	II	
GLYCERYL TRINITRATE	II	W.P. : — tablets containing not more than 1 mg — sustained-action preparations containing not more than 6.4 mg
GLYCLOPYRAMIDE	II	
GLYCOBIARSOL	II	
GLYCOPYRRONIUM BROMIDE	II	W.P. : U.D. not exceeding 1 mg *S.W. No. 3*
GLYCYCLAMIDE	II	
GLYHEXAMIDE	II	
GLYMIDINE SODIUM	II	
GLYOCTAMIDE	II	
GLYPARAMIDE	II	
GLYPINAMIDE	II	
GLYPROTHIAZOL	II	
GLYSOBUZOLE	II	
GRAMICIDIN	I	W.P. : — E.U. — lozenges
GRISEOFULVIN	I	E.U. : II
GUAMECYCLINE	I	
GUANETHIDINE	II	
GUANOCLOR	II	
GUANOXAN	II	

H

HACHIMYCINE	I	vaginal application : II
HALCINONIDE	I	
HALOPERIDOL	I	
HEDAQUINIUM CHLORIDE	II	
HEPTABARB*		
HEPTOBARBITAL*		
HEPTOLAMIDE	II	
HETACILLIN	I	
HEXAPROPYMATE	II	
HEXESTROL	I	E.U. : II
HEXETHAL*		
HEXOBARBITAL*		
HEXYLETHYLBARBITURIC ACID*		
HOMATROPINE	II	W.P. : U.D. not exceeding 0.5 mg *S.W. No. 3*
	I	ophthalmic preparations

HOMATROPINE METHYLBROMIDE	II	W.P. : U.D. not exceeding 2 mg *S.W. No. 3*
HOMOFENAZINE	II	
HYDRACARBAZINE	II	
HYDRALAZINE	II	
HYDRARGAPHEN	II	W.P. : E.U.
HYDROCHLOROTHIAZIDE	II	
HYDROCORTAMATE	I	
HYDROCORTISONE	I	
HYDROFLUMETHIAZIDE	II	
HYDROMADINONE	I	
HYDROXINDASATE	II	
HYDROXYCHLOROQUINE	II	W.P. : U.D. not exceeding 300 mg
HYDROXY-4-DIIODO-3,5-BENZOATE-n-BUTYL	I	
HYDROXYESTRONE	I	E.U. : II
HYDROXYPROGESTERONE	I	
HYDROXYSTENOZOLE	I	
HYDROXYZINE	II	W.P. : — tablets containing not more than 10 mg — suppositories containing not more than 50 mg
HYOSCINE	II	W.P. : U.D. not exceeding 0.3 mg *S.W. No. 3*
	I	ophthalmic preparations
HYOSCINE BUTYLBROMIDE	II	W.P. : U.D. not exceeding 10 mg *S.W. No. 3*
HYOSCINE METHOBROMIDE	II	W.P. : U.D. not exceeding 2.5 mg *S.W. No. 3*
HYOSCINE METHONITRATE	II	W.P. : U.D. not exceeding 3 mg *S.W. No. 3*
HYOSCYAMINE	II	W.P. : U.D. not exceeding 0.30 mg *S.W. No. 3*

I

IBUPROFEN	II	
IMICLOZAPINE	II	
IMIPRAMINE	I	
INDANOREX	I	
INDENESTROL	I	E.U. : II
INDOMETACIN	II	
INDOPINE	II	
INPROQUONE	I	
INTERMEDINE	I	
IODOPHENAZONE	II	W.P. : U.D. not exceeding 1 g *S.W. No. 2*
IPRINDOLE	I	
IPROCLOZIDE	I	
IPRONIAZIDE	I	
ISOAMINILE	II	W.P. : U.D. not exceeding 50 mg
ISOAMYLENGUANIDINE	II	
ISOCARBOXAZID	I	
ISOETARINE	II	
ISONIAZID	II	
ISOPRENALINE	II	W.P. : — E.U. : not more than 0.5 % — tablets containing not more than 20 mg
ISOPROMETHAZINE	II	
ISOPROPAMIDE IODIDE	II	W.P. : U.D. not exceeding 5 mg *S.W. No. 3*
ISOPYRIN	II	W.P. : U.D. not exceeding 500 mg *S.W. No. 2*
N.B. — INSULIN		W.P. even if used parenterally

J —

K

KANAMYCIN	I	
KETOPROFEN	II	
KITASAMYCIN	I	

L

LACTYLPHENETIDINE	II	W.P. : U.D. not exceeding 500 mg *S.W. No. 1*
LANATOSIDE C	II	
LANATOSIDES	II	
LAUDEXIUM	I	
LEVAMFETAMINE	I	
LEVARTERENOL	II	W.P. : topical use, including application to the skin
LEVISOPRENALINE	II	W.P. : tablets containing not more than 15 mg
LEVODOPA	I	
LEVOMEPROMAZINE	II	
LEVOPHACETOPERANE	I	
LEVOPROPICILLIN	I	
LEVOTHYROXINE SODIUM	II	
LINCOMYCIN	I	
LIOTHYRONINE	II	
LIVIDOMYCIN	I	
LOBELINE	II	W.P. : — E.U. — tablets containing not more than 3 mg — in liquid form containing a concentration not exceeding 0.5% w/v
LORAZEPAM	II	
LUCIMYCIN	I	
LUTUTRIN	I	
LYMECYCLINE	I	
LYNESTRENOL***	I	
LYPRESSIN	I	

M

MAFENIDUM**	I	
MANNOMUSTINE	I	
MAZINDOL	I	
MEBANAZINE	I	
MEBEVERINE	II	W.P. : U.D. not exceeding 100 mg
MEBOLAZINE	I	
MEBUTAMATE	II	
MECLOQUALONE	I	
MEDAZEPAM	II	
MEDROGESTONE	I	
MEDROXYPROGESTERONE	I	
MEDRYSONE	I	
MEFENAMIC ACID	II	W.P. : U.D. not exceeding 500 mg
MEFENOREX	I	
MEFRUSIDE	II	
MEGESTROL***	I	
MELENGESTROL	I	
MELITRACEN	I	
MELPHALAN	I	
MEPENZOLATE BROMIDE	II	W.P. : U.D. not exceeding 25 mg *S.W. No. 3*
MEPHENESIN	II	W.P. : ointments for E.U.
MEPHENOXALONE	II	

MEPHENTERMINE	I	
MEPHENYTOIN	II	
MEPROBAMATE	II	
MERALLURIDE	II	
MERBIURELIDIN	II	
MERCAPTOMERIN	II	
MERCAPTOPURINE	I	
MERCUDERAMIDE	II	
MERCUMATILIN SODIUM	II	
MERCUROPHYLLINE	II	
MERETHOXYLLINE	II	
MEROQUINOLAMIDE	II	
MERSALYL	II	
MERSALYL THEOPHYLLINE	II	
MESORIDAZINE	II	
MESTANOLONE	I	
MESTEROLONE	I	
MESTRANOL***	I	E.U. : II
MESULFAMIDE SODIUM**	I	
MESUXIMIDE	II	
METACYCLINE	I	
METAHEXAMIDE	II	
METAMFEPRAMONE	I	W.P. : U.D. not exceeding 50 mg
METAMPICILLIN	I	
METANDIENONE	II	W.P. : E.U. not exceeding 0.5%
METARAMINOL	II	
METASULFANILYL-BUTYL-CARBAMIDE	II	
METENOLONE	II	
METETHOHEPTAZINE	I	
METFORMIN	II	
METHALLATAL*		
METHALLENESTRIL	I	E.U. : II
METHAMPHETAMINE	I	
METHANDRIOL	II	
METHANTHELINIUM BROMIDE	II	W.P. : U.D. not exceeding 50 mg *S.W. No. 3*
METHAQUALONE	I	
METHARBITAL*		
METHAZOLAMIDE	II	
METHDILAZINE	II	W.P. : — U.D. not exceeding 5 mg — syrups containing not more than 0.4 mg/ml
METHEPTAZINE	I	
METHESTROL	I	E.U. : II
METHIOMEPRAZINE	II	
METHITURAL*		
METHOCARBAMOL	II	
METHOCIDIN	I	
METHOHEXITAL*		
METHOPROMAZINE	II	
METHOSERPIDINE	II	
METHOTREXATE	I	
METHOXAMINE	II	W.P. : nasal application not exceeding 0.25%
METHYCLOTHIAZIDE	II	
N-METHYLACETANILIDE	II	
METHYLANDROSTANDIOL	I	
N-METHYLBUTISOLE*		
METHYLCHLORESTRONE	I	E.U. : II
METHYLDOPA	II	
N,N-METHYLENE BIS-SULFACETAMIDE**	I	
METHYLEPHEDRINE	II	W.P. : U.D. not exceeding 30 mg

METHYLOESTRENOLONE	I	
METHYLPENTYNOL	II	
METHYLPENTYNOL CARBAMATE	II	
METHYLPHENIDATE	I	
METHYLPHENOBARBITAL*		
METHYLPREDNISOLONE	I	
METHYLTESTOSTERONE	I	
METHYLTHIOURACIL	II	
METHYPRYLON	II	
METHYSERGIDE	II	
METIAZINIC ACID	II	
METICILLIN	I	
METICRANE	II	
METOCLOPRAMIDE	II	
METOFENAZATE	II	
METOLAZONE	II	
METRIBOLONE	I	
METYLPERONE	I	
METYRAPONE	II	
MIKAMYCIN	I	
MIMBANE	II	
MINOCYCLINE	I	
MITHRAMYCIN	I	
MITOMYCIN	I	
MITOPODOZIDE	I	
MOPERONE	I	
MORAZONE	II	
MORPHOLINE	II	
MUCONOMYCINE A	I	E.U. : II

N

NAFCILLIN	I	
NAFTIDROFURYL	II	
NALOXONE	I	
NANDROLONE	I	
NAPHAZOLINE	II	W.P. : nasal application not exceeding 0.1%
NAPROXEN	II	
NARCOBARBITAL*		
NATAMYCIN	I	E.U. : II
PERCHLORATE SODIUM	II	
NEALBARBITAL*		
NEOARGENTARSPHENAMINE	II	
NEOARSPHENAMINE	II	W.P. : topical use
NEOCINCHOPHEN	II	
NEOMYCINS	I	W.P. : E.U.
NEOSTIGMINE BROMIDE	II	
NIALAMIDE	I	
NICOCODINE	II	
NICODICODINE	II	
NIDROXYZONE	II	W.P. : E.U.
NIFENAZONE	II	W.P. : — E.U. — U.D. not exceeding 500 mg *S.W. No. 2*
NIFLUMIC ACID	II	
NIFURATEL	II	W.P. : E.U.
NIFURETHAZONE	II	W.P. : E.U.
NIFURMERONE	II	W.P. : E.U.
NIFUROXAZIDE	II	W.P. : E.U.
NIFUROXIME	II	W.P. : E.U.
NIHYDRAZONE	II	W.P. : E.U.
NIRIDAZOLE	II	

NITRAZEPAM	II	
NITROFURANTOIN	II	W.P. : E.U.
NITROFURFURYLMETHYLETHER	II	W.P. : E.U.
NITROMETHAQUALONE	I	
NOPRYLSULFAMIDE**	I	
NORAMIDOPYRINE METHANESULFONATE SODIUM	II	W.P. : — E.U. — U.D. not exceeding 1 g *S.W. No. 2*
NORBOLETONE	I	
NORCODEINE	II	
DL-NOREPHEDRINE	II	W.P. : — U.D. not exceeding 50 mg — nasal application not exceeding 2%
NORETHANDROLONE	II	
NORETHISTERONE***	I	
NORETYNODREL***	I	
NORFENEFRINE	II	W.P. : U.D. not exceeding 10 mg
NORGESTREL***	I	
D-NORGESTREL***	I	
NORGESTRIENONE***	I	
NORTRIPTYLINE	I	
NORVINISTERONE	I	
NOVOBIOCIN	I	
NYSTATIN	I	W.P. : vaginal application

O

OCTATROPINE METHYLBROMIDE	II	
OESTROFURATE	I	E.U. : II
OESTROGENIC SUBSTANCES, conjugated	I	E.U. : II
OESTRONE	I	E.U. : II
OLEANDOMYCIN	I	
OLEANDRINE	II	
OPIPRAMOL	I	
ORAL-TURINABOL	I	
ORCIPRENALINE	II	W.P. : tablets containing not more than 20 mg
ORPHENADRINE	II	
ORTETAMINE	I	
OSTREOGRYCIN	I	
OUABAIN	II	
OXABOLONE CIPIONATE	I	
OXACILLIN	I	
OXANAMIDE	II	
OXANDROLONE	II	
OXAZEPAM	II	
OXITEFONIUM BROMIDE	II	
OXOGESTONE	I	
OXOLAMINE	II	W.P. : — U.D. not exceeding 50 mg — syrups containing not more than 1.5%
OXOMEMAZINE	II	W.P. : — U.D. not exceeding 10 mg — syrups containing not more than 1 mg/ml
OXOPHENARSINE	I	used as an anti-parasitic : II W.P. : topical use
OXPRENOLOL	II	
OXYCINCHOPHEN	II	W.P. : E.U.
OXYMESTERONE	II	
OXYMETHOLONE	I II	U.D. exceeding 5 mg U.D. not exceeding 5 mg
OXYPERTINE	II	
OXYPHENBUTAZONE	II	
OXYPHENCYCLIMINE	II	W.P. : U.D. not exceeding 10 mg *S.W. No. 3*

OXYPHENERIDINE	I	
OXYPHENONIUM BROMIDE	II	W.P. : U.D. not exceeding 5 mg *S.W. No. 3*
OXYTETRACYCLINE	I	
OXYTOCIN	II	W.P. : nasal preparations
P		
PANCURONIUM BROMIDE	I	
PAPAVERINE	II	W.P. : U.D. not exceeding 50 mg
PARA-AMINOSALICYLIC ACID	II	
PARAFLUTIZIDE	II	
PARALDEHYDE	II	
PARAMETHADIONE	II	
PARAMETHASONE	I	
PARAMETHYLAMPHETAMINE	I	
PARAOXONE	II	
PARATHIAZINE	II	W.P. : — U.D. not exceeding 25 mg — suppositories containing not more than 80 mg
PARGYLINE	I	
PAROMOMYCIN	I	
PAROXYPROPIONE	I	
PECAZINE	II	
PECILOCIN	I	
PEMOLINE	I	
PENAMECILLIN	I	
PENETHACILLINE	I	
PENFLURIDOL	I	
PENIMEPICYCLINE	I	
PENMESTEROL	I	
PENTAGESTRONE	I	
PENTAMETHONIUM BROMIDE	I	
PENTAZOCINE	I	
PENTHIENATE	II	W.P. : U.D. not exceeding 5 mg *S.W. No. 3*
PENTOBARBITAL*		
PENTOREX	I	
PERAZINE	II	
PERICIAZINE	II	
PERPHENAZINE	II	
PERPHENAZINE SULFOXIDE	II	
PETRICHLORAL	II	
PHANQUINONE	II	W.P. : U.D. not exceeding 50 mg
PHENACEMIDE	II	
PHENACETINE	II	W.P. : U.D. not exceeding 500 mg *S.W. No. 1*
PHENAGLYCODOL	II	
PHENALLYMAL*		
PHENANTRYLMETHYLHYDANTOIN	II	
PHENATINE	I	
PHENAZONE	II	W.P. : — E.U. — U.D. not exceeding 1 g *S.W. No. 2*
PHENBUTAMIDE	II	
PHENDIMETRAZINE	I	
PHENELZINE	I	
PHENERIDINE	I	
PHENETICILLIN	I	
PHENFORMIN	II	
PHENGLUTARIMIDE	II	
PHENICARBAZIDE	II	W.P. : U.D. not exceeding 500 mg *S.W. No. 1*

PHENINDIONE	II	
PHENMETRAZINE	I	
PHENOBARBITAL*		
PHENOXYBENZAMINE	II	
PHENOXYMETHYLPENICILLIN	I	
PHENPROBAMATE	II	
PHENPROCOUMON	II	
PHENPROMETHAMINE	II	W.P. : nasal application not exceeding 3 %
PHENSUXIMIDE	II	
PHENTERMINE	I	
PHENTOLAMINE	II	
PHENYLBUTAZONE	II	W.P. : E.U.
PHENYLDIBROMOMETHYL-METHYLHYDANTOIN	II	
PHENYTOIN	II	
PHLORETINE	I	E.U. : II
PHOLCODINE	II	W.P. : U.D. not exceeding 40 mg
PHTHALYLSULFACETAMIDE**	II	
PHTHALYLSULFAMETHIZOLE**	I	
PHTHALYLSULFATHIAZOLE**	II	
PILOCARPINE	II	W.P. : hair lotions containing not more than 1%
PIMOZIDE	II	
PIPAMAZINE	II	
PIPAMPERONE	I	
PIPENZOLATE BROMIDE	II	W.P. : U.D. not exceeding 5 mg *S.W. No. 3*
PIPERACETAZINE	II	
PIPERIDOLATE	II	W.P. : U.D. not exceeding 50 mg *S.W. No. 3*
PIPERIDYLETHYLBARBITURIC ACID*		
PIPEROXAN	II	
PIPERYLONE	II	W.P. : U.D. not exceeding 125 mg *S.W. No. 2*
PIPETHANATE	II	
PIPOBROMAN	I	
PIPOSULFAN	I	
PIPOTIAZINE	II	
PIPRADROL	I	
PIPROCURARIUM IODIDE	I	
PITUITARY POSTERIOR LOBE	II	W.P. : nasal preparations
PIVAMPICILLIN	I	
POLDINE	II	W.P. : U.D. not exceeding 2 mg *S.W. No. 3*
POLYESTRADIOL	I	E.U. : II
POLYMYXIN B	I	
POLYTHIAZIDE	II	
PORFIROMYCIN	I	
PRACTOLOL	II	
PREDNISOLONE	I	
PREDNISONE	I	
PREDNYLIDENE	I	
PREGNENOLONE	I	
PRENOXDIAZINE	II	
PRENYLAMINE	II	W.P. : U.D. not exceeding 60 mg
PRIMIDONE	II	
PRISTINAMYCIN	I	
PROBARBITAL SODIUM*		
PROBENECID	II	W.P. : U.D. not exceeding 500 mg
PROCAINAMIDE	II	
PROCARBAZINE	I	
PROCHLORPERAZINE	II	
PROCYCLIDINE	II	
PROCYMATE	II	

PRODILIDINE	II	
PROFENAMINE	II	
PROGESTERONE	I	
PROLACTINE	I	
PROLINTANE	I	W.P. : U.D. not exceeding 10 mg
PROMAZINE	II	
PROMESTRIENE	I	E.U. : II
PROMETHAZINE	II	W.P. : — suppositories containing not more than 50 mg — creams containing not more than 2% — syrups containing not more than 1 mg/ml — tablets containing not more than 25 mg
PRONETALOL	II	
PROPALLYLONAL*		
PROPANTHELINE BROMIDE	II	W.P. : U.D. not exceeding 15 mg *S.W. No. 3*
PROPETANDROL	I	
PROPICILLIN	I	
PROPIOMAZINE	II	
PROPRANOLOL	II	
PROPYLBARBITAL*		
PROPYLTHIOURACIL	II	
PROPYPHENAZONE	II	W.P. : — E.U. — U.D. not exceeding 1 g *S.W. No. 2*
PROPYROMAZINE BROMIDE	II	
PROTHIPENDYL	II	
PROTIONAMIDE	II	
PROTOANEMONIN	I	
PROTOKYLOL	II	
PROTRIPTYLINE	I	
PSEUDOEPHEDRINE	II	W.P. : — E.U. not exceeding 3% — U.D. not exceeding 60 mg
PUROMYCIN	I	
PYRANTEL	II	W.P. : U.D. not exceeding 250 mg
PYRAZINAMIDE	II	
PYRIDOSTIGMINE BROMIDE	II	
PYRITHYLDIONE	II	
PYROPHOS	I	
PYRROLIFENE	II	

Q

QUINACILLIN	I	
QUINESTRADOL	I	E.U. : II
QUINESTROL	I	E.U. : II
QUINETHAZONE	II	
QUINGESTANOL***	I	
QUINGESTRONE	I	
QUINIDINE	II	

R

RAUBASINE	II	
RAUWOLFIA, total alkaloids	II	
RENANOLONE	I	
RESCINNAMINE	II	
RESERPILINE	II	
RESERPINE	II	
RIDAZINE	II	
RIFAMIDE	I	
RIFAMPICIN	I	

RIFAMYCIN	I	
RISTOCETIN	I	
ROLITETRACYCLINE	I	
RONIDAZOLE	II	
RUFOCROMOMYCIN	I	
S		
SALAZOSULFADIMIDINE**	I	
SALAZOSULFAMIDE**	I	
SALAZOSULFAPYRIDINE**	I	
SALAZOSULFATHIAZOLE**	I	
SALBUTAMOL	II	
SARKOMYCINS	I	
SQUILL, isolated glycosides of	II	
SECBUTABARBITAL*		
SECOBARBITAL*		
SERUM GONADOTROPHIN	I	
SILANDRONE	I	
SODIUM STIBOCAPTATE	II	
SOMATOTROPHINE	I	
SPARSOMYCIN	I	
SPECTINOMYCIN	I	
SPIRAMYCIN	I	E.U. : II
SPIRONOLACTONE	I	
SPIROXASONE	II	
STANOZOLOL	II	
STENBOLONE	I	
STIBOPHEN	II	
STREPTODUOCIN	I	
STREPTOKINASE	II	
STREPTOMYCIN	I	
STREPTONIAZID	I	
STREPTONIGRIN	I	
STREPTOVARYCIN	I	
STRONTIUM SALT OF ASCORBOPHENYLBUTAZONE	II	
STROPHANTUS, isolated glycosides of	II	
STYRAMATE	II	
SUCCINYLSULFATHIAZOLE**	II	
SULFACARBAMIDE**	II	
SULFACETAMIDE**	I	
SULFACHLORPYRIDAZINE**	I	
SULFACHRYSOIDINE**	I	
SULFACINNAMINE**	I	
SULFACITINE**	I	
SULFACLOMIDE**	I	
SULFADIASULFONE SODIUM**	I	
SULFADIAZINE**	I	
SULFADICRAMIDE**	I	
SULFADIMETHOXINE**	I	
SULFADIMETHOXYTRIAZINE**	I	
SULFADIMIDINE**	I	
SULFADOXINE**	I	
SULFAETHIDOLE**	I	
SULFAETHYLPYRAZOLE**	I	
SULFAFURAZOLE**	I	
SULFAGUANIDINE**	II	
SULFALENE**	I	
SULFALOXIC ACID**	II	
SULFAMERAZINE**	I	
SULFAMETHIZOLE**	I	
SULFAMETHOXAZOLE**	I	

SULFAMETHOXYPYRIDAZINE**	I	
SULFAMETHYLTHIAZOL**	I	
SULFAMETOMIDINE**	I	
SULFAMETOXYDIAZINE**	I	
SULFAMETOYL**	I	
SULFAMIDOCHRYSOIDINE**	I	
SULFAMIDOMALEYL**	I	
SULFAMIDOPYRINE	II	W.P. : — E.U. — U.D. not exceeding 1 g *S.W. No. 2*
SULFAMONOMETHOXINE**	I	
SULFAMOXOLE**	I	
5-SULFANILAMIDO-1,3-DIMETHYL-2,6-DIOXO-4-IMINO HEXAHYDROPYRIMIDINE**	I	
SULFANILAMIDE**	I	
SULFANITRAN**	I	
SULFAPERIN**	I	
SULFAPHENAZOLE**	I	
SULFAPROXYLINE**	I	
SULFAPYRAZOLE**	I	
SULFAPYRIDINE**	I	
SULFARSIDE**	I	
SULFARSPHENAMINE	II	
SULFASOMIZOLE**	I	
SULFASUCCINAMIDE **	I	
SULFASYMAZINE**	I	
SULFATHIAZOLE**	I	
SULFATHIAZOL FORMALDEHYDE**	II	
SULFATHIOUREA**	I	
SULFATOLAMIDE**	I	
SULFISOMIDINE**	I	
SULFISOXAZOL DIETHANOLAMINE**	I	
SULFODIAMINE**	I	
SULFOMYXIN	I	
SULOCARBILATE	II	
SULPHABENZAMIDE**	I	
SULPHAMOPRINE**	I	
SULTIAME	II	
SURAMIN SODIUM**	I	
SUXAMETHONIUM CHLORIDE	I	
SUXETHONIUM CHLORIDE	I	

T

TALBUTAL*		
TAMOXIFEN	I	
TECLOTHIAZIDE	II	
TEMAZEPAM	II	
TERBUTALINE	II	
TESTOSTERONE	I	
TETRABARBITAL*		
TETRABENAZINE	II	
TETRACOSACTIDE	I	
TETRACYCLINE	I	
TETRAMISOLE	II	W.P. : U.D. not exceeding 500 mg
TETRYLAMMONIUM BROMIDE	I	
THIALBARBITAL*		
THIAMAZOL	II	
THIAMPHENICOL	I	
THIAZANOL	II	
THIAZINAMINE	II	W.P. : U.D. not exceeding 300 mg

THIAZOSULFONE**	I	
THIETYLPERAZINE	II	
THIOBARBITAL*		
THIOBUTOBARBITAL*		
THIOFURADENE	II	W.P. : E.U.
THIOHEXAMIDE	II	
THIOPENTAL SODIUM*		
THIOPORAN	II	
THIOPROPAZATE	II	
THIOPROPERAZINE	II	
THIORIDAZINE	II	
THIOSINAMINE	I	
THIOSTREPTONE	I	
THIOTEPA	I	
THYROGLOBULIN	II	
THYROTROPHIN	I	
THYROXINE	II	
TIAMIZIDE	II	
TIBOLONE	I	
TIEMONIUM IODIDE	II	W.P. : U.D. not exceeding 30 mg *S.W. No. 3*
TIFORMIN	II	
TIGESTOL	I	
TIOCARLIDE	II	
TIOGUANINE	I	
TIOMESTERONE	I	
TIOTIXENE	II	
TOBRAMYCIN	I	
TOFISOPAM	II	
TOLAZAMIDE	II	
TOLAZOLINE	II	W.P. : — E.U. — U.D. not exceeding 5 mg
TOLBOXANE	II	
TOLBUTAMIDE	II	
TOLPENTAMIDE	II	
TOLPRONINE	II	
TOLPYRRAMIDE	II	
TRANEXAMIC ACID	II	
TRANSCLOMIFENE	I	E.U. : II
TRANYLCYPROMINE	I	
TRENBOLONE	I	
TRENGESTONE	I	
TRESTOLONE	I	
TRETAMINE	I	
TRIAMCINOLONE ACETONIDE	I	
TRIAMCINOLONE HEXACETONIDE	I	
TRIAMCINOLONE	I	
TRIAMTERENE	II	
TRIAZIQUONE	I	
TRIBROMOETHANOL	II	
TRICHLORMETHIAZIDE	II	
TRICHLORMETHINE	I	
TRICLOFENOL PIPERAZINE	II	
TRICYCLAMOL CHLORIDE	II	
TRIDIHEXETHYL IODIDE	II	
TRIFLUOPERAZINE	II	
TRIFLUOREX	I	
TRUFLUPERIDOL	I	
TRIFLUPROMAZINE	II	
TRIHEXYPHENIDYL	II	
TRIIODOTHYROACETIC ACID	II	
TRIIODOTHYRONINE	II	

TRIMETHADIONE	II	
TRIMETOZINE	II	
TRIMIPRAMINE	I	
TRIOXYMETHYLMELAMINE	I	
TROLEANDOMYCIN	I	
TROPENZILINE BROMIDE	II	
TROPICAMIDE	II	
TUAMINOHEPTANE	II	W.P. : nasal drops
TUBOCURARINE CHLORIDE	I	
TYLOSIN	I	
TYROCIDIN	I	W.P. : — E.U. — lozenges
TYROTHRICIN	I	W.P. : - E.U. — lozenges

U

URAMUSTINE	I	
URETHANE	I	
UROKINASE	II	
USNIC ACID	I	

V

VALETHAMATE BROMIDE	II	W.P. : U.D. not exceeding 10 mg *S.W. No. 3*
VALNOCTAMIDE	II	
VANCOMYCIN	I	
VASOPRESSIN INJECTION	I	
VINBARBITAL*		
VINELASTINE	I	
VINCRISTINE	I	
VINLEUROSINE	I	
VINROSIDINE	I	
VINYLBITAL*		
VIOMYCIN	I	
VIRGINIAMYCIN	I	E.U. : II

W

WARFARIN	II	
WARFARINDEANOL	II	

X

XANTOCILLIN	I	

Y

YOHIMBINE	II	

Z

ZERANOL	I	