# **UNIVERSITY COLLEGE DUBLIN**



**UCD Internal Audit** 

Research Ethics Compliance Review (Animals)
Internal Control Questionnaire

No.	Audit Objective	Audit Test Questions / Procedures	Response / Result / Finding (Attach documentation if relevant)
1.	Awareness of Policies and Guidelines		
	Audit Objective is to confirm that staff and students have informed themselves of the relevant AREC Policies and Guidelines.	Have you read:  AREC 1 Operating Procedures and Guidelines  AREC2 Policy on the Use of Animals in Research and Teaching  AREC5 Policy on Breeding and Maintenance of GM Animals  AREC7 Policy on the Use of Post-Mortem Tissue from Animals	

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No.	Audit Objective	Audit Test Questions / Procedures	Response / Result / Finding (Attach documentation if relevant)
2.	Ethical Approval and Licences		
	Audit Objective is to confirm that all necessary ethical approvals and DOH licences are in place.	Obtain a copy of all ethical approvals.  Obtain a copy of any requests for amendments, revisions or extensions and check that these were approved by AREC.  Obtain all licences received from DOH in	
		relation to the work and check that all staff and students carrying out such work have a licence or are included on the licence.	

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No.	Audit Objective	Audit Test Questions / Procedures	Response / Result / Finding (Attach documentation if relevant)
3.	Training		
	Audit Objective is to ensure that all staff and	Have all staff and students performing	
	students are signed off as being competent to	licensed procedures attended a relevant	
	perform licensed procedures on animals before	training course (e.g. animal handling).	
	carrying them out unsupervised.	Obtain evidence of attendance /	
		certification.	
		Are these attendance / certification	
		records accessible in the event of an	
		inspection by DOH officials.	

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No.	Audit Objective	Audit Test Questions / Procedures	Response / Result / Finding (Attach documentation if relevant)
4.	Severity Banding		
	Audit Objective is to confirm the severity banding of animal work is kept under review to ensure that it remains within the limits approved / licensed for each research project.	Discuss with team leader.	

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No.	Audit Objective	Audit Test Questions / Procedures	Response / Result / Finding (Attach documentation/maps if relevant)
5.	Application of Humane Endpoints		
	Audit Objective is to confirm that records are	Discuss with team leader to establish if	
	maintained to ensure that any humane endpoints	records are maintained.	
	are applied in accordance with AREC recommendations.	Obtain copies of any records.	

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No.	Audit Objective	Audit Test Questions / Procedures	Response / Result / Finding (Attach documentation if relevant)
6.	Records for DOHC Licensing Regulations		
6.	Records for DOHC Licensing Regulations  Audit Objective is to ensure that information required under DOHC Licensing Regulations for each procedure is recorded and accessible.	Check from a sample of licensed procedures whether the following records, which are required under DOH licensing conditions, have been maintained:  a) the date on which the experiment was performed; b) the address of the establishment where the experiment was performed; c) the nature, purpose and duration of the experiment; d) the number of animals used in the experiment; e) the species and strain of each animal used in the experiment; f) whether the animal was obtained from a breeding establishment, a supplying establishment or another source; g) whether anaesthesia, analgesic or another method was used to relieve the animal's pain, distress or suffering and, if so, the type of anaesthesia, analgesic or method; h) whether the animal had been used in a previous experiment; i) whether the animal was kept alive, set free or killed at the end of the experiment;	
		j) if the animal was killing the final disposition of the carcass.	

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No.	Audit Objective	Audit Test Questions / Procedures	Response / Result / Finding (Attach documentation if relevant)
7.	Reporting of Adverse Events		
	Audit Objective is to ensure that any adverse	Discuss with team leader to ascertain if	
	events in the course of the animal research have	there have been any adverse events and, if	
	been reported to AREC.	so, whether these adverse events were	
		reported to AREC.	
		Obtain copy of any correspondence with	
		AREC in relation to adverse events.	
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