

Supplementary Table S1: PRISMA 2009 Checklist

Section/topic	#	Checklist item	Reported on page #
TITLE			
Title	1	Identify the report as a systematic review, meta-analysis, or both.	1
ABSTRACT			
Structured summary	2	Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number.	2
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of what is already known.	3
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	3-10
METHODS			
Protocol and registration	5	Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address), and, if available, provide registration information including registration number.	N/A
Eligibility criteria	6	Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale.	11
Information sources	7	Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched.	12
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	11
Study selection	9	State the process for selecting studies (i.e., screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis).	12
Data collection process	10	Describe method of data extraction from reports (e.g., piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators.	12
Data items	11	List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made	12
Risk of bias in individual studies	12	Describe methods used for assessing risk of bias of individual studies (including specification of whether this was done at the study or outcome level), and how this information is to be used in any data synthesis.	12
Summary measures	13	State the principal summary measures (e.g., risk ratio, difference in means).	12-13

Synthesis of results	14	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., I ²) for each meta-analysis.	13
Risk of bias across studies	15	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).	13
Additional analyses	16	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified.	13
RESULTS			
Study selection	17	Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram.	13
Study characteristics	18	For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations.	14-15
Risk of bias within studies	19	Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12).	14
Results of individual studies	20	For all outcomes considered (benefits or harms), present, for each study: (a) simple summary data for each intervention group (b) effect estimates and confidence intervals, ideally with a forest plot.	18-23
Synthesis of results	21	Present results of each meta-analysis done, including confidence intervals and measures of consistency.	23
Risk of bias across studies	22	Present results of any assessment of risk of bias across studies (see Item 15).	24
Additional analysis	23	Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]).	n/a
DISCUSSION			
Summary of evidence	24	Summarize the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., healthcare providers, users, and policy makers).	25
Limitations	25	Discuss limitations at study and outcome level (e.g., risk of bias), and at review-level (e.g., incomplete retrieval of identified research, reporting bias).	26
Conclusions	26	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	26
FUNDING			
Funding	27	Describe sources of funding for the systematic review and other support (e.g., supply of data); role of funders for the systematic review.	N/A

From: Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group (2009). Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement. PLoS Med 6(6): e1000097.doi:10.1371/journal.pmed1000097

For more information, visit: www.prisma-statement.org.

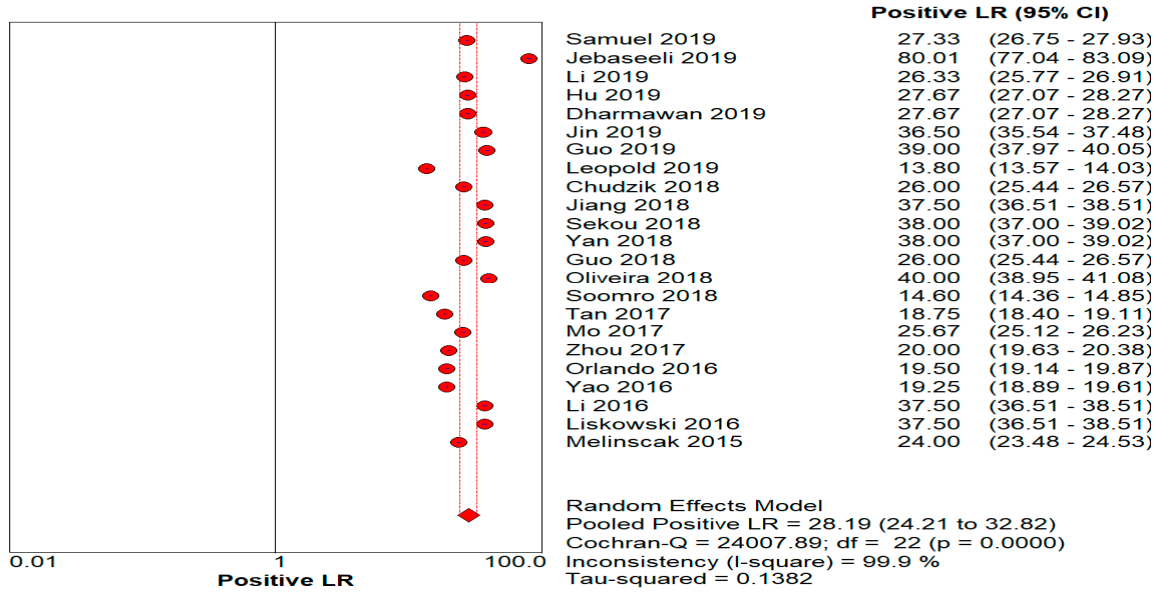
Table S2: Quality Assessment of Diagnostic Accuracy Studies-2 for Included Studies

Study	Risk of Bias				Applicability Concerns		
	Patients Selection	Index Test	Reference Standard	Flow and Timing	Patients Selection	Index Test	Reference Standard
Samuel 2019	😊	😊	?	?	😊	😊	?
Jebaseeli 2019	😊	😊	?	?	😊	😊	?
Li 2019	😊	😊	?	?	😊	😊	?
Hu 2019	😊	😊	?	?	😊	😊	?
Dharmawan 2019	😊	😊	?	?	😊	😊	?
Jin 2019	😊	😊	?	?	😊	😊	?
Guo 2019	😊	😊	?	?	😊	😊	?
Leopold 2019	😊	😊	?	?	😊	😊	?
Lin 2018	😊	😊	?	?	😊	😊	?
Chudzik 2018	😊	😊	?	?	😊	😊	?
Jiang 2018	😊	😊	?	?	😊	😊	?
Sekou 2018	😊	😊	?	?	😊	😊	?
Hajabdollahi 2018	😊	😊	?	?	😊	😊	?
Yan 20018	😊	😊	?	?	😊	😊	?
Guo 2018	😊	😊	?	?	😊	😊	?
Oliveira 2017	😊	😊	?	?	😊	😊	?
Soomro 2017	😊	😊	?	?	😊	😊	?
Tan 2017	😊	😊	?	?	😊	😊	?
Mo 2017	😊	😊	?	?	😊	😊	?
Zhou 2017	😊	😊	?	?	😊	😊	?
Dasgupta 2017	😊	😊	?	?	😊	😊	?
Sengur 2017	😊	😊	?	?	😊	😊	?
Orlando 2016	😊	😊	?	?	😊	😊	?
Yao 2016	😊	😊	?	?	😊	😊	?
Li 2016	😊	😊	?	?	😊	😊	?
Maji 2016	😊	😊	?	?	😊	😊	?
Lahiri 2016	😊	😊	?	?	😊	😊	?
Liskowski 2016	😊	😊	?	?	😊	😊	?
Fu 2016	😊	😊	?	?	😊	😊	?
Fu 2016	😊	😊	?	?	😊	😊	?
Melinscak 2015	😊	😊	?	?	😊	😊	?

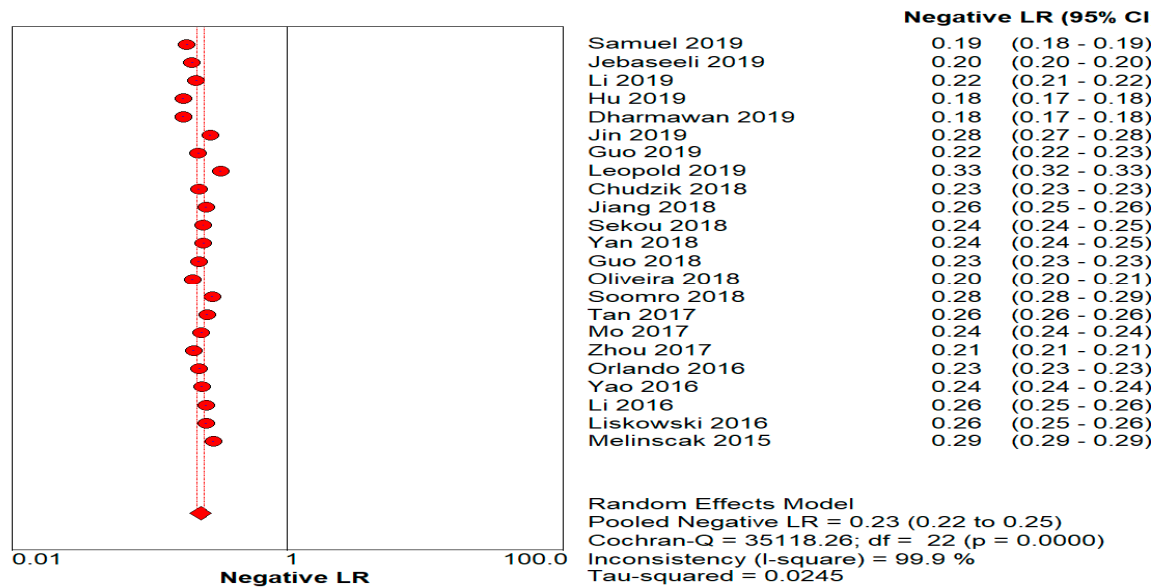
Note: 😊 = No risk bias; ? = Unclear risk bias; 😞 = High risk of bias

Supplementary figure F1-F4: Performance of DL in Retinal Vessel Segmentation:

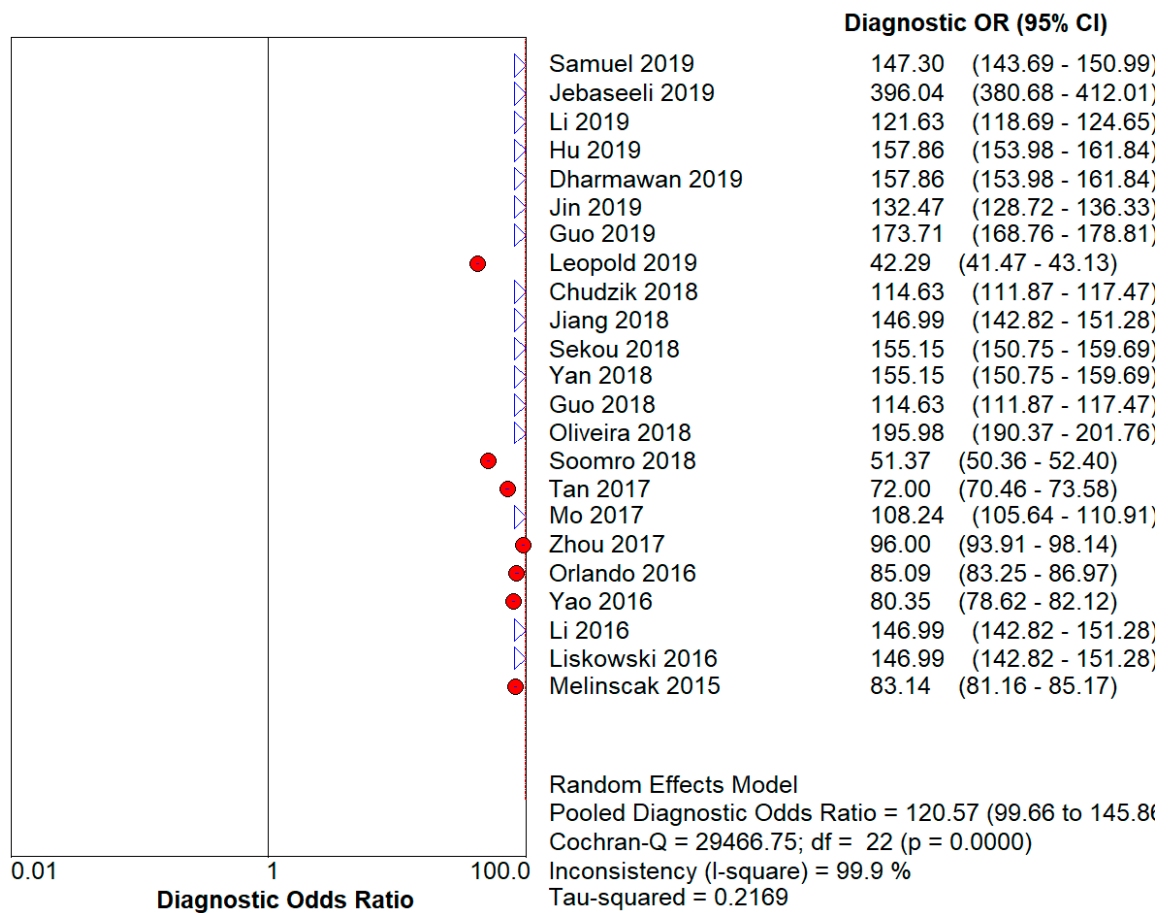
DRIVE database



A. Positive LR



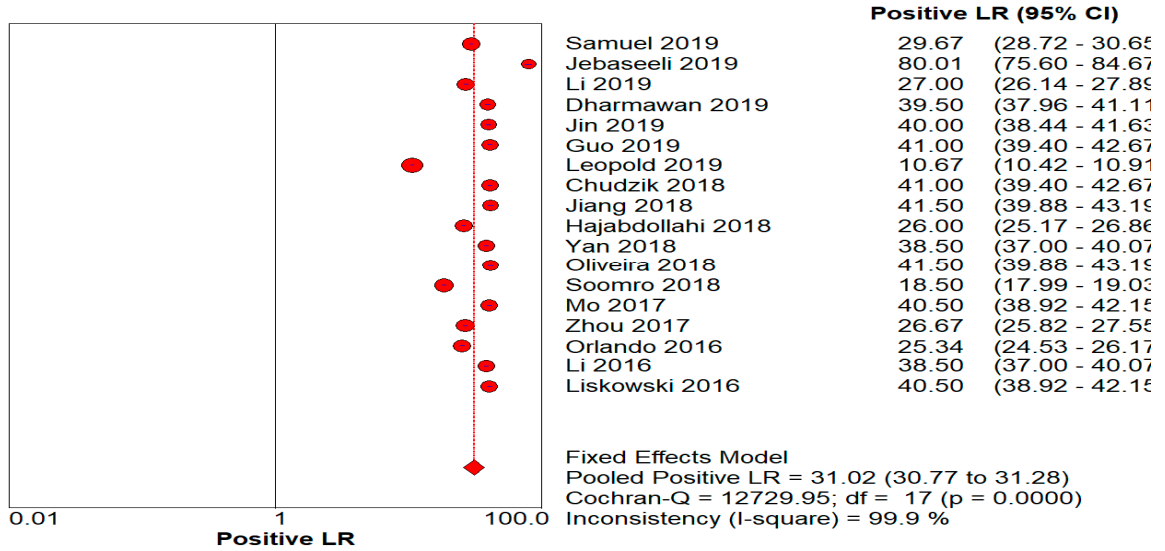
B. Negative LR



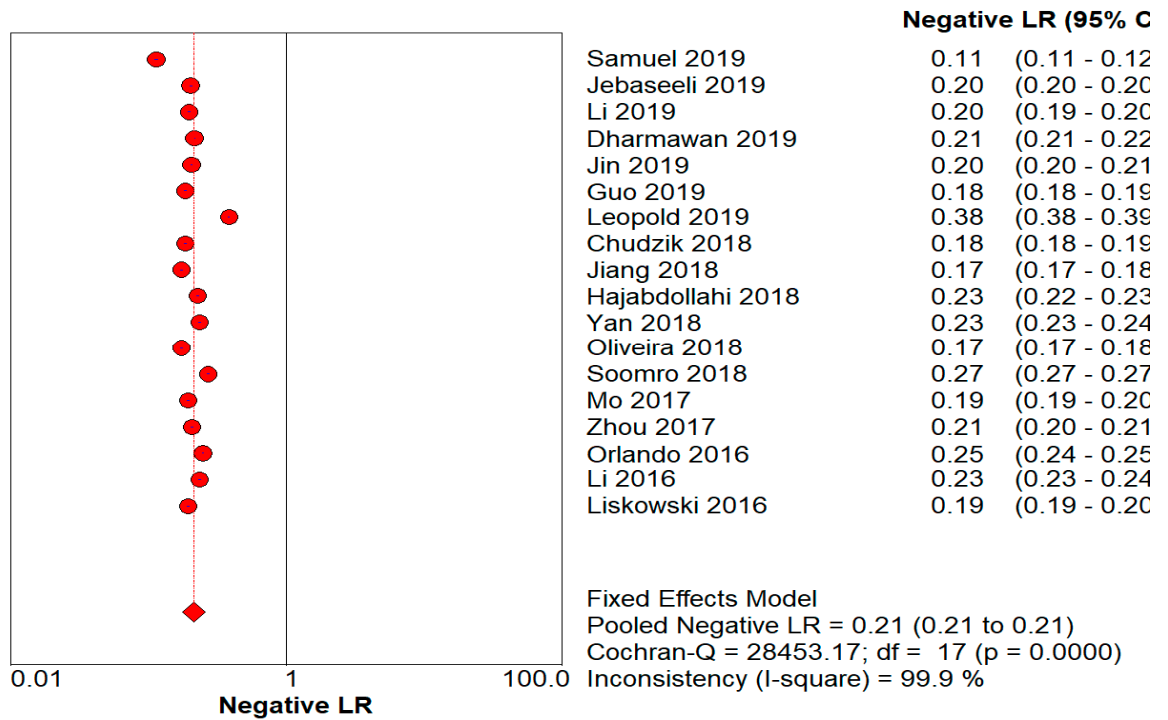
C. Diagnostic odds ratio

Figure F1: Performance of Deep learning in DRIVE database a) Positive LR b) Negative LR c) Diagnostic odds ratio

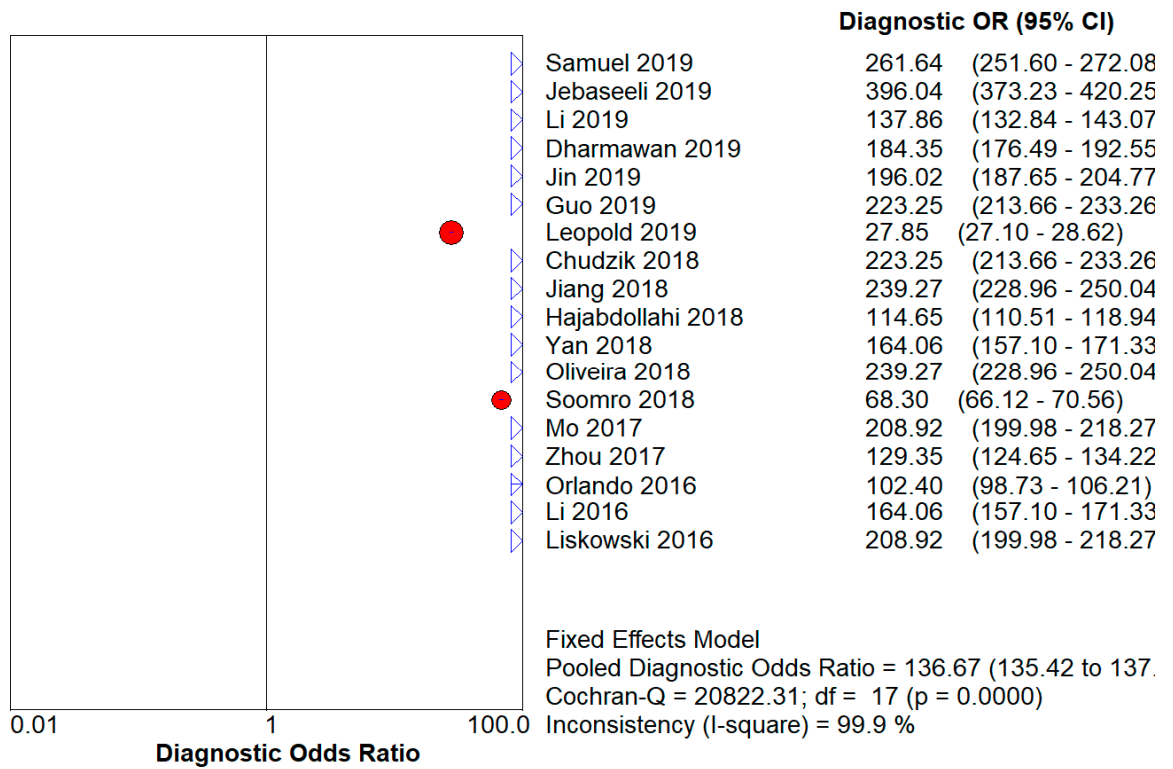
STARE DATABASE



A. Positive LR



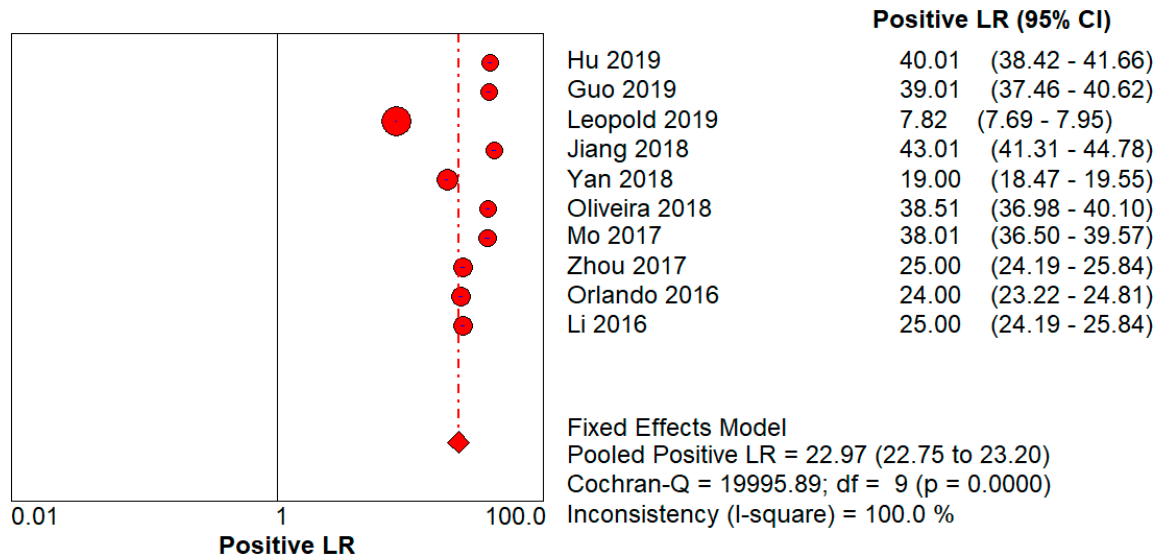
B. Negative LR



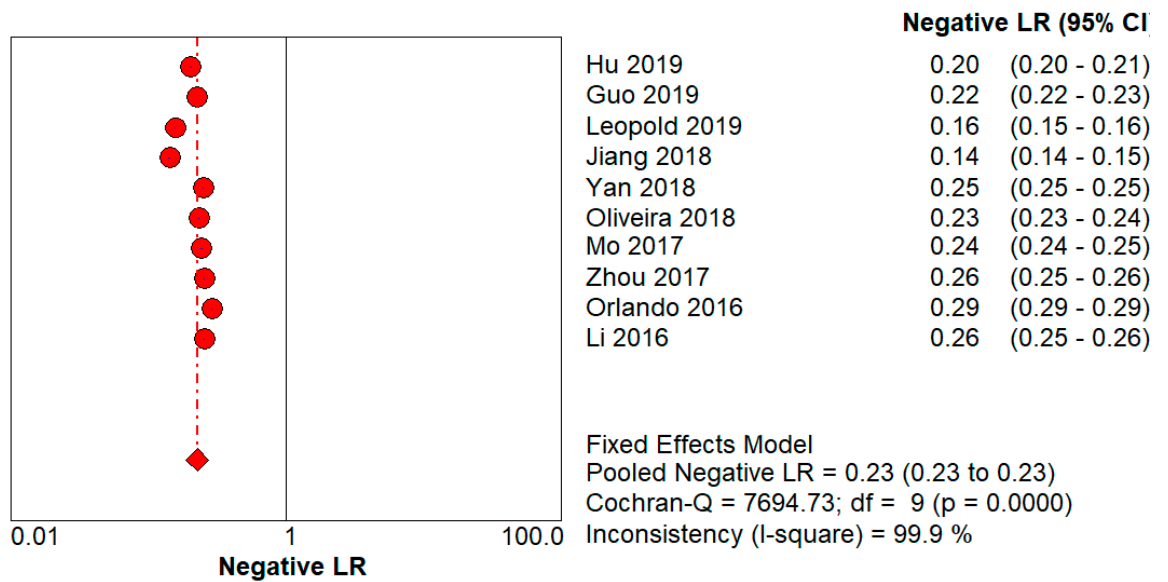
C. Diagnostic odds ratio

Figure F2: Performance of Deep leaning in STARE database a) Positive LR b) Negative LR c) Diagnostic odds ratio

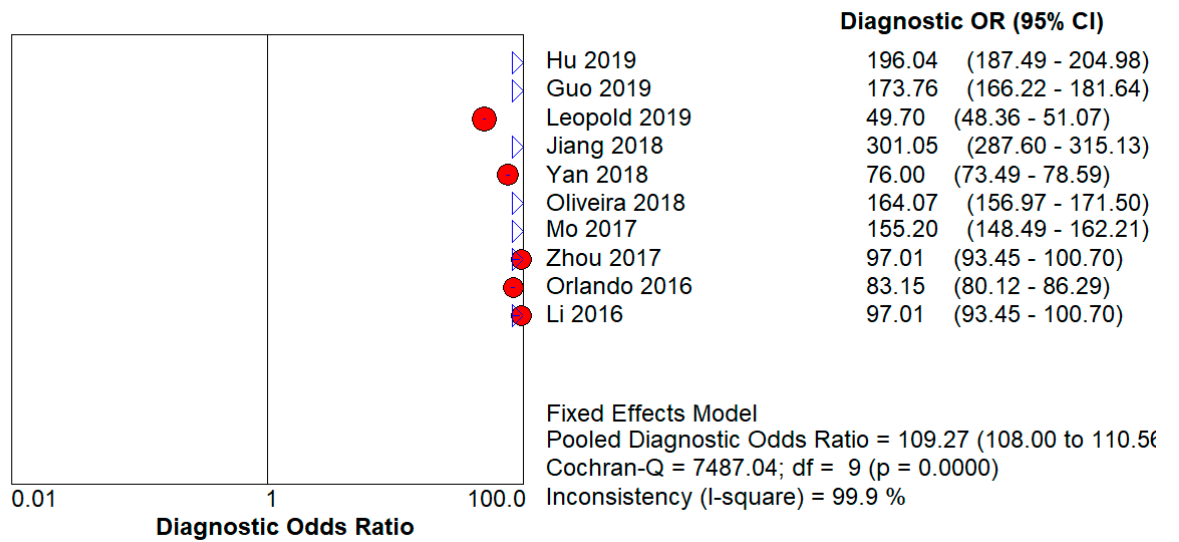
CHASE_DBI Database



A. Positive LR



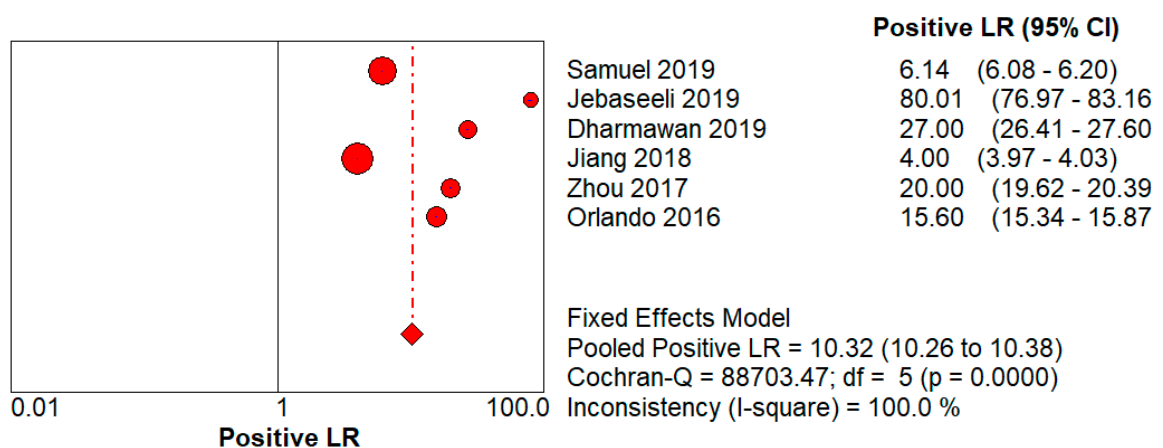
B. Negative LR



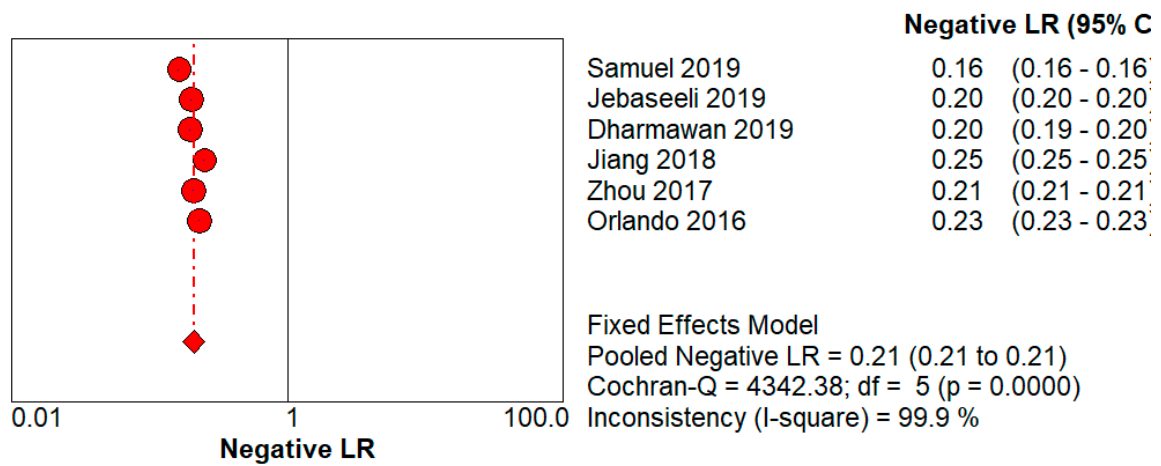
C. Diagnostic Odds Ratio

Figure F3: Performance of Deep learning in CHASE_DB1 database a) Positive LR b) Negative LR c) Diagnostic odds ratio

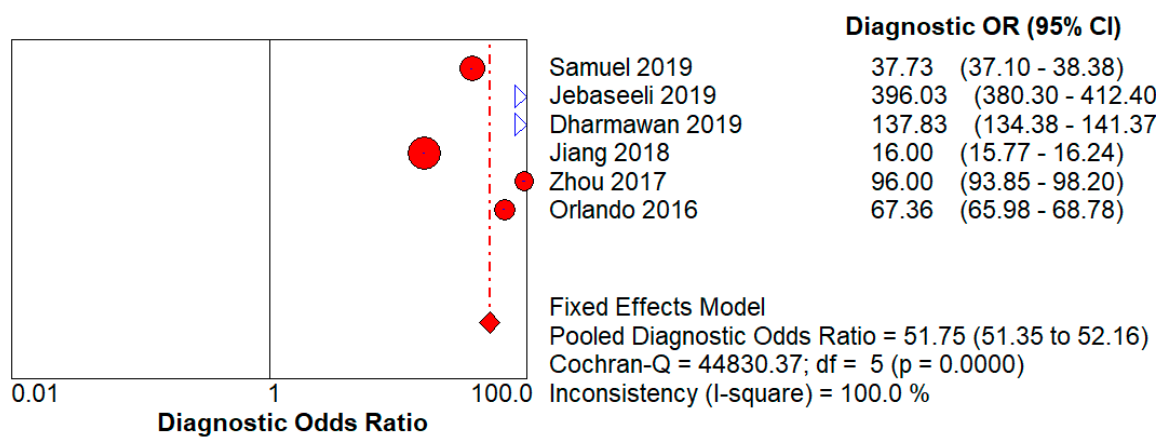
HRF database



A. Positive LR



B. Negative LR



C. Diagnostic Odds Ratio

Figure F4: Performance of Deep leaning in DRIVE database a) Positive LR b) Negative LR c) Diagnostic odds ratio