ISRCTN52062666 https://doi.org/10.1186/ISRCTN52062666

Maternal Acupuncture and Neonatal Abstinence Syndrome

Submission date	Recruitment status	[ ] Prospectively registered
09/09/2005	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
20/01/2006	Completed	[X] Results
Last Edited	**Condition category**	[ ] Individual participant data
24/07/2014	Mental and Behavioural Disorders	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Patricia Janssen

Contact details
University of British Columbia (UBC) Dept of Health Care and Epidemiology
5804 Fairvew Ave
Vancouver
Canada
V6T-1Z3
+1 604 875 2424, local 5414
pjanssen@interchange.ubc.ca

Additional identifiers

Protocol serial number
CAM 05-325

Study information

Scientific Title

Study objectives

Infants whose mothers have been randomized to receive standard care (methadone maintenance) in combination with daily acupuncture treatments versus standard care alone will require fewer days of treatment with morphine for neonatal abstinence syndrome.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Primary study design
Interventional

Study design
Randomised controlled trial

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Drug addiction

Interventions
Acupuncture using National Acupuncture Detoxification Association (NADA) protocol + standard care versus standard care alone.

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
Days of neonatal morphine treatment

Key secondary outcome(s)
1. Gestational age at birth
2. Rates of intrauterine growth restriction
3. Scores on the Neonatal Abstinence Syndrome Scale
4. Rates of admission to a level II or level III nursery and length of stay
5. Days to regain birthweight
6. Rates of apprehension by the Ministry of Children and Families prior to discharge from hospital

Completion date
01/08/2007

Eligibility

Key inclusion criteria

All women admitted to the Chemical Dependency Unit at BC Womens Hospital will be offered participation.

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
Female

Key exclusion criteria
Women who can neither read nor write English.

Date of first enrolment
01/08/2005

Date of final enrolment
01/08/2007

Locations

Countries of recruitment
Canada

Study participating centre
University of British Columbia (UBC) Dept of Health Care and Epidemiology
Vancouver
Canada
V6T-1Z3

Sponsor information

Organisation
Toronto SickKids Foundation (Canada)

ROR
https://ror.org/04374qe70

Funder(s)

Funder type
Charity

Funder Name
Toronto Sickkids Foundation

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	23/12/2012		Yes	No