

# **HSC Supplement to UNM Faculty Handbook Policy E40: Research Misconduct**

Title: HSC Supplement to UNM Faculty Handbook Policy E40: Research Misconduct							
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Owner(s) (Name and Title): Richard S. Larson, MD, PhD. Executive Vice Chancellor Vice Chancellor for Research	Revision Date:	Applies To: See applicability					

#### **PURPOSE**

The University of New Mexico's current Faculty Handbook Policy E40: Research Misconduct (FHB Policy E40), revised in 2002 and approved by the UNM Board of Regents on April 13, 2004, predates the issuance of the current Public Health Service (PHS) regulation (42CFR Part 93) dated June 16, 2005. FHB Policy E40 (Section 8.7) provides that any amendment to the Federal requirements in addressing research misconduct shall supersede the relevant portions of the UNM policy. UNM is committed to taking the appropriate steps to address the necessary updates to the FHB Policy E40 to meet the requirements of the current PHS regulations. However, given the time involved in addressing updates and obtaining approval of a Faculty Handbook policy, UNM HSC has implemented this supplement to the FHB Policy E40 to ensure UNM HSC compliance with the current PHS regulations. Although the UNM HSC remains governed by the University's policies, it has the authority to implement additional or more restrictive policies to meet the needs of its operations and all federal laws and regulations.

#### **APPLICABILITY**

FHB Policy E40, along with this supplement, are intended to carry out UNM HSC's responsibilities under the PHS regulations on Research Misconduct, 42 CFR Part 93. FHB Policy E40 and this supplement apply to allegations of research misconduct (as defined in FHB Policy E40), or in reporting research results involving:

- any individual who, at the time of the alleged research misconduct, was employed by, was an agent of, or was affiliated by contract or agreement with this institution; including, but not limited to, faculty, graduate/undergraduate students, staff, employees, contractors, visiting scholars, and any other member of the University's academic community and
- one or more of the following:
  - (1) PHS supported or non-PHS supported biomedical or behavioral research, research training or activities related to that research or research training, such as the operation of tissue and data banks and the dissemination of research information, (2) applications or proposals for PHS support or non-PHS support for biomedical or behavioral research, research training or activities related to that research or research training, or (3) plagiarism of research records produced in the course of research, research training or activities related to that research or research training. This includes any research proposed, performed, reviewed, or reported, or any research record generated from that research, regardless of whether an application or proposal resulted in a grant, contract, cooperative agreement, or other form of support.

These policies and procedures do not apply to authorship or collaboration disputes and apply only to



allegations of research misconduct that occurred within six (6) years of the date the institution or HHS received the allegation, subject to the subsequent use, health or safety of the public, and grandfather exceptions in 42 CFR § 93.105(b).

#### **POLICY STATEMENT**

This UNM HSC supplemental policy addresses omissions or areas that require additional clarification in *FHB Policy E40* in order to meet the current PHS regulations at 42 CFR Part 93. Section numbers refer to sections of *FHB Policy E40*. *Only sections requiring modifications or additions are listed*.

### 1. INTRODUCTION AND SCOPE

- Change title of section 1. From "INTRODUCTION AND SCOPE" to "INTRODUCTION"
- Eliminate last paragraph of section 1. INTRODUCTION AND SCOPE
- Address "scope" in new section titled APPLICABILITY (see below)

### 2. APPLICABILITY (new section)

FHB Policy E40, along with this supplement, are intended to carry out UNM HSC's responsibilities under the PHS Policies on Research Misconduct, 42 CFR Part 93. *FHB Policy E40* and this supplement apply to allegations of research misconduct (as defined below),, or in reporting research results involving:

- any individual who, at the time of the alleged research misconduct, was employed by, was
  an agent of, or was affiliated by contract or agreement with this institution; including, but
  not limited to, faculty, graduate/undergraduate students, staff, employees, contractors,
  visiting scholars, and any other member of the University's academic community and
- one or more of the following
  - (1) PHS supported or non-PHS supported biomedical or behavioral research, research training or activities related to that research or research training, such as the operation of tissue and data banks and the dissemination of research information, (2) applications or proposals for PHS support or non-PHS support for biomedical or behavioral research, research training or activities related to that research or research training, or (3) plagiarism of research records produced in the course of research, research training or activities related to that research or research training. This includes any research proposed, performed, reviewed, or reported, or any research record generated from that research, regardless of whether an application or proposal resulted in a grant, contract, cooperative agreement, or other form of support.

These policies and procedures do not apply to authorship or collaboration disputes and apply only to allegations of research misconduct that occurred within six years of the date the institution or HHS received the allegation, subject to the subsequent use, health or safety of the public, and grandfather exceptions in 42 CFR § 93.105(b).

### 3. GENERAL PRINCIPLES

- Replace section 3.3 with the following revised language:
  - 3.3 All faculty and staff will report observed, suspected, or apparent research misconduct in accordance with section 4.1 of this policy. Allegations may be made in writing, orally or anonymously and in all cases, must be sufficiently credible and specific. If an individual is unsure



whether a suspected incident falls within the definition of research misconduct, he or she may meet with or contact the Vice Chancellor for Research or HSC Research Integrity Officer (RIO) to discuss the suspected research misconduct informally, which may include discussing it anonymously and/or hypothetically. A copy of this policy shall be made available to the complainant.

- Add the following provision to section 3:
  - 3. 8. The institution will respond to each research misconduct allegation in a thorough, competent, objective and fair manner.
  - 3. 9. UNM HSC will ensure its deans, directors, chairs, and graduate advisors are reminded annually of the institution's policies and procedures on Research Misconduct including *FHB Policy E40* and this UNM HSC Supplement to *FHB Policy E40*. The HSC will also inform all faculty, students, and staff of (1) the need and importance of research integrity and (2) the importance of compliance with these policies and procedures.

#### 4. PRELIMINARY ASSESSMENT OF ALLEGATIONS

- Replace section 4.3 with the following revised language:
  - 4.3 Upon receiving an allegation of research misconduct, the Vice Chancellor for Research, or designee, shall conduct a preliminary assessment within seven (7) working days. The purpose of the preliminary assessment is to determine whether the allegation (1) is sufficiently credible and specific so that potential evidence of research misconduct may be identified, (2) whether the allegation falls within the definition of research misconduct and (3) whether it is within the jurisdictional criteria of this policy. An inquiry must be conducted if these criteria are met.

In conducting the preliminary assessment, the complainant, respondent, or other witnesses need not be interviewed and data need not be gathered beyond any that may have been submitted with the allegation, except as necessary to determine whether the allegation is sufficiently credible and specific so that potential evidence of research misconduct may be identified.

## 5. INQUIRY

• Replace section 5.2 with the following revised language:

# 5.2 Securing Research Records:

Prompt securing of the research records is in the best interest of both the respondent and UNM HSC. After determining that an inquiry will occur, the Vice Chancellor for Research will direct a process to obtain custody of all the research records and evidence needed to conduct the research misconduct proceeding, inventory the records and evidence and sequester them in a secure manner, except that where the research records or evidence encompass scientific instruments shared by a number of users, custody may be limited to copies of the data or evidence on such instruments, so long as those copies are substantially equivalent to the evidentiary value of the instruments. Sequestration of research records must occur on or before the date on which the respondent is notified of the allegation.



#### 6. INVESTIGATION

• Replace first sentence of section 6.4 Investigation Process with the following:

The Investigation Committee will pursue diligently all significant issues and leads discovered that are determined relevant to the investigation, including any evidence or additional instances of possible research misconduct, and continue the investigation to completion.

• Replace section 6.5 Investigation Report, paragraph 4 (beginning "The respondent will...") with the following revised language:

The respondent shall be given a copy of the draft investigation report for comment and, concurrently, a copy of, or supervised access to the evidence on which the report is based. The respondent will be allowed thirty (30) days from the date he/she received the draft report to submit comments. The respondent's comments must be included and considered in the final report. The complainant may be provided with those portions of the draft investigation report that address the complainant's role and opinions in the investigation, and the complainant will have thirty (30) days to submit any comments to the investigation committee. The report may be modified, as appropriate, based on the complainant's comments.

• Replace 6.6 Institutional Review and Determination, paragraph 4 with the following revised language:

Respondents may appeal the final determination to the University President. An appeal is limited to: (1) a claim of procedural error; and/or (2) a claim that the sanction imposed as a result of a finding of research misconduct is inappropriate.

• Replace 6.6 Institutional Review and Determination, paragraph 5 with the following revised language:

Except as to PHS funded research, the investigation shall be completed within 180 days of the first meeting of the Investigation Committee. However, for PHS sponsored research, unless an extension has been granted, the institution must submit the following to ORI within 120 days of the first meeting of the Investigation Committee: (1) a copy of the final investigation report with all attachments; (2) a statement of whether the institution accepts the findings of the investigation report; (3) a statement of whether the institution found misconduct and, if so, who committed the misconduct; and (4) a description of any pending or completed administrative actions against the respondent.

#### 8. OTHER CONSIDERATIONS

• Replace section 8.1.5 with the following language:

ORI shall be notified immediately, at any time during a research misconduct proceeding, if there is any reason to believe that any of the following conditions exist:

1. Health or safety of the public is at risk, including an immediate need to protect human or animal subjects;



- 2. HHS resources or interests are threatened;
- 3. Research activities should be suspended;
- 4. There is a reasonable indication of possible violations of civil or criminal law;
- 5. Federal action is required to protect the interests of those involved in the research misconduct proceeding;
- 6. The research misconduct proceeding may be made public prematurely and HHS action may be necessary to safeguard evidence and protect the rights of those involved; or
- 7. The research community or public should be informed.
- Replace section 8.5 Record Retention with the following language:

#### 8.5 Record Retention:

Records of the research misconduct proceeding will be maintained in a secure manner for 7 years after completion of any proceeding by the institution involving research misconduct allegation, or the completion of any ORI proceeding involving the allegation of research misconduct, whichever is later, unless custody of the records has been transferred to ORI or ORI has advised that the records no longer need to be retained. When it is determined that an investigation is not warranted, detailed documentation of the inquiry must be retained for at least 7 years after termination of the inquiry, so that ORI may assess the reasons why the institution decided not to conduct an investigation.

- Change sub-heading of section 8.3 from "Interim Administrative Action" to "Administrative Action"
- Add the following provision to section 8.3:

UNM HSC Officials shall ensure that administrative actions taken by the institution and ORI are enforced and shall take appropriate action to notify other involved parties such as sponsors, law enforcement agencies, professional societies, and licensing boards, of those actions.

#### **DEFINITIONS**

See UNM Faculty Handbook Policy E40

### **REFERENCES**

UNM Faculty Handbook Policy E40 PHS regulations at 42 CFR Part 93

### **RESPONSIBLITY**

This supplemental policy applies to any individual who, at the time of the alleged research misconduct, was employed by, was an agent of, or was affiliated by contract or agreement with the University of New Mexico Health Sciences Center (UNM HSC); including, but not limited to, faculty, graduate/undergraduate students, staff, employees, contractors, visiting scholars, and any other member of the UNM HSC.



## **RESOURCES AND TRAINING**

Resource/Department	Contact Information		
Vice Chancellor for Research	Richard S. Larson, MD, PhD  rlarson@salud.unm.edu  505-272-6950		
Research Integrity Officer	Catherine Penick <a href="mailto:cpenick@salud.unm.edu">cpenick@salud.unm.edu</a> 505-272-6950		
Compliance Hotline and Online Reporting	HSC Compliance Hotline 1-888-899-6092. Anonymous online reporting EthicsPoint		
Deans and Department Chairs	Consult UNM Directory		

## **DOCUMENT APPROVAL & TRACKING**

Item	Contact	Date		Approval		
Owner	Richard S. Larson, MD, PhD, Vice Chancellor for Research					
Consultant(s)	[Name, Title]					
Recommender(s)			N/A			
Committee(s)	Research Strategic Planning Committee		Yes			
HSC Legal Office	Ariadna Vazquez, Esq. Associate University Counsel	July 9, 2015		Yes		
Official Approver	Paul B. Roth, MD, MS Chancellor for Health Sciences CEO, UNM Health System Dean, School of Medicine			Yes		
Official Approver Signature	Date: July 9, 2015					
2nd Approver						
2nd Approver Signature (Optional)		Date:				
Policy Origination Date: 7/9/2015						

## **ATTACHMENTS**

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