

# CARDIOVASCULAR RESEARCH MSc



# Faculty of Life Sciences & Medicine Cardiovascular Division

# MSc in Cardiovascular Research TMSC1MTCVR Handbook for 2018/19

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# Introduction

Welcome to the MSc in Cardiovascular Research! The coming year will be challenging at times and occasionally you will find it difficult, but we trust that you will also find you will get a lot from it, that you will find it interesting, fun, and even inspirational. You will be allocated a personal Tutor, who will be able to advise you and guide you throughout the year.

### Disclaimer

This handbook was published in September 2016, and reflects the information available at that time. We have endeavoured to make this information as accurate as possible and any changes will be communicated on the Cardiovascular Research MSc site in KEATS. If there is any conflict between information on the web page and this handbook, then the web page should be taken as the authoritative source. If you should find any information that you consider to be out of date, inaccurate or misleading then please contact Dr Fraser (paul.fraser@kcl.ac.uk) with your reasons for considering the information to be inaccurate.

If you cannot find the information you require here it is likely to be in one of the other student handbooks.

Faculty Handbook.

Additional useful information relating to your programme and to student welfare can be found in the Faculty Handbook: <u>https://internal.kcl.ac.uk/lsm/students/pgt/handbook/index.aspx</u> <u>King's Student Handbook</u>

The university has an online student handbook, covering useful College level information about services for students: <u>http://www.kcl.ac.uk/aboutkings/quality/academic/myhandbook/index.aspx</u>

# **Communication and Contacts**

- → All communication about changes to the programme timetable and/or content will be sent to you via your King's email account and will also be notified on the programme's KEATS entry.
- ➔ To aid communication between members of the class a WhatsApp group will be set up early in the programme.
- → There is now a Facebook group that you are invited to join and communicate your comments.
- ➔ Two students will be elected to serve on the Staff Student Liaison Committee that meets once in each semester. The minutes of those meetings will be available on KEATS.

# **Programme Objectives**

At the end of the programme students will be able to:

- ➔ Evaluate and assimilate the scientific literature in a given subject area and to think critically about the results and methods.
- → Devise a hypothesis that can be tested experimentally.
- → Analyse data, appreciate the value of reproducibility of data and draw valid conclusions.
- → Collect data and apply appropriate methods to test a hypothesis.
- → Develop an ability to comprehend and synthesise complex information.
- → Organize a work-schedule, stick to deadlines, and prioritize activities.
- → Communicate clearly and effectively, both orally and through writing.

# Structure of the Cardiovascular Research MSc Programme

An MSc year needs to have 180 course credits cf. a BSc year of 120. This is made up of a research project of 90 cc (5-6 months), a Skills module of 30 cc, a Fundamental Cardiovascular Research Topics module of 30 cc and an Applied Cardiovascular Research Topics module of 30 cc.

Module name:		Module format:	СС
Cardiovascular Skills:	7MRV0012	Workshops & 2h Exam	30
Fundamental Cardiovascula	ır	9 Lectures & 9 Seminars	30
Research Topics:	7MRV0013	1 Essay + 3h Exam	
Applied Cardiovascular		8 Lectures & 8 Seminars	30
Research Topics:	7MRV0011	1 Essay + 3h Exam	
Cardiovascular Research		Laboratory project	
Project:	7MRV0015	Dissertation	60
	7MRV0016	Project performance	30
		Total:	180

The MSc Pass mark is 50%. A Merit is awarded for  $\ge$  60% and a Distinction for  $\ge$ 70%. Each module has to be passed.

# THE MODULES

#### CARDIOVASCULAR SKILLS (7MRV0012: 30 CC)

This module will consist of 2 elements: workshops for essential skills and advanced techniques. There are also sessions on the use of statistics and scientific interpretation that will be followed by an examination in May. A basic knowledge of statistics is required for this component, and a selfdirected teaching aid is available on KEATS for those who feel unsure of this.

**Lectures** & **Workshops** A workshop will consist of talks, with (where suitable) some interactive components such as demonstration, worked examples & hands-on experience. Currently the following topics are available: - tissue culture - Western blotting - Use of Animals – Experimental design and analysis - Scientific Interpretation - Leukocytes - Capillary Permeability - Flow cytometry – Skin – Histology - Heart failure - Artery stiffness – Animal Models of Heart Failure and Stroke – Angiogenesis – miRNA - Confocal Microscopy.

Assessment -Scientific Interpretation exam (55%: mandatory qualifying mark 40%) + 6 of the workshops (45%) assessed on a rolling basis. These 6 assessments will often consist of 4 to 6 short paragraphs in answer to questions that centre on the theoretical and practical aspects of the workshop. At least 5 of these must be passed at 50% with the 6<sup>th</sup> must be at least 40%.

#### LITERATURE BASED MODULES

#### FUNDAMENTAL CARDIOVASCULAR RESEARCH (7MRV0013: 30cc)

#### AND

#### Applied Cardiovascular Research (7MRV0011: 30cc)

These modules will address a number of topics that are the focus of current interest in the field of cardiovascular biology. The topics consist of introductory lectures to support student-led seminar sessions that consist of PowerPoint presentations of original research papers, with an internal expert present for guidance, followed by formalized question and answer sessions. The student questioners are instructed to prepare their questions in collaboration with the presenter to generate a good discussion of the science. Both the presenter and questioner will be awarded marks depending on the quality of the discussion they lead. The seminar reading lists and running order of presentations will be sent to the students by email and will be available on the programme's KEATS site.

Fundamental Cardiovascular Research	Staff
Endothelial control of vasculature;	R Siow; P Fraser
Endothelial Derived Hyperpolarization;	P Fraser; S Chapple
Capillary permeability;	P Fraser;
Leukocyte Transmigration;	A Ivetic; P Fraser
Thrombogenesis;	A Smith; P Fraser

Angiogenesis & Vascular Development; A Smith; P Fraser Control of Heart Rate & Rhythm; M Shattock; P Fraser Control of cardiac contraction; I Smyrnias; P Fraser Oxygen Sensing & Redox Signalling;

Assessment (30cc): Presentation and discussions (30%; mandatory qualifying mark 50%), 1h timed essay (15%; mandatory qualifying mark 40%) and a 3 question, 3 hour examination (55%; mandatory qualifying mark 40%) taken in May.

A Brewer; P Fraser

Applied Cardiovascular Research	Staff
Hypertension;	A Ferro; P Fraser
Atherogenesis;	C Shanahan; P Fraser
Acute coronary syndromes/MI;	D Perera; P Fraser
Heart Failure & Cardiac Remodelling;	I Smyrnias; P Fraser
Developmental origins of human disease	G Clough; P Fraser
Cardiac Regeneration – Stem Cells;	L Zeng; P Fraser
Diabetes and Cardiovascular Disease;	L Gnudi; P Fraser
Stroke;	P Fraser;
Aneurysms;	A Smith; P Fraser

Aneurysms;

Assessment (30cc): Presentation and discussions (30% core; mandatory qualifying mark 50%), 1 x 3,000 word essay after the Christmas vacation (15%; mandatory qualifying mark 40%) and on a 3 question, 3 hour examination (55%; mandatory qualifying mark 40%) taken in May.



Photo taken during the student-led seminar on aortic aneurysms in December 2016

#### **NOTES ON SEMINAR PRESENTATION**

The seminars are aimed to help you learn the subject matter by having a good discussion on and around the topic outlined by the reading list. It is important that certain ground rules are observed so that your presentations are understood and enjoyed by your audience.

Each paper should be presented in terms of:-

**AIMS** The authors will have had to justify the reasons for carrying out their experiments, you should mention these and place the work into some context.

**METHODS** The main methods should be **very** briefly outlined so that the audience can easily understand what really went on in the experiments. A simple diagram of the apparatus and/or the experimental procedures might be helpful, and is sometimes essential. **Do not be over elaborate**. **The methods for individual experiments are often best presented with the results.** 

**FINDINGS** The crux of the presentation. You should present carefully **selected** graphs, and possibly tables, to emphasise the points you (or the authors) wish to make. Any scientist publishing a paper realizes that the best way to get a message across is in terms of a graph, and often considerable care will have been taken in deciding how best to present the data. There is no need to present the full content of the paper, just those elements that you think are really important.

**DISCUSSION** Have the authors achieved their declared objectives? Are there possible alternative interpretations of their results? What is the net contribution to the subject of this piece of work? You will appreciate that you have to know the papers to a considerable depth to be able to present them well. It is not the intention that you read your notes. It is far, far better that you talk fluently to the class, and refresh your memory from time to time by glancing at a crib sheet. In that way whatever you say will be part of a conversation, and allow your personality to come across. A reading will be dry and stilted, far removed from ordinary human expression.

#### **DESIGNATED QUESTIONERS**

You must of course read the paper first! Your questions should be based on whether you think that the authors have really shown what they have claimed, could there be other alternative (and perhaps better!) explanations for their results, and even suggest a better experiment. You are encouraged to collaborate with the presenter and the others with an aim to generate as good a discussion as possible.

#### SEMINAR ASSESSMENT

Your performance in the seminars will be assessed equally on the quality of the presentation and on the quality of the discussion generated by intelligent and thoughtful questions. The presenter and the questioners are encouraged to work as a team to produce a good performance and will share the (marks) benefits of a good session. Free questions (those that are outside a designated presentation) are very much encouraged and will attract bonuses!

# CARDIOVASCULAR RESEARCH PROJECT (7MRV0015: 60 CC & 7MRV0016: 30 CC)

The project forms half of the programme and should be the equivalent of **6 months full-time work**. This means that you should spend around 4-5 months in the lab and the remaining time should be spent on reading the literature, data analysis and writing the dissertation. You are directed to view the Code of Practice for PGT Dissertations that can be found on the Faculty PGT Handbook at: <a href="https://internal.kcl.ac.uk/lsm/students/pgt/handbook/index.aspx">https://internal.kcl.ac.uk/lsm/students/pgt/handbook/index.aspx</a>

At the end of the project students should be able to:

- Design and carry out appropriate experiments to test a hypothesis.
- Interpret results and summarise main findings (conclusions).
- Carry out statistical analysis on data.
- Keep professional records of work done.
- Manage time effectively.
- Develop problem solving and trouble shooting skills.
- Have the ability to work easily and competently in a laboratory environment.
- Analyse data critically and effectively.
- Write up a research dissertation.

At the end of the project and assessment phases students should have developed research skills to a postgraduate standard.

#### **Project Supervision**

The project supervisor should spend time at the start of the project discussing the suggested topic, background reading, practical considerations, and timetabling. The supervisor will also take the necessary steps over Animal Licence, Ethical Committee permission, and safety regulations where relevant. You should ensure that you understand the details of each of these, as they have important legal and safety implications. Remember that **you are ultimately responsible for everything you do associated with the project, from your own safety, to proper attention to legal requirements over experimentation, and care of animals and human subjects.** The Home Office specifies that students must receive training in animal experimentation before applying for a Personal Licence for animal work. This training is organised by King's if necessary.

The project supervisors are asked to help in the following ways:

- encourage the student to plan the protocol and draw up the design of experiments.
- give assistance with learning how to calibrate, check and use equipment. You should understand the theory behind any apparatus used for your project work, not just its method of operation.
- give practical help initially during experiments, but thereafter encourage the student to work independently as much as possible.
- provide overall supervision of the student's work, with particular attention to regulations and safety.
- provide some key references, and suggest where to find recent information.
- give guidance on analysis and presentation of data and on the most appropriate statistical tests for the data generated.

- read and give constructive criticism of one, **and no more than two**, draft versions of the dissertation. The dissertation has to be the student's own work.
- give relevant information about the conduct of the project to the Examiners, e.g. any problems encountered, unavoidable delays, equipment faults, availability of subjects/animals, extent to which student worked alone or as part of team, extent of assistance required. Supervisors will be asked to complete report forms after reading the dissertation, and these will be forwarded to the Examiners.

#### **Project**

The starting date for the project depends on your other coursework commitments and timetable, but should be as early as possible.

#### Timetable:

Early October: Project Titles distributed. Discuss options with programme co-ordinator

Mid-October: e-mail project preference (4 projects) to the programme co-ordinator

#### (paul.fraser@kcl.ac.uk)

**Early November:** Announcement of preliminary allocation. Contact prospective supervisor and/or (if needed) discuss options with programme co-ordinator. Please provide a 1 page CV and short (no more than ½ page) statement to the programme co-ordinator <u>and</u> prospective supervisor describing why you wish to do this project.

Mid November: Confirm project choice & start literature search

**January-July 2019:** Experimental work on project. You should write up sections as you go along (Introduction, Methods, Results, Reference list), leaving only the Discussion and Abstract to complete after the end of experiments.

**End July 2019:** Draft dissertation to project supervisor for comment and correction. Your supervisor is only obliged to comment on one draft and will be instructed not to comment on more than 2 draft versions.

Mid-August 2019: Deadline for electronic submission of completed dissertation.

Early September 2019: Oral presentation – date to be confirmed

#### **Module structure**

### 1) Written dissertation

The total length of the dissertation, calculated as a word count using the "Tools/word count" option of the word-processing package **should be no more than 12,000 words** (including figures, tables and references within text but not the reference list). We will check the electronic version of your dissertation to ensure that the word count listed on the front cover sheet truly reflects the dissertation length: examiners will be asked not to read beyond the 12,000 word limit and submit their marks based on this upper limit. If your dissertation is overlong you will not gain credit for a thoughtful discussion that an examiner does not read.

A numbering system for Sections/Chapters helps. Pages should be numbered in sequence, and each Section should start on a fresh page. Figures and Tables look best if they are incorporated within the text, but they can be grouped together at the end of the appropriate Section if necessary. Each Figure or Table should have a number, a title, and a legend that describes its main features, so that it can be understood without reference to the text. In figures, show error bars (specify mean  $\pm$  SD or SEM), and *P* values, where relevant, using the convention \* <0.05, \*\* <0.01, \*\*\* <0.001. **Examiners pay particular attention to whether appropriate statistical analyses have been performed on the data – your statistics workshop should assist with this, but if in doubt ask your supervisor for advice.** In legends, give the 'n' number (specifying how many determinations of the parameter in how many treatment groups, animals or subjects). In Tables, give mean  $\pm$  SD or SEM and 'n', and *P* values where relevant.

#### The dissertation should consist of:

- Front cover sheet with details of number of words
- Title page with name of student and supervisor
- Abstract similar to a scientific paper (no more than one page) and divided into headed sections:
- Introduction & Aims; Methods; Results; Conclusion.
- Acknowledgements
- Abbreviations
- Table of contents
- List of tables and figures
- Introduction including a good review of the literature (appropriately referenced), showing brief
  historical development and recent work, and the chief questions to be addressed. Make sure that
  you give a balanced coverage, not just mentioning work from your host laboratory. If figures or tables
  are reproduced from published material, the source must be stated in the Figure legend and cited in
  the reference list.
- Aims it must be clear to the examiners what you are attempting to do in your project. It may be
  useful to give the aims as a series of bullet points with a brief outline of the approaches taken to
  address each aim.
- Methods given more fully than in a scientific paper, so that the examiners have a clear idea of what you did. It is useful to describe, in brief, the underlying principles of methods used. If you are involved in any *in vivo* experimental work on animals (even if you are not performing the procedures yourself) you MUST state the Home Office scheduled procedures that have been used. Similarly, dissertations that include work performed using human subjects MUST provide details of the ethical approval that was granted for the study.
- Results as in a paper, but you may include more examples of raw data. You should provide numerical data wherever possible, so if you obtain results in a non-quantifiable format (e.g. experimental traces, autoradiographs, etc.), you should attempt to quantify them (e.g. measurements of area under the curve, densitometric analyses, etc.). Wherever possible you should perform appropriate statistical analyses on the data obtained you should seek advice from your supervisor about the most appropriate statistical tests to apply to your data if necessary. The results text should describe the results presented, highlighting the main observations, and commenting on the statistical analyses.
- Discussion commenting intelligently and critically on the results obtained, and showing how they fit

in with the body of knowledge. Do not be afraid to criticise your own work, if you feel some parts are weak, and try to offer an explanation if your results are different from those of previous studies.

- Conclusion key points from the work, and suggestions for further study.
- Appendix the appendix contains information that you may not want to include in the main text such as large tables of raw data summarised in the Results section, checks on methods, DNA sequencing data, mathematical or theoretical considerations. You do not need to include an appendix if you do not have any additional information that has not been included in the Results section.
- References full details (title, year, journal, page numbers) of each reference cited in the text, in alphabetical order. If several different techniques are used, or different sub-projects done within the whole, it may be better to keep the sub-topics separate (e.g. by presenting Methods and Results of sub-project 1 together, then same for sub-project 2 etc.).

You will be expected to submit **two** copies of your dissertation to the Academic Centre in mid-August: the deadline will be confirmed by email in 2017. The examiners will both receive a copy that they will mark. The project supervisor who will be asked to complete an assessment form. Please note that the MSc examiners have requested that all dissertations must be submitted in a format that allows confirmation of word count and that can be scrutinised by plagiarism software if necessary. For this reason, you must submit an electronic copy of your dissertation (as well as the two paper copies). Please make sure that your electronic submission is labelled with your name, and hand it in at the same time as the paper copies of your dissertation.

### 60 cc will come from your written dissertation; mandatory qualifying mark 50%

2) Lab Performance: <u>15 cc will come from the supervisors assessment of your performance in the</u> <u>laboratory</u>; mandatory qualifying mark 50%. This will be based on your attendance, thoroughness in keeping records, ability to solve technical issues that come up and participation in laboratory and Divisional events.

#### 4) Oral Presentation

You will be expected to present a 10 minute PowerPoint presentation to provide an overview of your project followed by questions from internal and external examiners. Detailed instructions will be provided at a later stage. The Dissertation Abstract must be forwarded to the programme coordinator one week before the date of the presentation.

15 cc will come from your oral presentation; mandatory qualifying mark 40%

# Projects for 2018/19

The role of neutrophils in cardiovascular disease	Dr. Aleksandar Ivetic			
Are histone demethylases oxygen sensors?	Dr. Alison Brewer			
Seeking sense in antisense: Delineating the role of long non-coding RNA cell proliferation and apoptosis.	HIF1A-AS2 in endothelial <b>Dr. Anna Zampetaki</b>			
Investigating a novel role for the nuclear envelope LINC complex in cardi transduction	omyocyte mechano- Dr. Qiuping Zhang			
Do Vascular Abnormalities predict Acute Kidney Injury after CABG?	Prof. Cathy Shanahan			
Role of Nox4 in glycolysis of cardiac cells under stress	Dr Celio Santos			
Investigating the effect of Nrf2 on inflammation after myocardial infarction Dr. Daniel Bromage				
An experimental model of heart failure with preserved ejection fraction	Dr Min Zhang			
Deciphering the role of cyclin-dependent kinase-19 in pulmonary vascula	ar remodelling			
	Dr Olena Rudyk			
Determinants of coronary blood flow in patients with ischaemic heart failure Prof Divaka Perera				
Pathophysiology of Leaflet Thrombosis Following Trans-catheter Aortic V TAVI): An Investigational Study	/alve Implantation (PLT- Dr Tiffany Patterson			
Dietary nitrate and the nitrate-nitrite-NO pathway.	Dr Andrew Webb			
The role of inflammation in spinal cord injury following aortic aneurysm	repair			
	Prof Alberto Smith			
Activation of platelets by antiphospholipid antibodies	Prof Alberto Smith			
Comparison of the pro-angio/arteriogenic properties between adipose-d mesenchymal-derived stem cells	lerived and Prof Alberto Smith			
Stimulating the revascularising potential of monocytes isolated from pat ischaemia for use in cell therapy	ients with critical limb Prof Alberto Smith			
Can Sulforaphane afford protection against obesity and glucose intolera obese pregnancy?	nce associated with Dr Sarah Chapple			
Waveform Data Analysis.	Dr. Manasi Nandi			
Bioprinting of vascular endothelial and smooth muscle cells under oxyge vivo to improve clinical translation	n levels encountered in Prof Giovanni E. Mann			
Does p62 play an integral role in neuroinflammation?	Prof Giovanni E. Mann			
The role of Bilirubin in the regulation of ROS generation Impact of cardiovascular risk factors on trafficking IKca channels in endo	Dr Paul Fraser thelial cells			
	Dr Paul Fraser			
Mechanosensitive redox signalling in endothelial cells: Sirtuins and Nrf2 in vascular ageing				
	Dr Richard Siow			
Role of Nogo-B in distal collecting duct	Prof Luigi Gnudi			

# What past students said

"I would like to thank you for your help and guidance throughout the course. It certainly has been an enjoyable year" *Anjalee Chattersingh* 

"Thank you once again for everything this year, doing this master's has been one of the best experiences of my life." Puja Kapoor

"None of this would have been possible without your support throughout the year. You have been a fantastic course coordinator" *Nur Mousa* 

"I would like to reiterate my thanks for your guidance and support over the past year." *Ramith Gunawardena* 

"I had such a great experience throughout the year. I was motivated from the first day I entered the Waterloo campus, and people I met through the course were exceptional. I had so much genuine advice and guidance, and I am confident to say that I want to pursue my career in research in the future. Thank you very much for giving me an opportunity to study under your supervision." Sho Ito

"I really enjoyed the course. It has certainly been a valuable experience. Thank you for all the work you have done to make this course a success." Chris Seet

> This programme received 100% overall satisfaction in the Postgraduate Taught Experience Survey for the last 7 years