Welcome to

Veterinary Pharmacology, Toxicology and Therapeutics

227.305

Pharmacology - Why bother?

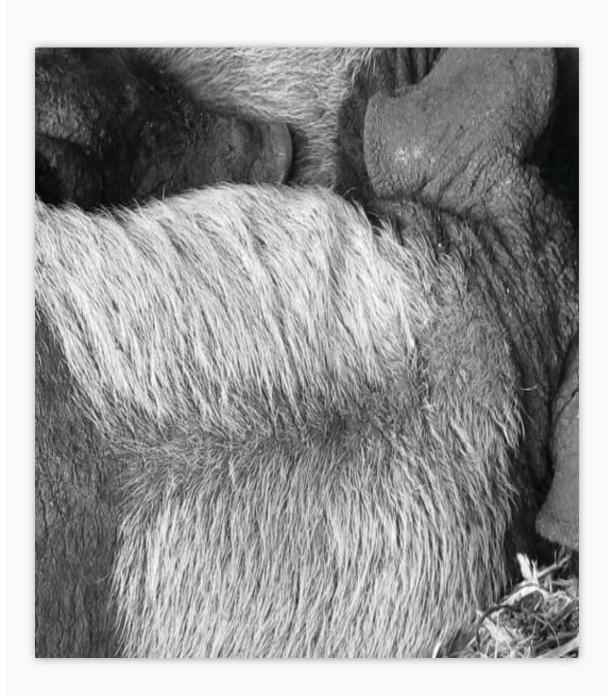
Treatment options

There are usually several things you can do, like:

- do nothing
- give drugs
- surgery
- change diet
- euthanasia
- all but the first involve drugs!

What do you need to know?







What do you need to know?

- history
- clinical exam findings
- differential list
- lab tests?
- diagnosis

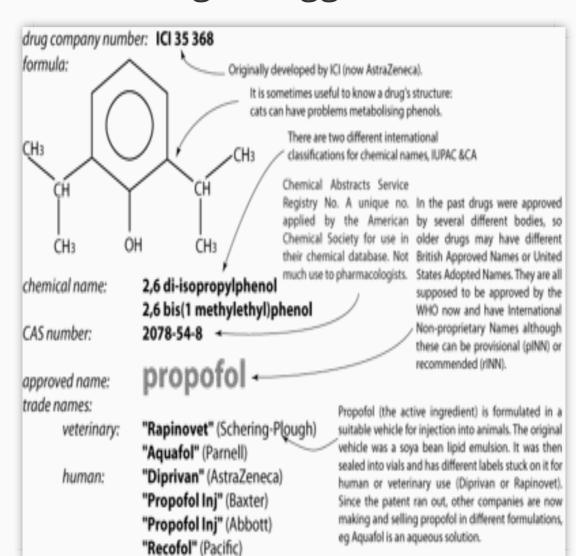
What do you need to know?

to treat the piglets?

- treatment objectives?
- drugs likely to be active?
- side effects & interactions?
- monitoring required?
- pharmacokinetics?
- dose?
- cost?
- do the benefits outweigh the risks?

Drug names

Don't get bogged down!



Active drugs?

- antiseptics
 - chlorhexidine
 - iodine

Active drugs?

- antibiotics
 - penicillins
 - narrow spectrum
 - broad spectrum
 - cephalosporins
 - tetracyclines
 - etc, etc

info you need to know

- antibiotics
 - penicillins
 - narrow spectrum
 - benzylpenicillin
 - Na benzylpenicillin
 - K benzylpenicillin
 - o procaine penicillin
 - benzathine penicillin
 - phenoxymethylpenicillin
 - broad spectrum
 - etc

info you need to know

pharmacokinetics

- penicillins
 - narrow spectrum
 - benzylpenicillin
 - Na benzylpenicillin
 - K benzylpenicillin
 - procaine penicillin
 - benzathine penicillin
 - phenoxymethylpenicillin
 - broad spectrum
- etc

Sources of info

- 1. scientific literature
- 2. textbooks
- 3. colleagues
- 4. www
- 5. drug companies plus this course!

resources Stream Calve Massey library



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Antimicrobial Susceptibility of Staphylococcus hyicus Isolated from Exudative Epidermitis in Pigs

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Exudative epidermitis or greasy pig syndrome is caused by the congulase-variable staphylococcal species Staphylococcus hyicus. Treatment of this disease is problematic because of the limited number of antimicrobial agents available for this purpose. Thirteen antimicrobial agents were evaluated for their activities against 100 S. bricus strains isolated from pigs with exudative epidermitis. Novobiocin was the most active compound tested, with an MIC for 90% of the strains tested (MIC,) of ≤0.06 µg/ml. Enrofloxacin, ampicillin, and celtiofur were the next most active compounds, with MIC_{se}s of 0.25, 0.5, and 1.0 µg/ml, respectively. However, 41.4% of the 99 strains tested were positive for β-lactamase production. The MIC_{ses} of erythromycin, tetracycline, and streptonycin were >32.0 µg/ml. Initial testing with sulfadiazine-trimethoprim yielded un MIC_{ss} of >64.0 μg/ml, but subsequent testing with thyuidine phosphorylase-supplemented medium yielded an MIC_{to} of 0.06 μg/ml. Both lincomycin and spectinomycin were relatively inactive against the S. hyicur strains tested, with MIC of >64.0 and >128.0 µg/ml, respectively. However, the combination of the two compounds at ratios of 1:2 (linconycin to spectinomycin) and 1:8 were more active, with MIC_{ss}s of 16.0 and 4.0 µg/ml, respectively. These results indicate that novobiocin and sulfadiazine-trimethoprim were the most active compounds tested against the S. Ayicus strains isolated from pigs with exudative epidermitis. Furthermore, the combination of lincomycin and spectinomycin was more active than the individual compounds against the strains tested.

Reading scientific papers to extract clinically useful info

Types of papers

- veterinary clinical trials
- human clinical trials
- basic science papers

Drug development

- synthesise compounds
- screen for activity on target
- screen for activity on other targets
- test in animal models
- pharmacokinetics
- animal toxicity testing
- phase 1 trials (healthy target species)
- phase 2 trials (sick target species)
- phase 3 trials (large nos. sick target species)
- (phase 4 trials post marketing surveillance)

"It has been clearly shown that drugs acting at the ORL1 receptor are good analgesics in isolated spinal cord preparations."

Faber et al., 1996, Br J Pharmac, 119, 189 – 190

reading papers

- Are the results valid?
 - randomisation?
 - all animals accounted for?
 - controls? dose? numbers?
- What are the results?
 - size of effect?
 - precision? (stats)
- How do the results compare to other studies?

reading papers

- Relevance to practice in NZ?
- Will the results help me in caring for my patients?

Phar-macolo-gy is fun!