

Supplementary table 2 Risk of bias assessment for enrolled retrospective studies by the modified Newcastle-Ottawa Scale[#]

Studies	Selection				Comparability		Exposure/Outcomes				Total score	Quality
	Item 1	Item 2	Item 3	Item 4	Item 1	Item 2	Item 1	Item 2	Item 3	Item 4		
Reynold 2015	1	1	1	0	0	1	0	0	1	1	7	Moderate
Teibel 2020	1	1	1	0	1	1	0	0	1	1	8	High
Tran 2021	1	1	1	0	1	1	1	1	1	1	9	High
Yan 2021	1	0	1	0	1	1	0	0	1	1	7	Moderate

[#] Items of the modified Newcastle-Ottawa Scale are listed below.

NEWCASTLE - OTTAWA QUALITY ASSESSMENT SCALE NON-RANDOMIZED STUDIES

Note: A study can be awarded a maximum of one star for each numbered item.

Selection

1) Representativeness of the exposed cohort

- a) truly representative of the average neonates requiring central line insertions in the NICU ✱
- b) somewhat representative of the average neonates requiring central line insertions in the NICU
- c) selected group of users, eg nurses, volunteers
- d) no description of the derivation of the cohort

2) Selection of the non-exposed cohort

- a) drawn from the same community as the exposed cohort ✱
- b) drawn from a different source
- c) no description of the derivation of the non-exposed cohort

3) Ascertainment of exposure

- a) secure medical record ✱
- b) structured interview
- c) written self-report
- d) no description

4) Demonstration that outcome of interest was not present at start of study

- a) yes ✱
- b) no

Comparability

Comparability of cohorts on the basis of the design or analysis

- 1) study controls for demographic characteristics including gestational age, sex, birth weight and ethnicity, *etc* ✱
- 2) study controls for characteristics of central lines including types of central lines and length of catheterization, *etc* ✱

Exposure/Outcomes

1) Ascertainment of exposure

- a) secure record with details ✱

- b) structured interview where blind to case/control status
 - c) interview not blinded to case/control status
 - d) written self-report or medical records
 - e) no description
- 2) Clear definition and report of outcomes
- a) yes, based on consensus clinical criteria ✱
 - b) yes, based on self-definition
 - c) no description
- 3) Was follow-up long enough for outcomes to occur
- a) yes (select an adequate follow up period for outcome of interest) ✱
 - b) no
- 4) Adequacy of follow up of cohorts
- a) complete follow up - all subjects accounted for ✱
 - b) subjects lost to follow up unlikely to introduce bias
 - c) subjects lost to follow up likely to introduce bias and no description of those lost
 - d) no statement