# Research

# Sample sizes of studies on diagnostic accuracy: literature survey

Lucas M Bachmann, Milo A Puhan, Gerben ter Riet, Patrick M Bossuyt

#### Abstract

#### Methods

**Objectives** To determine sample sizes in studies on diagnostic accuracy and the proportion of studies that report calculations of sample size.

**Design** Literature survey.

**Data sources** All issues of eight leading journals published in 2002.

**Methods** Sample sizes, number of subgroup analyses, and how often studies reported calculations of sample size were extracted.

**Results** 43 of 8999 articles were non-screening studies on diagnostic accuracy. The median sample size was 118 (interquartile range 71-350) and the median prevalence of the target condition was 43% (27-61%). The median number of patients with the target condition—needed to calculate a test's sensitivity—was 49 (28-91). The median number of patients without the target condition—needed to determine a test's specificity—was 76 (27-209). Two of the 43 studies (5%) reported a priori calculations of sample size. Twenty articles (47%) reported results for patient subgroups. The number of subgroups ranged from two to 19 (median four). No studies reported that sample size was calculated on the basis of preplanned analyses of subgroups.

**Conclusion** Few studies on diagnostic accuracy report considerations of sample size. The number of participants in most studies on diagnostic accuracy is probably too small to analyse variability of measures of accuracy across patient subgroups.

## Introduction

Estimates of sensitivity and specificity in small studies on diagnostic accuracy are usually imprecise, with wide confidence intervals. This makes it difficult to assess just how informative a test may be. Subgroup analysis is often needed because sensitivity and specificity may vary across patient subgroups, yet estimates are even less precise when subgroups are considered.<sup>1</sup> Investigators should calculate the sample size needed for sufficiently narrow confidence intervals at the planning stages of a study, as is common practice for randomised trials.<sup>2 3</sup> For example, if a diagnostic test requires a sensitivity of at least 90% for adequate decision making, the lower boundary of the 95% confidence interval should be at least 90%.

We hypothesised that studies of diagnostic accuracy rarely report considerations of sample size and tend to be small. We assumed that authors would state calculations of sample size if they had been performed. We investigated study sizes, the number of subgroup analyses, and how often studies on diagnostic accuracy reported calculations of sample sizes. Two reviewers independently screened all issues of the *BMJ*, *Lancet*, *New England Journal of Medicine*, and *JAMA* as well as four specialist journals (*Thorax, Gastroenterology, American Journal of Obstetrics and Gynecology*, and *European Journal of Pediatrics*) published in 2002 for studies on the accuracy of tests. From each full report we extracted data on the type of test(s) studied (table), study sizes, the number of subgroup analyses, and how often the studies reported calculations of sample size. We calculated 95% confidence intervals, medians, and interquartile ranges.

## Results

Fifty seven of 8999 articles reported test accuracy. Fourteen studies focused on a screening test and were excluded, which left 43 clinical studies for analysis. The median sample size was 118 (interquartile range 71-350) and the median prevalence was 43% (27-61%). The median number of patients with the target condition—needed to calculate a test's sensitivity—was 49 (28-91). The median number of patients without the target condition—needed to determine a test's specificity—was 76 (27-209).

Two of 43 studies (5%; 95% confidence interval 1.3% to 15.5%) reported a priori calculations of sample size, but no study reported that the sample size had been calculated on the basis of preplanned analyses of subgroups. Twenty articles (47%) reported results for subgroups of patients. The number of subgroups ranged from two to 19 (median four). Four studies used multivariable regression, but none used interaction terms.

## Discussion

In this survey of studies on diagnostic accuracy in eight major journals, only 4.7% of the studies reported that they considered sample size. Analysing small numbers of participants with and without the target condition usually yields imprecise estimates of overall diagnostic accuracy, and even less precise estimates of subgroups. For example, when the number of patients with the target condition is 49 the two sided 95% confidence interval of a sensitivity of 81% (40 true positives) is 68% to 91%.<sup>4 5</sup>

To ensure reasonably precise estimates of sensitivity and specificity investigators should consider sample sizes during the planning stages of the study. Investigators should calculate how precise the estimates of test accuracy should be for a particular diagnostic situation and report these calculations with confidence intervals. Arguably, sample size calculations are not important once data collection has been completed.<sup>2</sup> All that matters is the width of the confidence intervals. However, besides determining the minimum study size needed, calculations of sample size have another useful feature that remains important after the study has finished. These calculations require authors to Key features of 57 studies on accuracy of diagnostic tests published in eight major medical journals in 2002

First author	Type of test	Prevalence (%)	Sample size	Screening	Subgroup analysis	Multivariable analysis	Stratified reporting	Number o subgroups
Schneider	Imaging	2	8640	Yes	No	No	No	-
ilcher	Laboratory tests	0.5	8194	Yes	No	No	No	-
ahado-singh (1)	Laboratory tests	2	5641	Yes	Yes	Yes	No	2
ulasingam	Laboratory tests	3	4075	Yes	Yes	No	Yes	2
u	Physical examination	1	3710	Yes	No	No	No	-
intzileos	Imaging	2	3291	Yes	No	No	No	-
'asan	Laboratory tests	6	3177	Yes	Yes	Yes	Yes	4
ahado-singh (2)	Imaging	3	3003	No	Yes	Yes	No	5
elvachandran	History	4	2268	No	Yes	Yes	Yes	2
laisel	Laboratory tests	47	1586	No	No	No	No	_
enders	Laboratory tests	25	858	No	Yes	No	Yes	2
ïbble	Laboratory tests	44	602	No	Yes	No	Yes	2
		3	568	Yes	Yes	Yes	No	_
ahado-singh (3)	Laboratory tests							
zuma	Laboratory tests	6	561	Yes	Yes	No	Yes	3
eda	Physical examination	59	529	No	No	No	No	-
aing	History	21	458	Yes	No	No	No	-
chutter	Laboratory tests	55	412	No	No	No	No	-
/ang	Laboratory tests	50	394	No	Yes	No	Yes	2
luensterer	Imaging	6	386	No	No	No	No	-
havarria	Laboratory tests	7	378	No	No	No	No	-
ettenbacher	Imaging	17	350	No	Yes	No	Yes	3
ubin	Laboratory tests	39	342	No	No	No	No	_
uck	History	4	341	Yes	No	No	No	_
hezzi	Laboratory tests	3	306	No	No	No	No	_
	•	73	278					
liordan	History			No	Yes	Yes	No	19
im	Laboratory tests	36	251	No	Yes	No	Yes	2
ayssiere	History	5	242	Yes	Yes	Yes	No	2
′irkki	Laboratory tests	85	215	No	Yes	Yes	No	7
lemes	History	16	212	No	No	No	No	-
ughes	Laboratory tests	4	208	No	Yes	No	Yes	2
ouin	Other	43	199	No	No	No	No	-
ibeiro	Laboratory tests	85	177	No	Yes	No	Yes	2
liskin-Mashiah	Imaging	6	166	Yes	No	No	No	-
elan	Laboratory tests	27	139	No	No	No	No	_
Judkerk	Imaging	30	118	No	No	No	No	_
/ihm	Laboratory tests	58	113	No	No	No	No	_
IcManus	Other	64	110	Yes	No	No	No	
1cMahon	Physical examination	12	109	No	No	No	No	-
tiller	Other	6.5	107	No	No	No	No	-
ueholm	Imaging	69	106	No	No	No	No	-
ndrews	Imaging	53	100	No	No	No	No	-
oossens	Laboratory tests	32	97	No	Yes	No	Yes	4
eRoche	Laboratory tests	84	90	No	No	No	No	-
arang	Laboratory tests	39	80	No	No	No	No	-
arewood	Imaging	61	80	No	No	No	No	-
arsen	Other	75	79	No	Yes	No	Yes	2
/arke	Laboratory tests	41	71	No	No	No	No	_
ara	Imaging	66	60	No	No	No	No	-
erber	Laboratory tests	34	53	No	No	No	No	_
hmait	Imaging	85	53	No	No	No	No	_
	Laboratory tests							
eorgakoudi		64	44	No	No	No	No	-
agette	Other	79	42	No	Yes	No	Yes	3
arker	Imaging	*	33	No	No	No	No	-
dunsi	Laboratory tests	39	33	No	No	No	No	-
osmi	Imaging	53	32	No	No	No	No	-
roth	Other	41	29	No	No	No	No	-
atoh	Laboratory tests	61	23	No	No	No	No	-

This table provides information for both screening (excluded) and non-screening studies. \*Could not be determined.

think about the minimum precision needed for a test to be clinically meaningful. It is easier for readers to interpret reported confidence intervals if they have access to these data.

In conclusion, few studies on diagnostic accuracy report calculations of sample size. The number of participants in most studies on diagnostic accuracy is probably too small to analyse the variability of measures of accuracy across subgroups of patients.

Contributors: All members of the SUBIRAR (subjectivity rationality and reasoning) research collaboration (Klaus Eichler, Madlaina Scharplatz, and Johann Steurer, Horten Centre, University of Zurich, Switzerland, Ulrich Hoffrage, Max Planck Institute for Human Development and Cognition, Berlin, Germany; Alfons G Kessels, Hans Severens, Maastricht University, Germany; Khalid S Khan, University of Birmingham, UK; Jos Kleijnen, Centre for Reviews and Dissemination, University of York, UK) were involved in the design and critical review of the study. LMB, MAP, and GtR developed the protocol. LMB and MAP acquired the data. All authors interpreted the data and helped prepare the manuscript. LMB was guarantor.

Funding: LMB was supported by the Swiss National Science Foundation (grants 3233B0-103182 and 3200B0-103183). Competing interests: None declared.

# What is already known on this topic

To assess the minimum size needed for sufficiently narrow confidence intervals of sensitivity and specificity in study groups as a whole and in clinically relevant subgroups in particular, sample sizes should be considered at the planning stage of studies on test accuracy

#### What this study adds

Few studies on test accuracy report calculations of sample size

Overall size and subgroup size tend to be small in these studies, which leads to imprecise estimates of sensitivity and specificity

- Irwig L, Bossuyt P, Glasziou P, Gatsonis C, Lijmer J. Designing studies to ensure that estimates of test accuracy are transferable. *BMJ* 2002;324:669-71. 1
- 2 Schulz KF, Grimes DA. Sample size calculations in randomised trials: mandatory and
- mystical. Lancet 2005; 365:1348-53. Lijmer JG, Bossuyt PM, Heisterkamp SH. Exploring sources of heterogeneity in 3 systematic reviews of diagnostic tests. Stat Med 2002;21:1525-37.
- Pepe MS. The statistical evaluation of medical tests for classification and prediction. Oxford Statistical Science Series, Oxford University Press, 2003. www.fhcrc.org/science/labs/ 4 pepe/book/ (accessed 6 Apr 2006).
- Pepe MS. Study design and hypothesis testing. In: The statistical evaluation of medical tests for classification and prediction. New York: Oxford University Press, 2003:214-51. 5(Accepted 7 March 2006)

doi 10.1136/bmj.38793.637789.2F (Accepted 7 March 2006)

Division of Epidemiology and Biostatistics, Department of Social and Preventive Medicine, University of Bern, Switzerland Lucas M Bachmann senior research fellow

Horten Centre, University of Zurich, CH-8091 Zurich, Switzerland Milo A Puhan research fellou

Department of General Practice, Academic Medical Centre, 1105 AZ Amsterdam, Netherlands

Gerben ter Riet clinical epidemiologist

Department of Clinical Epidemiology and Biostatistics, Academic Medical Centre, Amsterdam, Netherlands

Patrick M Bossuyt professor

Correspondence to: L M Bachmann lucas.bachmann@evimed.ch

#### Amendment

This is version 2 of the paper. In this version the median number (interquartile range) of patients has been changed to 49 (28-91) and 76 (27-209) for patients with and without the target condition in the abstract and the results section.