1. Overview

SAHMRI’s vision is to conduct inspired research that will lead to better health outcomes. In order to achieve this vision, our researchers are committed to undertaking high-quality and innovative health and medical research that can be transformed into practical benefits for patients and the community.

SAHMRI is committed to the conduct of methodologically and ethically sound research that complies with the Principles, Responsibilities and Definitions of the Australian Code for the Responsible Conduct of Research (“the Code”), and other national guidelines in relation to values and community engagement.

1.1 Principles of responsible research conduct

As outlined in the Code, the Principles (P1-P8) that are the hallmarks of responsible research conduct are:

P1 - Honesty in the development, undertaking and reporting of research
• Present information truthfully and accurately in proposing, conducting and reporting research.

P2 - Rigour in the development, undertaking and reporting of research
• Underpin research by attention to detail and robust methodology, avoiding or acknowledging biases.

P3 - Transparency in declaring interests and reporting research methodology, data and findings
• Share and communicate research methodology, data and findings openly, responsibly and accurately.
• Disclose and manage conflicts of interest.

P4 - Fairness in the treatment of others
• Treat fellow researchers and others involved in the research fairly and with respect.
• Appropriately reference and cite the work of others.
• Give credit, including authorship where appropriate, to those who have contributed to the research.

P5 - Respect for research participants, the wider community, animals and the environment
• Treat human participants and communities that are affected by the research with care and respect, giving appropriate consideration to the needs of minority groups or vulnerable people.
• Ensure that respect underpins all decisions and actions related to the care and use of animals in research.
• Minimise adverse effects of the research on the environment.

P6 - Recognition of the right of Aboriginal and Torres Strait Islander peoples to be engaged in research that affects or is of particular significance to them
• Recognise, value and respect the diversity, heritage, knowledge, cultural property and connection to land of Aboriginal and Torres Strait Islander peoples.
• Engage with Aboriginal and Torres Strait Islander peoples prior to research being undertaken, so that they freely make decisions about their involvement.
• Report to Aboriginal and Torres Strait Islander peoples the outcomes of research in which they have engaged.

P7 - Accountability for the development, undertaking and reporting of research
• Comply with relevant legislation, policies and guidelines.
• Ensure good stewardship of public resources used to conduct research.
• Consider the consequences and outcomes of research prior to its communication.

P8 - Promotion of responsible research practices
• Promote and foster a research culture and environment that supports the responsible conduct of research.

The procedures outlined below are intended to provide a framework for ensuring monitoring and compliance with the Australian Code for the Responsible Conduct of Research by SAHMRI staff, students, partners and visiting researchers.

The procedures are based on the understanding that researchers associated with SAHMRI are committed to a culture of responsible conduct and that their work enhances the good name of SAHMRI and the profession to which they belong.

2. Responsibilities of SAHMRI researchers

SAHMRI researchers are required to comply with the following:
• Australian Code for the Responsible Conduct of Research (the Code);
• SAHMRI Procedures for Research Practice and any other relevant SAHMRI policies, procedures and guidelines as well as any applicable national guidelines including those covering ethical requirements for human and animal studies, requirements for confidentiality, occupational health and safety matters, and requirements for use of potentially hazardous agents; and
• NHMRC approved standards and guidelines on the NHMRC website which include, but are not limited to:
  • National Statement on Ethical Conduct in Human Research;
  • Guidelines Approved under Section 95A of the Privacy Act 1988;
  • Guidelines Under Section 95 of the Privacy Act 1988;
  • National Principles of Intellectual Property Management for Publicly Funded Research;
  • Guidelines for Genetic Registers and Associated Genetic Material;
  • Australian Code of Practice for the Care and Use of Animals for Scientific Purposes;
• **Guidelines to Promote the Wellbeing of Animals Used for Scientific Purposes - the assessment and alleviation of pain and distress in research animals.**

2.1 Research conduct

Research activity within SAHMRI must only be undertaken following appropriate education/training human and/or animal research ethics committee approval (see section 3.12). If required, approvals for the use of genetically modified organisms, carcinogenic/toxic chemicals and human/animal stem cells must also be obtained prior to commencement of research. Research associated with SAHMRI requires Governance compliance whereby researchers must include all relevant approval documentation associated with the research activity in SAHMRI’s research management system, Pure.

Researchers should adopt methods appropriate to the aims of the research and ensure that conclusions are justified by the results. Researchers are required to ensure that the ethics principles of research merit and integrity, justice, beneficence and respect are applied to human research and that the three R’s (Replacement, Reduction and Refinement) should be considered in all aspects of research involving animals.

In accordance with the Code, breaches and/or misconduct in research will be taken very seriously by SAHMRI (see section 3.13). A breach is defined as the failure to meet the principles and responsibilities of the Code and may refer to a single breach or multiple breaches. Research Misconduct is a serious breach of the Code which is also intentional or reckless or negligent.

2.2 Aboriginal and Torres Strait Islander peoples

Researchers associated with SAHMRI are required to ensure that they acknowledge, value and engage with Aboriginal and Torres Strait Islander peoples and communities and respect their legal rights and local laws, customs and protocols.

Researchers are expected to read the Code in conjunction with the following documents:

- **Ethical conduct in research with Aboriginal and Torres Strait Islander Peoples and communities: Guidelines for researchers and stakeholders** (NHMRC 2018),
- **Keeping research on track II** (NHMRC 2018) and;
- the **Guidelines for Ethical Research in Indigenous Studies** (Australian Institute of Aboriginal and Torres Strait Islander Studies 2012).

SAHMRI Researchers are also required to conduct research in accordance with the **South Australian Aboriginal Health Research Accord**, developed by Wardliparingga. The Accord has been developed through a series of consultations with Aboriginal Elders, organisations and community members. The South Australian Aboriginal Health Research Accord sets out nine principles by which Aboriginal health research in South Australia should be conducted. The principles within the Accord apply beyond Aboriginal Health research and the relevance of them relates to all research and
activities within SAHMRI and encourages a ‘culture of excellence’ in health and medical research.

**PRIORITIES:** Research should be conducted on priorities arising from and endorsed by the Aboriginal community to enhance acceptability, relevance and accountability.

**INVOLVEMENT:** The involvement of Aboriginal people and organisations is essential in developing, implementing and translating research.

**PARTNERSHIP:** Research should be based on the establishment of mutual trust, and equivalent partnerships, and the ability to work competently across cultures.

**RESPECT:** Researchers must demonstrate respect for Aboriginal knowledge, Aboriginal knowledge systems and custodianship of that knowledge.

**COMMUNICATION:** Communication must be culturally and community relevant and involve a willingness to listen and learn.

**RECIROCITY:** Research should deliver tangible benefits to Aboriginal communities. These benefits should be determined by Aboriginal people themselves and consider outcomes and processes during, and as a result of, the research.

**OWNERSHIP:** Researchers should acknowledge, respect, and protect Aboriginal intellectual property rights and ensure transparent negotiation of intellectual property use and benefit sharing.

**CONTROL:** Researchers must ensure the respectful and culturally appropriate management of all biological and non-biological research materials.

**KNOWLEDGE TRANSLATION AND EXCHANGE:** Sharing and translation of knowledge generated through research must be integrated into all elements of the research process to maximise impact on policy and practice.

### 2.3 Diversity and equity in health and medical research

SAHMRI supports diversity and equality within the Institution and is committed to ensuring that staff, researchers and students/trainees work in an environment free from discrimination, victimisation, harassment and bullying. For more information, please refer to the SAHMRI Equality in Employment Policy.

Additionally, SAHMRI acknowledges and is committed to the NHMRC gender equity principles, including the progression and retention of women in health and medical research. To support this, SAHMRI has implemented an Institution-wide Gender Equity Strategy.
2.4 Consumer and community engagement
SAHMRI researchers are encouraged to engage and facilitate consumer and community participation in their research with a commitment to transforming research into better health outcomes. This is a priority area for SAHMRI, evidenced by the establishment of the SAHMRI Consumer Engagement Committee. SAHMRI, in collaboration with Health Consumers Alliance SA, has prepared a Consumer and Community Engagement Framework which all staff should be familiar with. In addition, researchers are expected to be familiar with the Statement on Consumer and Community Participation in Health and Medical Research (NHMRC and Consumers’ Health Forum of Australia Inc. 2002). For more information, please contact Ms Alexandra Michelmore, Senior Project Officer, Consumer Engagement.

3. Guidelines regarding specific areas of responsible research conduct

3.1 Applying for funding
Grant applications, where SAHMRI is the Administering Institution are to be submitted to the funding body by the Research Office unless alternative arrangements have been made through prior consultation with the Research Office.

It is the responsibility of the applicant to contact the Research Office at least two weeks prior to the due date of any intended grant application to allow for adequate administration. For NHMRC funding applications, a signed Grant Application Cover Sheet must be provided to the Research Office before a review of the application can be undertaken.

Letters of support, signatures, referee reports and any other additional documentation required as part of a grant application are the responsibility of the applicant to arrange and ensure that they are included in the final version of the application to be submitted to the funding body.

The applicant is responsible for obtaining consent from other researchers (chief investigators and associate investigators) to be named on a grant application, if this is required. For NHMRC funding schemes, copies of the consents are to be provided to the Research Office before an application can be submitted.

All ethical approvals and/or safety clearances that are required for the research to be undertaken are the responsibility of the applicant to arrange. The relevant documentation must be provided to the Research Office and uploaded against the successful grant in Pure.

For NHMRC funded research, the chief investigator is required to complete ethics data at award in the NHMRC research grants management system (Sapphire) before payments can commence. This information is required to be certified by the Research Office.
3.2 Reporting to research funding bodies
The lead researcher or chief investigator receiving funding for a research activity or equipment item is the person responsible for reporting to the funding body, and as otherwise required. SAHMRI researchers receiving research grant funding from the NHMRC or any other funding body (government or private sponsor) must comply with reporting processes as specified by the funding body.

3.3 Involvement of external sites
When research is being undertaken at external sites to SAHMRI, including but not limited to hospital sites, supervisors must ensure that all researchers are aware of, and comply with, the appropriate confidentiality procedures and local codes of conduct. This is particularly important for students/trainees and early career researchers. Research Governance approval, through completion of a site-specific assessment form, must be obtained and recorded in Pure.

3.4 Ethical review of research applications
SAHMRI staff undertaking research involving animal experimentation, human participants (including surveys for research purposes), genetically modified organisms, carcinogenic/toxic chemicals or human or animal stem cells must have their research approved and monitored by a relevant authorised regulatory body. Additionally, the research activity cannot commence until the researchers involved have undertaken any required training.

In accordance with the provisions and processes identified by the relevant authorised regulatory body, all SAHMRI researchers are expected to:

- obtain relevant consents and approvals, including variations to research protocols, and maintain evidence of such approvals;
- comply and adhere with any relevant legislation, guidelines, conditions, policies and directives;
- comply with the requirement for monitoring and reporting on their research activities;
- provide relevant information about their research and research data, including progress reports, as requested; and
- notify the relevant authorised body immediately of any adverse events or experiences in their research that may be of concern in respect of ethical or safety matters.

Research Governance compliance requires all relevant approval documentation associated with the research activity be deposited into SAHMRI’s research management system, Pure. If researchers or other staff are unaware how to do this, they are to contact the Research Office.

Research must be approved and monitored by a relevant authorised body, as outlined below:

- proposals involving experimentation with animals on-site at SAHMRI are to be submitted to the SAHMRI Animal Ethics Committee. Where the research is conducted at a site external to SAHMRI, approval must be sought from the relevant Animal Ethics Committee;
proposals involving the participation of human subjects, or the use of human
tissue or blood, or those that involve participation of or may impact on
Indigenous peoples, are to be submitted to a relevant authorised Human
Research Ethics Committee;
proposals involving the use of clinical drugs in research trials must be submitted
to a relevant authorised regulatory Clinical Drug Trials Committee;
proposals involving biosafety matters including experimentation involving
 genetic modification or work with a genetically modified organisms;
carcinogens; toxic substances; and infectious substances; must be submitted
to the SAHMRI Biosafety Committee;
proposals involving radiation, where appropriate, must be submitted to the
SAHMRI Radiation Safety Officer; where a proposal may involve research with
significant hazards advice should be sought from the SAHMRI Director of
Research Support Services and, where appropriate, a relevant authorised
regulatory body.

3.5 Management of research materials, data and records
Researchers involved in any SAHMRI research activity must ensure that all laboratory
books and/or records, primary materials, research data (including electronic data) are
clear, accurate, complete, and retained in a safe, secure and appropriately referenced
form.

All researchers associated with SAHMRI research activities are expected to:
- Record their research (including experimental plan, design, data, results and
  outcomes) in an approved laboratory book. Research records should be
  reviewed and checked on a regular basis by the laboratory manager or
  supervisor;
- Record the location of the laboratory book/s, research data and primary
  materials (hard and electronic copies);
- Maintain and retain the laboratory book/s, research data and primary materials;
- Manage the storage of research data, data sources including any approvals
  granted, and primary materials;
- Capture and maintain their research activity in Pure for reporting purposes; and
- Maintain confidentiality of research data and primary materials, if required.

Whenever possible, original materials and data must be retained within the SAHMRI
Theme or Group in which they were generated and should be held for sufficient time
to allow access and reference. The process for the appropriate and safe disposal of
research data and primary materials when the specified period of retention has
finished should be decided in-conjunction with the relevant Research Manager.

When data are obtained from limited access databases, or via a contractual
arrangement, written indication of the location of the original data, or key information
regarding the database from which it was collected, must be retained by the
researcher and the responsible SAHMRI Theme or Group. Clinical trial data accessed
at SAHMRI must be in accordance with the SA Health Data Retention Policy and the
SAHMRI Information Security Management System Policy.
Where confidentiality provisions apply (for example, where the researchers or Institution have given undertakings to third parties, such as the subjects of the research), data must be kept in a way that reference to them by third parties can occur without breaching such confidentiality.

### 3.6 Supervision of research students and/or trainees
Supervisors must ensure appropriate induction and ongoing guidance, training/education and mentorship is provided to staff and students/trainees on the Code, SAMHRI’s principles for research practice, the management of research materials/data, records, applicable guidelines including those covering ethical guidelines and approvals for research studies, requirements for confidentiality, work health and safety matters, and requirements for the use of potentially hazardous agents. Where appropriate, supervisors may monitor staff and students/trainees conduct.

### 3.7 Intellectual property
Please refer to the SAHMRI Intellectual Property Policy (Policy Number 025).

### 3.8 Publication and dissemination of research findings
SAHMRI encourages its researchers to publish and disseminate their research to allow access by other researchers and the wider community. Publication and dissemination includes but is not limited to journal articles, correspondences, public announcements, conference and/or seminar presentations, advertising materials, research reports or other materials.

All reasonable steps must be taken to ensure that research findings, statistics, cited references, acknowledgements and public statements about research activities are complete, accurate and unambiguous. Where necessary, researchers are to take action to correct the record in a timely manner.

In order that SAHMRI’s achievements and investment in research receive appropriate acknowledgement and contribute to relevant measures of performance, SAHMRI is to be attributed as the affiliated Institution by all SAHMRI staff and students/trainees. This should be the case when the work was conducted at SAHMRI or elsewhere, even if the author has subsequently left the Institution. Joint publications of collaborative research projects must acknowledge all relevant institutions/organisations.

For publication and dissemination of research activities funded by the NHMRC, the acknowledgement requirements are outlined in the NHMRC Funding Agreement, Clause 20. Clause 20.2 indicates:

*Any material published in respect of a Research Activity must:*
  a. include the Grant Identification Number for the Research Activity; and
  b. specify that the contents of the published material are solely the responsibility of the Administering Institution, a Participating Institution or individual authors and do not reflect the views of NHMRC.

Clauses 20.3 and 20.4 indicate that the NHMRC logo must not be used without NHMRC’s prior written consent to that specific use of the logo, which the NHMRC my
give, refuse or revoke in its absolute discretion. This includes its font, colour, size and placement.

Published journal articles and other research outputs must be deposited into Pure (SAHMRI's research management system) so that they are accessible on the Pure Portal (which also serves as SAHMRI's open access institutional repository). In most cases this will be done automatically, but it is the responsibility of the researcher to ensure that all of their research outputs are captured in Pure as soon as possible following the date of publication.

3.9 Authorship
SAHMRI adopts the principles of the Code for attribution of authorship in the publication of research findings. Researchers should have made a significant contribution to the research and its output to be included as an author. All named authors must have agreed to be listed as an author. Authors should ensure that others who have contributed to the work are recognised in the research output and properly acknowledged. Individuals and/or organisations that provided facilities that contributed to the research activity should also be acknowledged in publication outputs.

When there is more than one author of a research output, one author should be nominated as executive author for the whole research output, and should take responsibility for record keeping regarding the research output.

Any part of an article critical to its main conclusion should be the responsibility of at least one author. An author’s role in a research output should be sufficient for that person to take public responsibility for at least that part of the output in the person’s area of expertise.

Disputes over authorship may arise (for example, regarding conflict of interest or authorship order). Where researchers are unable to reach mutual agreement on an issue of authorship, the following procedures may apply:

1. Any person involved in the dispute may seek advice from the Designated Person (Dr Leanne Sutherland) or Post-Graduate Student Coordinator (if involving Higher Degree Research students). A record of the dispute will be made in the Authorship Dispute Register, maintained by the SAHMRI Research Office.

2. Continuing disputes over authorship may be referred to the SAHMRI Theme Leader of the corresponding author for attempted resolution. Disputes involving co-authors from other institutions are to be handled by the institution of the corresponding author.

3. If the dispute remains unresolved after 30 days of acknowledgement, it may be referred to the SAHMRI Executive Director for determination. The Executive Director may engage an external arbitrator or mediator to assist in this process, although the final decision remains with the Executive Director.

4. Any determination made as part of a dispute resolution will be recorded in the Authorship Dispute Register but will not be considered as grounds for findings of research misconduct. However, proceeding to publication without agreement or formal determination of authorship following an acknowledged
dispute may be considered a breach of the Code or a case of research misconduct.

3.10 Peer review
SAHMRI recognises the importance of peer review and encourages its researchers to participate in the peer review process. Peer review should be undertaken and contributed to in a fair, rigorous and timely manner which maintains the confidentiality of the content.

3.11 Conflict of interest
Researchers have an obligation to disclose to their Manager and the Chief Operating Officer any conflict of interest which has the potential to influence research activities and/or outputs, this includes but is not limited to publications (for example, in authorship), public announcements, grant applications and applications for appointment and promotion. These may be a personal, financial or other interest which may represent an actual, potential or perceived conflict of interest. For more information on conflict of interest, please refer to the SAHMRI Code of Conduct Policy.

3.12 Confidentiality
If data of a confidential nature are obtained, for example from individual patient records or from certain questionnaires, confidentiality must be observed and researchers must not use such information for their own personal advantage or that of a third party. Confidentiality may also be necessary for a limited period in the case of contracted research or of non-contractual research, which is under consideration for patent protection. Confidentiality agreements may be required between SAHMRI, other Institutions, a researcher and/or a funding body of the research.

4. Research integrity
All SAHMRI staff, students, partners and visiting researchers must comply with the Code and associated SAHMRI Research Policies in relation to research integrity. Staff are encouraged to contact the SAHMRI Designated Officer in the first instance to discuss any concerns regarding a breach of the Code and/or research misconduct, please see the flowchart below for further detail. A concern, complaint or allegation may be made in writing or orally to the Designated Officer, confidentiality will be maintained throughout this process. Conflicts of interest must also be reported and will be managed.

In accordance with the Code, research breaches and/or research misconduct in will be taken very seriously by SAHMRI. A breach is considered to be the failure to meet the principles and responsibilities of the Code and may refer to a single breach or multiple breaches. Research Misconduct is a serious breach of the Code which is also intentional or reckless or negligent.

Concerns or allegations regarding breaches of the Code or research misconduct may include, but are not limited to, the following:
- fabrication of data: claiming results where none has been obtained;
• falsification of data, including changing records;
• plagiarism, including the direct copying of textual material, the use from other people without adequate attribution;
• inclusion of inaccurate or misleading information relating to a research activity;
• misleading ascription of authorship, including the listing of authors without their permission, attributing work to others who have not in fact contributed to the research, and the lack of appropriate acknowledgment of the work of a student/trainee or associate;
• other practices that seriously deviate from those commonly accepted within the research community for proposing, conducting or reporting research;
• an infringement of SAHMRI’s research policies and procedures on research conduct or other research related policies that is either intentional or caused by negligence;
• negligence, or failure to uphold commonly accepted standards in the conduct of research within the relevant field of research.

**Designated Officer**
SAHMRI has appointed a senior staff member from the Research Office as the Designated Officer. Any allegation of a breach of the Code and/or research misconduct should be made to the Designated Officer who will obtain the information required to undertake a preliminary assessment. The preliminary assessment will be undertaken by the Designated Officer and Assessment Officer/s within the Research Office. The outcome of the assessment will then be discussed with the Research Integrity Advisor and the appropriate next steps determined based on whether the allegation can or cannot be dismissed.

**Dr Leanne Sutherland** is the SAHMRI Designated Person.

**Research Integrity Advisor**
The Research Integrity Advisory can provide confidential advice to staff, students/trainees or other persons about:
• actions that might constitute a breach of the Code and/or misconduct;
• the rights and responsibilities of the complainant; and
• the procedures that will apply in the handling of allegations of breaches of the Code and/or misconduct at SAHMRI.

**Professor Maria Makrides** is SAHMRI’s Advisor in Research Integrity.

All records of complaints and/or allegations of a breach of the Code and/or research misconduct and any related correspondence (internal and external) will be securely located within the SAHMRI Research Office. Privacy in all matters will be upheld in accordance with the Guidelines Approved under Section 95A of the Privacy Act 1988 (2001) and the Guidelines Issued under Section 95 of the Privacy Act 1988 (2000).

An individual develops suspicion of a breach of the Code and/or research misconduct

Individual contacts the Designated Officer in the SAHMRI Research Office to raise their concern.

Designated Officer informs the Research Integrity Advisor and provides any information that the complainant has presented about the matter. Together they determine whether the complaint relates to a potential breach.

If a breach is identified and an investigation is to be undertaken:

Designated Officer and Assessment Officer undertake a preliminary assessment and determine:
- the need for further investigation;
- if the issue can be resolved without need for investigation;
- if the issue can be referred to other institutional processes; or
- if it is to be dismissed.

Designated Officer, Assessment Officer, and Research Integrity Advisor convene an internal investigation panel and make the appropriate notifications.

If research misconduct is established:

Designated Officer and Research Integrity Advisor will discuss with Responsible Executive Officer who will advise the person against whom the allegation was made and inform what disciplinary actions are to be taken.

If that person contests the finding of the internal investigation panel:

Designated Officer will refer the matter to an independent external research misconduct panel and will ensure that procedural fairness is afforded to the person against whom the allegation has been made.

External panel makes a decision taking into account results of internal finding plus submissions put by person against whom finding is made.

Decision communicated to the Executive Director, person who made allegation, person against whom allegation made, any funding body, SAHMRI Board.

If research misconduct is established:

External panel will make a recommendation as to the appropriate course of action which will be implemented by the Responsible Executive Officer.

Confidentiality will be maintained throughout this process. Conflicts of interest must also be reported and will be managed. Further detail regarding all components of this process can be found within the SAHMRI Research Procedures Policy.
Institutional Roles - Managing and Investigating Potential Breaches of the Code

<table>
<thead>
<tr>
<th>Role</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Responsible Executive Officer</strong> – Steve Wesselingh</td>
<td>A senior officer in an institution who has final responsibility for receiving reports of the outcomes of processes of assessment or investigation of potential or found breaches of the Code and deciding on the course of action to be taken.</td>
</tr>
<tr>
<td><strong>Designated Officer</strong> – Leanne Sutherland</td>
<td>A senior professional or academic institutional officer or officers appointed to receive complaints about the conduct of research or potential breaches of the Code and to oversee their management and investigation where required.</td>
</tr>
<tr>
<td><strong>Assessment Officer</strong> – Sarah Lawson/Yvette Growden</td>
<td>A person or persons appointed by an institution to conduct a preliminary assessment of a complaint about research.</td>
</tr>
<tr>
<td><strong>Research Integrity Advisor</strong> – Maria Makrides</td>
<td>A person or persons with knowledge of the Code and institutional processes nominated by an Institution to promote the responsible conduct of research and provide advice to those with concerns or complaints about potential breaches of the Code.</td>
</tr>
<tr>
<td><strong>Research Integrity Office</strong> – SAHMRI Research Office</td>
<td>Staff with responsibility for management of research integrity at an Institution.</td>
</tr>
<tr>
<td><strong>Review Officer</strong> – Collette Ordish</td>
<td>A senior officer with responsibility for receiving requests for a procedural review of an investigation of a breach of the Code.</td>
</tr>
</tbody>
</table>