

Haematology ESLIM_016_001

Purpose

For the determination of blood cell counts (white blood cells, red blood cells, platelets), haemoglobin measurement and the calculation of haematological indexes (mean cell volume, mean corpuscular haemoglobin and mean cell haemoglobin concentration). Haematology measurements are obtained using either the Beckman Coulter AcT Diff or Siemens Advia 2120.

 Standard Operating Procedure	Title: Analysis of Haematology samples	
	Doc. Number: ESLIM_016_001	Date Issued: 01/06/09

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2. Associated Documents:

ESLIM_027_001_blood_sample_handling_haematology

Beckman Coulter A^CT Diff operator manual

Siemens Advia 2120 operator manual

3. Notes:

- 3.1. The validity of results obtained from metabolic studies is largely dependent on methods of animal husbandry. It is of vital importance that individuals following this procedure are experienced and aware of the animal's welfare, and are familiar with the animal being tested, in order to reduce the anxiety levels of the animal prior to testing.
- 3.2. The majority of mouse metabolic studies are age/sex/strain dependent. It is important to keep these parameters comparable throughout a single experiment.
- 3.3. It is recommended that all metabolic experimentation is conducted at approximately the same time of day because physiological and biochemical parameters change throughout the day.
- 3.4. All samples should be considered as potentially hazardous

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4. Quality Control:

For the Beckman Coulter Act Diff see 4.1-4.5:

- 4.1. Each morning, all parameters are tested with blood “4C –ES Cell” control.

The 3 levels include:

Abnormal Low

Normal

Abnormal High

- 4.2. Controls are gently inverted eight times according to the manufacturer’s instruction before use.

- 4.3. Control values must be within three standard deviations, otherwise the measurement has to be repeated.

- 4.4. Controls can be stored at +4°C.

- 4.5. Control:

All control data are managed using the Biorad Unity Plus software that provides graphical reports (Levey-Jennings graphs, Youden diagram, and monthly cumulative histograms).

For the Siemens Advia 2120 Analyzer see 4.6-4.9

- 4.6. Each morning, all parameters are tested with fixed Testpoint control blood samples.

The 3 levels include:

Abnormal Low

Normal

Abnormal High

- 4.7. Controls are stored at 2-8°C and brought to room temperature on a roller mixer

before use



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4.8. Control values must be within the target range specified in the Advia 2120 software, otherwise the measurement has to be repeated.

4.9. All control data are managed using the Advia 2120 software that provides graphical reports.

5. Equipment:

5.1. A^CT diff (Beckman Coulter) or

5.2. Siemens Advia 2120 with multispecies software (Siemens Medical Solutions Diagnostics)

6. Supplies:

For the Beckman Coulter Act Diff see sections 6.1-6.2

6.1. Reagents:

A^CT Pak (ref 8448322 Beckman Coulter, France):

Reagent 1 = Diluent (balanced electrolyte solution)

Reagent 2 = Lytic reagent

6.2. Quality control:

Blood "4C -ES Cell" control (ref 7547114 Beckman Coulter) with 3 levels (abnormal low, normal and abnormal high).

For the Siemens Advia 2120 Analyzer see 6.3-6.4

6.3. Reagents:

Complete blood count timepac, Differential timepac, Perox sheeth solution, Universal rinse, Defoamer, Ez clean, Wash solution (Siemens Medical Solutions Diagnostics)

6.4. Quality control:

Testpoint controls (Siemens Medical Solutions Diagnostics) with 3 levels (Low, Normal and High).

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7. Procedure:

Summary of protocol:

- **Sample collection and storage**
- **Sample preparation**
- **Analysis**

7.1. Sample collection and storage:

7.1.1. Collect samples according to the blood and sampling procedures (refer to ESLIM_027_001).

7.1.2. Samples should be analysed within 2 hours after collection.

7.1.3. Volume:

12µl for the Beckman Coulter

200µl for the Siemens Advia 2120

7.1.4. Refer to section 4 to perform QC check

7.2. Sample preparation:

Immediately following sample collection put the blood samples (EDTA Microvette tubes) on a rotary agitator.

7.3. Analysis:

To perform the analysis: follow either the Siemens Advia 2120 operator manual or the Beckman ACT diff operator manual (pages 3-2 to 3-8)

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8. Parameters recorded:

The following parameters are required.

- White blood cell-count
- Red Blood Cell-count
- Haemoglobin
- Haematocrit
- Mean-cell-volume
- Mean-corpuscular-haemoglobin
- Mean-cell-haemoglobin-conc
- Platelets-count

9. Metadata recorded:

The following metadata is required.

- Equipment name (e.g. Haematology analyser)
- Equipment manufacturer (e.g. Siemens Medical Solutions Diagnostics)
- Equipment model (e.g. ADVIA 2120)
- Method of blood collection (e.g. retro-orbital)
- Date/Time of blood collection
- Anaesthesia used for blood collection (e.g. isoflourane)
- Day of measurement

The following metadata is optional.

- EMPReSSID for blood collection SOP
- Chip Card (Beckman analyser only)



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10. Supporting information:

There is no supporting information available for this SOP

11. History Review:

There is no history review available for this SOP

Parameters and Metadata

White blood cell count ESLIM_016_001_001 | v1.0

simpleParameter

Req. Analysis: false

Req. Upload: true

Is Annotated: true

Unit Measured: $10^3/\text{ul}$

Description: White_blood_cell_count

Red blood cell count ESLIM_016_001_002 | v1.0

simpleParameter

Req. Analysis: false

Req. Upload: true

Is Annotated: true

Unit Measured: $10^6/\text{ul}$

Description: Red_blood_cell_count

Haemoglobin ESLIM_016_001_003 | v1.0

simpleParameter

Req. Analysis: false

Req. Upload: true

Is Annotated: true

Unit Measured: g/dl

Description: Haemoglobin

Haematocrit ESLIM_016_001_004 | v1.0

simpleParameter

Req. Analysis: false

Req. Upload: true

Is Annotated: true

Unit Measured: %

Description: Haematocrit

Mean cell volume ESLIM_016_001_005 | v1.0

simpleParameter

Req. Analysis: false

Req. Upload: true

Is Annotated: true

Unit Measured: fL

Description: Mean_cell_volume

Mean corpuscular haemoglobin ESLIM_016_001_006 | v1.0

simpleParameter

Req. Analysis: false

Req. Upload: true

Is Annotated: true

Unit Measured: pg

Description: Mean_corpuscular_haemoglobin

Mean cell haemoglobin concentration ESLIM_016_001_007 | v1.0

simpleParameter

Req. Analysis: false

Req. Upload: true

Is Annotated: true

Unit Measured: g/dl

Description: Mean_cell_haemoglobin_concentration

Platelets count ESLIM_016_001_008 | v1.0

simpleParameter

Req. Analysis: false

Req. Upload: true

Is Annotated: true

Unit Measured: $10^3/\text{ul}$

Description: Platelets_count

Equipment name ESLIM_016_001_801 | v1.0

procedureMetadata

Req. Analysis: false

Req. Upload: true

Is Annotated: false

Description: Equipment_name

Equipment manufacturer ESLIM_016_001_802 | v1.0

procedureMetadata

Req. Analysis: true

Req. Upload: true

Is Annotated: false

Description: Equipment_manufacturer

Equipment model ESLIM_016_001_803 | v1.0

procedureMetadata

Req. Analysis: true

Req. Upload: true

Is Annotated: false

Description: Equipment_model

Method of blood collection ESLIM_016_001_804 | v1.0

procedureMetadata

Req. Analysis: true

Req. Upload: true

Is Annotated: false

Description: Method_of_blood_collection

EMPreSSID for blood collection SOP ESLIM_016_001_805 | v1.0

procedureMetadata

Req. Analysis: false

Req. Upload: false

Is Annotated: false

Description: EMPreSSID_for_blood_collection_SOP

Date/time of blood collection ESLIM_016_001_806 | v1.0

procedureMetadata

Req. Analysis: false

Req. Upload: false

Is Annotated: false

Description: DateTime_of_blood_collection

Anaesthesia used for blood collection ESLIM_016_001_812 | v1.0

procedureMetadata

Req. Analysis: true

Req. Upload: true

Is Annotated: false

Description: Anaesthesia_used_for_blood_collection

Date of measurement ESLIM_016_001_813 | v1.0

procedureMetadata

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

Description: Date_of_measurement

Chip card ESLIM_016_001_814 | v1.0

procedureMetadata

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

Description: Chip_Card
