

Supplementary Table 1. Time to first platelet response outcome among the included studies

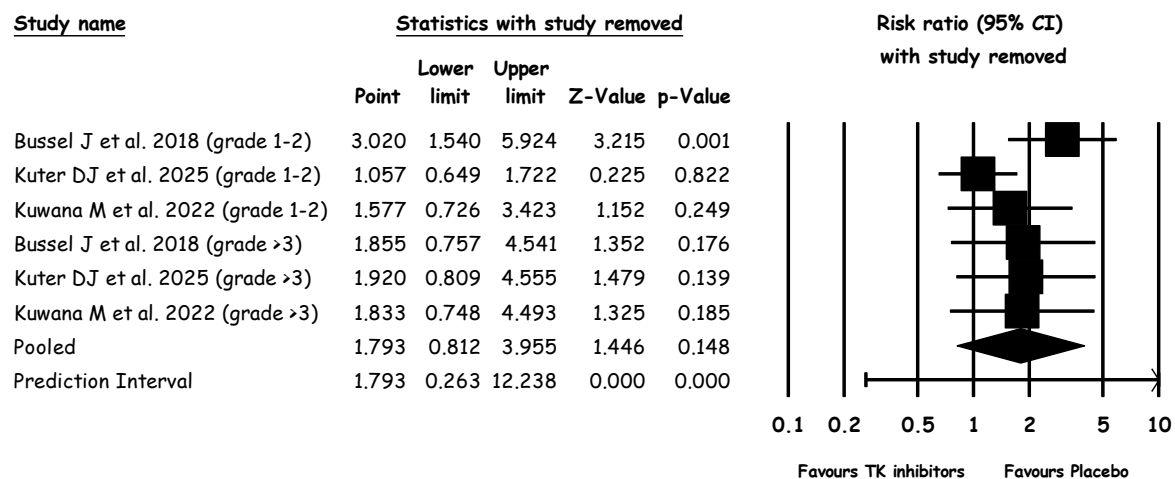
Study name	Time to First Platelet Response	
	TK inhibitors	Placebo
Bussel J et al. ^[15] 2018	Median time was 15 days for overall responders and 15.5 days for stable responders	Not reported
Hu Y et al. ^[16] 2024	Median time = 8 (8 – 12) days	Median time = 30 (24 – 46) days
Kuter DJ et al. ^[17] 2025	Median time = 36 days	Never achieved

TK inhibitors = tyrosine kinase inhibitors

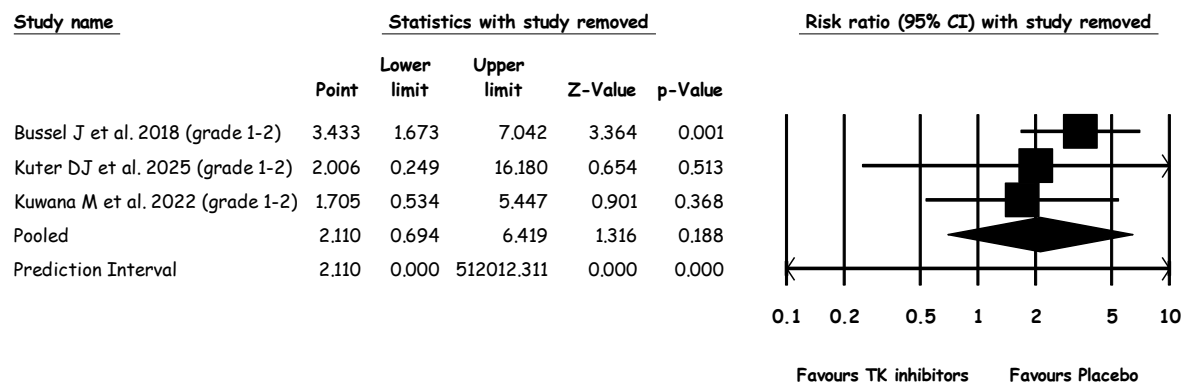
Supplementary Table 2. Bleeding events outcome among the included studies

Study name	Bleeding events	
	TK inhibitors	Placebo
Bussel J et al. ^[15] 2018	<ul style="list-style-type: none"> - Moderate or severe bleeding-related AEs occurred in 10/101 patients (9.9%) - Bleeding-related SAEs occurred in 4/101 patients (3.9%) 	<ul style="list-style-type: none"> - Moderate or severe bleeding-related AEs occurred in 8/49 (16%) patients - Bleeding-related SAEs occurred in 5/49 patients (10%)
Hu Y et al. ^[16] 2024	<ul style="list-style-type: none"> - Total bleeding events reduced significantly from 87/126 patients (69%) at baseline to 23/86 patients (27%) at week 24 	<ul style="list-style-type: none"> - Total bleeding events reduced significantly from 33/62 patients (53%) at baseline to 1/8 patients (13%) at week 24
Kuter DJ et al. ^[17] 2025	<ul style="list-style-type: none"> - Grade ≥ 2 bleeding events due to any cause were reported in 4% of patients 	<ul style="list-style-type: none"> - Grade ≥ 2 bleeding events due to any cause were reported in 12% of patients
Kuwana M et al. ^[18] 2023	<ul style="list-style-type: none"> - Bleeding events occurred in 3/22 patients (14%) 	<ul style="list-style-type: none"> - Bleeding events occurred in 2/12 patients (17%)

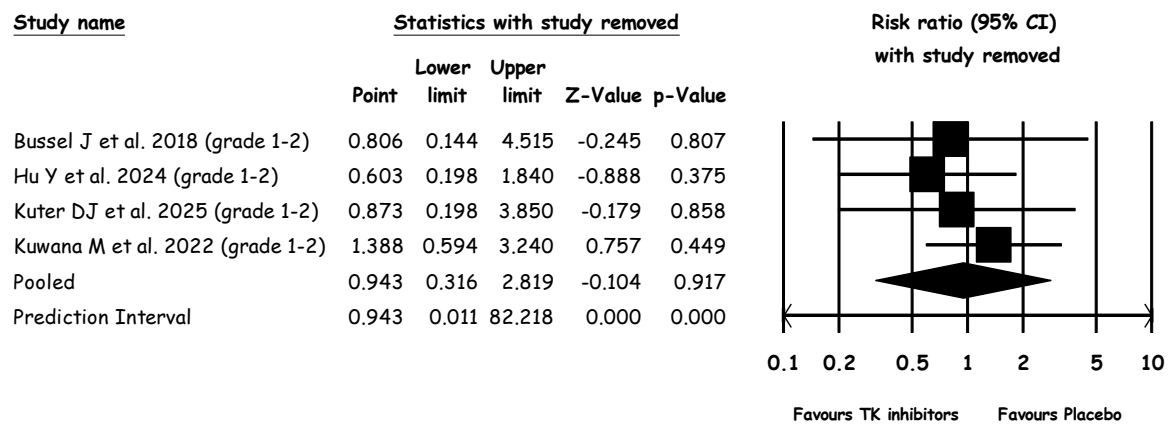
AEs = adverse events; SAEs = serious adverse events; TK inhibitors = tyrosine kinase inhibitors



Supplementary Figure 1. Sensitivity analysis by leave-one-out method for the overall diarrhea incidence outcome.



Supplementary Figure 2. Sensitivity analysis by leave-one-out method for the grade 1-2 diarrhea outcome.



Supplementary Figure 3. Sensitivity analysis by leave-one-out method for the grade 1-2 upper respiratory infection outcome.