Extrapolation of Adult Efficacy Data to Pediatric Patients

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Disclosure(s)

- No conflicts to disclose
- Off label drug use in neonates, treated in NICUs, is the current standard and therefore will be presented

Historical Drug "Development" in Children

Colic. diarrhea.
cholera & teething
alcohol (8.5%)
morphine (1/8 grain)



"BABY SERIEND"

CONTAINS EIGHT AND ONL-BALF PER CENT, ALCOHOL-ONE-FIGHTE CEATS SEL-PRITE OF NOFPENE IN EACH FLITD OF NE. J M.S. J. A. NOPP, Sole Prog. C. ROBERT NOPP, Mig. Chemist YORK, P.A., U. S. A., KING OF BABY SOOTHERS SPLENDID FOR Wind Colle, Griging in the Bowels, Diarrhoga, Cholera Infantum and Teething Troubles. Trial Size, 10c, Large, 25c.

: Trade Mark Registered

Teething
Deodorized
tincture of
opium (1.5%)

Deodorized Tinct. Opium 1 1-5 Per Cent.

TOTT'S

TEETHING

CORDIAL

Satisfies the Baby, pleases the Mother, gives rest to both.

Medication Use in NICUs – Pediatrix, Inc. Data for 2007: 72,647 Patients - Rate/1000 Discharges

Drug	Rank	Use
Gentamicin	1	822
Ampicillin	2	726
Surfactants	3	234
Caffeine	4	224
Furosemide	5	199
Vancomycin	6	177
Metoclopramide	7	82
<u>Fentanyl</u>	8	95
Dopamine	9	89
Midazolam	10	80
<u>Morphine</u>	11	71
Ranitidine	12	70
Cefotaxime	13	62
Phenobarbital	14	59
Indomethacin	15	54

Medication Use in NICUs, 2014

Drug	Rank	
Ampicillin	1	
Gentamicin	2	
Caffeine	3	
Vancomycin	4	
Beractant	5	
Furosemide	6	
<u>Fentanyl</u>	7	
Dopamine	8	
Midazolam	9	
Calfactant	10	
Metoclopramide	11	
Ranitidine	12	
Poractant alpha	13	
<u>Morphine</u>	14	
Cefotaxime	15	



Neonatal Clinical Pharmacology



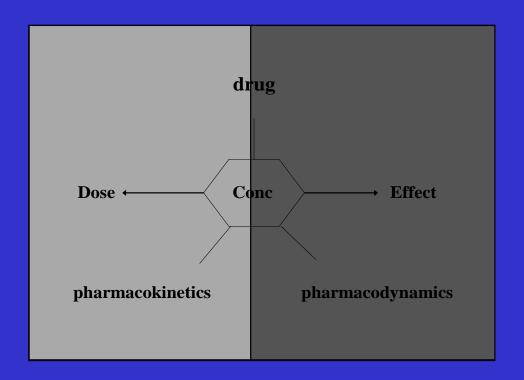








Role of Clinical Pharmacology



PK: what the body does to the drug: conc/time PD: what the drug does to the body: conc/effect

Extrapolation of Adult Efficacy Data to Pediatric Patients

Given the available data, is there a biological reason to believe that drugs with a well-established MOA (opioids, NSAIDs, acetaminophen, and local anesthetics) would be less effective (at similar concentrations) in pediatric patients less than 2 years of age compared to older children? If so, in which age group and what are the uncertainties? Assessment of PK and safety would be required in all age groups.

When Compared to Adults, Is Disease Progression and Response to Intervention Similar in Pediatrics?

Answer: No

NO EXTRAPOLATION

CONDUCT

- Adequate dose-ranging studies in children to establish dosing
- 2) Safety and efficacy trials at identified dose(s)

Similar Exposure-Response in Pediatrics and Adults?

Answer: Yes

Is the Drug (or active metabolite)
Concentration Measurable and Predictive
of Clinical Response?

Answer: Yes

FULL EXTRAPOLATION

CONDUCT

- 1) Adequate PK study to select dose(s) to achieve similar exposure in adults
- 2) Safety trials at identified dose(s)

Similar Exposure-Response in Pediatrics and Adults?

Is the Drug (or active metabolite)
Concentration Measurable and Predictive of Clinical Response?

Answers: No

Is There a PD Measurement That Can be Used to Predict Efficacy in Children?

Answer: Yes or No.

PARTIAL EXTRAPOLATION

CONDUCT

- 1) Adequate dose-ranging studies in children to establish dosing
- Safety and efficacy/PD trials at identified dose(s)

Extrapolation of Adult Efficacy Data to Pediatric Patients

Given the available data, is there a biological reason to believe that drugs with a well-established MOA (opioids, NSAIDs, acetaminophen, and local anesthetics) would be less effective (at similar concentrations) in pediatric patients less than 2 years of age compared to older children? If so, in which age group and what are the uncertainties? Assessment of PK and safety would be required in all age groups.

