

An open label phase I study in healthy subjects with blood group AB to investigate the safety, tolerability and efficacy of Uniplas™ LG

Submission date 22/12/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 10/01/2011	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 21/06/2011	Condition category Signs and Symptoms	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
UNI-111

Study information

Scientific Title

Study objectives

This is a trial in healthy subjects who have blood group AB to investigate the safety, tolerability and efficacy of Uniplas™ LG.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Local Ethics Committee (Ethikkommission der med.Uni.Wien und des Allg. Krankenhauses der Stadt Wien AKH) approved on the 12th November 2010 (ref: 779/2010)

Study design

Open-label non-randomised non-controlled phase I study

Primary study design

Interventional

Secondary study design

Non randomised controlled trial

Study setting(s)

Other

Study type(s)

Diagnostic

Participant information sheet

Not available in web format, please use the contact details below to request patient information material

Health condition(s) or problem(s) studied

Substitution of intentionally removed plasma

Interventions

Primary objective of this study is to investigate the safety and the tolerability of Uniplas™ LG, assessed by clinical and laboratory parameters with respect to subjects with blood group AB. IMP will be infused once and the subjects will be followed up until 3 months after administration of the IMP.

Intervention Type

Drug

Phase

Phase I

Drug/device/biological/vaccine name(s)

Uniplas™ LG

Primary outcome measure

Haemoglobin (Hb), measured at baseline, less than or equal to 30 minutes before and less than 5 minutes post plasmapheresis, 15 minutes and 2 hours post-transfusion, 24 hours and 7 days post-plasmapheresis and 3 months after administration of IMP.

Secondary outcome measures

1. Parameters of haemolysis: haptoglobin, free Hb, indirect bilirubin
2. Complement activation: CH50, C3c, C4
3. Circulating immune complexes (CIC): IgG, IgA, IgM
4. DAT (direct antiglobulin test)
5. Isoagglutinines (in case of a positive DAT)
6. Haematology: RBC count, WBC count, platelets, Hct, Hb
7. Standard safety lab (Clinical chemistry): sodium (Na⁺), potassium (K⁺), calcium (Ca²⁺), creatinine, ALAT, gamma-glutamyl transferase (gGT), total protein (TP)
8. Haemostatic Panel I: aPTT, PT, Fbg
9. Haemostatic Panel II: FII, FV, FVII, FVIII, FIX, FX, FXI, Protein C, Protein S, plasmin inhibitor
10. Urine analysis: WBC, nitrite, pH, protein, glucose, ketones, urobilinogen, bilirubin, blood/Hb
11. Changes in viral status over the study period: anti-HIV-1/2, HBsAg, anti-HBc, anti-HCV, anti-CMV, anti-HAV, anti-Parvovirus B19
12. Overall tolerability, AE monitoring, vital signs including body temperature

Measured at baseline, less than or equal to 30 minutes before and less than 5 minutes post plasmapheresis, 15 minutes and 2 hours post-transfusion, 24 hours and 7 days post-plasmapheresis and 3 months after administration of IMP.

Overall study start date

01/10/2010

Completion date

01/04/2011

Eligibility

Key inclusion criteria

1. Signed written informed consent
2. Subject must be capable to understand and comply with all relevant aspects of the study protocol
3. Blood group AB
4. Healthy male or female subjects greater than or equal to 18 years of age
5. Female subject must have a negative pregnancy test (human chorionic gonadotropin [HCG]-based assay)
6. Female subject must apply sufficient methods of contraception
7. Subject must have no clinically relevant abnormalities in medical history and general physical examination
8. A standard health insurance must be in place for the subject

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

4 to 6 (Study no longer recruiting: Last patient out on 28/03/2011)

Key exclusion criteria

1. Pregnancy or lactation
2. Subject got tattoos within the last 3 months
3. Subject was treated therapeutically with FFP, blood or plasma-derived products in the previous 6 months
4. Angiotensin converting enzyme (ACE)-inhibitors
5. Subject has a history of severe hypersensitivity to blood products or plasma protein
6. History of angioedema
7. History of coagulation disorder or bleeding disorder and any known abnormality affecting coagulation, fibrinolysis or platelet function
8. Any other clinically relevant history of disease
9. Subject has clinically significant abnormal laboratory values
10. Subject has IgA deficiency
11. Seropositivity for hepatitis B surface antigens (HBsAg), hepatitis C virus (HCV), human immunodeficiency virus (HIV-1/2) antibodies
12. Symptoms of a clinically relevant illness within 3 weeks before Visit 2
13. Subject has a history of or a suspected drug or alcohol abuse
14. Participation in another clinical study within the past 4 weeks

Date of first enrolment

01/10/2010

Date of final enrolment

01/04/2011

Locations**Countries of recruitment**

Austria

Study participating centre

Oberlaaerstrasse 235

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Sponsor information

Organisation

Octapharma AG (Switzerland)

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Sponsor type

Industry

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Funder(s)

Funder type

Industry

Funder Name

Octapharma AG (Switzerland)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

