




Supplementary Materials

Table S1. Searching strategy

Pubmed (OVID)	EMBASE
<ol style="list-style-type: none">1. exp Hepatitis B2. acute.tw.3. 1 and 24. randomized controlled trial.pt.5. controlled clinical trial.pt.6. lamivudine7. nucleoside8. 4 or 5 or 6 or 79. 3 and 8	<ol style="list-style-type: none">1. hepatitis B2. acute3. 1 and 24. randomized controlled trial5. controlled clinical trial6. lamivudine7. nucleoside8. 4 or 5 or 6 or 79. 3 and 8

Table S2. Summary of findings for the main comparison. Lamivudine vs no intervention for acute hepatitis B virus infection

Lamivudine compared to standard of care for acute hepatitis B infection. **Setting: inpatients**

Certainty assessment							№ of patients		Effect		Certainty
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	lamivudine	standard of care	Relative (95% CI)	Absolute (95% CI)	
Undetectable HBV DNA											
5	randomised trials	serious	serious	not serious	not serious	none	148/173 (85.5%)	192/223 (86.1%)	OR 0.96 (0.54 to 1.70)	5 fewer per 1000 (from 91 fewer to 52 more)	 1, 2 Low
Seroconversion HBsAg											
5	randomised trials	serious	serious	not serious	not serious	none	95/173 (54.9%)	136/223 (61.0%)	OR 0.54 (0.33 to 0.90)	152 fewer per 1000 (from 270 fewer to 25 fewer)	 1, 2 Low
Mortality											
4	randomised trials	serious	serious	not serious	not serious	none	7/158 (4.4%)	15/207 (7.2%)	OR 0.54 (0.21 to 1.36)	32 fewer per 1000 (from 56 fewer to 24 more)	 1, 2 Low

CI: confidence interval; OR: odds ratio; HBV: Hepatitis B Virus; HBsAg: Hepatitis B surface Antigen; SOC: Standard of Care

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.


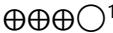
Very low quality: We are very uncertain about the estimate.

¹ The risk of bias in the trials was high (downgraded by 1 level for risk of bias).

² Moderate heterogeneity in results across studies (downgrade the quality of evidence for this outcome by 1 level).

Table S3. Summary of findings. Entecavir vs Lamivudine for acute hepatitis B virus infection

Entecavir compared to Lamivudine for acute hepatitis B infection **Setting:** inpatients


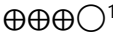
Certainty assessment							№ of patients		Effect		Certainty	
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	[Entecavir]	[Lamivudine]	Relative (95% CI)	Absolute (95% CI)		
Seroconversion HBsAg												
1	randomised trials	serious	not serious	not serious	not serious	none	16/69 (23.2%)	11/21 (52.4%)	OR 3.64 (1.31 to 10.13)	276 more per 1,000 (from 67 more to 394 more)	 Moderate	
Reduction viral load												
1	randomised trials	serious	not serious	not serious	not serious	none	14/21 (66.7%)	56/69 (81.2%)	OR 0.46 (0.16 to 1.38)	147 fewer per 1,000 (from 404 fewer to 44 more)	 Moderate	

CI: confidence interval; OR: odds ratio; HBsAg: Hepatitis B surface Antigen

¹ The risk of bias in the trials was high (downgraded by 1 level for risk of bias).

Table S4. Summary of findings. Entecavir vs no intervention for acute hepatitis B virus infection

Entecavir compared to standard of care for acute hepatitis B infection **Setting:** inpatients

Certainty assessment							№ of patients		Effect		Certainty	
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	[Entecavir]	[standard of care]	Relative (95% CI)	Absolute (95% CI)		
Seroconversion HBVsAg												
1	randomised trials	serious	not serious	not serious	not serious	none	11/21 (52.4%)	43/110 (39.1%)	OR 1.71 (0.67 to 4.38)	132 more per 1,000 (from 90 fewer to 347 more)	 Moderate	
Reduction viral load												
1	randomised trials	serious	not serious	not serious	not serious	none	11/21 (52.4%)	102/110 (92.7%)	OR 0.16 (0.05 to 0.50)	256 fewer per 1,000 (from 538 fewer to 63 fewer)	 Moderate	

CI: confidence interval; OR: odds ratio; HBsAg: Hepatitis B surface Antigen

¹ The risk of bias in the trials was high (downgraded by 1 level for risk of bias).