NON-TECHNICAL SUMMARY

Protection after heart attack or stroke

Project duration

5 years 0 months

Project purpose

- (a) Basic research
- (b) Translational or applied research with one of the following aims:
  - (i) Avoidance, prevention, diagnosis or treatment of disease, ill-health or abnormality, or their effects, in man, animals or plants

Key words

No answer provided

Animal types | Life stages
-------------|-----------------
Mice         | adult, pregnant, juvenile, neonate, embryo
Rats         | adult

Retrospective assessment

The Secretary of State has determined that a retrospective assessment of this licence is not required.

Objectives and benefits
Description of the projects objectives, for example the scientific unknowns or clinical or scientific needs it's addressing.

What is the aim of this project?

The aim of this project is to understand how heart and brain tissue can be saved from damage following a heart attack or stroke.

Potential benefits likely to derive from the project, for example how science might be advanced or how humans, animals or the environment might benefit - these could be short-term benefits within the duration of the project or long-term benefits that accrue after the project has finished.

Why is it important to undertake this work?

Heart attack and stroke are the leading cause of death worldwide and present a huge burden on health systems such as the NHS. These diseases occur when an artery supplying either the heart muscle or the brain respectively is blocked, disrupting the delivery oxygen and nutrients as well as the removal of waste products. This inevitably leads to death of the tissue that is without blood supply, which can lead to both severe long-term disability and increased mortality in patients.

What outputs do you think you will see at the end of this project?

This project will develop and test novel protective drugs for heart attack and stroke, in order to pave the way towards the use of these compounds in patients. The information gathered from this project will allow us to see how these substances affect the heart or brain and what doses and treatment regimens are most likely to be effective. We will also examine whether lifestyle changes (such as diet and exercise) or pre-existing diseases alter their effectiveness.

We aim to publish in high-ranking journals in order to inform the scientific community. I also regularly take part in public engagement activities, such as Pint of Science, University-wide Science festival, "Naked Scientist" and appear in the popular press in order to inform the general public about the progress and new findings of our research.

Ultimately, the above results and achievements will pave the way towards the use of novel compounds in patients with heart attack and stroke.

What will be the impact of this proposed work on humans / animals / the environment in the short-term (within the duration of the project), in the medium-term and the long-term (which may accrue after the project is finished)?

Initially, our fellow scientists will benefit from our findings about the underlying mechanisms of health and disease in heart attack and stroke. We publish in prestigious journals and are active members of many scientific societies with regular participation at meetings.
In the second line, we anticipate bringing at least one of our novel compounds towards use in patients with heart attack or stroke within the next 5 years. Drug development is a long and complicated process, but we are confident that this project will enable us to move forward to a first-in-man study on this timescale.

How will you maximise the outputs of your work?

My group is very well connected, not only locally, but also on a national and international level. The quality of our animal work has made my group one of the leading groups in the country investigating damage during heart attack. We regularly publish our results, including negative outcomes or unsuccessful approaches.

My group and my collaborators are very active on various outreach channels, including twitter, popular press, radio, and websites. The general public has access to all our findings and we widely share our knowledge. We have a large number of collaborators world-wide, which work together with us to maximise our findings and make translation towards patient care more likely.

Species and numbers of animals expected to be used

- Mice: 4400
- Rats: 800

Predicted harms

Typical procedures done to animals, for example injections or surgical procedures, including duration of the experiment and number of procedures.

Describe, in general terms, the procedures animals will undergo, eg injections, surgical procedures. Include the typical number of procedures individual animals will undergo and the likely duration of suffering.

Typically either purpose-bred genetically-altered animals are used or genetically normal mice will be used. The breeding protocols follow standard breeding procedures.

In a subset of animals certain conditions such as exercise, administration of substances or alteration of diet will be applied in order to mirror more closely the conditions in humans. This includes pre-existing lifestyle (exercise or diet) or pre-existing medication.

Finally, animals will be subjected to surgical procedures to induce either a heart attack or a stroke. These final procedures will be performed under terminal anaesthesia and the animal will not be allowed to recover.
Expected impacts or adverse effects on the animals - for example, pain, weight loss, inactivity or lameness, stress, or abnormal behaviour - and how long those effects are expected to last.

Overall, the animals will only suffer mild discomfort and no lasting harm. Most of the interventions, such as exercise or dietary intervention, are not expected to have any discomfort and certainly no harm to the animals. Rarely, very mild discomfort will occur.

The surgical procedures will be performed solely under confirmed general anaesthesia and the animal will not be allowed to recover. We further aim to perform any interventions, such as imaging or administration of substances under the terminal anaesthesia. Only rarely it is necessary to do this beforehand.

Expected severity categories and the proportion of animals in each category, per species.

What are the expected severities and the proportion of animals in each category (per species)?

- Protocol 1: mild
- Protocol 2, 3: non-recovery
- Protocol 4, 5: mild
- Protocol 6, 7: moderate
- Protocol 8, 9, 10: mild

What will happen to the animals at the end of the study?

- Kept alive

Replacement

State what non-animal alternatives are available in this field, which alternatives you have considered and why they cannot be used for this purpose.

Why do you need to use animals to achieve the aim of your project?

Information about the distribution of the drugs, effects on the heart and brain, or blood pressure and exercise tolerance cannot be answered using cell-based models.

The novel drugs which will be tested with this license have already undergone extensive testing in suitable non-regulated models in cell culture, isolated cells or isolated organ preparations. Therefore, we already know most of the mechanisms of action and have information about possible toxicity, dosing and the most effective treatment regimen. However we need to confirm efficacy in animal models before we can translate these compounds to human use. Furthermore, these animal models have to mimic the
situation of patients as close as possible. This can be done by an alteration of diet, exercise, application of pre-existing medication, and the use of pre-diseased genetically-altered models.

**What was your strategy for searching for non-animal alternatives?**

We are already using isolated organ applications for testing in the heart (Langendorff preparation). This cannot be done with the brain.

**Why were they not suitable?**

Cell lines cannot mimic important conditions, such as exercise, and do not have crucial confounding elements, such as circulation or the blood brain barrier.

There have been decades of research trying to find a suitable cell-based model of ischaemia (lack of oxygen to tissue)/reperfusion (re-introduction of oxygen and blood flow to tissue), but so far none have been found.

**Reduction**

**Explain how the numbers of animals for this project were determined. Describe steps that have been taken to reduce animal numbers, and principles used to design studies. Describe practices that are used throughout the project to minimise numbers consistent with scientific objectives, if any. These may include e.g. pilot studies, computer modelling, sharing of tissue and reuse.**

**How have you estimated the numbers of animals you will use?**

The number of animals is determined by previous experience with an identical methodological approach under a different license. Furthermore, we will use power calculations based on our previous data to calculate numbers for each individual experimental setting.

**What steps will you take to reduce animal numbers? Where applicable, what principles will you use to design experiments?**

I use the NC3R Experimental Design Assistant (EDA) in order to thoroughly design the studies.

In addition, I am on an EU committee (COST Action “Cardioprotection”) which aims to standardise ischaemia/reperfusion experiments throughout Europe considering the 3Rs as well as highest scientific standards. Within this consortium, my group is one of the reference groups for acute heart attack in mice.

**What other measures apart from good experimental design will you use to minimise numbers?**
Wherever possible, we try to assess as many parameters in a single animal as possible and reduce the numbers in one treatment group to an absolute minimum that the statistical analysis will allow us as determined by power analysis. This includes performing sophisticated imaging techniques which allow us to gain much additional information about the heart attack or stroke in a single animal.

**Refinement**

Give examples of the specific measures (e.g., increased monitoring, post-operative care, pain management, training of animals) to be taken, in relation to the procedures, to minimise welfare costs (harms) to the animals. Describe the mechanisms in place to take up emerging refinement techniques during the lifetime of the project.

Why are the animals, models and methods you will use the best to meet your objectives? Why will your approach cause the least pain, suffering, distress or lasting harm?

The present project will use either wild-type mice and rats or genetically-altered mice. The genetically-altered strains are lacking important mechanistic elements which helps us to understand the mechanisms of how we can protect against heart attack or stroke. We are already breeding these genetically-altered lines under a different license and the animals do not show any clinical signs.

The main aim of the license is to test protective effects in heart attack and stroke, and as both models will be performed under terminal anaesthesia they are therefore free of any suffering.

The alteration of diet or exercise and administration of substances are the most refined models used to determine these important factors which can influence the outcome of heart attack and stroke. There will not be any lasting harm and only mild discomfort for the animals due to these models.

Why can’t you use a less sentient animal, (for example at an immature stage, a less sentient species or using terminally anaesthetised animals)?

Mice and rats are the most useful species from which it is possible to obtain relevant and meaningful physiological and pathophysiological information. Especially non-invasive imaging such as MRI and PET cannot be performed in smaller species. Furthermore, mice allow genetic manipulations in order to more specifically study the underlying mechanisms.

What are you going to do to refine the procedures (for example increased monitoring, post-operative care, pain management, training of animals) to minimise the welfare costs (harms) to the animals?

We closely monitor the animals throughout the experiments. This includes the use of sophisticated heart and brain function monitors, such as ECG and blood flow monitor (doppler), as well as temperature control. If anaesthesia is applied, the efficacy is monitored closely throughout the procedure. Should any anaesthesia or surgical problem occur which would potentially harm the animal, the experiment will be terminated and the animal humanely killed.
In protocols or procedures where anaesthesia is not used, animals will be monitored for any deviation from normal health and behaviour and should any signs appear animals will be killed immediately.

The animals are allowed to acclimatise to either the exercise protocol or the dietary changes.

**What published best practice guidance will be followed to ensure experiments are conducted in most refined way?**

We will follow the NC3Rs Experimental Design Assistant (EDA) see https://www.nc3rs.org.uk/experimental-design as well as the following guidelines:

- for pilot studies (https://www.nc3rs.org.uk/conducting-pilot-study);
- ARRIVE guidelines, https://www.nc3rs.org.uk/arrive-guidelines;
- IMPROVE guidelines for stroke models: https://www.nc3rs.org.uk/news/improve-ing-animal-welfareexperimental-stroke-research;
- published guidelines to assist with planning animal research and testing, such as the PREPARE guidelines: http://journals.sagepub.com/doi/full/10.1177/0023677217724823

**How will you ensure you continue to use the most refined methods during the lifetime of this project?**

I receive regular updates via the Biological Service of the University of Cambridge about news and advances in the 3Rs.

**Explain the choice of species and the related life stages**

Since the effects on hemodynamic parameters and the elimination/distribution of compounds are essential parameters of a pharmacotherapy, isolated organ systems cannot be used. It is not possible to gain this information in vitro since the whole-body system needs to be intact due to the influences that administration, distribution, metabolism and excretion have on the availability and efficacy of the compound.

Furthermore, changes in exercise levels, diet or pre-existing diseases cannot be mimicked without animals.

Rodents are the smallest possible species where interventions, such as ligation of a coronary artery or occlusion of a brain artery in order to induce a heart attack or stroke respectively are possible.

In addition, rodents offer the possibility to use genetically-altered models in order to mirror patient conditions (such as pre-existing diseases) more closely.