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LAMPIRAN

1. Tabel hasil telaah krtitis artikel 1

Eksperimental Treatment With Favipiravir for COVID-19: Open-Label Control Study

JBI Critical Appraisal Checklist for Quasi-Experimental Studies (non-randomized experimental studies)

Reviewer : DA,SS

Author : Qingxian Cai, et al

	525		NO	UNCLEAR	NA
NO	Question				
1	Is it clear in the study what is the 'cause'				
	and what is the 'effect' (i.e. there is no				
	confusion about which variable comes				
	first)?	,			
2	Were the participants included in any				
	comparisons similar?				
3	Were the participants included in any	$\sqrt{}$			
	comparisons receiving similar				
	treatment/care, other than the exposure				
	or intervention of interest?	,			
4	Was there a control group?	V			
5	Were there multiple measurements of the				
	outcome both pre and post the				
	intervention/exposure?		,		
6	Was follow up complete and if not, were				
	differences between groups in terms of				
	their follow up adequately described and				
	analyzed?	,			
7	Were the outcomes of participants				
	included in any comparisons measured in				
	the same way?	,			
8	Were outcomes measured in a reliable				
	way?	,			
9	Was appropriate statistical analysis used?	V			

2. Tabel hasil telaah krtitis artikel 2

Favipiravir versus Arbidol for COVID-19: A Randomized Clinical Trial

JBI Critical Appraisal Checklist for Randomized Controlled Trials

Reviewer : DA,SS

Author : Chang chen, et al

		YES	NO	UNCLEAR	NA
NO					
1	Was true randomization used for assignment of participants to treatment groups?	V			
2	Was allocation to treatment groups concealed?				
3	Were treatment groups similar at the baseline?				
4	Were participants blind to treatment assignment?		V		
5	Were those delivering treatment blind to treatment assignment?		V		
6	Were outcomes assessors blind to treatment assignment?		1		
7	Were treatment groups treated identically other than the intervention of interest?	√			
8	Was follow up complete and if not, were differences between groups in terms of their follow up adequately described and analyzed?		√		
9	Were participants analyzed in the groups to which they were randomized?	V			
10	Were outcomes measured in the same way for treatment groups?	1			
11	Were outcomes measured in a reliable way?	√			
12	Was appropriate statistical analysis used?	$\sqrt{}$			
13	Was the trial design appropriate, and any deviations from the standard RCT design (individual randomization, parallel groups) accounted for in the conduct and analysis of the trial?	V			

3. Tabel hasil telaah kritis artikel 3

Clinical Course of a Crtically ill Patient with Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2)

JBI Critical Appraisal Checklist for Case Reports

Reviewer : DA,FN

Author : Nozomi takahashi, et al

		YES	NO	UNCLEAR	NA
NO					
1	Were patient's demographic characteristics clearly described?	1			
2	Was the patient's history clearly described and presented as a timeline?	V			
3	Was the current clinical condition of the patient on presentation clearly described?		√		
4	Were diagnostic tests or assessment methods and the results clearly described?	V			
5	Was the intervention(s) or treatment procedure(s) clearly described?			√ 	
6	Was the post-intervention clinical condition clearly described?			√ 	
7	Were adverse events (harms) or unanticipated events identified and described?		V		
8	Does the case report provide takeaway lessons?	$\sqrt{}$			

4. Tabel hasil telaah kritis artikel 4

Neuroleptic Malignant Syndrome in Patient with COVID-19

JBI Critical Appraisal Checklist for Case Reports

Reviewer : DA,FN

Author : Mitsuhito soh, et al

		YES	NO	UNCLEAR	NA
NO					
1	Were patient's demographic characteristics clearly described?		V		
2	Was the patient's history clearly described and presented as a timeline?	1			
3	Was the current clinical condition of the patient on presentation clearly described?	$\sqrt{}$			
4	Were diagnostic tests or assessment methods and the results clearly described?		√		
5	Was the intervention(s) or treatment procedure(s) clearly described?		√		
6	Was the post-intervention clinical condition clearly described?			√ 	
7	Were adverse events (harms) or unanticipated events identified and described?			V	
8	Does the case report provide takeaway lessons?	1			

5. Perhitungan nilai persetujuan tinjauan sistematis

	Pengamat 1/ reviewer 1							
		Ya Tidak Jumlah						
	Ya	Α	В	a + b				
Pengamat 2/	Tidak	С	D	c + d				
reviewer 2	Jumlah	a + c	b + d	a+b+c+d=				
				n				

Ket:

Persetujuan= (a + d)/n x 100%

Cohen's Kappa= (Po-Pe)/(1-Pe)

Po=(a+d)/n

Pe = [(a + c)(a + b)/n + (b + d)(c + d)/n]/n

Kategori nilai Cohen's Kappa (Altman, 1991):

 $0.00 \le k \le 0.20 = Rendah$

 $0.21 \le k \le 0.40 = Lumayan$

 $0.41 \le k \le 0.60 = Cukup$

 $0.61 \le k \le 0.80 = Baik$

0.81≤ k ≤ 1.00 = Sangat baik

a. Skrining Tahap I (TIAB screening)

	Pengamat 1/ reviewer 1						
		Include Exclude Jumlah					
B	Include	26	5	31			
Pengamat 2/	Exclude	5	121/124	126/129			
reviewer 2	Jumlah	31	126/129	157/160			

Persetujuan =
$$(a + d)/n \times 100\%$$

= $(26+124)/160 \times 100\%$
= $0.937 (0.937\%)$
Cohen's Kappa = $(Po-Pe)/(1-Pe)$
= $(0.936-0.683)/(1-0.683)$
= $0.798 (Baik)$
Po = $(a + d)/n$
= $(26+121)/157=0.936$
Pe = $[(a + c)(a + b)/n + (b + d)(c + d)/n]/n$
= $[(31)(31)/157 + (126)(126)/157]/157$
= $6,12 + 101,12/157$
= 0.683

b. Skrining Tahap II (Full Text Screening)

	Pengamat 1/ reviewer 1							
		Include Exclude Jumlah						
	Include	4	0	4				
Pengamat 2/	Exclude	0	22	22				
reviewer 2	Jumlah	4	22	26				

Persetujuan =
$$(a + d)/n \times 100\%$$

= $(4+22)/26 \times 100\%$
= $1 (100\%)$
Cohen's Kappa = $(Po-Pe)/(1-Pe)$
= $(1-0.739)/(1-0.739)$
= $1,00 \text{ (Sangat baik)}$
Po = $(a + d)/n$
= $(4+22)/26=1$
Pe = $[(a + c)(a + b)/n + (b + d)(c + d)/n]/n$
= $[(4)(4)/26 + (22)(22)/26]/26$
= $0.615 + 18,615/26$
= 0.739