

Table S3: Qualitative evaluation of textbook information missing in the German Wikipedia.

Textbook information not present in Wikipedia (keywords)	Evaluation by lecturers (A-F)						sum	didactic relevance (yes if sum > 4)
	(1= didactically relevant; 0 = didactically not relevant)							
	A	B	C	D	E	F		result
<b>Acetylsalicylic acid</b>								
Acute coronary syndrome	1	1	1	1	1	1	6	yes
Migraine attack	1	1	1	0	1	0	4	no
Inhibition of endothelial prostacyclin synthesis	1	1	0	0	1	1	4	no
CNS side effects (high doses)	0	1	0	0	1	0	2	no
<b>Alendronic acid</b>								
Osteoclast uptake	1	1	1	1	0	1	5	yes
<b>Allopurinol</b>								
Isomere of hypoxanthin	1	0	1	0	0	1	3	no
Oral application	0	1	0	0	1	1	3	no
<b>Amoxicillin</b>								
H. influenzae infections	0	1	1	1	0	1	4	no
High oral resorption when compared to Ampicillin	1	0	1	1	1	1	5	yes
<b>Atropine</b>								
Iridolysis	0	0	0	0	1	1	2	no
Mydriasis	1	1	0	1	1	1	5	yes
50% renal elimination	0	0	0	0	0	0	0	no
<b>Benzylpenicillin</b>								
Neurotoxicity (high doses)	1	1	0	0	1	0	3	no
<b>Biperiden</b>								
IND: M. Parkinson	1	1	1	1	1	1	6	yes
<b>Bromocriptine</b>								
Increased GH incretion	1	0	1	0	0	1	3	no
Pleural fibrosis	0	0	0	1	1	1	3	no
<b>Caffeine</b>								
Increase of catecholamine levels	0	1	1	0	1	1	4	no
Interactions with uric acid metabolism	0	0	0	0	0	0	0	no
Tremor	1	1	1	0	1	1	5	yes
Seizures	1	1	0	1	1	1	5	yes
Anxiety	0	1	0	0	0	1	2	no
Increased gastric acid production	1	1	1	0	0	1	4	no
<b>Candesartan</b>								
Long duration of action	1	1	0	0	1	0	3	no
<b>Carbamazepine</b>								
Use-dependent block of neuronal discharges	1	0	1	1	0	1	4	no
<b>Cefepime</b>								
IND: Pseudomonde infections	1	1	1	1	1	1	6	yes
IND: infections with gram negative bacteria	1	1	1	1	1	1	6	yes
IND: infections with gram positive bacteria	1	1	0	1	1	1	5	yes
<b>Ciclosporin</b>								
IND: autoimmune disorders, rheumatoide arthritis	1	0	1	1	1	1	5	yes
Variable oral bioavailability	1	1	1	1	0	1	5	yes
Predominant hepatic biotransformation	1	1	0	1	1	1	5	yes
Predominant hepatic elimination	0	0	0	0	0	1	1	no
Hyperglycemia	0	0	0	1	0	1	2	no
<b>Ciprofloxacin</b>								
Inhibition of topoisomerase II	1	1	1	0	0	1	4	no
Renal elimination	1	0	0	0	1	1	3	no
CNS adverse effects	1	0	0	0	1	0	2	no
<b>Clarithromycin</b>								
Alternative antibiotic for patients with penicillin allergy	1	1	0	1	0	1	4	no
<b>Cyclophosphamide</b>								
Immunosuppressive effect via inhibition of leucocytes	1	0	1	1	1	1	5	yes
<b>Diazepam</b>								
Dose-dependent anxiolytic, sedative, myotonolytic or anti-convulsant	1	1	1	1	1	1	6	yes
conjugation of hydroxyl group during biotransformation	0	0	0	0	0	0	0	no
Reduced / slowed elimination in liver disease	1	0	1	1	0	1	4	no
Floppy infant syndrom	1	1	0	1	1	1	5	yes
Reduced libido	0	0	0	0	0	1	1	no
<b>Domperidone</b>								
IND: migraine attacks	0	1	0	0	1	1	3	no
Reflux	1	0	0	0	1	0	2	no
<b>Doxazosin</b>								
Tachycardia	1	1	0	0	1	1	4	no
Increased heart failure incidence	0	1	1	1	1	1	5	yes

<b>Doxycycline</b>								
IND: Proprii bacteria infections	0	1	0	1	0	1	3	no
IND: Yersinia infections	0	0	0	1	0	1	2	no
IND: Protozoa infections	0	0	0	1	0	1	2	no
IND: Lymphgranuloma venerum	0	0	0	0	0	1	1	no
Penetrates blood-placenta-barrier	0	1	0	1	1	1	4	no
Variable penetration of blood-brain-barrier	0	0	0	0	0	1	1	no
Plasma protein binding 90%	0	0	0	0	0	0	0	no
Onycholysis	0	0	0	0	0	1	1	no
Increase of intracranial pressure	0	0	0	0	0	1	1	no
Esophageal mucositis	0	1	0	1	0	0	2	no
<b>Estradiol</b>								
IND: hormone therapy in menopause	1	1	1	1	1	1	6	yes
Binding to estradiol response element	1	0	0	1	1	1	4	no
Suppression of ovulation	1	1	0	1	1	1	5	yes
Higher oral dose	0	1	0	0	0	0	1	no
Hepatic elimination	1	1	0	1	1	0	4	no
Elimination product estriol	0	0	0	0	0	1	1	no
Glucuronidation, sulphation	0	1	0	1	0	1	3	no
Role of renal elimination	1	1	0	0	0	1	3	no
Transdermal application	1	1	1	1	1	1	6	yes
Thrombosis	1	1	1	1	1	1	6	yes
Endometrial carcinoma	1	1	1	1	1	1	5	yes
Edema	1	1	1	1	0	1	5	yes
Cardiovascular complications	1	1	1	1	1	1	6	yes
<b>Ethanol</b>								
Increase in blood pressure	0	1	0	0	1	0	2	no
Hypoglycemia	1	1	1	0	1	0	4	no
Increased diuresis	1	1	0	1	0	1	4	no
Decreased sexual potency	1	1	0	0	0	1	3	no
Distribution in total body water	0	1	0	1	1	1	4	no
Elimination increased in alcoholics	1	1	0	0	0	1	3	no
CYP450 metabolism	0	1	0	1	0	0	2	no
Energy source	1	1	0	1	1	1	5	yes
<b>Flucloxacillin</b>								
Effectiveness against penicilline-sensitive bacteria	0	1	0	1	1	1	4	no
Oral application	1	1	0	1	1	1	5	yes
Half-life 30 bis 60 min	0	0	0	0	0	0	0	no
High plasma protein binding	0	0	0	0	0	0	0	no
<b>Fluconazole</b>								
High water solubility	0	1	0	0	0	0	1	no
Application oral or i.v.	1	1	0	1	1	1	5	yes
Oral bioavailability > 90%	0	1	0	1	0	1	3	no
High liquor concentration	1	1	0	1	1	1	5	yes
Half-life 30 h	1	1	0	1	0	0	3	no
Mainly unmodified renal elimination	1	1	0	0	1	1	4	no
Dose adaption in renal insufficiency	1	1	1	1	1	1	6	yes
<b>Flumazenil</b>								
Half-life 1 h	1	1	0	0	0	0	2	no
<b>Furosemide</b>								
Effective if GFR <30 ml/min	1	1	1	1	1	1	6	yes
Increased thrombosis risk	1	1	1	1	1	1	6	yes
<b>Gentamicin</b>								
Intracellular site of action	0	0	0	0	0	1	1	no
No penetration of blood-liquor-barrier	1	1	1	0	1	1	4	no
Low plasma protein binding	0	1	0	0	0	0	1	no
No metabolism	1	1	0	0	0	1	3	no
Renal elimination by glomerular filtration	1	1	0	1	1	1	5	yes
Reduced elimination in renal insufficiency	1	1	1	1	1	1	6	yes
CI: pregnancy	1	1	0	1	1	1	4	no
<b>Glyceryl trinitrate</b>								
Duration of action 15-30 min	1	1	1	0	1	1	5	yes
<b>Goserelin</b>								
Regulation of GnRH-receptors	0	1	0	1	1	1	4	no
Reduction of sex hormone levels	1	1	1	1	1	1	6	yes
Hot flush	1	1	1	1	1	1	6	yes
<b>Heparin, unfractionated</b>								
Tissue irritation at injection site	0	0	0	0	0	1	1	no
<b>Hydrochlorothiazide</b>								
Hypercalciuria	0	1	0	1	1	1	4	no
Onset of action after 1-2 h	0	1	0	0	1	0	2	no
Loss of action if renal function is reduced	0	1	1	1	1	1	5	yes
<b>Ibuprofen</b>								
Oxidation as biotransformation process	0	0	0	0	0	0	0	no
<b>Imipenem</b>								
Beta-lactamase stable	1	1	1	1	1	1	6	yes
<b>Insulin lispro</b>								
Duration of action 2 h	1	1	0	0	1	1	4	no
Max. drug action after 40 min	1	1	0	0	1	1	4	no

<b>Isoflurane</b>								
MAC-value 1.2%	0	1	0	0	1	0	2	no
Metabolisation 0.2%	0	1	0	1	0	1	3	no
Quick onset of drug action	1	1	0	0	1	1	4	no
<b>Isoniazid</b>								
High tissue distribution	1	1	0	1	1	1	5	yes
<b>Lamotrigine</b>								
IND: focal seizures	1	1	1	1	0	1	5	yes
IND: Lennox-Gastaut-syndrome	0	0	0	1	1	0	2	no
<b>Levodopa</b>								
Metabolization to dopamin by dopa-decarboxylase	1	1	1	1	1	1	6	yes
Loss of efficacy after longtime use	1	1	0	1	1	1	5	yes
<b>Lithium</b>								
Hyperthyreotic metabolic state	0	1	0	0	0	1	2	no
Therapeutic effect after 1-2 weeks	1	1	1	1	1	1	6	yes
Prophylactic effects after 6-12 months	1	1	1	1	1	1	6	yes
Maximal plasma concentration after 2 h	0	1	0	0	0	0	1	no
Euthyret struma	1	1	1	1	1	1	6	yes
ECG abnormalities	1	1	0	1	1	1	5	yes
Renal diabetes insipidus	1	1	0	1	1	1	5	yes
<b>Loperamide</b>								
Inhibition of propulsive peristalsis	1	1	1	1	1	1	6	yes
Reduction of fluid loss	1	1	1	1	1	1	6	yes
CNS AE in children	1	1	1	1	1	1	6	yes
<b>Metformin</b>								
Unmodified renal elimination	1	1	1	0	1	1	5	yes
Half-life 2-5 h	0	1	0	0	0	0	1	no
Caution if used in elderly patients	1	1	1	1	0	0	4	no
<b>Methotrexate</b>								
Direct inhibition of enzymes	1	1	0	0	1	0	3	no
Drug resistance	0	1	0	0	1	0	2	no
<b>Metoclopramide</b>								
Antiemetic effect	1	1	1	1	1	1	6	yes
Increased tonus of lower esophagus sphincter	1	1	0	1	1	1	5	yes
Higher risk for AE in children	1	1	0	1	1	0	4	no
<b>Metoprolol</b>								
Blood pressure reduction after weeks	1	1	0	0	1	1	4	no
<b>Metronidazole</b>								
HP triple therapy	1	1	1	0	0	0	3	no
<b>Mirtazapine</b>								
Agitated depression	1	0	0	0	0	0	1	no
<b>Morphine</b>								
Sedation	1	1	1	1	1	1	6	yes
Oral bioavailability 15%	1	1	0	1	1	1	5	yes
Hepatic glucuronidation	1	0	0	0	0	0	1	no
Half-life: several hours	1	0	1	0	0	1	3	no
AE: increased GI-tonus	1	1	1	1	1	1	6	yes
AE: bradycardia	0	1	0	1	1	1	4	no
AE: miosis	1	1	0	1	1	1	5	yes
AE: effect on hypophyseal hormones	0	0	0	1	1	1	3	no
AE: delayed gastric emptying	0	1	0	1	1	1	4	no
AE: spastic constipation	1	1	1	1	1	1	6	yes
CI: lung diseases	0	0	0	0	1	0	1	no
Use during birth	1	0	0	1	1	1	4	no
<b>Naloxone</b>								
Use in neonatal medicine	0	1	0	0	1	0	2	no
Half-life approx. 1 h	1	1	1	0	1	0	4	no
<b>Nifedipine</b>								
Rapid onset of action	1	1	1	1	1	1	6	yes
Oral bioavailability 50%	0	1	0	0	0	0	1	no
Half-life 2 h	1	1	0	0	0	0	2	no
<b>Nicotine</b>								
Intoxication: excitation, followed by depolarisation block	1	1	0	1	1	1	5	yes
Absorption pH-dependent	0	1	0	1	0	0	2	no
Elimination: 10% unmodified renal elimination	0	1	0	1	0	0	2	no
Increase of plasma lipids	1	0	0	1	1	1	4	no
GI-ulcera	1	0	0	0	1	1	3	no
<b>Norepinephrine</b>								
$\alpha_2$ -mediated inhibition of noradrenalin release	1	1	1	0	1	1	5	yes
Effect on $\beta_2$ receptors	0	1	0	1	1	1	4	no
Renale elimination	1	1	0	0	0	1	3	no
<b>Omeprazole</b>								
Duration of action until re-synthesis of proton pump	1	1	1	0	1	1	5	yes
Duration of action > half-time (irreversible inhibition of proton pump)	1	1	0	0	1	1	4	no
Hypergastrinemia	1	0	0	0	1	0	2	no

<b>Ondansetron</b>								
Co-medication with steroids results in increased efficacy	0	1	1	1	1	1	5	yes
Half-life 3.5 h	0	0	0	0	0	0	0	no
Duration several hours	0	0	0	1	1	1	3	no
<b>Pancuronium</b>								
IND: intoxications with elevated muscle tension	1	0	0	0	1	1	3	no
<b>Perchlorate</b>								
Aplastic anemia	1	1	1	1	1	0	5	yes
<b>Physostigmine</b>								
Carbamylation of ACh-esterase	0	0	1	0	0	0	1	no
<b>Pilocarpine</b>								
Accommodation errors	1	1	0	1	1	1	5	yes
Reduced twilight vision	0	1	0	1	1	1	4	no
Systemic actions	0	0	0	0	1	1	2	no
<b>Prednisolone</b>								
IND: leucemia	1	1	1	1	1	0	5	yes
Decreased cytokine synthesis	1	1	1	1	1	1	6	yes
Inhibition of all inflammation phases	1	0	1	1	1	1	5	yes
Negative feedback mechanisms	1	1	1	1	1	1	6	yes
Affinity to mineral corticoid receptor	1	1	1	1	1	1	6	yes
Oral application	0	1	0	0	1	1	3	no
Glucuronidation, sulphatation	0	1	0	0	0	0	1	no
Sodium retention, potassium excretion via mineral corticoid receptor action	1	1	1	0	1	1	5	yes
Muscle weakness	1	1	0	1	1	1	5	yes
Euphoria	1	1	1	1	1	1	6	yes
Adrenal cortex atrophy	1	1	1	1	1	1	6	yes
Glaucoma, cataract	0	1	1	1	1	1	5	yes
Cl: osteoporosis	1	1	1	1	1	1	6	yes
Cl: GI-ulcers	1	1	1	1	1	1	6	yes
Cl: hypertonus	1	1	1	0	1	1	5	yes
Cl: diabetes mellitus	1	1	0	1	1	1	5	yes
Cl: infectious diseases	1	1	1	1	1	1	6	yes
Cl: pregnancy	1	1	0	1	1	1	5	yes
<b>Propofol</b>								
Drug action after 1 min	0	1	0	0	1	1	3	no
Duration of action 10 min	1	1	0	0		1	3	no
Half-time 30 min	0	1	0	0		0	1	no
Hepatic metabolisation	1	1	0	0	0	1	3	no
<b>Pyrazinamide</b>								
Inhibition of fatty acid synthesis by pyrazincarbonate acid	0	0	0	0	0	0	0	no
Inhibition of mycolic acid synthesis	1	1	1	1	1	1	6	yes
High tissue distribution	0	1	0	0		1	2	no
<b>Ranitidine</b>								
Dizziness	1	0	0	0	1	1	3	no
<b>Rifampicin</b>								
Application p.o.	0	1	0	0	0	1	2	no
Plasma protein binding 80%	0	1	0	0	0	0	1	no
High organ distribution	1	1	0	1	0	1	4	no
Half-life 2-5 h	0	1	0	0	0	0	1	no
After longtime use reduction of half-life	0	1	0	1	1	1	4	no
GI-tract side effects	0	1	0	0	1	1	3	no
Adverse effects of the skin	1	1	0	0	1	1	4	no
Neurological side effects	1	0	0	0	0	1	2	no
Flu symptoms	0	1	0	0	1	1	3	no
Use in pregnancy	0	1	0	0	1	1	3	no
Use during lactation	0	1	0	0	1	1	3	no
<b>Suxamethonium</b>								
Blockade of action potentials	1	1	0	0	1	1	4	no
Longlasting paralysis	1	1	0	1	1	0	4	no
<b>Tazobactam</b>								
IND: Severe infections	0	1	0	0	1	1	3	no
<b>Thiamazole (Methimazole)</b>								
Inhibition of iodine incorporation to tyrosines	1	1	1	0	1	1	5	yes
Delayed thyreostatic effect	1	1	1	1	1	1	6	yes
<b>Valproic acid</b>								
IND: focal seizures	1	0	1	1	0	1	4	no
Inhibition of T-type-calcium channels	0	0	0	0	0	1	1	no
High plasma protein binding	1	1	0	1	0	0	3	no
<b>Vancomycin</b>								
No penetration of outer membrane in gram negative bacteria	0	0	1	1	1	1	4	no
No hemodialysis	0	0	0	1	0	0	1	no
Irritation of veins after i.v. application	0	1	0	1	1	1	4	no