

## Application summary

### Application title

*This should be the name of the MA/MSc course.*

### Proposed duration of funding (months)

### Proposed start date

### Name of administering organisation

*Please enter the name of the organisation where you intend to hold the award. If your application is successful, this is the organisation that will be responsible for administering the award.*

### Lead applicant's address at administering organisation

Department/Division

Organisation

Street

City/Town

Postcode/Zipcode

Country

*Please enter the address where you will be based at the administering organisation. If your application is successful, this is the address that will be used in the award letter.*

Please confirm that this is the only application for the Humanities and Social Science Master's Studentship scheme being submitted from your prospective host organisation this year.

*The relevant authority/Master's programme contact within the university should be able to confirm this to you.*

## Lead applicant

### Lead applicant details

Full Name

<b>Department</b>	
<b>Division</b>	
<b>Organisation</b>	
<b>Address Line 1</b>	
<b>City/Town</b>	
<b>Postcode</b>	
<b>Country</b>	
<b>Telephone No.</b>	
<b>Email Address</b>	

<b>ORCID iD</b>	
<b>ORCID iD</b>	

*Lead applicants must add their ORCID iD. Find out more about ORCID on our website.*

**Career history (current/most recent first)**

<b>From</b>	<b>To</b>	<b>Position</b>	<b>Organisation</b>

*Please provide details of your current position (if applicable) and all previous posts held, listing most recent first.*

Skills gained relevant to this application

**Education/training**

<b>From</b>	<b>To</b>	<b>Qualification</b>	<b>Subject</b>	<b>Organisation</b>

*Please provide details of relevant education/training, listing the most recent first.*

<b>Clinical status</b> Do you have a medical/veterinary degree?	
--	--

*Please note that this includes dental and clinical psychology degrees.*

Please specify

Are you clinically active?	
----------------------------	--

What is your specialty?

Please choose your specialty from the dropdown list – if it is not on the list, select 'Other' and specify.

Please specify

**Career breaks**

Have you had any career breaks or periods of part-time work, for example parental or long-term sick leave?

We encourage applications from researchers who have taken career breaks, and wish to ensure that any such breaks are duly taken into account when considering your track record. Please state when and for what period of time you took a break, or were working on a part-time basis.

Please provide details

Do you wish to undertake this award part time?

If you wish to undertake this award part time, please contact the Trust to discuss your requirements.

**University Degrees/Diplomas**

Give subject, university, dates of attendance and class of degree if already graduated, or most recent year's classification and marks.

Name of current or most recent academic tutor. Please specify their department and organisation.

Recommendation from current tutor, or another appropriate academic referee if you are not presently undertaking a course of study (500 words maximum).

## Sponsors

### Sponsor

*The sponsor must be based at the administering organisation.*

1

<b>Sponsor</b>	
<b>Full Name</b>	
<b>Department</b>	
<b>Division</b>	
<b>Organisation</b>	
<b>Address Line 1</b>	
<b>City/Town</b>	
<b>Postcode</b>	
<b>Country</b>	
<b>Telephone No.</b>	
<b>Email Address</b>	

*The sponsor must be based at the administering organisation.*

<b>Title of current post</b>

<b>Recommendation from prospective Head of Department (or equivalent)</b> The recommendation should include a brief description of the course/programme and an outline of what is unique about it, as well as any synergies with research taking place in the host department (500 words maximum).
---

## Resubmissions

Is this a resubmission of an application submitted to the Wellcome Trust within the last 24 months?	
---	--

*Applicants must contact the Wellcome Trust before resubmitting an application.*

Please describe how this application differs from the original (200 words max.)
---

## Research summary

**Research summary**  
Please provide a summary of your proposed research, including key goals (200 words max.)

*This will be used as a short form 'abstract' and is necessary to enable the Trust to classify your proposal by subject area. The summary should be as complete as possible within the word limit, and should include key words which best describe the proposal to enable text searching.*

**Lay summary**  
Please provide a summary of your proposed research that people who may not be familiar with the subject can understand. We may edit your summary and then use it to describe your research on our website and elsewhere.

*You don't need to oversimplify your research, but try to explain it as clearly as possible. You should write in the first person ("I" and "we") and structure your summary in this order:*

- *background to the research problem*
- *your approach*
- *expected impact of your work.*

**Example of a lay summary**  
*Complete diagnostic autopsies (CDA) remain the gold standard for determining cause of death, but performing them in low- and middle-income countries (LMICs) is challenging. Facilities are inadequate, skilled staff scarce and public acceptance low. A minimally invasive autopsy (MIA) procedure involving organ-directed sampling has been proposed as an alternative. Oxford University Clinical Research Unit (OUCRU) is evaluating the use of MIA in Vietnam, but the method's ultimate effectiveness will depend on its public reception. The public view on post mortem examinations and consent for them are complex and under-researched. I will use interviews, focus groups and participant observations to assess the practice and perceptions of autopsy in Vietnam and Nepal. I will investigate socio-cultural factors surrounding these perceptions and explore ethical barriers preventing autopsy uptake. I will try to determine whether MIA may be more acceptable than traditional forms of post mortem. I will then work alongside clinicians to develop more culturally sensitive and appropriate methods of obtaining consent to autopsy.*

## Details of proposal

Motivation for choosing this subject and this organisation (250 words max.)

--

Provisional dissertation topic (500 words max.)

Outline briefly the career you hope to pursue (250 words max.)

## Outputs management and sharing

Will the proposed research generate outputs of data, software, materials or intellectual property that hold significant value as a resource for the wider research community?	
<p><i>As set out in our Data, Software and Materials Management and Sharing Policy, all Wellcome-funded researchers are expected to manage their research outputs in a way that will achieve the greatest health benefit, maximising the availability of research data, software and materials with as few restrictions as possible. If your proposed research is likely to generate significant outputs - data, software, materials and/or intellectual property - that will hold clear value as a resource for others in academia or industry, you are required to provide an outputs management plan.</i></p> <p><i>Our guidance on developing an outputs management plan sets out the circumstances under which such a plan is required and gives an overview of what you should consider.</i></p> <p><i>Your plan should be clear, concise, proportionate and focus specifically on how outputs will be identified, managed and then used to advance potential health benefits. You should set out and address clearly the following:</i></p> <p><i>1) For significant data, software and materials outputs</i></p> <ul style="list-style-type: none"><li><i>(i) What significant outputs will your research generate?</i></li><li><i>(ii) When do you intend to share these outputs?</i></li><li><i>(iii) Where will you make these outputs available?</i></li><li><i>(iv) How will they be discovered and accessed by others?</i></li><li><i>(v) Are limits on sharing required?</i></li><li><i>(vi) How will these outputs be preserved?</i></li></ul> <p><i>2) For intellectual property outputs</i></p> <ul style="list-style-type: none"><li><i>(i) What IP will your research generate?</i></li><li><i>(ii) How will you protect this IP?</i></li><li><i>(iii) How will the IP be used to achieve health benefits?</i></li><li><i>(iv) Provide the name and contact details for the person in your organisation (e.g. Technology Transfer Officer or Business Development executive) who can act as a point of contact for Wellcome in connection with the protection and</i></li></ul>	

commercialisation of this IP.

3) Describe any resources that you will need to deliver your outputs management plan.

Please note that regardless of whether or not a plan is required, you must ensure that data and original software underlying published research findings are accessible at the point of publication.

Which approach do you intend to use to maximise the impact of your significant research outputs to improve health and benefit the wider research community?

Please provide an outputs management plan. Ensure this describes any significant data, software, materials or intellectual property outputs, their management, and resources required (refer to guidance).

(700 words max.)

Please refer to guidance next to the above question: 'Will the proposed research generate outputs of data, software, materials or intellectual property that hold significant value as a resource for the wider research community?'

## Public engagement

Do you have plans for engaging with the non-academic public about your work?

*The Wellcome Trust is committed to engaging with society about the research it supports. We aim to foster mutual trust and understanding and place science within a societal, historical and cultural context. Further information is available on the Wellcome Trust's website.*

*We expect those researchers who receive funding from the Wellcome Trust to help support an environment within which science can flourish by informing, consulting and collaborating with the non-academic public.*

Please provide a brief outline of your public engagement plans.  
(250 words max.)

*Describe your plans to engage the non-academic public about your work beyond press and media activity. Engagement that is essential for the ethical conduct of the research, such as patient information leaflets or community advisory boards, should be part of your research methodology and included within your main research costs.*

Please note that we provide support for Wellcome Trust funded researchers to engage with the non-academic public. Do you wish to receive information about training, funding and other public engagement opportunities?

## Location of activity

Will the funded activity take place at more than one location?

*It is important that we are able to track the countries and organisations where research activity is taking place and the approximate proportion of the funds that will be spent at each location.*

*You should list any locations where you will be conducting research or redirecting funds outside of the administering organisation. This includes, but is not limited to, anywhere in receipt of indirect funding, fieldwork sites, and time spent working in another institution/laboratory. This does not include conference attendance.*

*Salary costs, if requested, should be attributed to the employing organisation.*

For each location, select the country and, where applicable, state the organisation (please include the administering organisation). Indicate the approximate percentage of the total funds that will be spent in each location, entering zero for locations where no significant funds will be spent. Salary costs, if requested, should be attributed to the employing organisation.

Country	Organisation	Percentage of funds

## Costs requested

Please select the currency in which you wish to apply.

*It is expected that costs within the application will be submitted in the currency which, in the view of the applicant(s), best enables the activity to be undertaken. In the majority of cases, the currency specified is likely to be the local currency. Where this is not the case, please explain the reasons for selecting the chosen currency.*

*Please refer to the Wellcome Trust's website for further information regarding selecting a currency.*

*If at any point, the Wellcome Trust is unable to award in the currency requested, discussions will be held with the administering organisation to decide whether an alternative currency should be used. If you have any concerns that the currency you would like to request may not be readily available, please contact the Wellcome Trust by e-mailing: [grantpayments@wellcome.ac.uk](mailto:grantpayments@wellcome.ac.uk).*

Is the selected currency your local currency?

What is your local currency?

Please state clearly the reasons for requesting costs in the selected currency and the exchange rate used (100 words max.)

--

<b>Studentship fees</b>	
Are you requesting studentship fees?	
<p>Please provide details of your studentship fees. The Trust reserves the right to query and challenge the costs requested, where it considers it appropriate and necessary to do so.</p> <p>Please note that the Wellcome Trust provides a standard stipend for Master's students which will be added if your application is successful.</p> <p>Grants are cash-limited at the point of award.</p>	

**Studentship fees**

Description	Total
<p>State the annual amount of approved university and college fees (if appropriate) at the UK/EC student rate in the description, and specify the total amount of fees requested.</p> <p>International fees can only be requested by applicants from low- and middle-income countries.</p>	

**Research involving human participants, human biological material and identifiable data**

Does your project involve human participants, human biological material, or identifiable/potentially identifiable data?	
<p>The following notes relating to 'Research involving human participants, human biological material and identifiable data' are intended to provide guidance and advice in completing the form, rather than a comprehensive review of the legal and regulatory environment in which the application is made.</p> <p>The World Health Organization defines research with human subjects as "any social science, biomedical, behavioural, or epidemiological activity that entails systematic collection or analysis of data with the intent to generate new knowledge, in which human beings i) are exposed to manipulation, intervention, observation, or other interaction with investigators either directly or through alteration of their environment, or ii) become individually identifiable through investigator's collection, preparation, or use of biological material or medical or other records."</p> <p>The Wellcome Trust policy position on research involving human participants can be found on the Wellcome Trust website (<a href="http://www.wellcome.ac.uk/wellcome-trust-policy-position-research-involving-human-participants">www.wellcome.ac.uk/wellcome-trust-policy-position-research-involving-human-participants</a>)</p> <p>Ethical approval (usually from the appropriate National Health Service (NHS) research ethics committees) is required for all Wellcome Trust funded research involving human participants, biological samples or personal data. Personal data, in the context of the 1998 Data Protection Act (Section 3.2, and Annex 3), comprise information about living people who can be identified from the data, or from combinations of the data and other information which the person in control of the data has, or is likely to have in future. Any use of personal data or biological samples, relating to living or dead persons, should conform to MRC guidelines available at: <a href="http://www.mrc.ac.uk/documents/pdf/personal-information-in-medical-research/">http://www.mrc.ac.uk/documents/pdf/personal-information-in-medical-research/</a> and <a href="http://www.insight.mrc.ac.uk/2015/02/12/getting-the-best-out-of-biological-samples/">http://www.insight.mrc.ac.uk/2015/02/12/getting-the-best-out-of-biological-samples/</a>.</p> <p>Approval from other regulatory bodies such as the Human Fertilisation and Embryology Authority (HFEA) or the Gene Therapy Advisory Committee in the UK should also be sought where necessary, e.g. research involving human embryos may require a licence from the HFEA (please refer to <a href="http://www.hfea.gov.uk">www.hfea.gov.uk</a> for more information). If your proposal involves research on gene therapy which requires regulatory approval, approval should be sought from your Local Research Ethics Committee, the University's Genetic Manipulation Committee, the Gene Therapy Advisory Committee and the</p>	

Medicines and Healthcare products Regulatory Agency (MHRA).

The organisation must ensure that ethical approval is in place at all relevant times during the project. For research carried out at multiple sites, ethics committee approval must cover each site.

Where the research, or part of the research, is to be performed outside the UK, independent ethics review must be obtained. For research involving people living in low and middle income countries, see the Wellcome Trust's website ([www.wellcome.ac.uk/funding/managing-grant/guidance-notes-research-involving-people-low-and-middle-income-countries](http://www.wellcome.ac.uk/funding/managing-grant/guidance-notes-research-involving-people-low-and-middle-income-countries)).

Please confirm that you have read the Trust's guidance on the feedback of health-related findings in research and that you are in the process of considering your approach to this.

The Wellcome Trust's guidelines on the feedback of health-related findings in research can be found on the Trust's website.

Please state by whom and when the ethics of the project has been, or will be, reviewed and specify any other regulatory approvals that have been obtained, or will be sought.

We reserve the right to see relevant approval documents at any point during the lifetime of the grant, in accordance with our policy position on research involving human participants.

The following notes relating to 'Research involving human participants, human biological material and identifiable data' are intended to provide guidance and advice in completing the form, rather than a comprehensive review of the legal and regulatory environment in which the application is made.

The World Health Organization defines research with human subjects as "any social science, biomedical, behavioural, or epidemiological activity that entails systematic collection or analysis of data with the intent to generate new knowledge, in which human beings i) are exposed to manipulation, intervention, observation, or other interaction with investigators either directly or through alteration of their environment, or ii) become individually identifiable through investigator's collection, preparation, or use of biological material or medical or other records."

The Wellcome Trust policy position on research involving human participants can be found on the Wellcome Trust website ([www.wellcome.ac.uk/wellcome-trust-policy-position-research-involving-human-participants](http://www.wellcome.ac.uk/wellcome-trust-policy-position-research-involving-human-participants))

Ethical approval (usually from the appropriate National Health Service (NHS) research ethics committees) is required for all Wellcome Trust funded research involving human participants, biological samples or personal data. Personal data, in the context of the 1998 Data Protection Act (Section 3.2, and Annex 3), comprise information about living people who can be identified from the data, or from combinations of the data and other information which the person in control of the data has, or is likely to have in future. Any use of personal data or biological samples, relating to living or dead persons, should conform to MRC guidelines available at: <http://www.mrc.ac.uk/documents/pdf/personal-information-in-medical-research/> and <http://www.insight.mrc.ac.uk/2015/02/12/getting-the-best-out-of-biological-samples/>.

Approval from other regulatory bodies such as the Human Fertilisation and Embryology Authority (HFEA) or the Gene Therapy Advisory Committee in the UK should also be sought where necessary, e.g. research involving human embryos may require a licence from the HFEA (please refer to [www.hfea.gov.uk](http://www.hfea.gov.uk) for more information). If your proposal involves research on gene therapy which requires regulatory approval, approval should be sought from your Local Research Ethics Committee, the University's Genetic Manipulation Committee, the Gene Therapy Advisory Committee and the Medicines and Healthcare products Regulatory Agency (MHRA).

The organisation must ensure that ethical approval is in place at all relevant times during the project. For research carried out at multiple sites, ethics committee approval must cover each site.

Where the research, or part of the research, is to be performed outside the UK, independent ethics review must be obtained. For research involving people living in low and middle income countries, see the Wellcome Trust's website ([www.wellcome.ac.uk/funding/managing-grant/guidance-notes-research-involving-people-low-and-middle-income-countries](http://www.wellcome.ac.uk/funding/managing-grant/guidance-notes-research-involving-people-low-and-middle-income-countries)).

In the course of your project, do you propose to use facilities within the National Health Service (NHS) or to involve patients being cared for by the NHS?

By agreeing to fund work which requires NHS support, the Wellcome Trust is agreeing to abide by the Statement of Partnership on Non-commercial R&D in the NHS in England (and the corresponding statements in Northern Ireland, Scotland, and Wales). Researchers must therefore meet the obligations of the Partnership and may not carry out any research until the NHS has given its consent.

The Research Governance Framework for Health & Social Care, published by the Department of Health in England can be downloaded from the Department of Health website <http://www.dh.gov.uk/health/category/research>. Please note that the Wellcome Trust cannot act as sponsor.

Is a formal sponsor required for the project, for example under the Medicines for Human Use (Clinical Trials) Regulations or the Research Governance Framework for Health and Social Care and equivalent guidance?

Please indicate which organisation(s) has/have agreed to fulfil this role. Please note that the Wellcome Trust cannot act as sponsor.

## Risks of research misuse

Please confirm that you have considered whether your proposed research could generate outcomes that could be misused for harmful purposes.

In preparing research proposals, the Wellcome Trust wishes to encourage applicants and their administering organisations to consider carefully any ethical, safety or security implications associated with the research, including any risks that the potential outcomes could be misused for harmful purposes. Such purposes would include actions which lead to harm to humans, animals or the environment - including terrorist misuse.

Where there are judged to be **tangible** (i.e. real and non-hypothetical) risks that the proposed research **will itself** generate outcomes that could be misused to cause harm, organisations should take appropriate steps to monitor the research as it proceeds and minimise these risks. The Wellcome Trust recognises that most research could conceivably generate results that might hypothetically be misused at some point in the future, and is not asking applicants to appraise these kinds of remote and hypothetical risks.

Applicants should refer to the Wellcome Trust's position statement on managing risks of research misuse [www.wellcome.ac.uk/funding/managing-grant/managing-risks-research-misuse](http://www.wellcome.ac.uk/funding/managing-grant/managing-risks-research-misuse).

Have you identified any tangible risks of this type?

Please briefly describe these risks and the steps that you and your organisation will take to manage

them  
(250 words max.)

## Freedom to operate/conflicts of interest

**Describe any freedom to operate issues or potential conflicts of interest that have been identified or that might arise and how these will be or have been addressed.**

In particular, please consider the following:

- Do any of the individuals involved in the project hold any consultancies or equities in, or directorships of, companies or other organisations that might have an interest in the results of the proposed research?
- Will the proposed research use technology, materials or other inventions that are subject to any patents or other form of intellectual property protection?
- Will any element of the research be subject to agreements with commercial, academic or other organisations, including arrangements with collaborators named in the grant application, that might lead to intellectual property issues or restrictions?

(350 words max.)

*Please describe any freedom to operate issues or potential conflicts of interest that may affect your ability to carry out the proposed research and/or to comply with the Trust's grant conditions.*

*Where the proposed research, in whole or in part, is subject to agreements with commercial, academic or other organisations, e.g. Materials Transfer Agreements, the Wellcome Trust will expect a written assurance from the administering organisation that the terms of any such agreement do not conflict with the Trust's grant conditions, particularly in relation to the publication of research and the granting of research rights.*

*Please refer to the Wellcome Trust's website for our policy on the relationship between Trust-funded researchers and commercial entities: [www.wellcome.ac.uk/funding/managing-grant/policy-relationships-between-trust-funded-researchers-and-commercial-organisations](http://www.wellcome.ac.uk/funding/managing-grant/policy-relationships-between-trust-funded-researchers-and-commercial-organisations).*

*Details of our policy on intellectual property can be found in our Grant Conditions [www.wellcome.ac.uk/funding/managing-grant/grant-conditions](http://www.wellcome.ac.uk/funding/managing-grant/grant-conditions).*

*Applicants should disclose all relevant information pertinent to their grant proposal, including proprietary information where appropriate, in order to provide the most comprehensive picture of their proposed research.*

*If no issues have been identified, please enter N/A.*

## Wellcome Trust supported facilities

Will the research project be based in one of the following Wellcome Trust supported facilities:

- the Wellcome Trust Sanger Institute
- a Wellcome Trust Centre
- an Africa and Asia Programme
- the Francis Crick Institute?

*We are interested to find out about Trust-funded research projects based in these facilities and wish to collect this data for information purposes.*

Please specify

Sample