Clinical Chemistry UCDLA_CBC_003

Purpose

Clinical chemistry determines biochemical parameters in plasma including enzymatic activity, specific substrates and electrolytes.

Ontological description: MP:0001545 – blood physiology abnormalities.

Experimental Design

• Minimum number of animals: 7M + 7F

• Age at test: Week 59

• Sex: We would expect the results of this test to show sexual dimorphism

Equipment

- 1. Clinical chemistry analyser
- 2. Vortex
- 3. Refrigerated centrifuge
- 4. Eppendorf tubes
- 5. Pipettes (200-1000 ul)

Procedure

Set up the clinical chemistry analyser and perform QC analyses of the control reagents in accordance with the equipment guidelines.

Sample collection and preparation:

- a. Collect the appropriate volume of blood required (160-200l of plasma), for the clinical chemistry analyser being used for assessment, preferably using a tube with an anticoagulant with the relevant blood collection procedure (see IMPC protocol Blood collection by retro-orbital puncture). Time of day for collection is in the morning, starting no earlier than 07:30.
- b. Keep whole blood samples on ice until centrifugation. Centrifuge for 10 minutes at 5000 x g in a refrigerated centrifuge set at 8°C. If plasma samples cannot be analysed immediately, keep them in the fridge until analysis.
- c. Analysis of samples is optimally done on the day of collection. When not possible the plasma samples can be stored at 2-8°C. If samples require storage for > 48 hours,

- freeze plasma at -20 °C in single aliquots. All samples are allowed to come to room temperature prior to analysis.
- d. Use plasma samples undiluted or diluted to a ratio of 1:2 with deionised water if the volume is insufficient.
- e. Plasma samples that were frozen or stored in the fridge should be vortexed briefly and centrifuged again at ~5000 x g for 2-3 minutes immediately prior to analysis. If necessary, remove fibrin clots using a wooden applicator.

Analysis:

Samples that produce results that lie outside the linear range for a specific assay have to be re-tested. In some cases it may be necessary to dilute samples with water to bring test results into range.

Notes

Blood collection for Clinical Chemistry and Hematology is usually performed as a non-fasting, terminal procedure but can be performed as a non-terminal procedure under certain circumstances. Mice from the terminal procedure may be used for subsequent gross pathology and other procedures included in terminal assessments. Whole blood (for Hematology) and plasma (for Clinical Chemistry) require different collection tubes so two independent samples are required from each mouse.

The information about the date of the experiment, that is the date when the measurement is performed, is an important parameter which is to be submitted in the Experiment xml file (dateOfExperiment="2013-02-28").

Dilution. Dilution of blood is highly discouraged, but is allowed when the total necessary amount is not obtained. If dilution is necessary then the assays should be done in one run.

Hemolysis. Two fields currently exist to capture metadata information about the hemolysis status in the clinical chemistry plasma samples. The first is the LIH Hemolysis severity score which can only be performed by clinics who run one of the Beckman Coulter AU-series of analysers. Such clinics are encouraged to capture and submit the hemolysis score of the LIH test in this field. Clinics who do not have an AU analyser are encouraged to use the second /alternative field which is simply titled Hemolysis. Simply enter "slight", "moderate", or "marked" based on whether the sample is visibly haemolysed or not. Provision of this information is not compulsory and it is suggested that any clinic completes at least one field or the other (not both).

Data QC

- 1. Plasma samples must be free of Fibrin clots in order to be analysed.
- 2. Badly haemolysed samples should be discarded.
- 3. Each morning, all parameters are tested with control sera (see ESLIM_015_001_Annex_3: Controls for biochemistry on AU400). Some parameters are tested with control serum level 1 (Beckman Coulter System Reagent, ODC0003)

- and control serum level 2 (Beckman Coulter System Reagent, ODC0004), which consists of lyophilised human plasma with a normal and a pathological concentration. Other parameters are tested with specific controls from other suppliers.
- 4. Controls are thawed and vortexed before utilisation and loaded according to the analyser's display. Control values must lie within the acceptable range indicated by the manufacturer, otherwise the specific tests must be recalibrated and specific measurements repeated. Controls can be stored in 200l aliquots of control sera frozen at -20°C for up to 1 month.

Metadata and examples

Metadata	Example
Equipment ID	ID of the machine used when more than 1 is used having same model and manufacturer. E. g. machine 1, machine 2, machine Minnie, machine Mickey Mouse, etc.
Equipment manufacturer	Manufacturer of the equipment. E.g. Olympus Diagnostics.
Equipment model	Model of the equipment. E.g. AU400
Blood collection tubes	The tubes used for blood collection. E.g. Sarstedt Li-Heparin gel tubes or Kabe Labortechnik Lithium heparin coated tubes.
Anaesthesia used for blood collection	The drug used for anaesthesia during blood collection. E. g. Isofluorane.
Method of blood collection	Concise description of the method used for blood collection. E.g. retro-orbital puncture.
Anticoagulant	Anticoagulant drug used for blood collection. E. g. Li-Heparin.
Samples kept on ice between collection and analysis	Yes/No.

Storage temperature from blood collection till measurement	E.g. 2°C
Sample status	Indicate if the sample were frozen (analysis on the same day of collection not possible) or fresh (analysis on the same day of collection). E.g Fresh/Frozen.
Plasma dilution	Dilution is highly discouraged but if necessary indicate here. E.g. "No dilution" or 1:2. Note that results submitted to DCC are assumed to be already corrected for any dilutions made.
ID of blood collection SOP	ID of the protocol followed for blood collection. Can be a center specific protocol. E.g. ESLIM_024_001.
Date and time of blood collection	Time of day for collection is in the morning, starting no earlier than 07:30. E.g. Year, month, day, time.
Date of measurement	The day of blood analysis. Year, month, day.
Hemolysis status	If no AU analyser score is provided, indicate here the gauged degree of hemolysis. E.g. slight/moderate/marked.
Blood collection experimenter ID	An ID of any format to be used coherently both inside the same procedure and for all procedures indicating the experimenter who collected the blood. E.g. Harw_001, or 1/2/3.
Blood analysis experimenter ID	An ID of any format to be used coherently both inside the same procedure and for all procedures indicating the experimenter who analyzed the blood. E.g. Harw_001, or 1/2/3.
Date equipment last calibrated	Most recent date in which the equipment (or any part of) used in the procedure was subject to a calibration event.
Date and time of sacrifice	The date and time when the mouse is sacrified.

Parameters and Metadata

ID of blood collection SOP UCDLA_CBC_045_001 | v1.1

procedureMetadata

Req. Analysis: false Req. Upload: true Is Annotated: false

Options: PHENO_CBC, ESLIM_024_001, sop.inv.019, RIKENMPP_004a_003, sop.inv.063,

Free fatty acids UCDLA_CBC_026_001 | v1.4

simpleParameter

Req. Analysis: false Req. Upload: false Is Annotated: true

Unit Measured: mmol/l

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Lactate dehydrogenase UCDLA_CBC_022_001 | v1.2

simpleParameter

Req. Analysis: false Req. Upload: false Is Annotated: true

Unit Measured: U/I

Cholesterol ratio UCDLA_CBC_058_001 | v1.0

simpleParameter

Req. Analysis: false Req. Upload: false Is Annotated: true Derivation: div('UCDLA_CBC_015_001', 'UCDLA_CBC_016_001') Glucose UCDLA CBC 018 001 | v1.5 simpleParameter Req. Analysis: false Req. Upload: true Is Annotated: true Unit Measured: mg/dl Sodium UCDLA_CBC_001_001 | v1.3 simpleParameter Req. Analysis: false Req. Upload: false Is Annotated: true Unit Measured: mmol/l

Total protein UCDLA_CBC_006_001 | v1.2

simpleParameter

Req. Analysis: false	Req. Upload: true	Is Annotated: true
Unit Measured: g/l		
HDL-cholesterol UC simpleParameter	DLA_CBC_016_001 v1.4	
Req. Analysis: false	Req. Upload: true	Is Annotated: true
Unit Measured: mg/dl		
Blood collection ex	(perimenter ID ucdl./	A CBC 049 001 Lv1 1
procedureMetadata		
Req. Analysis: false	Req. Upload: true	Is Annotated: false
Total cholesterol U	CDLA CBC 015 001 Lv1.4	
simpleParameter	<u> </u>	
Req. Analysis: false	Req. Upload: true	Is Annotated: true
Unit Measured: mg/dl		

Blood analysis experimenter ID UCDLA_CBC_051_001 | v1.0

procedureMetadata

Reg. Analysis: false Reg. Upload: true Is Annotated: false UIBC (unsaturated iron binding capacity) UCDLA_CBC_024_001 | v1.0 simpleParameter Req. Analysis: false Req. Upload: false Is Annotated: true Unit Measured: umol/l Date equipment last calibrated UCDLA_CBC_050_001 | v1.2 procedureMetadata Req. Analysis: false Req. Upload: false Is Annotated: false

Lipase UCDLA_CBC_021_001 | v1.1

simpleParameter

Req. Analysis: false Req. Upload: false Is Annotated: true

Unit Measured: U/I		
Sample status UCDL procedureMetadata	_A_CBC_043_001 v1.1	
Req. Analysis: false	Req. Upload: true	Is Annotated: false
Options: Fresh, Frozen, Fresh	h and frozen,	
Anticoagulant UCDL procedureMetadata	A_CBC_038_001 v1.1	
Req. Analysis: false	Req. Upload: true	Is Annotated: false
Options: No, Lithium Heparin	, Sodium Heparin, Heparine,	
Samples kept on ic A_CBC_042_001 v1.1 procedureMetadata	ce between collection	on and analysis ucdl
Req. Analysis: true	Req. Upload: true	Is Annotated: false
Options: Yes, No,		

Storage temperature from blood collection till measurement UCDLA_CBC_041_001 | v1.3

procedureMetadata

Req. Analysis: true Req. Upload: true **Is Annotated:** false Unit Measured: C Options: 2, 18-22, 4, -80, Fasting UCDLA_CBC_057_001 | v1.0 procedureMetadata Req. Analysis: true Req. Upload: true Is Annotated: false **Options:** Sixteen hours before bleeding, No, Four hours before bleeding, Equipment model UCDLA_CBC_035_001 | v1.0 procedureMetadata

Req. Analysis: true Req. Upload: true Is Annotated: false

Options: Integra 400 Plus, AU 480, 7020, JCA-BM6070, JCA-BM2250 (Advia 2400), AU 400, UniCel 600 Pro, AU 680, Hitachi 917, Cobas,

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Sample type UCDLA_CBC_056_001 | v1.0

procedureMetadata

Req. Analysis: true Req. Upload: true Is Annotated: false

Options: Plasma, Serum,

Chloride UCDLA_CBC_003_001 | v1.4

Req. Analysis: false Req. Upload: false Is Annotated: true

Unit Measured: mmol/l

simpleParameter

Date and time of blood collection UCDLA CBC 046 001 | v1.2

procedureMetadata

Req. Analysis: false Req. Upload: true Is Annotated: false

Alkaline phosphatase UCDLA_CBC_014_001 | v1.2

simpleParameter

Req. Analysis: false Req. Upload: true Is Annotated: true

Unit Measured: U/I		
Creatine kinase uci	OLA CRC 028 004 Lv4 2	
simpleParameter	DLA_CBC_028_001 V1.2	
Req. Analysis: false	Req. Upload: false	Is Annotated: true
Unit Measured: U/I		
Hemolysis status UprocedureMetadata	JCDLA_CBC_048_001 v1.1	
Req. Analysis: false	Req. Upload: false	Is Annotated: false
Options: Slight, Marked, Mod	erate, None,	
Albumin UCDLA_CBC_simpleParameter	007_001 v1.2	
Req. Analysis: false	Req. Upload: true	Is Annotated: true
Unit Measured: g/l		

Anesthesia used for blood collection UCDLA_CBC_036_001 | v1.0

procedureMetadata

Req. Analysis: true Req. Upload: true Is Annotated: false

Options:

Injection narcosis with Ketamine (110mg/kg)/Xylazine (11mg/kg)/ Antipamezole (Antisedan, 1mg/kg),

Gas anaesthesia with Isofluorane.

Injection narcosis with Ketamine (137mg/kg)/Xylazine (6.6mg/kg),

Injection narcosis with Sodium Pentobarbital (Euthatal),

Injection narcosis with Ketamine (110mg/kg)/Xylazine (11mg/kg),

Injection narcosis with Sodium Pentobarbital (Pentobarb, 0.1ml),

Injection narcosis with Tribromoethanol (Avertin),

Injection narcosis with Ketamine (100mg/kg)/ Xylazine (10mg/kg)/Antipamezole (Antisedan, 1mg/kg),

Injection narcosis with Ketamine (100mg/kg)/Xylazine (10mg/kg), No,

Injection narcosis with Sodium Pentobarbital (Somnopentyl),

Difficult bleed UCDLA CBC 055 001 | v1.0

procedureMetadata

Req. Analysis: false Req. Upload: false Is Annotated: false

Options: No, Yes,

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Date and time of sacrifice UCDLA CBC 040 001 LV1.1

	Req. Upload: true	
Fructosamine UCDLA simpleParameter	A_CBC_020_001 v1.2	
Req. Analysis: false	Req. Upload: false	Is Annotated: true
Unit Measured: umol/l		
Magnesium UCDLA_ConsimpleParameter	CBC_054_001 v1.5	
Req. Analysis: false	Req. Upload: false	Is Annotated: true
Unit Measured: mg/dl		
Iron UCDLA_CBC_011_00 simpleParameter	01 v1.5	
Req. Analysis: false	Req. Upload: false	Is Annotated: true
Unit Measured: mg/dl		

Calcium UCDLA_CBC_009_001 | v1.5

simpleParameter

Reg. Analysis: false Reg. Upload: true Is Annotated: true Unit Measured: mg/dl Thyroxine UCDLA_CBC_053_001 | v1.2 simpleParameter Req. Analysis: false Req. Upload: false Is Annotated: true Unit Measured: ug/dl Urea (Blood Urea Nitrogen - BUN) UCDLA_CBC_004_001 | v1.5 simpleParameter Req. Analysis: false Req. Upload: true Is Annotated: true Unit Measured: mg/dl

Creatinine UCDLA_CBC_005_001 | v1.5

simpleParameter

Req. Analysis: false Req. Upload: true **Is Annotated:** true Unit Measured: mg/dl Transferrin UCDLA_CBC_031_001 | v1.2 simpleParameter Reg. Analysis: false Reg. Upload: false Is Annotated: true Unit Measured: mg/dl Reagent manufacturer UCDLA_CBC_059_001 | v1.0 procedureMetadata Req. Analysis: true Req. Upload: false Is Annotated: false Options: Wako and Sekisui, Beckman Coulter, Microgenics,

Uric acid UCDLA_CBC_029_001 | v1.2

simpleParameter

Req. Analysis: false Req. Upload: false Is Annotated: true

Unit Measured: umol/l

Ferritin UCDLA_CBC_03 simpleParameter	30_001 v1.3	
Req. Analysis: false	Req. Upload: false	Is Annotated: true
Unit Measured: ng/ml		
Total bilirubin UCDLA simpleParameter	A_CBC_008_001 v1.4	
Req. Analysis: false	Req. Upload: true	Is Annotated: true
Unit Measured: mg/dl		
Alanine aminotrans	sferase ucdla_cbc_01	3_001 v1.2
Req. Analysis: false	Req. Upload: true	Is Annotated: true
Unit Measured: U/I		

Glycerol UCDLA_CBC_027_001 | v1.4

simpleParameter

Req. Analysis: false Req. Upload: false Is Annotated: true

Unit Measured: mmol/l

Equipment manufacturer UCDLA_CBC_034_001 | v1.0

procedureMetadata

Req. Analysis: true Req. Upload: true Is Annotated: false

Options: Hitachi, Roche, Beckman Coulter, JEOL (Siemens), Cobas, Olympus Diagnostics,

Alpha-amylase UCDLA_CBC_023_001 | v1.2

simpleParameter

Req. Analysis: false Req. Upload: false Is Annotated: true

Unit Measured: U/I

Method of blood collection UCDLA_CBC_037_001 | v1.0

procedureMetadata

Req. Analysis: true Req. Upload: true Is Annotated: false

Options: Heart puncture, Ta	ail vein, Retro-orbital pund	cture, Jugular vein, Cardia	ac puncture,

C-reactive protein UCDLA_CBC_032_001 | v1.0

simpleParameter

Req. Analysis: false Req. Upload: false Is Annotated: true

Unit Measured: mg/l

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Phosphorus UCDLA_CBC_010_001 | v1.6

simpleParameter

Req. Analysis: false Req. Upload: true Is Annotated: true

Unit Measured: mg/dl

Blood collection tubes UCDLA_CBC_039_001 | v1.1

procedureMetadata

Req. Analysis: false Req. Upload: false Is Annotated: false

Options: Kabe Labortechnik Lithium heparin coated tubes, Eppendorf 1.7ml, Kabe Labortechnik 1000ul Lithium Heparin, Sarstedt Li-Heparin gel tubes, BD Microtainer Lithium Heparin Tube, TERUMO CAPIJECT Lithium heparin coated tubes,

Greiner MiniCollect Lithium Heparin 1ml,			
globin A1c (HbA1c)	UCDLA_CBC_052_001 v1.		
Req. Upload: false	Is Annotated: true		
CDLA_CBC_025_001 v1.3			
Req. Upload: false	Is Annotated: true		
	Req. Upload: false		

BD Microtainer Lithium Heparin/PST Gel Blood Tube.

Aspartate aminotransferase UCDLA_CBC_012_001 | v1.2

simpleParameter

Req. Analysis: false Req. Upload: true Is Annotated: true

Unit Measured: U/I

Triglycerides UCDLA_CBC_017_001 | v1.4 simpleParameter Req. Analysis: false Req. Upload: true Is Annotated: true Unit Measured: mg/dl Sample dilution UCDLA_CBC_044_001 | v1.2 procedureMetadata Req. Analysis: false Req. Upload: true Is Annotated: false **Options:** Neat plasma, 1:3, Yes (by Equipment, automatically), 1:2, 1:5, 1:4, Neat serum, Varies. Potassium UCDLA_CBC_002_001 | v1.3 simpleParameter Req. Analysis: false Req. Upload: false Is Annotated: true Unit Measured: mmol/l

LIH (Hemolysis Severity - available on AU analysers) UCDLA

_CBC_019_001 | v1.3

simpleParameter

Req. Analysis: false Req. Upload: false Is Annotated: false

Equipment ID UCDLA_CBC_033_001 | v1.0

procedureMetadata

Req. Analysis: false Req. Upload: true Is Annotated: false