

#### **CMHD Pathology Core**

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# **CMHD Pathology Report**

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ReportID: Report Date: March 06, 2013

Pathologist: H. Adissu



#### **Mouse Genetics Project**

Wellcome Trust Sanger Institute Wellcome Trust Genome Campus Hinxton, Cambridge CB10 1SA UK

email:

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**CMHD LabID: N12-1515** 

#### **Relevant History:**

Phenotype:

abnormal snout morphology asymmetric snout short snout decreased body length decreased body weight decreased total body fat amount decreased body weight abnormal radius morphology abnormal ulna morphology abnormal cranium morphology abnormal tooth morphology partial lethality

# AnimalID: M00583718 Histopathology Findings:

# kidney (MA:0000368)

# **Histopath Description:**

Nearly half of the renal parenchyma is replaced by a large cyst; the renal parenchyma surrounding the cyst is mildly compressed.

#### **Morphological Diagnosis:**

**Duration:** Chronic; **Distribution:** Unilateral; **Severity:** severe; **MPATH Diagnosis:** hydronephrosis MPATH:635

# **Definitive Diagnosis:**

Hydronephrosis with mild parenchymal atrophy.

#### **Histopathology Comments:**

Hydronephrosis is usually caused by an ascending obstructive urinary lesion; this is not evident in the examined sections.



Kidney, hydronephrosis, 1.25x



Kidney, normal, 1.25x

# liver (MA:0000358)

#### **Histopath Description:**

hepatic lipidosis is not present

# **Definitive Diagnosis:**

No hepatic lipidosis

# **Histopathology Comments:**www.cmhd.ca/pathology/reports/histopathology\_report\_wtsi.asp?ID=94926735

No hepatic lipidosis consistent with poor growth and weight gain

# pancreas (MA:0000120)

# **Histopath Description:**

There are multifocal areas of necrosis within multiple lobules. There are occasional polymorphonuclear cells within the necrotic parenchyma.

# **Morphological Diagnosis:**

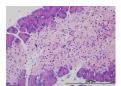
Distribution: multifocal; Severity: moderate;

#### **Definitive Diagnosis:**

Pancreatic coagulative necrosis

#### **Histopathology Comments:**

This is uncommon lesion; the pathogenesis is not clear.



Pancreas, necrosis,

#### skin (MA:0000151)

#### **Histopath Description:**

There are multifocal serocellular crusts overlying a focal hyperplastic epidermis. There is mild perivascular dermatitis.

#### **Morphological Diagnosis:**

**Duration:** Chronic; **Distribution:** Focal; **Severity:** mild; **MPATH Diagnosis:** inflammation MPATH:212

#### **Definitive Diagnosis:**

Focal serocellular exudate and epidermal hyperplasia (epidermitis)

#### brain (MA:0000168)

# **Histopath Description:**

Mild hydrocephalus.

# **Morphological Diagnosis:**

Severity: mild; MPATH Diagnosis: hydrocephalus MPATH:639

#### **Definitive Diagnosis:**

Mild hydrocephalus of the lateral ventricles

# AnimalID: M00604074 Histopathology Findings:

# liver (MA:0000358)

# **Histopath Description:**

hepatic lipidosis is not present

# **Definitive Diagnosis:**

No hepatic lipidosis

#### **Histopathology Comments:**

No hepatic lipidosis consistent with poor growth and weight gain

# heart (MA:0000072)

#### **Histopath Description:**

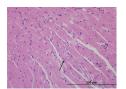
at the base of the left ventricle, there are large myocardiocytes with gigantic vesicular nuclei (Anitschkow cells). Some of these cells have vacuolated cytoplasm. There is mild dissaray of myocardial fibers.

# **Morphological Diagnosis:**

**Distribution:** multifocal; **Severity:** mild; **MPATH Diagnosis:** cardiomyopathy MPATH:615

#### **Definitive Diagnosis:**

Myocardial degeneration, mild



Left ventricle, degeneration, 40x

#### brain (MA:0000168)

#### **Histopath Description:**

Mild hydrocephalus.

**Morphological Diagnosis:** 

Severity: mild; MPATH Diagnosis: hydrocephalus MPATH:639

**Definitive Diagnosis:** 

Mild hydrocephalus of the lateral ventricles

# AnimalID: M00583721 Histopathology Findings:

liver (MA:0000358)

#### **Histopath Description:**

hepatic lipidosis is very minimal affecting approximately 1% of heaptocytes

# **Definitive Diagnosis:**

Minimal hepatic lipidosis

#### **Histopathology Comments:**

Minimal hepatic lipidosis is consistent with poor growth and weight gain

# brain (MA:0000168)

# **Histopath Description:**

Mild hydrocephalus.

# **Morphological Diagnosis:**

Severity: mild; MPATH Diagnosis: hydrocephalus MPATH:639

#### **Definitive Diagnosis:**

Mild hydrocephalus of the lateral ventricles

# AnimalID: M00583719 Histopathology Findings:

# kidney (MA:0000368)

# **Histopath Description:**

There is multifocal lymphocyte and macrophage infiltrates within the renal pelvis

# **Morphological Diagnosis:**

**Distribution:** Multifocal to coalescing; **Severity:** mild;

#### **Definitive Diagnosis:**

Chronic pyelitis

### liver (MA:0000358)

#### **Histopath Description:**

hepatic lipidosis is not present

# **Definitive Diagnosis:**

No hepatic lipidosis

# **Histopathology Comments:**

No hepatic lipidosis consistent with poor growth and weight gain

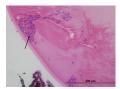
#### lens (MA:0000275)

#### **Histopath Description:**

There are clusters of hypertrophic lens cells within the lens ate the anterior aspect.

### **Morphological Diagnosis:**

MPATH Diagnosis: cataract MPATH:29



lens, cataract, note clusters of lens cells within the lens substance, 40x

#### retina (MA:0000276)

# **Histopath Description:**

Involving one eye, there are clusters of external nuclear structures within the internal plexiform layer.

# **Morphological Diagnosis:**

Distribution: Focal; Severity: mild;

#### **Definitive Diagnosis:**

Retinal dysplasia

# **Report Summary and Recommendation:**

Absence of hepatic lipidosis in this line is consistent with decreased body weight, decreased total body fat amount, decreased body weight documented in clinical phenotyping. The significance of the pancreatic lesion in one of the mice (M00583718) is unceratin. Mild inflammation of the renal pelvis was noted in two mice (M00583718 and M00583719); the former had unilateral hydronephrosis. The lesion suggests unilateral obstruction proximal to the urinary bladder. Cataract was observed in one mouse (M00583719). Microscopic examination of single sections is not adequate to confirm the skeletal and facial morphodeviants.