

1 *Supplementary material*

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## 3 Supplementary Table 1: Quality Assessment For Case Cohort Studies

Selection (MAX 4 stars)	
1	<p>Representativeness of the exposed cohort</p> <p>a) truly representative of the average _____ (describe) in the community</p> <p>b) somewhat representative of the average _____ in the community</p> <p>c) selected group of users eg nurses, volunteers</p> <p>d) no description of the derivation of the cohort</p>
2	<p>Selection of the non exposed cohort</p> <p>a) drawn from the same community as the exposed cohort</p> <p>b) drawn from a different source</p> <p>c) no description of the derivation of the non exposed cohort</p>
3	<p>Ascertainment of exposure</p> <p>a) secure record (eg surgical records)</p> <p>b) structured interview</p> <p>c) written self report</p> <p>d) no description</p>
4	<p>Demonstration that outcome of interest was not present at start of study</p> <p>a) yes</p> <p>b) no</p>
Comparability (MAX 2 stars)	
5	<p>Comparability of cohorts on the basis of the design or analysis</p> <p>a) study controls for _____ (select the most important factor)</p> <p>b) study controls for any additional factor</p>
Outcome (MAX 3 stars)	
6	<p>Assessment of outcome</p> <p>a) independent blind assessment</p> <p>b) record linkage</p> <p>c) self report</p> <p>d) no description</p>

7	<p>Was follow-up long enough for outcomes to occur</p> <ol style="list-style-type: none"> <li>yes (select an adequate follow up period for outcome of interest)</li> <li>no</li> </ol>
8	<p>Adequacy of follow up of cohorts</p> <ol style="list-style-type: none"> <li>complete follow up - all subjects accounted for</li> <li>subjects lost to follow up unlikely to introduce bias - small number lost - &gt; ____ % (select an adequate %) follow up, or description provided of those lost)</li> <li>follow up rate &lt; ____ % (select an adequate %) and no description of those lost</li> <li>no statement</li> </ol>

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6 Supplementary Table 2: Quality Assessment For Cross-Sectional Studies

Selection (MAX 5 stars)	
1	<p>Representativeness of the cases:</p> <ul style="list-style-type: none"> <li>a) Truly representative of the population (consecutive or random sampling of cases). 1 score</li> <li>b) Somewhat representative of the average population (non-random sampling) . 1 score</li> <li>c) Selected demographic group of users. 0 score</li> <li>d) No description of the sampling strategy. 0 score</li> </ul>
2	<p>Sample size:</p> <ul style="list-style-type: none"> <li>a) Justified and satisfactory. 1 score</li> <li>b) Not justified. 0 score</li> </ul>
3	<p>Non-Response rate</p> <ul style="list-style-type: none"> <li>a) <b>The response rate is satisfactory (<math>\geq 95\%</math>).</b> 1 Score</li> <li>b) The response rate is unsatisfactory (<math>&lt; 95\%</math>), or no description. 0 Score</li> </ul>
4	<p>Ascertainment of the screening/surveillance tool:</p> <ul style="list-style-type: none"> <li>a) Validated screening/surveillance tool. 2 scores</li> <li>b) Non-validated screening/surveillance tool, but the tool is available or described. 1 score</li> <li>c) No description of the measurement tool. 0 score</li> </ul>
Comparability (MAX 1 star)	
5	<p>The potential confounders were investigated by subgroup analysis or multivariable analysis.</p> <ul style="list-style-type: none"> <li>a) The study investigates potential confounders. 1 score</li> <li>b) The study does not investigate potential confounders. 0 score</li> </ul>
Outcome (MAX 3 stars)	
6	<p>Assessment of the outcome:</p> <ul style="list-style-type: none"> <li>a) Independent blind assessment. 2 scores</li> <li>b) Record linkage. 2 scores</li> <li>c) Self report. 1 score</li> <li>d) No description. 0 score</li> </ul>

7	<p>Statistical test:</p> <ul style="list-style-type: none"><li data-bbox="398 249 1319 339">a) The statistical test used to analyze the data is clearly described and appropriate. 1 score</li><li data-bbox="398 361 1319 449">b) The statistical test is not appropriate, not described or incomplete. 0 score</li></ul>
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8      **Supplementary Table 3: Case Cohort Studies Quality Assessment.**  
 9      Points  $\geq 7$  were considered as “good”, 2 to 6 points were considered  
 10     as “fair”, and  $\leq 1$  point was considered as “poor” quality.

Study	1	2	3	4	5	6	7	8	Total
Bhaskar et al.	1	1	1	0	2	1	1	1	8
Devarbhavi et al.	1	0	1	0	0	1	1	1	5
Diaz-Quijano et al.	1	1	1	1	0	1	1	1	7
Djossou et al.	1	1	1	1	2	1	1	1	9
Fernández et al.	1	1	1	0	2	1	0*	0*	6
Gayathri et al.	1	1	1	1	2	1	1	1	9
Hsieh et al.	1	0	1	1	0	1	1	1	6
Huang et al.	1	1	1	1	2	1	1	1	9
Lam et al.	1	0	1	0	0	1	1	1	5
Lee et al.	1	1	1	0	2	1	1	1	8
Lee et al.	1	1	1	0	2	1	1	1	8
Leo et al.	1	1	1	1	2	1	1	1	9
Marois et al.	1	1	1	1	2	1	1***	0	8
McBride et al.	1	0	1	0	0	1	1	1	5
Nguyen et al.	1	1	1	0	2	0	1	1	7
Park et al.	1	1	1	0	1	1	1	0	6
Phakhounthong et al.	1	1	1	0	NA	1	1	1	6
Pongpan et al.	1	NA	1	0	NA	1	1	1	5
Pongpan et al.	1	NA	1	0	2**	1	1	1	7
Potts et al.	1	1	1	1	2	1	1	1	9
Sachdev et al.	1	NA	1	0	0	1	1***	1	5
Srisuphanunt et al.	1	1	1	0	1	1	1	1	7
Suwarto et al.	1	NA	1	0	0	1	1	1	5
Suwarto et al.	1	NA	1	0	0	1	1	1	5

Tangnarak et al.	1	NA	1	0	0	1	1	1	5
Md-Sani et al.	1	NA	1	0	2	1	1	1	7
Pang et al.	1	1	1	1	1	1	1	1	8

\* Information not available in study

\*\* differences between severity groups shown to be insignificant

\*\*\* follow up length not defined

14

15 **Supplementary Table 4: Cross-sectional Studies Quality**

16 **Assessment.** Points  $\geq 7$  were considered as “good”, 2 to 6 points were  
17 considered as “fair”, and  $\leq 1$  point was considered as “poor” quality.

Study	1	2	3	4	5	6	7	Total
Chi et al.	1	1	1	1	1	1	1	7
Yang et al.	1****	1	1	2	1	2	1	9

18 \*\*\*\* convenience sampling from five government and three private hospitals

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20 Supplementary Figure 1: Most sensitive for mortality in adults - Weighted risk scoring [27]<sup>1</sup>

<u>Variable</u>	<u>Score</u>
Platelet $<50 \times 10^9$ cells/L (initial)	0 or 1
Leukocytosis (during stay)	0 or 2
Gastrointestinal bleed $<72$ h	0 or 2
Haemoconcentration (during stay)	0 or 2
<b>Total</b>	<b>0 to 7</b>

*Score  $\geq 2$ : High mortality risk*

21

22 Supplementary Figure 2: Most specific for severity in children - BDSS formula [10]

$$\text{BDSS} = -1.297 + 4.234 \times (\text{Pulse Pressure in mmHg}) + 1.284 \times (\text{Mucosal bleed: 1 or 0}) + 0.489 \times (\text{Third-spacing fluid loss: 1 or 0})$$

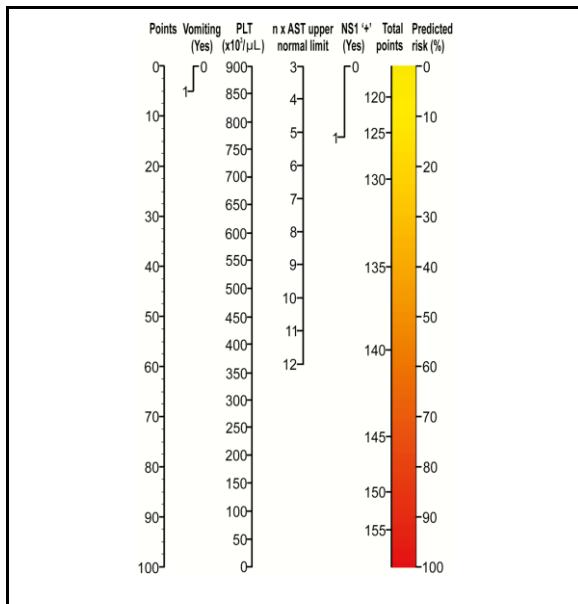
*BDSS > 0.9285: Severe dengue*

23

24 Supplementary Figure 3: Most sensitive for severity in children - Nomogram of prognostic  
25 model [16]<sup>2</sup>

<sup>1</sup> Licence CC BY 4.0 (assessed from <https://www.mdpi.com/2077-0383/7/11/396> on 5 October 2024)

<sup>2</sup> Licence CC BY-NC-ND 4.0 (assessed from <https://academic.oup.com/cid/article/64/5/656/2747462> on 5 October 2024)



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27 Supplementary Figure 4: Most specific for mortality in children - Vasopressor-Ionotrope  
 28 Score (VIS) formula [13]

$$\text{VIS} = \text{Dopamine } (\mu\text{g kg}^{-1} \text{ min}^{-1}) + \text{Dobutamine } (\mu\text{g kg}^{-1} \text{ min}^{-1}) + 100 \times \text{Epinephrine } (\mu\text{g kg}^{-1} \text{ min}^{-1}) + 100 \times \text{Norepinephrine } (\mu\text{g kg}^{-1} \text{ min}^{-1}) + 10 \times \text{Vasopressin } (\text{U kg}^{-1} \text{ min}^{-1}) + 10 \times \text{Milrinone } (\mu\text{g kg}^{-1} \text{ min}^{-1})$$

*VIS ≥ 22.5: High mortality risk*

29

30 Supplementary Figure 5: Most specific for severity in adults - Warning Signs [20]<sup>3</sup>

A combination of 3 of the Warning Signs for Severe Dengue as follows:

- 1. Abdominal Pain**
- 2. Mucosal Bleeding**
- 3. Persistent Vomiting OR Change in Haematocrit**

31

32 Supplementary Figure 6: Most sensitive for severity in adults - Warning Signs [20]

At least 1 of the Warning Signs for Severe Dengue as follows:

- 1. Abdominal pain**
- 2. Persistent Vomiting**

<sup>3</sup>Licence CC BY 2.0 (assessed from <https://bmccinfectdis.biomedcentral.com/articles/10.1186/1471-2334-13-498> on 5 October 2024)

- 3. **Mucosal bleeding**
- 4. **Clinical fluid accumulation**
- 5. **Hepatomegaly >2cm**
- 6. **Increase Haematocrit** (and concurrent reduced platelets)

33

34 Supplementary Figure 7: Most specific for severity in adults - Weighted risk scoring [32]

<u>Variable</u>	<u>Score</u>
Age $\geq$ 65-year-old	0 or 1
Leukocytosis (WBC $>10 \times 10^9$ )	0 or 2
<b>Total</b>	<b>0 to 3</b>
<i>Score <math>\geq</math> 1: High risk of severe dengue</i>	

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36 Supplementary Figure 8: Most specific for mortality in adults - Qualitative integer scoring  
37 system [35]

<u>Variable</u>	<u>Score</u>
Age $\geq$ 65-year-old	0 or 1
Systolic Blood Pressure $<90$ mmHg	0 or 1
Haemoptysis	0 or 1
Diabetes mellitus	0 or 1
Chronic bedridden	0 or 1
<b>Total</b>	<b>0 to 5</b>
<i>Score <math>\geq</math> 3: High mortality risk (45.5%)</i>	

38

39 Supplementary Figure 9: Probability of Intensive Care Unit (ICU) admission (only study with  
40 reported Sensitivity and Specificity) [36]<sup>4</sup>

<sup>4</sup> Licence CC BY 2.0 (assessed from <https://bmccinfectdis.biomedcentral.com/articles/10.1186/s12879-014-0649-2> on 5 October 2024)

**Probability of ICU requirement ( $P_{index}$ )** =  $0.106 \times x_1 + 0.004 \times x_2 + 0.326 \times x_3 -$

10.601

$P_{index} \geq -1.4$ : High risk of ICU admission

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