Nurses and Families a Partnership in Research







- O Nurses and families
- Resources and active recruitment
- O Bedside nurses involvement in research

What is Informed Consent?

CompetentSufficient InformationVoluntary



Cultural Considerations

• What does the family need

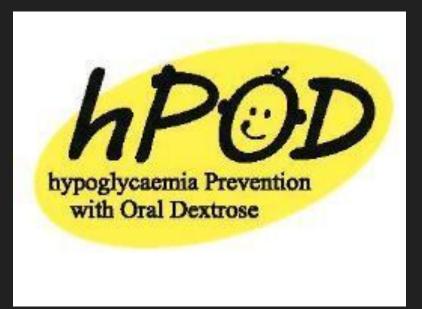
Clinical Trials Pamphlet

• Background

- O Aim
- What participation involves
- O Eligibility
- O Significance

Active Recruitment

hPOD-Hypoglycaemia Prevention with Oral Dextrose



To determine if oral 40% dextrose gel shortly after birth will prevent low blood sugars in babies at risk of hypoglycaemia, reducing admission to a NICU/SCBU

hPOD-Eligibility

- $O \ge 35$ weeks
- O Large for gestational age
- Small for gestational age
- O Infants of diabetic mothers
- Mother intending to breast feed
- No indication for admission to SCBU/NICU

hPOD-Method

Gel dosage 0.5mls/kg
Study gel administered at 1 hour of age
Gel administered via buccal mucosa
Standard blood sugar monitoring

hPOD- Recruitment

O Diabetic Clinic

- O Antenatal Ward
- Womens Assessment Clinic/ Delivery Suite

ProVIDE- Protein in IVN: Impact on Devlopment

The ProVEDe Study

Protein IVN: Impact on Development

To establish if giving extra protein in the first week of life, improves growth and brain development in extremely low birth weight babies

ProVIDE-Eligibility

- **O** <1000g
- O UAC
- Admitted to a NICU <24 hours
- No abnormalities; affecting inborn error of metabolism
- Singleton or twin pregnancy

ProVIDE-Method

- Randomised < 24 hours
- O 5 days
- Blood testing on day 1 and day 5
- O Intermittent growth measurements
- Follow up at 2 years

ProVIDE- Recruitment

Antenatal Ward
NICU post delivery
Waiver of consent for 24 hours

GLOW-Glucose in Well Babies



To describe the normal blood sugar concentrations in healthy term babies, during the first five postnatal days.

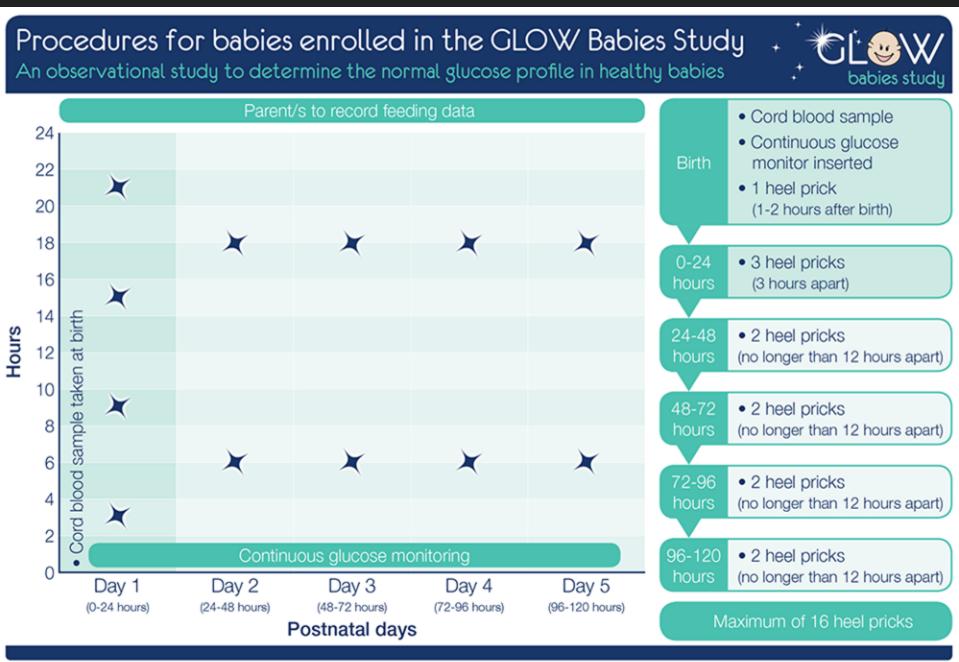


Inclusion criteria

- Babies Singleton
 - 37 to 42 completed weeks' gestation
 - Appropriate weight for gestation age (birthweight \geq 10th centile and \leq 90th centile)
 - Living within the greater Hamilton area
- Mothers Body Mass Index > 18.5 or ≤ 30 kg/m² before 20 weeks

Exclusion criteria

- **Babies** Apgar score < 7 at five minutes of age
 - Skin conditions preventing the attachment of the continuous glucose monitor
 - Unwell for any reason
- Mothers
 Diabetes, drug dependency, or using medications that may affect the baby's blood glucose concentration e.g. steroids



TM11/15











Meet the + CLOW team







Ohttps://www.facebook.com/GlowBabiesStudy

GLOW Facebook Page

"...We really enjoyed the visits and the chats, it became part of our normal routine and we will miss them. To not only be part of a study to help others but to receive the best care possible from birth was a luxury and provides us with reassurance just from the visits that we had a healthy baby.

The team gave us all the time and information we needed and the whole experience was relaxing, such lovely people..."

"....I didn't mind taking part in the study at all, it was super easy and I slept through nearly every heal prick..."

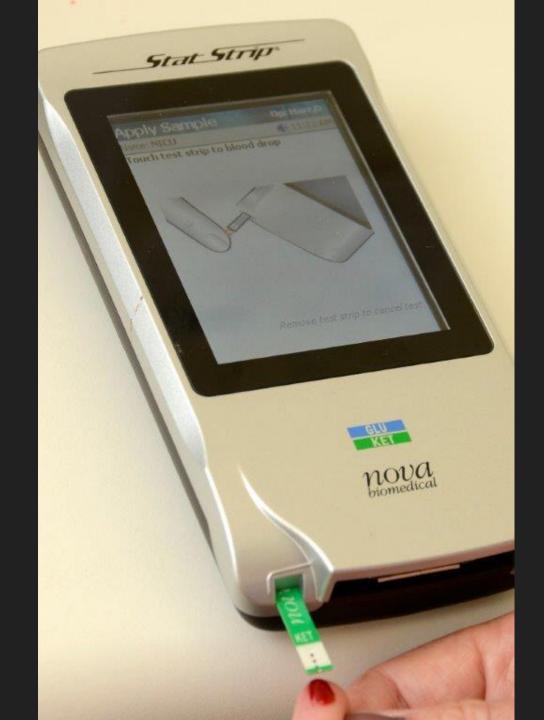
ROCKET-Reliability of Point of Care Ketone Trial



To determine the accuracy of the Point of Care Testing analyser to obtain results for ketones

ROCKET-Eligibility





ROCKET-Bedside Nurse

- O Discussion with family
- O Consent
- O Blood testing
- Case report form
- Publish and present results

Nurses and Research

Discuss studies
Consents
Data Collection
Support

What can you take back to your unit?

- Research is not scary
- Engage nurses in research
- Empower nurses in the process



What informed consent involves
Clinical trials pamphlet
Review current studies
Nurses role

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Questions?