A study to assess Clinician-perceived failure rates of commonly used ACTs

Interviewer's Name:	Code:
Region:	Code:
Health Facility Name:	Code:
Date of Interview:/	
Clinician-perceived ACT Treatment Failure despite having received an Artemisinin Comb artemisinin resistance.	
SECTION A: DEMOGRAPHICS	
1. Gender: [1] Male [2] Female	
2. Age (in completed years):	
3. Education Level:	
[1] Certificate	7. Health Facility Type:
[2] Diploma	[1] Public
[3] Bachelor	[2] Private Not-for-Profit
[4] Masters	[3] Private for-Profit
[5] Other (specify)	
	8. Health Facility Status:
4. Professional experience (Years):	[1] Hospital
E If less than 1 year in O4 state	[2] Health Centre III
5. If less than 1 year in Q4 , state number of completed months	[3] Health Centre II
0.00	[4] Private Clinic
6. Professional Cadre:[1] Physician	[5] Pharmacy
[2] Medical Officer	[6] Drug Shop
[3] Pharmacist	[7] Other, (specify)
[4] Nurse	
[5] Clinical Officer	
[6] Pharmacy Technician	
[7] Other (specify)	I

SECTION B: PERCEIVED ACT FAILURE

Ple	ase, complete the questionnaire by indicating the appropriate responses.
1.	What is the approximate number of malaria-patients you see per day?
	In the use of ACTs in treating uncomplicated malaria, have you ever encountered any treatment failure(s) in your malaria-patients? [1] Yes [2] No
3.	Have you suspected any ACT treatment failure in the <u>past 4 weeks</u> ? [1] Yes [2] No
4.	If YES to Q3 , how many cases of ACT treatment failure?
	Have you received patient-complaints of ACT treatment failure in the <u>past 4</u> weeks? [1] Yes [2] No
6.	If YES to Q5 , how many patient-complaints of ACT treatment failure?
7	Briefly describe the most recent case of ACT treatment failure you have
	encountered providing information on patient age, brand of ACT involved, clinical outcome & action taken; e.t.c.
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	outcome & action taken; e.t.c. Have you reported any ACT treatment failure(s) in the past 6-months? (<i>Please</i>
8.	Outcome & action taken; e.t.c. Have you reported any ACT treatment failure(s) in the past 6-months? (<i>Please tick one</i>)
8.	outcome & action taken; e.t.c. Have you reported any ACT treatment failure(s) in the past 6-months? (<i>Please tick one</i>) [1] Yes [2] No (<i>Skip to 13</i>) If <i>YES</i> to Q8, to whom have you reported the most recent ACT treatment

11. What motivates you to report ACT treatment failure(s)?				
12. Do you get feedback on the Ar [1] Yes [2] No	CT treatment failure(s) you report?			
13. Do you feel that circumstance failure to ACTs?	s in your setting make it difficult to report treatment [1] Yes [2] No			
14. Explain your response to Q13				
15. What can be done to improve setting?	the reporting of treatment failure to ACTs in your			
appropriate)	ed ACTs at your health facility? (<i>Please tick all</i>			
Brand	Coartem			
D-Artepp (GPSC)	Lumartem			
Artequin	Malfan			
Combiart	Artem			
Ridmal	Arexel			
Glumac	Lonart			
Duocotecxin	Lumether			
P-Alaxin	Lumaren			
Artefan	Other, (specify)			
17. Which ACT brand(s) have you of tick all appropriate)	observed to result in treatment failure? (Please			
Brand	Coartem			
D-Artepp (GPSC)	Lumartem			
Artequin	Malfan			
Combiart	Artem			
Ridmal	Arexel			
Glumac	Lonart			
Duocotecxin	Lumether			
P-Alaxin	Lumaren			
Artefan	Other, (specify)			

18.Do you think ACT [1] Yes	resistance is a gr [2] No	rowing concern nationally? [9] Don't Know
19. If yes to Q18, brie	afly describe why?	
20. Do you think AC ⁻ [1] Yes	T resistance is a g [2] No	rowing concern in your institution? [9] Don't Know
21. If yes to Q20 , brie	ofly describe why?	
SECTION C: DRUG	FACTORS RELATE	ED TO ACT FAILURE
22 Do you think the	polor of an ACT a	ould load to poor nationt compliance haves
treatment failure?		ould lead to poor patient compliance hence [2] No
23. Briefly describe w	yhy giving eyemple	267
		
24. Do you think the t treatment failure?		ould lead to poor patient compliance hence [2] No
05. Priofly describe w	thy giving avample	
25. Briefly describe w	my giving example	55 f
26. Do you think the s	size of an ACT tab	olets could lead to poor patient compliance
hence treatment f		
27. Briefly describe w	hy giving example	es?
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28. Do you think the number of tablets swallowed could lead to poor patient compliance hence treatment failure? [1] Yes [2] No
29. Briefly describe why giving examples?
30. Do you think that inadequate information about an ACT could lead to patient misuse of the drug hence treatment failure? [1] Yes [2] No
31. Briefly describe why giving examples?
32. Do you think the dosing frequency of an ACT could lead to poor patient compliance hence treatment failure? [1] Yes [2] No
33. Briefly describe why giving examples?
34. What other factors in the practice of <u>patients</u> are responsible for the poor response of patients to ACT? (<i>Please briefly outline</i>)
35. What other factors in the practice of <u>clinicians</u> are responsible for the poor response of patients to ACT? (<i>Please briefly outline</i>)

We appreciate your time taken to respond to this questionnaire. Thank you

This a collaborative study between
National Drug Authority and Makerere University Department of Pharmacology and therapeutics.