Supplementary Table 4: Adverse events.

	ASMQ	AL	DP	AL
Number of patients	**n = 59**	**n = 59**	**n = 62**	**n = 56**
Anemia	15(25.4%)	9(15.3%)	7(11.3%)	10(17.9%)
Diarrhea	0	0	1(1.6%)	0
Abdominal pain	1(1.7%)	0	1(1.6%)	0
Gastroenteritis	1(1.7%)	0	2(3.2%)	0
Tinea capitis	5(8.5%)	3(5.1%)	4(6.5%)	4(7.1%)
Respiratory tract infection	3(5.1%)	15(25.4%)	12(19.4%)	9(16.1%)
Rash	2(3.4%)	1(1.7%)	0	3(5.4%)
Dizziness	1(1.7%)	0	0	0
Fatigue	0	1(1.7%)	0	0
Headache	1(1.7%)	1(1.7%)	5(8.1%)	1(1.8%)
Fever	0	0	1(1.6%)	0
Nausea	1(1.7%)	0	0	0

Some of the adverse events that occurred in study participants. None of the study participant vomited, all adverse events were mild, at low frequency and resolve spontaneously.