Infant Feeding –
allergies, intolerances,
& the evidence;
hardly ever black & white

DAA conference May 2011

030511

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Aim of this workshop

 To give you a treatment pathway that enables you to practice with confidence.

 How do we do this with current evidence available?

- Research Evidence + Clinical Practice = Better Practice
- Separating the black & white from the grey



Most diet therapy is based on:

- Known physiology of the condition
 - as that knowledge increases diet therapy benefits
 e.g diabetes, inborn errors of metabolism
- Known content of foods
 - related to condition e.g. various nutrients

... We know the physiology of some symptoms eg reflux, but *that physiology* does not help us decide which foods to change

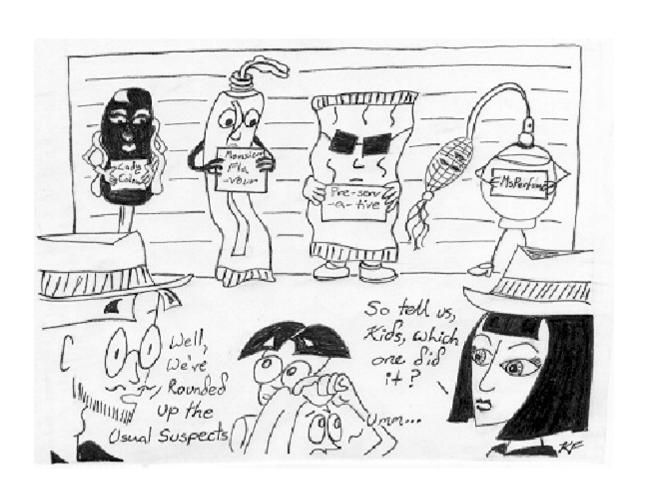
We need the physiology of the mechanism whereby those symptoms arise - unknown

- In allergy we have the mechanism & allergy tests do inform practice with clinical trials
- In FODMAPS we have a mechanism & a hypothesis - re what foods affect gut function
- In food intolerance we do not have the physiology of the mechanism - it is known to be pharmacological & to be complex

FI is 'adverse reaction to foods' i.e. toxicology Q: What foods are poisonous?

- What foods are 'poisonous' is not clear check in nutrition texts
- Remember: "There are no poisonous substances, only poisonous doses" &
 "One man's meat is another man's poison" e.g favism broad beans a genetic group
- Q: content of suspect substances in foods

The usual suspects



What are the suspect substances?

- Present Ho salicylates, amines, glutamates and many additives - have DBPCT
 - but analysis of salicylate did not correspond to patient tolerance e g apples, onions
 - but many families have aspirin sensitivity
 - complicated as amount varies with variety, amount of ripening & season
 - Q: natural benzoates, other compounds
 - Q: natural flavours my Ho

Levels of 'werry bad'!



More grey areas

- All is <u>not known</u> re *suspect substances*
- Q: role of whole foods eg wheat & milk
- UK Ho 'few foods diet' so takes longer
- Q: not "Is it allowed?" > "Is it tolerated?"
- Physiology in the individual is not known
- My Ho sulphotransferases varies over life, in family members, & with Total Body Load
- Q: Are body smells intermediary metabolites?

Adverse reactions > individual diet

 "The aim of the diet therapy is to find the diet therapy" for each individual

- Evidence has a history thinking over 35 yrs
- Evidence has a present current Ho each patient is a new case study
- Evidence has a **future** genetics, immune sys interacting with pharmacology
- Next 35 yrs enjoy new info for me!!

Considered hypothesis, test, review,

then

```
next hypothesis

test

review

then

next hypothesis

test

review

then

next hypothesis

test

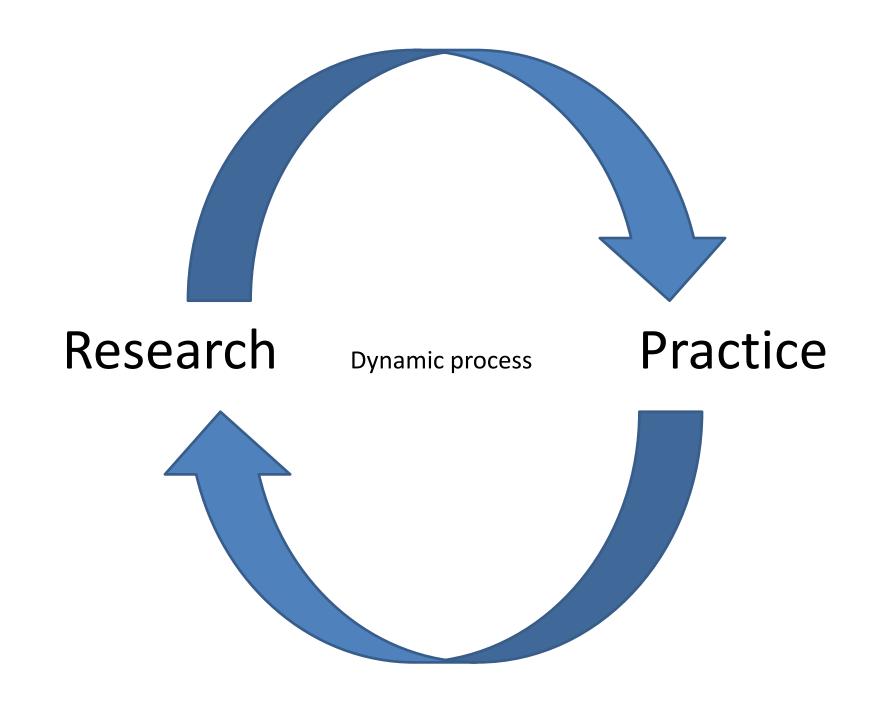
review

then

next hypothesis

test
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review



Practice informs evidence

- Tolerance adds to salicylate data. See history
- Chems "aggravate the underlying condition"
- Smells matter, & Flavour in food
- Family Sensitivity History gives data
- Total Body Load relevant environment
- Individual differences eg amine tolerance
- Info from different ages and symptoms
 All inform practice and research

The total body load



Practice > description of the group

- In practice diet practice is based on the understanding of the physiology and the food content eg FODMAPS
- Or on understanding of eg gut flora eg use of antibiotics and pro and prebiotics
- Q: Who do you choose for which diet treatment?
- In Food Sensitive can use knowledge of sample

Food intolerant	FODMAPS	Probiotics
Foods usually causing reactions	Different foods usually causing reactions	Effect of usual food not relevant
Multi system symptoms	Variety of gut symptoms	Those reported to benefit
Family Sensitivity History & aspirin		? Genetic diffs
Smell sensitive pain sensitive		
Supersensitive to sensory input		
Supertasters Cravings w'thdrawl		
Individua diffs		

The family sensitivity history



Treatment options for FI and Reflux

- From evidence:-
- Reflux thickening of feeds
- Allergy allergy tests basis of practice
- Adverse reactions exclude suspect foods
- Now DBPCTs exclude suspect chemicals
- New buzz word > 'personalised medicine' >>>
 "personalised diet therapy'
- Diet is for patient <u>not</u> for particular symptoms

In animal care

- With the best care
- With a homogenous sample genetically

2% do not thrive

 How does this apply to the genetic variety in human babies?

To Treatment pathway

- Each baby is a product of its environment internal, metabolic and external
- Each baby has its own variety of symptoms physical, sleep, mood, activity etc
- Each baby has its own variety of food intolerances
- Each will add evidence to add to your evidence based practice.

Diet investigation resources

- My Self Help book: Are You Food Sensitive? available from <u>www.FoodIntolerancePro.com</u>
- Dietitians Manual in professional pack
- Fussy Babies > Fussy Baby
- Dietitians Association of Australia DINER
- RPAH handbook <u>www.allergy.net.au</u>
- Dietitians at a hospital near you, or listed in the phone book in private practice.