TOP 5 > VETERINARY TRENDS/ETHICS & HUMAN ANIMAL BOND

Top 5 Tips for Reading a Veterinary Scientific Study

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Results from scientific research can help provide optimal care. Daily decisions should lead to effective diagnostic procedures and therapeutic interventions with optimal risk:benefit ratios. It is therefore important to be able to select and evaluate scientific literature relevant to the field and/or patient. Critical evaluation aids in identification of strengths and weaknesses of a study and its relevance and validity in the clinic¹; relevant information for critical evaluation is typically found in the methods and results sections.

The following are the authors' top 5 tips for evaluating a scientific study.

TOP 5 TIPS FOR READING A SCIENTIFIC STUDY

- 1. Determining Relevance
- 2. Identifying Study Design
- 3. Considering Potential Bias
- 4. Identifying Appropriate Results
- 5. Evaluating Literature Guides

<u>1</u> Determining Relevance

For ease of application, the circumstances (eg, patient and condition being evaluated) and available resources (eg, therapies, equipment, required skills) in the study should be similar to those in the clinic. Other important considerations include compliance with local legal requirements, ethics, and wishes of the pet owner.² For example, the study may recommend a surgical procedure, but surgery may not be the best course of action in a patient with high anesthetic risk; therefore, the recommendations from the study are less applicable to the case.

2 Identifying Study Design

Different study designs are best suited to address different questions (*Table 1*).³ Regardless of whether the study concerns diagnostics, therapeutic procedures or treatments, disease prevention, transmission, or another type of research, the evidence can be ranked based on the methodology used.⁴ Methodology most likely to minimize systematic errors is considered to have the least bias, and methodology with the most systematic errors results in the most bias. Systematic errors are not based on chance; they are the result of problems with the study design or methods used to obtain data.⁵ Increased levels of bias increase the likelihood of distorted results. Clinical decision-making should be based on evidence from the least-biased, but most applicable, study design available.

TABLE 1COMMON STUDY TYPES1,2

Study Type*	Description
Meta-analysis	 Quantitative statistical analysis combining data from several studies conducted as part of a systematic literature review
Systematic review	 Method of collating and summarizing information from all published articles addressing a particular question Follows defined and rigorous methods to search and select literature, assess quality, and make conclusions
Randomized controlled trial	 Intervention study used to assess the effect of a treatment or intervention Study subjects randomly allocated to either an intervention or a control group (ie, no treatment, placebo, current best treatment) Ideally, everyone involved in the study should be blinded so no one knows which treatment each patient received
Cohort study	 Prospective or retrospective study in which exposed and unexposed groups (cohorts) are observed over a period of time Outcome (eg, disease) is measured at the end of the study period Can identify risk factors associated with disease and estimate incidence
Diagnostic test validation study	 Used to establish the usefulness of diagnostic tests for specific purposes Patients are tested using both tests (often a new test compared with an accepted test), which are then compared to establish the sensitivity, specificity, and likelihood ratios of the new/repurposed test
Case-control study	 Retrospective study comparing patients with (ie, cases) and without (ie, controls) the disease of interest and often carried out using clinical notes recorded by a veterinary clinic after interaction the pet owner Patient histories are examined to identify risk factors for the disease
Cross-sectional study	 Looks at a sample of the population at a single point in time, most commonly to determine the prevalence of a certain disease; can also include questionnaire-based studies
Case series	 Description of the presentation, diagnosis, treatment, and outcome of several patients with the same disease or syndrome Typically, there are no disease-free patients for comparison, and any differences in management are not considered
Case report Expert opinion	 Description of a single case Can be an individual or a group of experts and provides some evidence Useful when no other scientific, research-driven information is available

Consensus	 Although consensus statements based on peer-reviewed literature
Statement	can be relatively robust, those that rely solely on expert opinion
	without attempts to gather opinions objectively (eg, using specific
	frameworks, such as the Delphi approach) may be less reliable

*Study types are listed in order from lowest to highest risk for bias.

3 Considering Potential Bias

The level of evidence (eg, meta-analysis, randomized clinical trial, case report, expert opinion or experience) in a study indicates how prone it is to bias.⁶ The quality of the study, which is determined by an appraisal process, can also be an indicator of bias. Appraisal should include investigating factors, such as the type, age, sex, and number of study participants (ie, sample size); enrollment criteria; definitions of conditions; which and how examinations were performed; how participants were allocated to different groups; and whether the study was blinded (ie, humans involved in the study did not know which treatment group participants were allocated to). Assessment for potential bias should also include whether clear inclusion and exclusion criteria are given and whether outcome measures are reasonable and relevant. Questions asked during the appraisal process depend on the study design and research question of interest (*Tables 2-4*).

A common flaw in veterinary publications is not including a sufficient number of subjects or samples to draw robust conclusions.⁷ A sample size calculation or reasoning regarding the number of participants included in the study should be included in the methods, results, and discussion sections.

In some studies, patients with a specific disease are enrolled without a clear case definition or a documented diagnostic procedure (including which test results indicated physiologic/not physiologic conditions) that can be used to confidently identify the presence of disease, making it difficult to draw a conclusion. How participants are allocated to different groups is important because unequal distribution (eg, including more severely ill patients in one group compared with another) can distort outcomes.⁸ In addition, a lack of blinding can influence recorded outcomes, as researchers may be (consciously or unconsciously) biased in the conduction or interpretation of a study when they know which patients received specific treatments.

TABLE 2

Step 1: Evider	The level	
	 Meta-analysis (statistical combination of the results of several studies) Clinical trial Case report 	5 points 3 points 2 points 1 point
	Expert opinion or experience	I
Step 2: Additi	onal quality criteria (regarding corresponding evidence level)	
Meta-	• Literature search is exhaustive and reproducible	2 points
analysis	 Included trials are clinically comparable 	4 points
	 Included trials are of high quality (ie, randomized, controlled, blinded) 	2 points 2 points
	 Results are discussed objectively and critically, including questions regarding comparability and bias 	
Clinical trial	• Trial comprises a sufficient number of participants or samples, including a sample size calculation to identify the appropriate number of participants or samples	2 points
	• Essential information (eg, number included, breed, age, sex, inclusion criteria, housing) is given regarding participants	1 point
	 Trial is composed of an adequate control group 	3 points
	Trial is randomized	1 point
	• Trial is blinded	1 point
	• Examinations and interventions are described in detail, and	1 point
	 results are presented completely Adequate statistical procedures are used, and any data that are 	1 point
	incomplete or missing are documented	1 point
	Results are discussed critically	1 point 1 point
	References are extensive and current	-
Case report	• Essential information (eg, number included, breed, age, sex, inclusion criteria, housing) is given regarding participants	2 points
	 Examinations and interventions are described in detail 	2 points
	 Results are discussed critically 	2 points
	References are extensive and current	1 point
Expert	Results are discussed critically	1 point
opinion or experience	 References are extensive and current 	1 point
-	a tion of points for an overall score good; 12-10 = good; 9-7 = satisfactory; 6-4 = adequate; 3-2 = = fail	

TABLE 3 EVALUATION GUIDE FOR RESEARCH ON DIAGNOSTIC TESTS

Study design	• Disease/condition to be tested is clearly defined		1 point
	• Clear, defined test results indicating physiologic/not physiologic		2 points
	conditions		1 point
	• Clear inclusion and exclusion criteria for participants or samples		
	are reported		1 point
	• Appropriate number of participants or samples are included		1 point
	 Procedures are described in detail 		2 points
	• Study is blinded		
Test	 Test is compared with an acknowledged gold standard 		1 point
characteristics	 Sensitivity and specificity of the test are given 		2 points
	 Repeatability (same result obtained when test is repeated) is 		1 point
	good		1 point
	 Possible biases or other problems of the test 		
	(preanalytic/analytic) are discussed		
Practical	• Quality of the test results are discussed in context with other		1 point
relevance	diagnostic tools for the given disease or condition		
	• Applicability and reliability of the test are discussed objectively		1 point
Summation of points for an overall score			
15-13 = very good; 12-10 = good; 9-7 = satisfactory; 6-4 = adequate; 3-2 = inadequate;			
1 = fail			

TABLE 4

EVALUATION GUIDE FOR LITERATURE REVIEWS

Literature	• Literature search was conducted systematically via databases		4 points
search and	and is well documented		
inclusion	 Search terms used are documented 		2 points
	• More literature was searched in reference lists of acquired		1 point
	articles (eg, hand searching*)		2 points
	• Inclusion and exclusion criteria for papers are well-documented		
Assessment	 Quality of each paper is assessed systematically 		4 points
	 Findings and conclusions are discussed objectively 		2 points
Summation of points for an overall score			
15-13 = very good; 12-10 = good; 9-7 = satisfactory; 6-4 = adequate; 3-2 = inadequate;			
1 = fail			

*Hand searching is the examination of reference lists of included studies in order to identify other relevant citations.

<u>4</u> Identifying Appropriate Results

Presentation of crucial information (eg, patient age and medical history, case or control definitions, diagnoses) should be examined.¹ Lack of clarity on whether specific aspects of study design, methods, and results were not considered or just not reported by authors,³ and journal word count restrictions leading to fewer details provided, can cause difficulties

for the reader. Special attention should be given to conclusions, as they may be based on weak or absent scientific data or go beyond the stated research question.³

Transparent reporting is crucial; several reporting guidelines have been developed to ensure important details are not missed.⁹ Quality of a study decreases when critical appraisal is not possible due to poor reporting.

Authors should include whether ethical approval was sought, as well as conflicts of interest and sources of research funding, as these may influence study design or interpretation.¹

5 Evaluating Literature Guides

Several tools are available to guide critical assessment of a scientific study, including evaluation guides,¹⁰ which can help assess key features and quality. Summarizing rating points can provide confidence in recommendations based on study results, although additive scores may not always be necessary. Some tools are included here, but these do not cover all possible scientific research approaches, and some items may not be helpful or easily assessed in every case.

The study design should be determined first, so the appropriate evaluation guide can be chosen. Additional criteria, including information content and objectivity, should be assessed using the appropriate guide. Determining whether the right statistics have been used may be challenging, but focusing on the design features that should be present can help determine study quality.

Conclusion

Critical reading of scientific studies can be time-consuming, but practice can make it easier to decide whether a study is relevant and valid and can be applied in the clinic.¹ Identifying limitations can help results be interpreted and applied appropriately.

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Suggested Reading

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