

SCB-219M

Preliminary Phase 1 Data

December 29th, 2023

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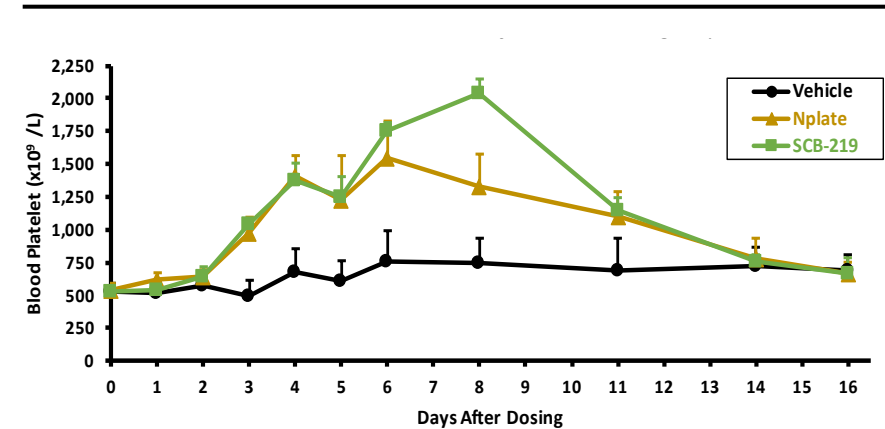
SCB-219M (TPO Mimetic Bispecific-Fc)

- ✓ SCB-219M is a novel fusion protein (TPO mimetic bispecific-Fc) in Phase 1 Clinical Testing
- ✓ Initially targeted to treat **Chemotherapy-Induced Thrombocytopenia (CIT)**

Potential Significant Differentiation & Advantages Compared to Commercially-Available Native TPO-Based Therapy in China

- ✓ **Potent & Durable Efficacy:** SCB-219M may potentially overcome reduced efficacy observed for native TPO therapy due to anti-drug antibodies (ADA)
- ✓ **More Convenient Dosing:** SCB-219M's longer half-life may enable it to achieve a more convenient dosing regimen compared to both native TPO-based therapy and Nplate (romiplostim)
- ✓ **Blockbuster Market Potential:** Product sales for native TPO-based therapy (TPIAO) in China reached **over RMB 3 billion** in 2022
- ✓ Opportunities for near-term **value creation via development & commercial partnerships in China and globally** for SCB-219M to be evaluated

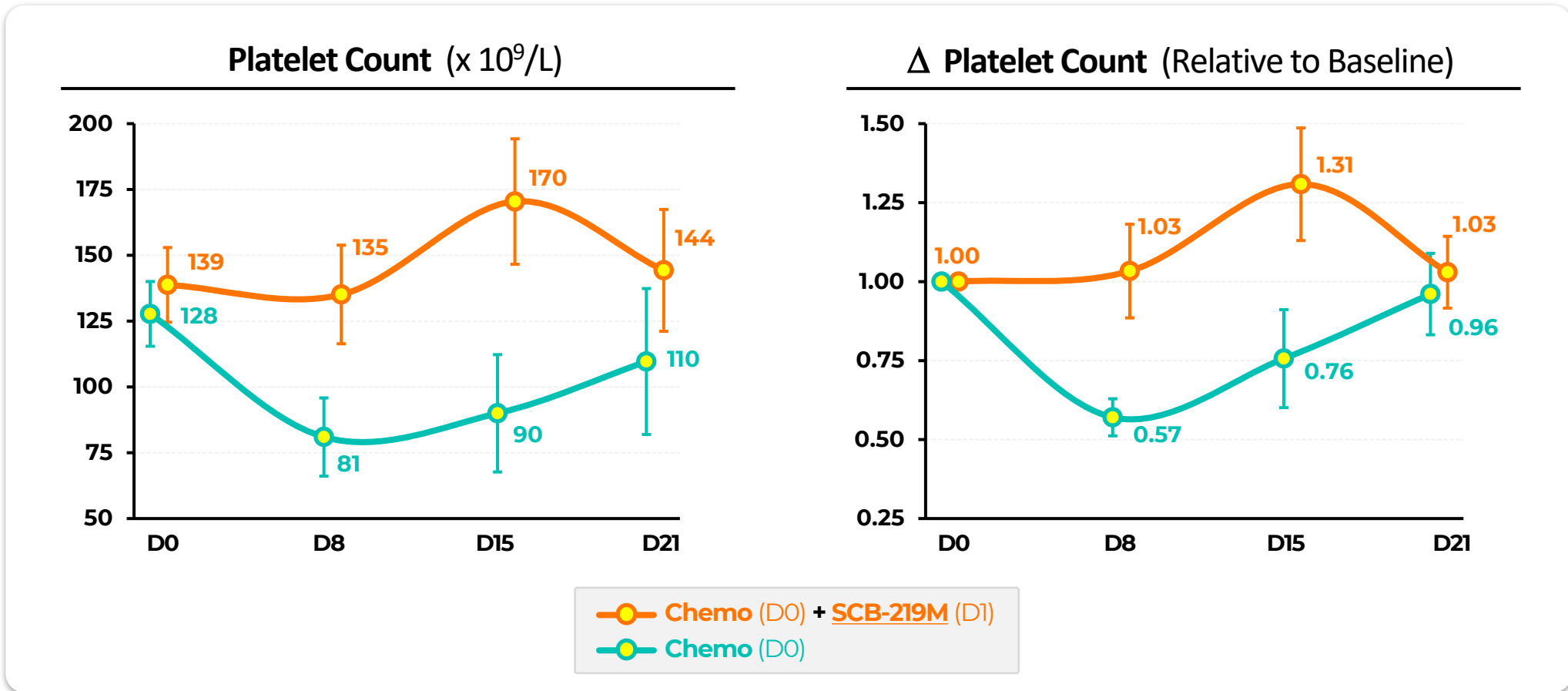
Blood Platelet Level in SD Rats



Phase 1 Clinical Trial Data Readout in Chemotherapy-Induced Thrombocytopenia (CIT) Announced in **DEC-2023**

Preliminary Phase 1 Data: *Efficacy*

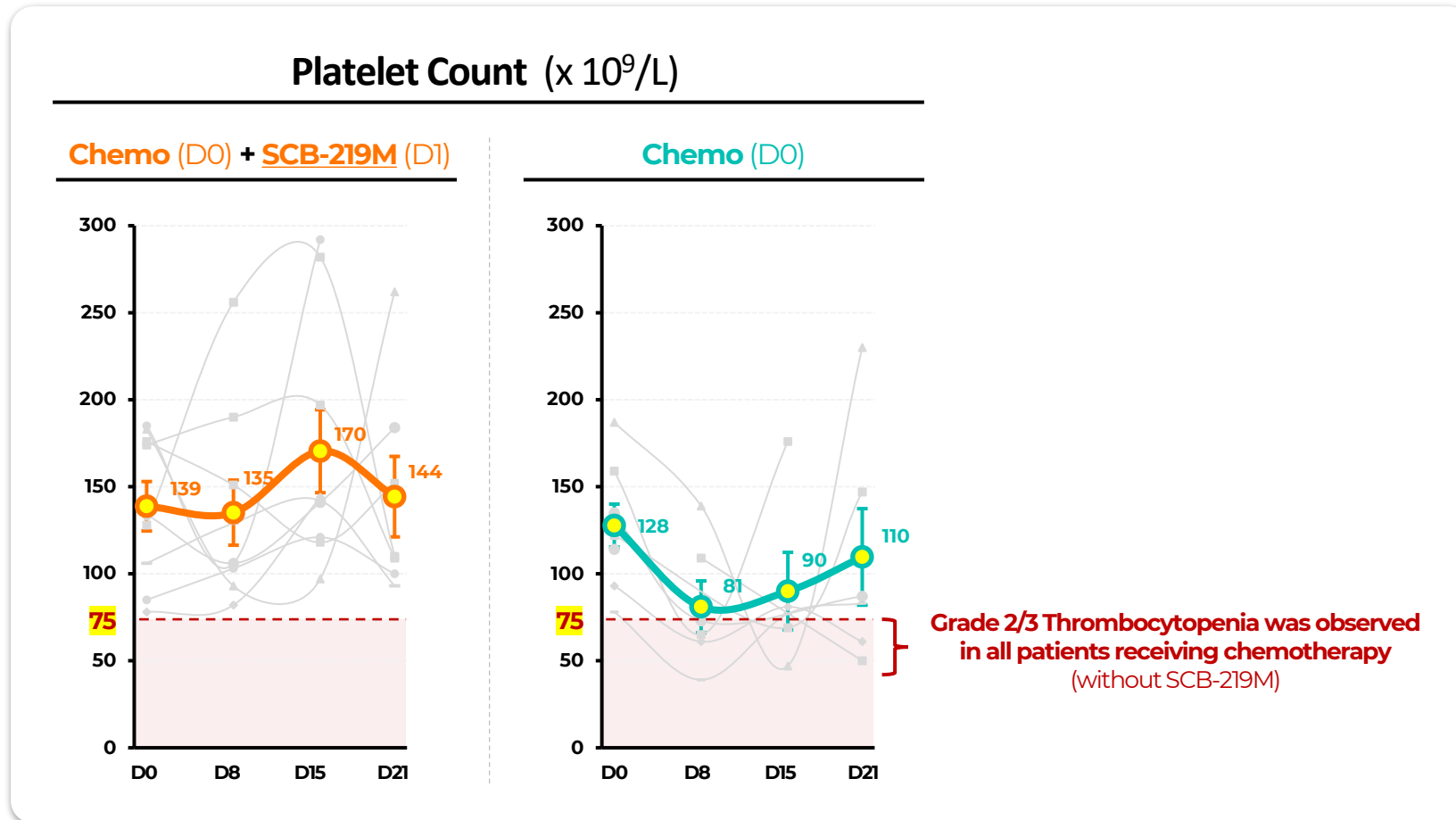
- ✔ Significant platelet count maintenance/recovery observed in CIT patients following chemotherapy (on Day 0) plus **a single dose of SCB-219M** (on Day 1)
- ✘ Compared to chemotherapy-alone (without SCB-219M), platelet counts dropped by >40% versus baseline (in the same patients prior to study enrollment)



Note: Preliminary SCB-219M Phase 1 results in 9 CIT patients (data not final and subject to change). Chemotherapy infusion administered on Day 0 (D0), and SCB-219M administered subcutaneously on Day 1 (D1). Mean values ± Standard errors (SE) shown (where available).

Preliminary Phase 1 Data: *Efficacy*

- ✓ **All CIT patients** enrolled maintained platelet counts $>75 \times 10^9/L$ at 1-week following chemotherapy (on Day 0) plus **a single dose of SCB-219M** (on Day 1), with **durable responses** through at least 3-weeks
- ✗ In comparison, following **chemotherapy-alone** (without SCB-219M) in the same patients prior to enrolling into the trial, all patients observed platelet counts drop to $<75 \times 10^9/L$ (i.e. Grade 2 or 3 thrombocytopenia) between one and three weeks



Note: Preliminary SCB-219M Phase 1 results in 9 CIT patients (data not final and subject to change). Chemotherapy infusion administered on Day 0 (D0), and SCB-219M administered subcutaneously on Day 1 (D1). Mean values \pm Standard errors (SE) shown (where available). Grey lines represent individual CIT patients.

Preliminary Phase 1 Data: *Summary*

Durable Preliminary Efficacy & PK Profile

- ✓ Durable preliminary efficacy and pharmacokinetic profile observed for SCB-219M are potentially supportive of dosing intervals ≥ 2 -weeks
- ✓ If further confirmed, this profile could enable convenient dosing of SCB-219M synchronized with any given patient's chemotherapy regimen, typically 2-3 weeks per cycle.
 - In contrast to current standard of care biologic treatments for CIT in China requiring daily injections⁽¹⁾ and globally requiring weekly injections⁽²⁾

Safety Profile

- ✓ Favorable safety and tolerability profile for SCB-219M observed to-date
- ✓ No serious adverse events (SAEs)
- ✓ No dose-limiting toxicity (DLT)

Phase Ib trial evaluating repeated dosing of SCB-219M in CIT and cancer therapy-induced thrombocytopenia (CTIT) patients is planned to initiate in 2024

Note: Preliminary SCB-219M Phase 1 results in 9 cancer patients (data not final and subject to change).

(1) TPIAO (3SBio; <https://ypk.39.net/666055/manual>).

(2) Nplate; romiplostim (<https://doi.org/10.3324/haematol.2020.251900>; <https://doi.org/10.1200/JCO.18.01931>).

Thank You!