

**Supplemental digital content 3. Non-fatal serious adverse events reported from administration of dose 1 up to study end (Total vaccinated cohort)**

<b>Group</b>	<b>n</b>	<b>Serious adverse event description</b>
<6S	1	Malaria and septicemia
	1	Malaria and gastroenteritis
<6NS	4	Gastroenteritis
	2	Malaria and gastroenteritis
	1	Malaria and pneumonia
	1	Urinary tract infection
7-11S	1	Gastroenteritis
	1	Septicemia
7-11NS	2	Gastroenteritis
	1	Malaria and bronchitis
	1	Enteritis
12-23S	1	Osteomyelitis
	1	Malaria and pneumonia
12-23NS	1	Malaria and septicemia
	1	Malaria and gastroenteritis

**Footnote:** <6S, infants with SCD enrolled at 8–11 weeks of age, primed with 3 doses and boosted with PHiD-CV + DTPw-HBV/Hib + OPV vaccines at 2, 3, 4 and 9–10 months of age; <6NS, infants without SCD enrolled at 8–11 weeks of age, primed with 3 doses and boosted with PHiD-CV + DTPw-HBV/Hib + OPV vaccines at 2, 3, 4 and 9–10 months of age; 7-11S, children with SCD enrolled at 7–11 months of age, primed with 2 doses and boosted with PHiD-CV; 7-11NS, children without SCD enrolled at 7–11 months of age, primed with 2 doses and boosted with PHiD-CV ; 12-23S, children with SCD enrolled at 12–23 months of age, vaccinated with 2 doses of PHiD-CV; 12-23NS, children without SCD enrolled at 12–23 months of age, vaccinated with 2 doses of PHiD-CV; n, number of children reporting the serious adverse event.