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Early effect of agomelatine on general interest in outpatients suffering major depressive disorder

Submission date
24/03/2011

Recruitment status
No longer recruiting

Registration date
04/05/2011

Overall study status
Completed

Last Edited
18/04/2018

Condition category
Mental and Behavioural Disorders

[X] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

Plain English summary of protocol
Not provided at time of registration and not expected to be available in the future

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
CL3-20098-083

Study information

Scientific Title
Early effect of agomelatine on general interest in outpatients suffering major depressive disorder: a parallel group, randomised, double-blind, multicentre study

Study objectives
To assess the early effect of agomelatine on general interest in outpatients suffering from major depressive disorder

Please note tht as of 26/11/2012, the anticipated end date for this trial was updated from 30/09/2012 to 30/03/2013.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Ethics approval was obtained before recruitment of the first participants

Study design
Parallel group randomised double-blind multicentre study

Primary study design
Interventional

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Depression

Interventions
Therapeutic oral doses of agomelatine and therapeutic oral doses of selective serotonin reuptake inhibitors (SSRI) for 12 weeks, a randomised double-blind period

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Agomelatine

Primary outcome(s)
General interest score obtained from the Visual Analogue Scale (VAS) reflecting the item 13 of the QIDS-SR 16 scale

Key secondary outcome(s))
To provide additional efficacy, safety and tolerability data on agomelatine

Completion date
30/03/2013

Eligibility

Key inclusion criteria
1. Male or female outpatients
2. Aged between 18 and 65 years (inclusive) at the time of selection
3. Fulfilling Diagnostic and Statistical Manual of Mental Disorders 4th edition, Text Revision (DSM-IV-TR) criteria for current major depressive episode (MDE) < = 12months, of moderate to severe intensity
4. Major depressive episode diagnosis documented using the brief structured Mini-International Neuropsychiatric Interview (M.I.N.I.)
5. Quick Inventory of Depressive Symptomatology Self-Report (QIDS-SR16) item 13 (General interest) > = 2
6. Hamilton Rating Scale for Depression (HAM-D-17) total score > = 22
7. Clinical Global Impression - Severity (CGI-S) (Severity of illness) > = 4 (moderately to severely ill)
8. Requiring an antidepressant treatment

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Lower age limit
18 years

Sex
All

Key exclusion criteria
1. Patient fulfilling DSM-IV-TR criteria for a MDE of mild intensity or severe episode with psychotic features,catatonic features or with duration <4 weeks
2. All types of depression other than MDD, according to DSM-IV-TR criteria
3. Marked suicidal intent and/or known suicidal tendencies for the current episode defined as a score of 4 at the item 3 of the HAM-D-17 and/or in the investigator's opinion based on the patient's medical history, previoussuicide attempts, quality of social and familial support
4. Pregnancy, breastfeeding or possibility of becoming pregnant during the study and without an effective contraception
5. Hepatic impairment
6. Severe renal insufficiency

Date of first enrolment
23/05/2011

Date of final enrolment
30/03/2013

Locations

Countries of recruitment

Romania

Study participating centre
Spitalul Clinic de Neuropsihiatrie Craiova
Craiova
Romania
200317

Sponsor information

Organisation
Institut de Recherches Internationales Servier (France)

ROR
https://ror.org/034e7c066

Funder(s)

Funder type
Industry

Funder Name
Institut de Recherches Internationales Servier (France)

Results and Publications

Individual participant data (IPD) sharing plan
The datasets generated during and/or analysed during the current study will be available upon request from https://clinicaltrials.servier.com if a Marketing Authorisation has been granted after 1st January 2014.

IPD sharing plan summary
Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Basic results				No	No