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Comparison of Library Publishing Workflows by Open Access Model

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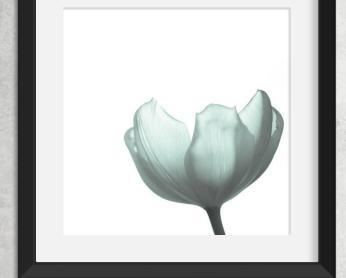
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Comparison of Library Publishing Workflows by Open Access Model

Sue Ann Gardner CHALLENGES OF CONTEMPORARY PUBLISHING CONFERENCE Lublin, Poland 2023.05.25



Library publishing

Population of an institutional repository is a form of publishing

PDF-based publishing: Workflow steps

Manuscript Intellectual property Editing Review and develop content Copyedit Proofread

Production Typeset Graphics Identifiers/Metadata

Published work



Publishing: Type of activity

Human relations Legal Quality assurance Production | Discoverability Post-production

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PDF-based publishing: Types of activity

Human relations	Legal	Quality assurance	Production Discoverability	Post- production
Acquire content	Contract	Develop content	Typeset text	Marketing
Develop content	Determine permissions	Fact check	Add graphics	Preservation
Review content	Select license	Copyedit	Lay out pages	Metrics
	Graphics permissions	Correct proofs	Metadata/ identifiers	
			Create other formats	

PDF-based publishing: Workflow steps

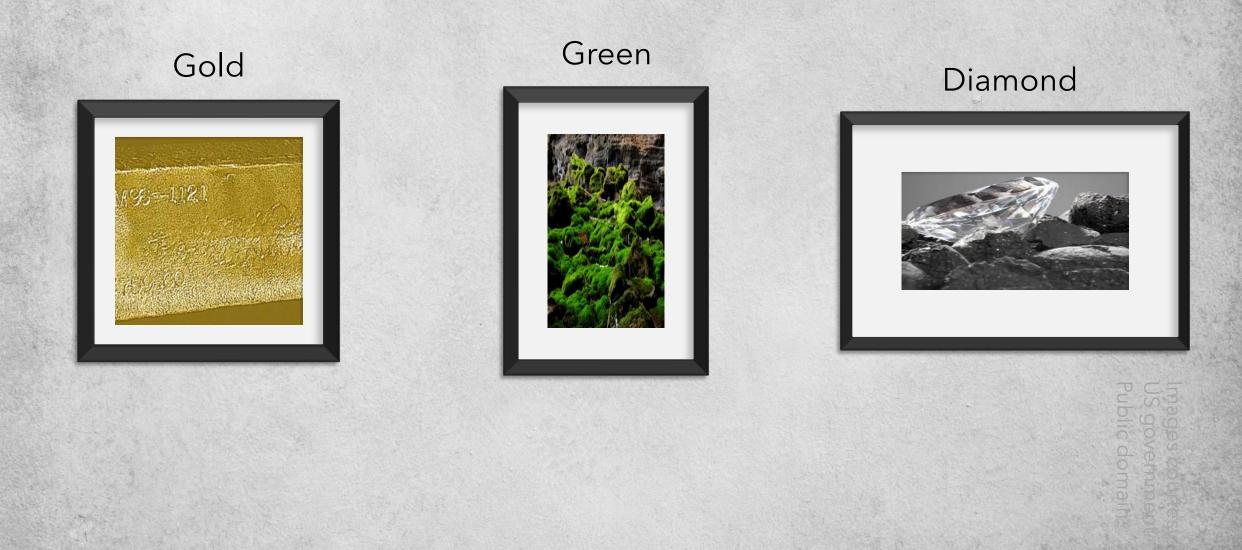
Publishing activity	Type of activity
Acquire content	Human relations
Determine permissions	Legal
Choose license	Legal
Contract	Legal
Develop and review manuscript	Quality assurance Human relations
Copyedit	Quality assurance
Typeset text	Production
Select and/or create graphics	Production Legal
Lay out pages (text + graphics)	Production
Make final corrections	Quality assurance
Generate completed text	Production
Generate and affix identifiers	Production Discoverability
Post electronic version/Metadata	Production Discoverability
Create other formats/instances	Production
Marketing	Post-production
Preservation	Post-production
Capture and analyze use-metrics	Post-production



Comparison of Workflows

Gold | Green | Diamond

University of Nebraska-Lincoln Libraries **Open access models, shorthand**



Open access models, working definitions at UNL

Gold open access (OA) means scholarship that is **made available open immediately** after publication in a commercially-published journal (may be for-profit or not-for-profit) after paying an **open access charge**

Green OA means **republishing** work that was published previously and is available often behind a paywall (but not always) and is then **put into our repository** in a copyright-legal form

Diamond OA means work that we **publish originally** in the University of Nebraska–Lincoln Libraries

These are not standard definitions but help me discuss OA models with authors.

GOLD Original publishing	GREEN Republishing	DIAMOND Original publishing
Advise and guide author	Acquire content (post-print manuscript)	Acquire content (original manuscript)
	Determine permissions	Choose permissions and license
		Develop and review manuscript
		Copyedit
	Typeset text	Typeset text
		Select and/or create graphics
	Lay pages out (text + graphics)	Lay pages out (text + graphics)
	Make final corrections	Make final corrections
	Add identifiers and metadata	Add identifiers and metadata
Post/point to content online	Post content online	Post content online
Post-production	Post-production	Post-production



Extra steps for original publishing (compared to republishing)

Choose permissions and license, contract Develop and review content Copyedit Select/create graphics

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GOLD Original publishing

Advise and guide author

Post or point to published content



Gold OA vignette

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ACQUISITION

GREEN Republishing	DIAMOND Original publishing
Acquire content	Acquire content
(post-print manuscript)	(original manuscript)

Post-print manuscript



93 Transduction of the Geomagnetic Field as Evidenced from

94 Alpha-band Activity in the Human Brain

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Jacob N. H. Abrahams², Sam E. Bernstein⁵, Ayumu Matani⁴, Shinsuke Shimojo^{1,3*}, & Joseph L.
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 103

105 Abstract

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Accepted Manuscript

eNeuro

Magnetoreception, the perception of the geomagnetic field, is a sensory modality well-106 107 established across all major groups of vertebrates and some invertebrates, but its presence in humans has been tested rarely, yielding inconclusive results. We report here a strong, specific 108 human brain response to ecologically-relevant rotations of Earth-strength magnetic fields. 109 110 Following geomagnetic stimulation, a drop in amplitude of EEG alpha oscillations (8-13 Hz) occurred in a repeatable manner. Termed alpha event-related desynchronization (alpha-ERD), 111 112 such a response has been associated previously with sensory and cognitive processing of external 113 stimuli including vision, auditory and somatosensory cues. Alpha-ERD in response to the magnetic field was triggered only by horizontal rotations when the static vertical magnetic 114 field was directed downwards, as it is in the Northern Hemisphere; no brain responses were 115 elicited by the same horizontal rotations when the static vertical component was directed up-116 wards. This implicates a biological response tuned to the ecology of the local human population, 117 118 rather than a generic physical effect. Biophysical tests showed that the neural response was sensitive to static components of 119 120 the magnetic field. This rules out all forms of electrical induction (including artifacts from the electrodes) which are determined solely on dynamic components of the field. The neural re-121

122 sponse was also sensitive to the polarity of the magnetic field. This rules out free-radical 'quan-123 tum compass' mechanisms like the cryptochrome hypothesis, which can detect only axial align-124 ment. Ferromagnetism remains a viable biophysical mechanism for sensory transduction and

provides a basis to start the behavioral exploration of human magnetoreception.

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Original manuscript

Basic methods and protocols for molecular techniques in parasite diagnostics

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Molecular systematics, i.e., the use of DNA sequences to address a variety of questions on the identity, species boundaries, and relationships of organisms has now become a powerful and useful approach that complements or even supplants traditional systematics based on morphology. A perusal of the literature on parasite systematics suggests that most of our recent understanding and hypotheses of parasite identification and relationships have been, and continue to be, obtained through the application of molecular methods (e.g., Olson et al., 2003; Nadler et al., 2010). This review will attempt to summarize some key protocols in molecular systematics as are used for studying helminth parasites.

Collection of specimens:

The first step in doing molecular systematics is the proper recovery of helminths from the host. Although the specimens used for DNA extraction and subsequent processing need not be handled in the same gentle manner as specimens for morphological studies, they should be collected live, cleaned in 0.6% saline or PBS (phosphate buffered saline) by gentle pipetting or agitation in a petridish to wash off adhering debris, and then fixed for subsequent processing. Specimens that are going to be used for DNA work should be fixed directly in 95% or 100% ethanol, making sure that the ethanol does not contain denaturing agents such as ketones, aldehydes, methanol or kerosene, which are harmful to DNA. A careful reading of the label on the ethanol bottle will tell you what denaturing agents were used. Often, commercially available 95% ethanol is preferred because it may not contain any denaturing agents. Isopropanol can be allowed as a denaturing agent. The sample should be stored in ethanol in a cryovial or in a similar suitable vial, and should be kept chilled in a regular freezer (at -20° C) if possible or in a regular refrigerator (approximately 4° to 8° C) until use. As a cautionary note, formalin is very harmful for DNA work and the worms being used for DNA analysis should not be brought in contact with formalin.

Note: Each time a sample of worms is collected with the intention of doing molecular work, a small subsample of worms from the same batch should also be separately fixed for a corresponding voucher sample to confirm the identity of the worms being studied. These specimens should be fixed by the proper techniques that will allow good stained whole mounts to be produced and be suitable for histology and scanning electron microscopy. For certain helminths (tapeworms, trematodes, nematodes), using hot (steaming) 5% or 10% neutral buffered formalin is an easy way of producing relaxed and well fixed specimens for subsequent stained whole-mounts. If a fume hood or proper ventilation is not available, killing helminths with hot PBS (or saline) and then placing them in unheated fixatives (formalin, FAA, etc.) will suffice for producing adequate stained whole mounts, but worms fixed in this way are not suitable for histology and not ideal for SEM work.

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	Marketing		
Preservation	Preservation		
Capture/analyze use-metrics	Capture/analyze use-metrics		



Green OA vignette

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Diamond OA vignette

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	Copyedit			
Typeset text	Typeset text			
	Select/create graphics			
Lay pages out (text + graphics)	Lay pages out (text + graphics)			
Quick proof corrections	Careful proof corrections			
Generate and affix identifiers	Generate and affix identifiers			

Words of Encouragement

If you are already republishing, do not be intimidated by original publishing



Dziękuję Questions? Ideas?

Mam nadzieję, że spodoba ci się dalsza część konferencji

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