

Investigating the effects of repeated once-daily administration of the non-hormonal Neurokinin 1,3 receptor antagonist NT-814 on sex hormone levels in healthy women

Submission date 14/09/2020	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 18/09/2020	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 13/08/2021	Condition category Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Endometriosis is a condition where small pieces of the womb lining (the endometrium) are found outside the womb. Uterine fibroids are non-cancerous growths that develop in or around the womb. The ideal treatment for endometriosis and uterine fibroids would lower the female sex hormone estrogen, whilst avoiding the menopausal flushing and bone loss associated with current treatments. Substance P and neurokinin B bind to the neurokinin (NK) receptors 1 and 3, respectively, to control reproductive hormone levels. NT-814 is a new, non-hormonal treatment which blocks the NK 1,3 receptors. However, the effects of NT-814 on reproductive hormone levels in healthy women is currently unknown. Current treatments for hormone-driven disorders, such as endometriosis and uterine fibroids, are limited by side effects and effectiveness. The aim of this study is to assess the effectiveness and safety of NT-814 in healthy women.

Who can participate?

Healthy women aged 18-45 with regular menstrual cycles, not taking any medications or hormonal contraception

What does the study involve?

Healthy women attend for two consecutive menstrual cycles. No treatment is given in Cycle 1. During Cycle 2, women are randomly allocated to receive placebo (dummy drug) or NT-814 40 mg, 80 mg or 120 mg for up to 21 days. In each cycle, blood samples are taken on days 3/4, 9/10, 15/16 and 21/22 to measure blood reproductive hormone levels.

What are the possible benefits and risks of participating?

Regular monitoring (using blood samples, clinical observations and ECG) are used throughout to maintain participant safety.

Where is the study run?

Quotient Sciences (USA)

When is the study starting and how long is it expected to run?
February 2019 to September 2019

Who is funding the study?
KaNDy Therapeutics (USA)

Who is the main contact?
Dr Steve Pawsey
steve.pawsey@nerretherapeutics.com

Contact information

Type(s)

Public

Contact name

Dr Steve Pawsey

Contact details

KaNDy Therapeutics
Stevenage Biosciences Catalyst
Stevenage
United Kingdom
SG1 2FX
+44 (0)1438 906960
steve.pawsey@nerretherapeutics.com

Type(s)

Scientific

Contact name

Dr Steve Pawsey

Contact details

KaNDy Therapeutics
Stevenage Biosciences Catalyst
Stevenage
United Kingdom
SG1 2FX
+44 (0)1438 906960
steve.pawsey@nerretherapeutics.com

Additional identifiers

EudraCT/CTIS number

Nil known

IRAS number

ClinicalTrials.gov number

Nil known

Secondary identifying numbers

814-1-05

Study information

Scientific Title

A randomized, single-blind, placebo-controlled study of the effects of repeat doses of NT-814 on oestradiol and other sex hormone concentrations in healthy pre-menopausal female volunteers

Study objectives

NT-814 is a dual NK1,3 receptor antagonist and therefore has the theoretical potential to reduce GnRH pulsatility by blocking the endogenous effects of NKB and substance P on the reproductive axis. This would lower LH levels and subsequently estradiol concentrations in healthy pre-menopausal women.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 22/05/2019, Institutional Review Board Advarra (6940 Columbia Gateway Drive, Suite 110, Columbia, MD 21046, USA; +1 (0)410 884 2900; adviser@advarra.com), ref: Pro00034218

Study design

Single-centre phase 1 randomized single-blind placebo-controlled study

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Reproductive hormone levels in women with hormone-driven disorders

Interventions

This is a Phase 1, randomized, single-blind, placebo-controlled study designed to determine the effects of NT-814 (40, 80, 120, 160 mg once daily) on GnRH pathway hormones in healthy female subjects.

Thirty-two healthy women attend for two consecutive menstrual cycles. In each cycle, blood samples are taken on days 3/4, 9/10, 15/16 and 21/22 to measure serum reproductive hormone levels, and plasma NT-814 levels (Cycle 2 only). No treatment is given in Cycle 1 (baseline). During Cycle 2, participants are randomized by block randomization to receive placebo or NT-814 40 mg, 80 mg or 120 mg (n=8 per group) for up to 21 days.

Intervention Type

Drug

Phase

Phase I

Drug/device/biological/vaccine name(s)

NT-814: a novel, non-hormonal NK1,3 antagonist

Primary outcome measure

GnRH pathway hormones measured using serum LH, FSH, oestradiol and progesterone concentrations on Days 3/4, 9/10, 15/16 and 21/22 in both Cycle 1 (baseline) and Cycle 2 (treatment)

Secondary outcome measures

1. The safety of NT 814 measured using clinical laboratory assessments (haematology and biochemistry), vital signs (BP and HR), and recording of treatment-emergent adverse events on Days 3/4, 9/10, 15/16 and 21/22 in both Cycle 1 (baseline) and Cycle 2 (treatment)
2. The PK-PD relationship of NT-814 measured using NT-814 plasma concentration levels during Cycle 2 Days 3/4, 9/10, 15/16 and 21/22
3. Menstrual cycle length measured using the difference in cycle length (in days) between Cycle 1 and Cycle 2

Overall study start date

01/02/2019

Completion date

30/09/2019

Eligibility

Key inclusion criteria

1. Healthy, female, aged 18 to 45 years inclusive (age at time of informed consent)
2. Have regular (approximately) monthly menstrual periods
3. Be able and willing to understand and comply with the requirements of the study and give written informed consent
4. Have a body mass index (BMI) in the range 18.0 to 32.0 kg/m² and body weight not less than 40.0 kg
5. Be judged to be in good health, based on the results of medical history, physical examination, vital signs, 12-lead ECG, and clinical laboratory findings
6. Not be pregnant and not lactating, with no intention of pregnancy during study treatment
7. Agree to use two acceptable methods of birth control, one of which a barrier method with spermicide for the duration of participation in the study and 30 days after the last dose of study medication

Participant type(s)

Healthy volunteer

Age group

Adult

Lower age limit

18 Years

Sex

Female

Target number of participants

32 women

Total final enrolment

33

Key exclusion criteria

1. Been previously enrolled in this study or any other study with NT-814
2. Have clinically significant findings on physical examination at screening
3. Have any relevant medical history, in particular: liver or renal insufficiency, cholecystectomy, significant cardiac, vascular, pulmonary, gastrointestinal, endocrine, neurologic, haematologic, rheumatologic, psychiatric, or metabolic disturbances, any inflammatory illness or any other condition that the Investigator considers should exclude the subject
4. Undergone bilateral oophorectomy and/or hysterectomy
5. Have positive serology for any of hepatitis B, hepatitis C, or human immunodeficiency virus (HIV)
6. Have any serum biochemistry and full blood count outside the normal reference ranges and considered by the Investigator to be of clinical significance, assessed at screening and on Cycle 1 Day 21/22. Haemoglobin must not be less than 11.0 g/dL at either screening or Day 21/22
7. Have Stage 2 or higher hypertension (supine/semi-recumbent systolic blood pressure [SBP] >160 mmHg; diastolic blood pressure [DBP] >100 mmHg or Stage 1 hypertension (supine/semi-recumbent SBP 140-160 mmHg; DBP 90-100 mmHg associated with indication for treatment i.e. evidence of end-organ damage, diabetes or a 10 year cardiovascular risk, estimated using a standard calculator, e.g. <https://qrisk.org/three/>, greater than 20%. Measurements are based on the mean of duplicate values recorded at least 2 minutes apart and are assessed at screening, on Cycle 1 Day 21 or 22 and pre-dose on Cycle 2 Day 1 or 2
8. Have clinically relevant abnormal 12 lead ECG, including QTcF >450 msec, QRS interval >120 msec, PR interval >220 msec, assessed at screening, on Cycle 1 Day 21 or 22 and pre-dose on Cycle 2 Day 1 or 2; if any value are out of range, two repeated assessments are permitted at that time point and the value based on the mean of the triplicate measure
9. Have a history of any grade of drug or alcohol use disorder according to Diagnostic and Statistical Manual of Mental Disorders 5 criteria (or later edition if applicable) within 6 months before Screening or have a positive test result(s) for drugs of abuse (opiates including methadone, cocaine, amphetamines, methamphetamines, cannabinoids, barbiturates, and benzodiazepines) at screening, Cycle 1 Day 21 or 22, or Cycle 2 Day 1 or 2
10. Have a clinically significant acute illness within 7 days before first study drug administration
11. Been drinking, on average, more than 6 cups of coffee or other caffeinated beverages daily (where each cup of coffee or beverage contained approximately 120 mg caffeine)

12. Been smoking more than an average of 5 cigarettes (or equivalent) per day
13. Use any non-permitted prior prescription, over-the-counter or herbal medication
14. Have a history of clinically significant drug and/or food allergies, particularly known allergy to any of the excipients used in the study medications
15. Receive an investigational drug (including vaccines) or use an investigational medical device within 3 calendar months before the first dose of study medication or currently enrolled in an investigational study
16. Have major surgery within 2 calendar months before first dose of study medication, or surgery planned during the time the subject is expected to participate in the study
17. Donate blood or experienced acute loss of a significant amount of blood within 3 calendar months before first dose of study medication
18. Have psychological and/or emotional problems that would render the informed consent invalid or limited the ability of the subject to comply with the study requirements
19. Have any condition for which, in the opinion of the Investigator, participation would not have been in the best interest of the subject or that could prevent, limit, or confound the protocol-specified assessments

Date of first enrolment

06/06/2019

Date of final enrolment

16/07/2019

Locations

Countries of recruitment

United States of America

Study participating centre**Quotient Sciences**

3898 NW 7th St

Miami

United States of America

FL 33126

Sponsor information

Organisation

KaNDy Therapeutics

Sponsor details

Stevenage Bioscience Catalyst

Gunnels Wood Road

Stevenage

United Kingdom

SG1 2FX

+44 (0)1438 906 960
info@kandytherapeutics.com

Sponsor type
Industry

Funder(s)

Funder type
Industry

Funder Name
KaNDy Therapeutics

Results and Publications

Publication and dissemination plan
Planned publication in a high-impact peer-reviewed journal

Intention to publish date
01/12/2020

Individual participant data (IPD) sharing plan
Based on the sponsor's data transparency policy (<https://www.clinicalstudydatarequest.com/Study-Sponsors/Study-Sponsors-Bayer.aspx>), individual patient-level data will not be published for this type of study (single centre, small patient number) in order to avoid the risk for re-identification of participants.

IPD sharing plan summary
Not expected to be made available

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol file	version V1.0	10/05/2019	08/10/2020	No	No
Results article		13/07/2021	13/08/2021	Yes	No