## A. In vivo therapeutic efficacy study

## Sample collection and treatment: Subjects enrolled on D0, administered CQ (N = 125) Withdraws (n = 5): Loss to follow up (n = 13)Completed 28-day follow up, administered PQ (n = 107) Pv+ on D0 only (n = 74, 74 isolates) Pv+ on D0, 2 only (n = 15, 30 isolates) Pv+ on D0, 2, 7 only (n = 1, 2 isolates) Pv+ on D0, 2, 7, Rec (n = 1, 3 isolates) PQ completed (n = 1) Pv+ on D0, 2, Rec (n = 2, 6 isolates) PQ completed (n = 2) Pv+ D0 and Rec (n = 14, 28 isolates) PQ completed (n = 7)PQ not completed (n = 7)Genotype processing: Selected for aenotyping (D0, n = 33; D2, n = 19; DRec, n = 17) Pvmdr1 sequencing (n = 69) Failed to amplify (D2, n = 19; DRec, n = 2) 958M/Y976/1076L (D0, n = 33; DRec, n = 14) 958M/Y976/F1076 (DRec, n = 1) MS fragment analysis (n = 50)

Failed to amplify (D2, n = 19) Full allelic haplotypes (D0, n = 33: DRec. n = 17)

## B. In vitro drug assay

## Sample collection and treatment:

