Supplemental Table 3.1 Data extraction sheet for RCTs

| | | | Public | cation and study | details | | | | | | | |
|--------------------------|------------|--------------|---------------------|------------------|--------------------------|---------|---------------|--|--|--|--|--|
| Authors | | | | | | | | | | | | |
| Year of | | | | | | | | | | | | |
| publication | | | | | | | | | | | | |
| Country | | | | | | | | | | | | |
| Study design | | | | | | | | | | | | |
| Number of | | | | | | | | | | | | |
| participants | | | | | | | | | | | | |
| Groups | | | Experimental | | Control | | | | | | | |
| | | plat | elet count thres | hold | platelet count threshold | | | | | | | |
| | | | $(\times 10^{9}/L)$ | | (×10 ⁹ /L) | | | | | | | |
| | | | | | | | | | | | | |
| Clinical characteristics | | | | | | | | | | | | |
| | Experin | nental | Experimental | Experimental | Control | Control | Control | | | | | |
| | median (or | | IQR (or SD) | total | median (or | IQR (or | total | | | | | |
| | mean) | | | | mean) | SD) | | | | | | |
| GA(w) | | | | | | | | | | | | |
| BW (g) | | | | | | | | | | | | |
| Platelet | | | | | | | | | | | | |
| count | | | | | | | | | | | | |
| $(\times 10^{9}/L)$ | | | | | | | | | | | | |
| Number of | | | | | | | | | | | | |
| platelet | | | | | | | | | | | | |
| transfusions | | | | | | | | | | | | |
| | | |] | Primary outcom | ies | | | | | | | |
| | | Experimental | | Experimental | Control | | Control | | | | | |
| | | event | | total | event | | total | | | | | |
| In-hospital mortality | | | | | | | | | | | | |
| or major bleeding | | | | | | | | | | | | |
| events | | | | | | | | | | | | |
| | | | | Bleeding episod | es | | | | | | | |
| IVH | | | | | | | | | | | | |
| ICH | | | | | | | | | | | | |
| PH | | | | | | | | | | | | |
| Frank rectal b | leeding | | | | | | | | | | | |
| Other bleeding | | | | | | | | | | | | |
| | | | S | econdary outco | mes | | | | | | | |
| | | | | Major morbidit | y | | | | | | | |
| | | Experimental | | Experimental | Control event | | Control total | | | | | |
| | | | event | total | | | | | | | | |
| PDA | | | | | | | | | | | | |
| | | | | | | | | | | | | |

| BPD | | | | | | | | | | | | | |
|------------------------|----------------|--|--------------|---------------|---------|---------------|---------|--|--|--|--|--|--|
| Sepsis | | | | | | | | | | | | | |
| NEC | | | | | | | | | | | | | |
| ROP | | | | | | | | | | | | | |
| Other outcome measures | | | | | | | | | | | | | |
| LOS (days) | Experimental | | Experimental | Experimental | Control | Control | Control | | | | | | |
| | median (or | | IQR (or SD) | total | median | IQR (or | total | | | | | | |
| | mean) | | | | (or | SD) | | | | | | | |
| | | | | | mean) | | | | | | | | |
| | | | | | | | | | | | | | |
| Adverse effects of | f Experimental | | Experimental | Control event | | Control total | | | | | | | |
| transfusion | event | | total | | | | | | | | | | |
| | | | | | | | | | | | | | |
| Other information | | | | | | | | | | | | | |
| Type and dose of | | | | | | | | | | | | | |
| platelet component | | | | | | | | | | | | | |
| Any sponsorship or | | | | | | | | | | | | | |
| funding | | | | | | | | | | | | | |

IQR: interquartile range; SD: standard deviation; GA: gestational age; BW: birth weight; IVH: intraventricular haemorrhage; ICH: intracranial haemorrhage; PH: pulmonary haemorrhage; BPD: bronchopulmonary dysplasia; NEC: necrotizing enterocolitis; PDA: patent ductus arteriosus; NEC: necrotizing enterocolitis; ROP: retinopathy of prematurity; LOS: length of stay