DRUG CARDS DAILY

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NAME(S): Generic: digoxin (di JOKS in) | Brand: Digox, Digitalis, Lanoxicaps

PHARMACOLOGIC & THERAPEUTIC CLASS: Antiarrhythmic Agent | Cardiac Glycoside | Ionotrope

DOSAGE FORM & STRENGTH: Tabs: 62.5 mcg, 125 mcg, 187.5 mcg, 250 mcg | Sol: 0.05 mg/mL | Inj: 0.25 mg/mL

INDICATION(S) & DOSING(S): ADULTS (NOTE: Use lean body weight (LBW) for dose calculations. Doses are adjusted based on levels. The IV formulation is preferred to IM.)

1. Heart Failure with reduced Ejection Fraction (HFrEF):

- 0 Tx range is b/t 3.4-5.1 mcg/kg/dose PO qd. IM/IV dosing is between 2.4-3.6 mcg/kg/dose.
- O A loading dose of 10-15 mcg/kg PO or 8-12 mcg/kg IM/IV divided into 3 doses. 50% is given first, then 25% given in 6-8 hours and the last 25% in 6-8 hours.
- 2. Atrial Fibrillation:
 - O Tx range is b/t 3.4-5.1 mcg/kg/dose PO qd. IM/IV dosing is b/t 2.4-3.6 mcg/kg/dose for rate control.
 - O Loading dose is b/t 10-15 mcg/kg PO OR 8-12 mcg/kg IM/IV divided into three doses. 50% given initially. 25% after 6-8 hours. Then the remaining 25% after 6-8 hours.
- OFF LABEL | Paroxysmal Supraventricular Tachycardia (PSVT) conversion: Refer to HfrEF dosing.

INDICATION(S) & DOSING(S): PEDIATRICS (Weight based dosing based on LBW)

- 1. HFrEF:
 - O Premature neonates: Tx range is <u>4.7-8 mcg/kg/day PO</u> divided q12h OR b/t <u>3.8-6.2 mcg/kg/day IM/IV</u> divided q12h. Loading dose is b/t <u>20-30 mcg/kg PO</u> OR b/t <u>15-25 mcg/kg IM/IV</u> divided into 3 doses with 50% initially, 25% 4-8 hours later, and the final 25% given 4-8 hours later. Dosing based on LBW.
 - O Full-term neonates: Tx range is 7.5-11.3 mcg/kg/day PO divided q12h OR b/t 6-9 mcg/kg/day IM/IV divided q12h. Loading dose is b/t 25-35 mcg/kg PO OR b/t 20-30 mcg/kg IM/IV divided into 3 doses with 50% initially, 25% 4-8 hours later, and the final 25% given 4-8 hours later. Dosing based on LBW.
 - 0 1 month to 2 years: Tx range is <u>11.3-18.8 mcg/kg/day PO</u> divided q12h OR b/t <u>9-13 mcg/kg/day IM/IV</u> divided q12h. Loading dose is b/t <u>35-60 mcg/kg PO</u> OR b/t <u>30-50 mcg/kg IM/IV</u> divided into 3 doses with 50% initially, 25% <u>4-8 hours later</u>, and the final 25% given 4-8 hours later. Dosing based on LBW.
 - O 2 years to 5 years: (Note the dec from prev age range) Tx range is 9.4-13.1 mcg/kg/day PO divided q12h OR b/t 7.6-10.6 mcg/kg/day IM/IV divided q12h. Loading dose is b/t 30-45 mcg/kg PO OR b/t 25-35 mcg/kg IM/IV divided into 3 doses with 50% initially, 25% 4-8 hours later, and the final 25% given 4-8 hours later. Dosing based on LBW.
 - 5 years to 10 years: (Note the dec from prev age range) Tx range is <u>5.6-11.3 mcg/kg/day PO</u> divided q12h OR b/t <u>4.6-9 mcg/kg/day IM/IV</u> divided q12h. Loading dose is b/t <u>20-35 mcg/kg PO</u> OR b/t <u>15-30</u> mcg/kg IM/IV divided into 3 doses with 50% initially, 25% 4-8 hours later, and the final 25% given 4-8 hours later. Dosing based on LBW.
 - >10 years: (Note 1. The dec from prev age range; 2. Freq is qd instead of divided q12 dosing; 3. Load dose freq is now 6-8h instead of 4-8h) Tx range is <u>3.4-5.1 mcg/kg/day PO</u> qd OR b/t <u>2.4-3.6 mcg/kg/day</u>
 <u>IM/IV</u> qd. Loading dose is b/t <u>10-15 mcg/kg PO</u> OR b/t <u>8-12 mcg/kg IM/IV</u> divided into 3 doses with 50% initially, 25% 6-8 hours later, and the final 25% given 6-8 hours later. Dosing based on LBW.



• OFF LABEL | Atrial Fibrilation and PSVT conversion: Similar to HfrEF w/ changes in dosing ranges. Please refer to most current literature in case there were range changes. Same are ranges as follows: Premature neonates, Full-term neonates, 1 month to 2 years, 2 years to 5 years, 5 years to 10 years, & >10 years.

MECHANISM OF ACTION & PHARMACOLOGY

• MOA: Na/K ATPase pump is inhibited → Increase of sodium intracellularly → Increased influx of calcium → increased contractility and improved baroreflex sensitivity. Specific to HF there is an increase in contractility and possible improvement in baroreflex sensitivity. Regarding Supraventricular arrhythmias there is an increased refractory period, a decrease of conduction velocity, some positive inotropic effects, and a decrease ventricular rate. | Absorbed in the upper small intestine. It is passive and non-saturable. | Metabolized in stomach via hydrolysis. Only ~16% through the liver. | 50-70% excreted in the urine. | The onset of action is b/t 1-2 hours when taken orally and 5-60 minutes if taken intravenously. | Time to peak is 1-3 hours. | Duration of action is b/t 3-4 days. | Half-life elimination in neonates is between 61-170 hours. Infants 18-25 hours. Children 18-36 hours. Adults 36-48 hours. | ~25% protein bound.

SPECIAL POPULATIONS & CONSIDERATIONS

 Diet: Make sure to maintain a healthy amount of dietary potassium in order to decrease the risk of hypokalemia. Low potassium increases risk of toxicity from digoxin | Renally Impaired: If eGFR is b/t 10-50 give b/t 25-75% of usual dose q24-36h. If eGFR is <10 give b/t 10-25% usual dose q48h. | Some caution in pts to be aware of are pts w/ ventricular fibrillation, mycarditis, acute MI, hypokalemia, hypocalcemia, renal impairment, AV block, bradycardia, and thyroid disease. | Digoxin Toxicity: S/sx are anorexia, n/v, visual changes, and usually associated w/ levels >2 ng/mL. Risk factors are pts w/ low body weight, advanced age, renal impairment, and low K+/Ca++/Mg++ | Pregnancy/Lactation: May use. Harm not expected. No known risk. Limited data.

SIDE EFFECTS

- **Common**: Dizziness, headache, n/v/d, bradycardia, weakness, confusion, palpitations, depression, & more.
- **Serious**: Intenstinal ischemia, delirium, thrombocytopenia, severe bradycardia, and AV block.

DRUG INTERACTIONS

- Some considerations made when accessing possible DI: P-gp substrate, meds that delay gastric emptying, binding to anion/cation exchange resin/polymer, bradycardia, & meds that shorten QT interval.
- Some drug(s) of note to be avoided: clonidine, neomycin, succinylcholine, and fingolimod.
- Some drug(s) of note that warrant caution/adjustments: acarbose, aloe, amiodarone, azithromycin, atropine, colestipol, carvedilol, esomeprazole, doxycycline, indomethacin, ibuprofen, warfarin, and many others.

MONITORING PARAMETERS

- Drug specifics: In HF the tx drug levels are b/t 0.5-0.9 ng/mL. In Afib b/t 0.8-2 ng/mL. Digoxin serum concentrations should be drawn approximately 6-8h after last dose. Anything >2 ng/mL is considered a toxic level of the medication.
- General: Serum drug levels, HR, electrolytes, & Cr.

PATIENT COUNSELING INFORMATION

- Digoxin is commonly used in the treatment of **Heart Failure** and **Chronic Atrial Fibrillation** by reducing strain on the heart and helping to maintain a strong/steady/normal heartbeat.
- Taken w/ or w/o food with maintenance dosing typically being once daily unless otherwise specified by Pbr.



- Foods high in fiber and medications such as cholestyramine, cholestipol, or psyllium will decrease drug absorption so digoxin should be taken at least 2 hours before or 2 hours after.
- Medication should be taken regularly at the same time each day.

REFERENCE(S) & RESOURCE(S)

- 1. <u>https://online.epocrates.com/drugs/19710/digoxin/Monograph</u>
- 2. https://www.drugs.com/ppa/digoxin.html
- 3. https://www.webmd.com/drugs/2/drug-4358/digoxin-oral/details



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DRUG CARDS D A I L Y

Monday at 7 am EST (6 am CST, 4 am PST)

HEY NEW GRAD!

So you landed that perfect job offer or got the perfect match for your PGY1 and now the <u>ONLY</u> thing standing in your way is passing the NAPLEX and MPJE.

Here are some NAPLEX & MPJE prep recommendations to help you do everything you can to **pass the first time!**

HEY STUDENT!

When I was P1 one of the best pieces of advice I got from those before me was to use a NAPLEX Prep book while learning each topic.

This helps focus your learning and the repetition helps to retain info and indirectly prepare you for the NAPLEX

