

# The Harwell Frozen embryo and sperm archive (FESA)

<https://www.neurodegenerationresearch.eu/survey/the-harwell-frozen-embryo-and-sperm-archive-fesa/>

## Name of resource

The Harwell Frozen embryo and sperm archive (FESA)

## Name of Principal Investigator

Title Dr

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## Summary

The MRC's frozen embryo and sperm archive holds a collection of mouse models of human disease that are distributed to the wider scientific community. We also hold some mouse models that have been frozen for our own researchers that are not available to the community.

### 1a. The resource holds animal models relevant to the study of the following neurodegenerative diseases

Motor neurone diseases

Prion disease

Huntington's disease

Neurodegenerative disease in general

Parkinson's disease

Spinocerebellar ataxia (SCA)

Spinal muscular atrophy (SMA)

### 1b. The resource holds:

Animals

Frozen embryos

Frozen sperm

### 2a. The resource acts as a centre for access and distribution to external groups (who are not the PIs of the resource)

## 2b. Procedures and rules for access

Access independent of collaboration with PI

Local/ regional access

National access

International access

Access to industry

Charge for retrieval

## 3a. Does the resource develop animal models for external groups

1

## 3b. Types of models provided

Not applicable

## 4a This activity is supported as:

Independent of collaboration

## 4b. The supplied material deposited in a central repository

2

## 5a Disease models available

| Disease | Species | Available to external user (Y/N) | Full phenotypic character (Y/N or partial) | Phenotypes                             | Genotypes or other subtypes |
|---------|---------|----------------------------------|--|--|-----------------------------|
| HD      | Mouse   | N                                | Partial                                    | Short life span/tremors                | Hets                        |
| Prion   | Mouse   | Y                                | Y  | Suceptibility to disease               | Hets/Homs                   |
| SMA     | Mouse   | Y                                | Partial                                    | Neuropathology                         | Homs                        |
| PD      | Mouse   | Y                                | Partial                                    | Olfactory memory and motor control     | Het                         |
| MND     | Mouse   | Y                                | Y  | Mice show dystrophic changes in mucscl | Hets/Homs                   |
|         |         |                                  |  | histopathology                         |                             |

## 5b. Other models/phenotypes available through the resource relevant to neurodegenerative conditions

| No. of models     | Available to external users | Full phenotypic characterisation available (Y/N or partial) | Nature of phenotype |
|-------------------|-----------------------------|---|---------------------|
| Low grip strength | 3                           | Y   | Partial             |
| 2                 | Y                           | Partial   | Memory deficit      |
| 2                 | Y                           | Partial   | Muscle tremors      |
| Numerous          | Y                           | Partial   | Various phenotypes  |

## 6. European or international consortia or networks to which the resource is linked

**Jackson Laboratories, USA**International mouse knockout consortium

European Mutant Mouse Archive

International Mouse Knockout Consortium

**7a. Maintenance of the resource is dependent on continued funding**

1

**7b. End date of current funding period**

N/A

**7c. Expected lifespan of the resource (in years)**

1000

**7d. Other plans affecting future use**

FESA is a core funded MRC resource but it is also supported in part by the funds we receive from the EU through our membership of the EMMA consortium. The current EMMA grant will come to an end Dec 2012.