BMJ Global Health

Antimalarial stocking decisions among medicine retailers in Ghana: implications for quality management and control of malaria

Adams Osman,¹ Fiifi Amoako Johnson,² Simon Mariwah,³ Daniel Amoako-Sakyi,⁴ Samuel Asiedu Owusu,⁵ Martins Ekor,⁴ Heather Hamill ⁽ⁱ⁾, ⁶ Kate Hampshire⁷

ABSTRACT

To cite: Osman A, Amoako Johnson F, Mariwah S, *et al.* Antimalarial stocking decisions among medicine retailers in Ghana: implications for quality management and control of malaria. *BMJ Glob Health* 2023;**6**:e013426. doi:10.1136/ bmjgh-2023-013426

Additional supplemental material is published online only. To view, please visit the journal online (http://dx.doi.org/10. 1136/bmjgh-2023-013426).

Received 14 July 2023 Accepted 12 August 2023

Check for updates

© Author(s) (or their employer(s)) 2023. Re-use permitted under CC BY. Published by BMJ.

For numbered affiliations see end of article.

Correspondence to

Professor Heather Hamill; heather.hamill@sociology.ox. ac.uk

Global health efforts such as malarial control require efficient pharmaceutical supply chains to ensure effective delivery of quality-assured medicines to those who need them. However, very little is currently known about decision-making processes within antimalarial supply chains and potential vulnerabilities to substandard and falsified medicines. Addressing this gap, we report on a study that investigated decision-making around the stocking of antimalarial products among private-sector medicine retailers in Ghana. Licensed retail pharmacies and over-the-counter (OTC) medicine retail outlets were sampled across six regions of Ghana using a twostage stratified sampling procedure, with antimalarial medicines categorised as 'expensive,' 'mid-range,' and 'cheaper,' relative to other products in the shop. Retailers were asked about their motivations for choosing to stock particular products over others. The reasons were grouped into three categories: financial, reputation/ experience and professional recommendation. Reputation/ experience (76%, 95% CI 72.0% to 80.7%) were the drivers of antimalarial stocking decisions, followed by financial reasons (53.2%, 95% Cl 48.1% to 58.3%) and recommendation by certified health professionals (24.7%, 95% CI 20.3% to 29.1%). Financial considerations were particularly influential in stocking decisions of cheaper medicines. Moreover, pharmacies and OTCs without a qualified pharmacist were significantly more likely to indicate financial reasons as a motivation for stocking decisions. No significant differences in stocking decisions were found by geographical location (zone and urban/rural) or outlet (pharmacy/OTC). These findings have implications for the management of antimalarial quality across supply chains in Ghana, with potentially important consequences for malaria control, particularly in lower-income areas where people rely on low-cost medication.

INTRODUCTION

Well-governed pharmaceutical supply chain networks are integral to health systems and a critical enabler of universal health coverage (UHC), helping to ensure timely access to quality, safe and effective essential medicines.

WHAT IS ALREADY KNOWN ON THIS TOPIC

⇒ Malarial control requires efficient pharmaceutical supply chains to ensure effective delivery of quality-assured medicines to those who need them. However, very little is known about decision-making processes within antimalarial supply chains and potential vulnerabilities to substandard and falsified (SF) medicines.

WHAT THIS STUDY ADDS

- ⇒ We confirm the uneven distribution of licensed medicine outlets across Ghana; rural areas and the less affluent north of Ghana have much poorer access to essential medicines.
- ⇒ Private-sector medicine outlets are motivated to stock quality medicines rather than by purely financial considerations.
- ⇒ Financial considerations were most important for the cheapest products and for outlets without a pharmacist.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

- ⇒ Private-sector outlets are crucial to the delivery of primary care to underserved communities.
- ⇒ All medicine outlets must have trained staff to make informed stocking decisions.
- ⇒ Ensuring access to affordable quality-assured antimalarials is key in both the fight against SF medicines and in moving towards genuine universal health coverage.

However, the proliferation of substandard and falsified (SF) medicines in low-income and middle-income countries (LMICs) suggests weaknesses within supply chains that threaten global health efforts such as malaria control and the progress towards UHC.^{1–3} Africa has the highest reported prevalence of poor-quality medicines, with antimalarials (19.1%) being the most affected drug class.³ Like any other African country, poor-quality medicines in Ghana are a major health threat, but the

BMJ

challenge has more to do with substandard than falsified products, where studies on the prevalence of substandard medicine products reported varied results. For example, Wilson *et al*⁴ and Tivura *et al*,⁵ respectively. found that 37% and 35% of antimalarials were substandard fueled by uncontrolled/managed borders, high drug demand, corruption, limited political will, poor coordination among institutions/countries, poverty, and illiteracy.⁶

We have observed that national and international efforts to curb the SF medicine menace largely focus on the public sector. However, in many LMICs, public, private and informal medicine distribution networks intertwine to form complex, fragmented and poorly regulated supply chains.^{7 8} Many of the world's poor continue to source essential medicines from private-sector (including 'informal') outlets, owing to the inaccessibility of public facilities and frequent medicine stock-outs.⁹ Existing literature suggests that private-sector actors might sometimes behave (deliberately or otherwise) in ways that undermine supply chain robustness with potentially dire consequences for affordability, quality (SF issues) and availability of medicines.¹⁰⁻¹⁴ Indeed, there is now a significant body of evidence showing that stock-outs, cost and imbalances in demand and supply drive the emergence of SF medicines.^{15 16}

This paper reports findings from a wider research project tracking decision-making through antimalarial supply chains in Ghana, from retail outlets up to manufacturers and importers. The focus here is on the retail outlets: the 'last mile' of medicine supply, whose decisions may directly impact on availability, quality, type, affordability of medicines to households and efficient flow of medicines to areas needed.^{10–13} Thus, the study sought to determine whether, in addition to quality considerations, retailers are influenced by other factors such as the financial cost of medicine and recommendations from health practitioners or suppliers. We predict that outlets in more economically constrained contexts will be more driven by financial imperatives and have less capacity to consider likely drug quality (based on outlets self-assessment of brand reputation, recommended efficacy by health specialists and personal experience rather than active ingredients), than those in more economically favourable contexts. More specifically, we hypothesise that:

- ► (H1) Outlets in more remote and poorer areas of the country (i.e., northern, rural) are more likely to be driven by financial considerations and less likely to be influenced by considerations of likely quality in antimalarial stocking decisions, compared with those located in more densely populated and wealthier areas of the country (i.e., southern/central, urban).
- ► (H2): Over-the-counters (OTCs) are more likely to be driven by financial considerations and less likely to be influenced by considerations of quality in antimalarial stocking decisions, compared with pharmacies because they are mostly in rural areas and low-income areas.

- BMJ Glob Health: first published as 10.1136/bmjgh-2023-013426 on 21 September 2023. Downloaded from http://gh.bmj.com/ on February 8, 2024 by guest. Protected by copyright
- (H3) For cheaper medicines, financial considerations are likely to be more important and quality less important in antimalarial stocking decisions, compared with more expensive medicines.

Conceptual issues: antimalarial supply chain and SF medicines

After two decades of massive gains in the fight against malaria, recent data are warning of advances tapering-off. An estimated 14 million more cases and 47 000 more malaria deaths occurred in 2020 relative to 2019; an increase that persists even after adjusting for COVID-19-induced disruption of malaria services.¹⁷ Important global malaria strategy milestones for 2020 have been missed, and there is a risk of missing the 2030 target entirely if bold, immediate and smart actions are not taken to restore and improve gains.

National malaria control strategies for many endemic countries hinges on preventive chemotherapies such as intermittent preventive treatment of malaria in pregnancy, perennial malaria chemoprevention and seasonal malaria chemoprevention.¹⁸¹⁹ Other preventive chemotherapies used depend on country-specific needs and may include post-discharge malaria chemoprevention for children at risk of malaria and recently treated for severe anaemia, and intermittent preventive treatment of malaria for school-aged children. The recommended medicines for these programmes rely on the pharmaceutical supply chains (sometimes including private sector actors) to reach the target populations. Thus, it is important for stakeholders to thoroughly understand the antimalarial supply chain specifically and safeguard it from vulnerabilities that promote the proliferation of SF antimalarials.

Ghana's pharmaceutical supply chain resembles that of other LMICs in some ways but the local business environment, client preferences and policy initiatives such as the National Health Insurance Scheme (drugs are either subsidised or completely covered) are key distinguishing features. To date, very few studies have looked at the antimalarial supply chain in Ghana and none explores the relationship between various actors in the supply chain and the emergence of SF antimalarials.²⁰ The ongoing threat of malaria and the limitations of current scholarship on antimalarial supply chains in Ghana suggests our study is timely and relevant in addressing a key threat to malaria control gains.

METHODS

Study context

Ghana is a West Africa country divided into 16 administrative regions across 3 main agro-ecological zones (see figure 1), with a population of about 30.8 million.²¹ Ghana's economy is classified as lower-middle income but with high multidimensional poverty (income, energy, food, formal education, health), persistent



Figure 1 Sampled areas per regions in Ghana.

especially in rural areas and northern/savannah part of the country. $^{\rm 22\,23}$

Ghana's climate (temperature: 24°C–32°C, rainfall: 780mm–2160 m annually) and ecology (forest, dry equatorial, and savannah) make it highly susceptible to malaria and some other vector borne diseases. Malaria is responsible for over 38% of outpatient visits and 27.3% of health facility admissions,²⁴ and for about 25% of under-5 deaths in Ghana.²⁵ Ghana's health system faces multiple challenges: limited healthcare facilities, uneven spatial distribution with concentration in the south (forest and coastal zones), limited health personnel, an inefficient health insurance scheme and high cost of medicines.^{26 27} Medicine distribution and sales are mainly privately owned, with limited quality control measures, which have implications for the proliferation of SF medicines in the country.⁴

Study design

The data for the study are derived from a survey conducted in 2021 as part of the 'STREAMS' project (Strengthening private-sector Medicine Systems to tackle the persistence of poor-quality medicines in Africa), funded by the UK Medical Research Council. The research was codesigned by all authors at a workshop in Accra in March 2020. Data were collected from licensed retail pharmacies and OTC medicine retail outlets across Ghana to explore decision-making around the stocking of antimalarials. The study design was descriptive in nature, with the aim of describing the characteristics of sampled retail outlets and determining the factors influencing decisions to stock specific types and brands of antimalarials. Comparisons were made between outlets in different locations (border/non-border; rural/urban and across regions). Specifically, the study adopted a case series design, where

AMA, Accra Metropolitan Area; KMA, Kumasi M

Table 1

Zone

Forest

Coastal

Total

Savannah

_									
1	Number of registered	and sampled	pharmacies and (DTCs					
				Number of registered			No of sampled		
	Border/non-border	Rural/urban	District	Pharmacies	OTCs	Total	Pharmacies	OTCs	Total
1	Border	Urban	Bawku Municipal	5	43	48	5	15	20
	Non-border	Rural	Central Gonja	2	22	24	2	18	20
	Border	Rural	Bia East	2	26	28	2	18	20
	Non-border	Urban	KMA	422	421	843	10	10	20
	Border	Rural	Ketu South	9	80	89	9	11	20
	Non-border	Urban	AMA	623	314	937	10	10	20
				1063	906	1969	38	82	120
a i au	mple of retail out e basic understand n Ghana (given th nd to provide data tive studies.	tlets were st ling about st le current pa to inform fu	udied to help ocking of anti- aucity of infor- ture nationally	(location, products (assisted po minimise i	type, siz (see belor ersonal in non-samp	e, person w) were r nterviewin ling error	nel, etc.) an ecorded usir ng tool (Kob s.	d antii ig a co: oToolB	malarial mputer- Box), to
procedure ge stratified sampling design was adopted to nat the data collected were robust and guar- fficient use of resources. The sampling frame list of pharmacies (2879) and OTCs (12249) from the Ghana Pharmacy Council as of 2019. rst stage of the sampling process, the country fied into the three zones—Savannah/Northern				The individual responsible for stocking decisions in each outlet was asked to list all the oral medicines currently in stock for treating acute malaria in adults, and then to identify three of their best-selling products: one more expensive, one relatively cheap, and one 'mid-range'. (Note that these were relative assessments rather than absolute—that is, an 'expensive' product in one outlet might be a 'shaap' one in another). For each selected					

a small sample of retail outlets were st provide the basic understanding about st malarials in Ghana (given the current pa mation) and to provide data to inform fu representative studies.

Sampling procedure

A two-stage stratified sampling design w ensure that the data collected were rol anteed efficient use of resources. The sa was the list of pharmacies (2879) and obtained from the Ghana Pharmacy Cour At the first stage of the sampling proces was stratified into the three zones-Savan (Savannah, Northern, North East, Upper East and Upper West regions), Forest/Middle (Western North, Ashanti, Ahafo, Eastern, Oti, Bono East and Bono regions) and Coastal/Southern (Western, Central, Greater Accra and Volta regions). These ecological zones are the statistical aggregation and reporting zones used by the Ghana Statistical Service.

At the second stage, the zones were stratified into border and non-border districts, because previous works in Africa and Ghana suggested that proximity to an international border (especially 'leaky' borders) promotes smuggling of substandard drugs.^{5 28} In each zone, one border and one non-border district were randomly sampled, ensuring that at least one district is as predominantly rural and the other urban (table 1).

Pharmacies and OTCs in each selected district were independently listed with unique identifiers. Using simple random sampling, 20 outlets (10 pharmacies and 10 OTCs) were selected from each district. In districts where the number of pharmacies were less than 10, all pharmacies were selected and supplemented with OTCs to achieve a total of 20 outlets.

Data collection

A team of research assistants were deployed to each district to collect data from the sampled outlets. The position of each outlet was recorded using a handheld Global Positioning System (GPS) device, enabling superimposition of relevant secondary data. Data on the outlet

s in each currently d then to ne more d-range'. her than ne outlet might be a 'cheap' one in another). For each selected product, we recorded brand name, generic name, API, retail price, manufacturer, country of origin, expiry date and other details from the packages. Retailers were then asked to provide information on the source of each product (supplier, purchase price, quantity purchased and date) before explaining the reasons for choosing to stock that particular product. The latter question was asked in an open-ended way and respondents were encouraged to list all the considerations pertaining to each product. The interviewing tool was precoded with a series of possible responses that were identified from the pilot stage (plus an 'other' category"), but these were not shared with respondents, to avoid biasing responses.

To test the hypotheses above, we began by grouping the 'reasons' responses into two broad categories: financial and quality related. In practice, it can be very difficult to discern the quality of a medicine from visual inspection alone, so people often make use of proxies such as reputation, recommendation or personal experience.^{29 30} On this basis, we further subdivided the 'quality' responses into those based on personal experience or reputation (i.e., non-professional) and those based on recommendation from a 'professional' source. This resulted in three categories, operationalised as follows. Financial was coded '1' if the respondent mentioned low price, profitability, or free delivery as considerations; otherwise, it was coded '0'. Experience/reputation was coded '1' if the respondent mentioned customer demand, positive brand

reputation, customers' report of good previous experience, or personal (shop attendant) positive experience. Finally, professional recommendation was coded '1' if the respondent mentioned recommendation by government, health professionals or suppliers; often prescribed by doctors; or public advertisement. In relation to the hypotheses above, the latter two categories (reputation/ experience and professional recommendation) were used as a proxy for considerations of quality, as opposed to decisions driven primarily by economic motivations.

The primary covariates (ecological zone, rural–urban location, type of outlet and category of antimalarials) were selected based on the hypothesises of the study, while the confounders were selected based on the literature,³¹ and included border/non-border location, number of staff, average number of customers per day, years of operation and availability of qualified pharmacist. The confounders were included in the analysis to examine the true associative effects of the hypothesised (primary) variables.

Data analysis

Frequency tables were used to describe the general characteristics of the sampled pharmacies and OTCs. Cross-tabulation was used to examine how the reasons for stocking vary by the primary and confounding factors, using χ^2 tests to assess statistically significant differences. Binary logistic regression was employed to examine the primary and confounding factors that were statistically significantly associated with the reasons for stocking anti-malarials. To prevent model overfit, only confounders significant at the 5% level were retained in the model.

Patient and public involvement statement

Patients or the public were not involved in the design, or conduct, or reporting, or dissemination plans of our research.

RESULTS

Background characteristics of the sampled outlets

Table 2 shows the background characteristics of the sampled outlets. Data were collected from 38 pharmacies (31.7%) and 82 OTCs (68.3%) located in rural (40.2%) and urban (60.8%) areas of Ghana. There were equal proportions (33.3%) of outlets in each of the three ecological zones (Southern, Middle, and Northern). Retail outlets were engaged in selling diverse antimalarials which were categorised into more expensive (38.7%), mid-range (34.4%) and cheaper (26.9%) based on cost. Based on the exchange rate at the time of fieldwork (US\$1=GH¢7.6), the retail prices of the 'more expensive' medicines ranged from US\$0.39 to US\$8.16 (median US\$1.97), while 'mid-range' products ranged from US\$0.26 to US\$3.42 (median US\$1.05), and 'cheaper' drugs ranged from US\$0.26 to US\$1.97 (median US\$0.66).

Bivariate analysis of the reasons for stocking antimalarials by outlet characteristics

As noted above, retailers' decisions regarding which antimalarials to stock were categorised into financial,

Table 2 Characteristics of sampled retail outlets						
Characteristics	Frequency	Per cent				
Type of outlet						
Pharmacy	38	31.7				
OTC	82	68.3				
Place of location						
Urban	73	60.8				
Rural	47	39.2				
Ecological zone						
Southern	40	33.3				
Middle	40	33.3				
Northern	40	33.3				
Border/Non-Border						
Border	60	50				
Non-Border	60	50				
Price bracket (multiple response)						
Cheaper	100	26.9				
Mid-range	128	34.4				
More expensive	144	38.7				
NO of staff						
One	73	60.8				
Two	28	23.3				
Three or more	19	15.8				
NO of customers per day						
Less than 25	39	32.5				
25–60	37	30.8				
More than 60	44	36.7				
Years of operation						
Less than 5 years	31	25.8				
5–10 years	23	19.2				
More than 10 years	66	55.0				
Has qualified pharmacist						
Yes	119	50.7				
No	25	49.3				
OTC, over the counter.						

experience/reputation and professional recommendation (table 3). Overall, the most widely cited set of factors influencing decision-making were personal experience and reputation, mentioned in 76.3% of cases (95% CI 72.0% to 80.7%). This was followed by financial reasons (53.2% of cases, 95% CI 48.1% to 58.3%), and then professional recommendation (24.7% of cases, 95% CI 20.3% to 29.1%).

The analysis (table 3) revealed that financial reasons for stocking antimalarials were not dependent on ecological zone, rural/urban location, or type of outlet (pharmacy or OTC). However, price bracket did make a difference. For 'cheaper' products, financial reasons were mentioned T-1-1- 0

Background characteristics	Financial % (95% Cl)	Experience/reputation % (95% CI)	Professional recommendation % (95% CI)
Overall	53.2 (48.1 to 58.3)	76.3 (72.0 to 80.7)	24.7 (20.3 to 29.1)
Primary factors			
Ecological zone	P value=0.456	P value=0.314	P value=0.047
Southern	48.7 (39.4 to 57.9)	81.4 (74.2 to 88.6)	16.8 (9.9 to 23.7)
Middle	56.7 (48.0 to 65.3)	74.0 (66.4 to 81.7)	26.0 (18.3 to 33.6)
Northern	53.8 (45.3 to 62.3)	74.2 (66.8 to 81.7)	30.3 (22.4 to 38.2)
Location	P value=0.123	P value=0.101	P value=0.062
Urban	50.4 (44.2 to 56.6)	73.8 (68.3 to 79.3)	21.8 (16.6 to 26.9)
Rural	58.9 (50.2 to 67.6)	81.5 (74.6 to 88.3)	30.6 (22.5 to 38.8)
Type of outlet	P value=0.208	P value=0.168	P value=0.611
Licensed pharmacy	49.4 (41.4 to 57.3)	72.7 (65.7 to 79.8)	23.4 (16.7 to 30.1)
Licensed OTC	56.0 (49.4 to 62.6)	78.9 (73.5 to 84.3)	25.7 (19.9 to 31.5)
Price bracket of antimalarial	P value=0.000	P value=0.314	P value=0.018
Cheaper	82.0 (74.4 to 89.6)	80.0 (72.1 to 87.9)	21.0 (13.0 to 29.0)
Mid-range	54.7 (46.0 to 63.3)	78.1 (70.9 to 85.3)	18.8 (12.0 to 25.5)
More expensive	31.9 (24.3 to 39.6)	72.2 (64.9 to 79.6)	32.6 (25.0 to 40.3)
Confounders			
Border/non-border location	P value=0.030	P value=0.201	P value=0.856
Border	58.8 (51.8 to 65.9)	79.1 (73.3 to 85.0)	25.1 (18.9 to 31.4)
Non border	47.6 (40.4 to 54.8)	73.5 (67.1 to 79.9)	24.3 (18.1 to 30.5)
No of staff	P value=0.364	P value=0.089	P value=0.100
One	54.0 (46.8 to 61.2)	79.7 (73.9 to 85.5)	21.9 (16.0 to 27.9)
Two	57.4 (47.4 to 67.5)	68.1 (58.6 to 77.6)	33.0 (23.4 to 42.5)
Three or more	47.3 (36.9 to 57.6)	78.0 (69.5 to 86.6)	22.0 (13.4 to 30.5)
No of customers per day	P value=0.650	P value=0.649	P value=0.278
Less than 25	54.2 (44.1 to 64.2)	74.0 (65.1 to 82.8)	27.1 (18.1 to 36.0)
25–60	56.2 (47.3 to 65.1)	76.0 (68.4 to 83.7)	28.1 (20.1 to 36.1)
More than 60	50.7 (42.7 to 58.6)	78.9 (72.4 to 85.5)	20.4 (14.0 to 26.8)
Year of operation	P value=0.450	P value=0.830	P value=0.537
Less than 5 years	50.0 (40.6 to 59.4)	76.4 (68.4 to 84.3)	23.6 (15.7 to 31.6)
5-10 years	50.5 (40.3 to 60.8)	78.5 (70.1 to 86.9)	29.0 (19.8 to 38.3)
More than 10 years	56.8 (49.3 to 64.3)	75.1 (68.6 to 81.7)	23.1 (16.7 to 29.4)
Has qualified pharmacists	P value=0.026	P value=0.275	P value=0.132
Yes	50.6 (45.1 to 56.2)	77.4 (72.8 to 82.1)	23.2 (18.5 to 27.9)
No	66.1 (54.3 to 78.0)	71.0 (59.6 to 82.4)	32.3 (20.5 to 44.0)

A substant data substant a la stata la substant a

OTC, Over The Counter.

in 82.0% of cases (95% CI 74.4% to 89.6%), compared with 54.7% (95% CI 46.0% to 63.3%) for 'mid-range' and 31.9% (95% CI 24.3% to 39.6%) for 'more expensive' products. Regarding the confounders, significant differences were found in mentions of financial considerations between border/non-border locations (p=0.03) and between outlets with/without a qualified pharmacist present (p=0.026). In outlets close to international borders, financial considerations were mentioned in

58.0% of cases (95% CI 51.8% to 65.9%), compared with 47.6% of cases (95% CI 40.4% to 54.8%) in non-border areas. Further, 66.1% (95% CI 54.4% to 78.0%) of outlets without a pharmacist present cited financial reasons for stocking decisions, compared with 50.6% (95% CI 45.1% to 56.2%) of those with a pharmacist present.

No significant differences in the proportions citing experience/reputation were found in either the primary factors (ecological zone, rural–urban locatio, and outlet

Table 4 Factors associated with reasons for stocking antimalarials							
	Financial OR (95% CI)	Experience/reputation OR (95% CI)	Professional recommendation OR (95% CI)				
Ecological zone							
Southern	1.00	1.00	1.00				
Middle	1.45 (0.82 to 2.58)	0.60 (0.32 to 1.12)	1.67 (0.88 to 3.18)				
Northern	1.06 (0.57 to 1.96)	0.55 (0.28 to 1.06)	1.89 (0.97 to 3.68)				
Location							
Urban	1.00	1.00	1.00				
Rural	1.39 (0.78 to 2.46)	1.69 (0.9 to 3.2)	1.38 (0.76 to 2.5)				
Type of outlet							
Licensed pharmacy	1.00	1.00	1.00				
Licensed OTC	1.14 (0.67 to 1.94)	1.19 (0.69 to 2.05)	0.92 (0.52 to 1.6)				
Price bracket							
Cheaper	1.00	1.00	1.00				
Mid-range	0.22 (0.11 to 0.42)**	0.85 (0.44 to 1.64)	0.84 (0.43 to 1.63)				
More expensive	0.08 (0.04 to 0.15)**	0.63 (0.33 to 1.17)	1.72 (0.94 to 3.15)				
Confounders							
Has qualified pharmacists							
Yes	1.00						
No	1.99 (1.02 to 3.87)*						
*p<0.01, **p<0.05. OTC, over the counter.							

type or price bracket) or any of the confounders. By contrast, 'professional recommendation' was significantly more likely to be mentioned in relation to 'more expensive' antimalarials 32.6% (95% CI 25.0% to 40.3%), compared with products that were 'mid-range' (18.8% of cases, 95% CI 12.0% to 40.3%) or 'cheaper' 21.0% of cases, 95% CI 13.0% to 29.0%). 'Professional recommendation' was also more likely to be cited as a driver of decision-making in the Northern part of Ghana (30.3% of cases, 95% CI 22.4% to 38.2%) and in the Middle Belt (26.0% of cases, 95% CI 18.3% to 33.6%), compared with the Southern zone (16.8% of cases, 95% CI 9.9% to 23.7%).

Multivariate analysis of factors associated with antimalarial stocking decisions

Table 4 shows the primary factors hypothesised to be associated antimalarial decisions and confounding factors found to be statistically important (p<0.05) in the bivariate analyses above. Although the direction of effects was hypothesised for financial motivations, this only reached statistical significance for one of the primary factors (price bracket) and one of the confounders (presence of a pharmacist). Financial considerations were significantly more likely to factor into decision-making for cheaper products, followed by mid-range, and least likely to matter for more expensive medicines (p<0.01). Presence of a pharmacist was found to be associated with a lower likelihood of financially driven stocking decisions compared with outlets with no pharmacist present (p<0.05). None of the hypothesised and confounding factors were associated with either personal experience/ reputation or professional recommendation in the multi-variate analysis.

Disaggregated analysis (online supplemental appendix 1) revealed that profitability considerations were significantly more likely to factor into decisionmaking for outlets in rural areas (p<0.01), and border outlets likely to consider low price antimalarial (p<0.05). Personal positive experience was significantly likely to be considered in antimalarial stocking decisions in the rural areas. Customer previous experience was likely considered in the northern ecological zone but positive brand less likely in the same ecological zone.

DISCUSSION

Private-sector pharmaceutical supply chains are integral to the fight against SF and the success of global health initiatives like the attainment of UHC and malaria control milestones. Cognisant of the complex nature of private sector pharmaceutical supply chains in LMICs and the vital role that medicine retail outlets play in ensuring that good quality antimalarials get to patient,³² this study investigated the reasons behind antimalarial stocking decisions at the medicine retail outlet level in a country with a fragile health system and suboptimal regulatory structures (although better than many other countries in the Region).

The emergence of personal experience and reputation (one group of proxies for quality) as the overall leading set of motivations driving antimalarial stocking decisions in this study is quite instructive. Although international stakeholders are gradually warming up to the reality that the private sector play a vital role in malaria control, there is still some amount of scepticism about the financial motivations of the sector.^{33 34} Such scepticism is perhaps fueled by the overly commercial orientation of privatesector retail outlets and studies suggesting that pricing and profitability are the main drivers of stocking practices of retail outlets.³⁵ Thus, our finding that perceived quality is apparently more important overall than financial considerations in antimalarial stocking decisions in Ghana is refreshing. However, financial motivations did emerge as the second most popular driver of stocking decisions overall.

In considering variation between different types/locations of outlets and products, the rejection of our hypothesis (H1) that outlets in poorer and more remote areas are likely to prioritise profit over quality of antimalarials is noteworthy. Despite a palpable rural-urban dichotomy in Ghana and substantial socioeconomic differences between zones and region,²⁹ outlets in the rural and more northern areas appeared not to be any more motivated by financial considerations than their counterparts in relatively affluent areas. This finding is intriguing considering the dire economic situation in rural parts of Ghana, with outlets operating on small margins and many people unable to afford anything but the cheapest medicines, where one might expect that economic imperatives would be particularly pressing. Other work suggests that outlets in rural areas are often operated by persons from those communities who are highly trusted and widely known, and thus may be particularly motivated to ensure their clients receive effective treatment wherever possible.³⁶⁻³⁸

The impact of financial considerations on stocking decisions came to the fore in two other situations: price bracket and presence/absence of a pharmacist. However, H2 was not accepted as there were no significant differences in financial consideration stocking decisions between OTCs and pharmacies. Outlets' financial considerations were influenced stocking decisions for 'cheaper' compared with 'more expensive' antimalarials supporting H3 (For cheaper medicines, financial considerations are likely to be more important and quality). This is perhaps unsurprising: lower-cost medicines tend to be purchased by the poorest within a community, for whom pricing is likely to be particularly sensitive, thereby incentivising retailers to select the very cheapest products. In other words, it may be that selecting primarily on the basis of perceived quality is a 'luxury' than can only be entertained in less economically constrained circumstances. The Global Fund's Affordable Medicines Facility for Malaria (AMFm) project, which provided a subsidy for the purchase of ACTs (artemisinin combination

therapies, currently recommended by the WHO for firstline treatment of acute malaria) was a direct response to meet the demand for efficacious, low-cost in LMICs.³⁹ Evidence from Ghana suggests that the AMFm was effective in making Quality-Assured ACTs readily available, even in rural medicine outlets.⁴⁰ Similarly, government subsidies were reported to have improved access to antimalarials and reduced SF medicines by 50% in rural Uganda.⁴¹ Our finding that financial considerations still drive the stocking of lower-cost antimalarials suggests that self-sustaining initiatives fashioned after AMFm might be important in curbing the SF medicine threat.

Second, the presence of a qualified pharmacist was associated with a lower influence of financial considerations in stocking decisions, compared with outlets without a pharmacist. Other studies have suggested that the unavailability of trained pharmacists at outlets promotes 'irrational' dispensing but very little is known about the impact on stocking decisions. If the person responsible for stocking decisions lacks the necessary pharmacological knowledge, they may rely more on a pricing heuristic. This is important because pharmacists are more likely to be present in large, urban, upscale pharmacies than in small, rural, OTC outlets.

By contrast, none of our hypotheses regarding the impact of outlet location (zone or rural/urban) or outlet type (pharmacy/OTC) on antimalarial stocking decisions were supported. Some previous studies in LMICs⁴² have reported differences between the two, claiming that pharmacies are more loyal to products and rent-seeking, and are thus more likely to stock products based on their personal interest. However, such tendencies were not evident in our study. It may be that the sample sizes used in our study were too small and lacked sufficient statistical power (in most cases, the direction of the effect was as predicted but did not reach statistical significance), or it may be that there really are no differences in these respects. As noted above, it would be useful in the future to scale-up this study to be nationally representative.

CONCLUSIONS

Several findings from this study have practical implications for SF medicines and malaria control in Ghana. First, through the sampling, our study confirms the uneven distribution of licensed medicine outlets across Ghana, with those living in urban areas and in the relatively affluent southern parts of the country having much better access (especially to pharmacies) than those living in more rural areas and in the less affluent north. While this is not a new finding, it continues to affect access to essential medicines like antimalarials for large parts of the population. Second, broadly speaking, private-sector medicine outlets across the country were stocking antimalarials based on the price bracket of the drug. It is clearly time that policy-makers recognised the reality that private-sector outlets perform a crucial role in delivering primary care to underserved communities, often doing

the best they can be in difficult circumstances, and work to support their efforts.

Third, in two specific sets of circumstances, financial considerations came more to the fore: for the cheapest products and for outlets without a pharmacist present. These cases present potential supply-chain vulnerabilities, where price may trump considerations of quality. It is notable that both are more likely to affect disadvantaged rural communities, who are less likely to be able to afford anything but the lowest-cost medicines and who may be less likely to have access to an outlet staffed by a qualified pharmacist. Ensuring that all medicine outlets have staff with adequate training to make informed stocking decisions (whether or not they are fully qualified pharmacists) and ensuring that access to affordable qualityassured antimalarials continues in a sustainable manner in the post-AMFm era are therefore key priorities in both the fight against SF medicines and in moving towards genuine UHC.

Author affiliations

¹Department of Geography Education, University of Education, Winneba, Ghana ²Department of Population and Health, University of Cape Coast Faculty of Social Sciences, Cape Coast, Ghana

³Department of Geography and Regional Planning, University of Cape Coast Faculty of Social Sciences, Cape Coast, Ghana

⁴School of Medical Sciences, University of Cape Coast, Cape Coast, Ghana

⁵Directorate of Research, Innovation and Consultancy, University of Cape Coast, Cape Coast, Ghana

⁶Department of Sociology, Oxford University, Oxford, UK

⁷Department of Anthropology, Durham University, Durham, UK

Twitter Simon Mariwah @smariwah, Daniel Amoako-Sakyi @dasakyi, Samuel Asiedu Owusu @sakowusu and Heather Hamill @DrHeatherHamill

Contributors AO, FAJ and SM designed the study, managed the survey, analysed the data, and drafted the paper. DA-S designed the data and drafted the paper. SAO managed the survey and drafted the paper. ME analysed the data and drafted the paper. HH is Co-I on the MRC grant that funded the research, designed the study, drafted the pape, r and is responsible for the overall content as guarantor. KH is PI on the MRC grant that funded the research, designed the data, drafted the paper, and is responsible for the overall content as guarantor.

Funding This work was supported by the Medical Research Council (MRC) UK grant number MR/T022132/1 $\,$

Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not applicable.

Ethics approval This study involves human participants and was approved by the University of Cape Coast Institutional Review Board UCC1RB/EXT/2020/21, and by the Department of Anthropology Durham University ANTH-2020-07-07T17. Participants gave informed consent to participate in the study before taking part.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available on reasonable request.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution 4.0 Unported (CC BY 4.0) license, which permits others to copy, redistribute, remix, transform and build upon this work for any purpose, provided the original work is properly cited, a link to the licence is given, and indication of whether changes were made. See: https://creativecommons.org/licenses/by/4.0/.

ORCID iD

Heather Hamill http://orcid.org/0000-0002-0301-7057

REFERENCES

- 1 Study on the public health & socioeconomic impact of SF medical products. Geneva World Health Organisation; 2017.
- 2 WHO. WHO global surveillance and monitoring system for substandard and falsified medical products. Geneva: World Health Organization, 2017b. Available: https://apps.who.int/iris/handle/ 10665/326708
- 3 Ozawa S, Evans DR, Bessias S, *et al.* Prevalence and estimated economic burden of substandard and falsified medicines in low- and middle-income countries: a systematic review and meta-analysis. *JAMA Netw Open* 2018;1:e181662.
- 4 Wilson BK, Kaur H, Allan EL, et al. A new handheld device for the detection of falsified medicines: demonstration on falsified artemisinin-based therapies from the field. Am J Trop Med Hyg 2017;96:1117–23.
- 5 Tivura M, Asante I, van Wyk A, et al. Quality of artemisinin-based combination therapy for malaria found in Ghanaian markets and public health implications of their use. *BMC Pharmacol Toxicol* 2016;17:48.
- 6 McCabe A, Seiter A, DiackA, et al. Private sector pharmaceutical supply and distribution channels in Africa: a focus on Ghana, Malawi and Mali. World Bank, 2011. Available: https://openknowledge. worldbank.org/handle/10986/13590
- 7 Mackintosh M, Tibandebage P, Karimi Njeru M, et al. Rethinking health sector procurement as developmental linkages in East Africa. Soc Sci Med 2018;200:182–9.
- 8 Seidman G, Atun R. Do changes to supply chains and procurement processes yield cost savings and improve availability of pharmaceuticals, vaccines or health products? A systematic review of evidence from low-income and middle-income countries. *BMJ Glob Health* 2017;2:e000243.
- 9 Gautam CS, Utreja A, Singal GL. Spurious and counterfeit drugs: a growing industry in the developing world. *Postgrad Med J* 2009;85:251–6.
- 10 Minzi OMS, Moshi MJ, Hipolite D, *et al*. Evaluation of the quality of amodiaquine and sulphadoxine/pyrimethamine tablets sold by private wholesale pharmacies in Dar Es Salaam Tanzania. *J Clin Pharm Ther* 2003;28:117–22.
- 11 Syhakhang L, Lundborg CS, Lindgren B, et al. The quality of drugs in private pharmacies in Lao PDR: a repeat study in 1997 and 1999. *Pharm World Sci* 2004;26:333–8.
- 12 Bate R, Tren R, Mooney L, et al. Pilot study of essential drug quality in two major cities in India. PLoS One 2009;4:e6003.
- 13 Bate R, Coticelli P, Tren R, et al. Antimalarial drug quality in the most severely malarious parts of Africa - a six country study. PLoS One 2008;3:e2132.
- 14 Pisani E, Nistor A-L, Hasnida A, *et al.* Identifying market risk for substandard and falsified medicines: an analytic framework based on qualitative research in China, Indonesia, Turkey and Romania. *Wellcome Open Res* 2019;4:70.
- 15 Newton PN, Bond KC, 53 signatories from 20 countries. COVID-19 and risks to the supply and quality of tests, drugs, and vaccines. *Lancet Glob Health* 2020;8:e754–5.
- 16 WHO. World malaria report 2021. Geneva World Health Organisation; 2021. Available: https://www.who.int/teams/global-malariaprogramme/reports/world-malaria-report-2021
- 17 Darteh EKM, Buabeng I, Akuamoah-Boateng C. Uptake of intermittent preventive treatment in pregnancy for malaria: further analysis of the 2016 Ghana malaria indicator survey. *J Public Health* (Berl) 2021;29:967–78.
- 18 Tangena J-A, Hendriks CMJ, Devine M, et al. Indoor residual spraying for malaria control in sub-Saharan Africa 1997 to 2017: an adjusted retrospective analysis. *Malar J* 2020;19:150.
- 19 Bossert TJ, Bowser DM, Amenyah JK. Is decentralization good for logistics systems? Evidence on essential medicine logistics in Ghana and Guatemala. *Health Policy Plan* 2007;22:73–82.
- 20 Ghana Statistical Service. Population and Housing Census General Report. Accra Ghana Statistical Service; 2021.

BMJ Global Health

- 21 Ahmed A, Gasparatos A. Multi-dimensional energy poverty patterns around industrial crop projects in Ghana: enhancing the energy poverty alleviation potential of rural development strategies. *Energy Policy* 2020;137:111123.
- 22 Zhang C, Rahman MS, Rahman MM, et al. Trends and projections of universal health coverage indicators in Ghana, 1995-2030: a national and subnational study. *PLoS One* 2019;14:e0209126.
- 23 2014 Annual Report: National Malaria Control Programme. Accra Ministry of Health, Ghana; 2015.
- 24 Yankson R, Anto EA, Chipeta MG. Geostatistical analysis and mapping of malaria risk in children under 5 using point-referenced prevalence data in Ghana. *Malar J* 2019;18:67.
- 25 Agyemang-Duah W, Peprah C, Peprah P. Factors influencing the use of public and private health care facilities among poor older people in rural Ghana. *J Public Health (Berl)* 2020;28:53–63.
- 26 Kwarteng A, Akazili J, Welaga P, et al. The state of enrollment on the national health insurance scheme in rural Ghana after eight years of implementation. Int J Equity Health 2019;19:4.
- 27 Walker EJ, Peterson GM, Grech J, et al. Are we doing enough to prevent poor-quality antimalarial medicines in the developing world?BMC Public Health 2018;18:630.
- 28 Hampshire K, Hamill H, Mariwah S, et al. The application of signalling theory to health-related trust problems: the example of herbal clinics in Ghana and Tanzania. Soc Sci Med 2017;188:109–18.
- 29 Hamill H, Hampshire K, Mariwah S, et al. Managing uncertainty in medicine quality in Ghana: the cognitive and affective basis of trust in a high-risk, low-regulation context. Soc Sci Med 2019;234:112369.
- 30 Soares CM, Nascimento B, Chaves LA, et al. Public procurement of medicines: scoping review of the scientific literature in South America. J Pharm Policy Pract 2019;12:33.
- 31 Goodman C, Kachur SP, Abdulla S, et al. Retail supply of malariarelated drugs in rural Tanzania: risks and opportunities. *Trop Med Int Health* 2004;9:655–63.
- 32 Awor P, Wamani H, Bwire G, et al. Private sector drug shops in integrated community case management of malaria, pneumonia, and diarrhea in children in Uganda. Am J Trop Med Hyg 2012;87:92–6.

- 33 Bennett A, Avanceña ALV, Wegbreit J, et al. Engaging the private sector in malaria surveillance: a review of strategies and recommendations for elimination settings. *Malar J* 2017;16:252.
- 34 Mayora C, Kitutu FE, Kandala N-B, et al. Private retail drug shops: what they are, how they operate, and implications for health care delivery in rural Uganda. BMC Health Serv Res 2018;18:532.
- 35 Baffoe G, Zhou X, Moinuddin M, et al. Urban-rural linkages: effective solutions for achieving sustainable development in Ghana from an SDG interlinkage perspective. Sustain Sci 2021;16:1341–62.
- 36 Okai GA, Abekah-Nkrumah G, Asuming PO. Perceptions and trends in the use of community pharmacies in Ghana. *J Pharm Policy Pract* 2019;12:25.
- 37 Hampshire K, Mariwah S, Amoako-Sakyi D, *et al.* "It is very difficult in this business if you want to have a good conscience": pharmaceutical governance and on-the-ground ethical labour in Ghana. *Glob Bioeth* 2022;33:103–21.
- 38 Tougher S, ACTwatch Group, Ye Y, et al. Effect of the affordable medicines facility--malaria (AMFm) on the availability, price, and market share of quality-assured artemisinin-based combination therapies in seven countries: a before-and-after analysis of outlet survey data. *Lancet* 2012;380:1916–26.
- 39 Freeman A, Kwarteng A, Febir LG, et al. Two years post affordable medicines facility for malaria program: availability and prices of anti-malarial drugs in central Ghana. J Pharm Policy Pract 2017;10:15.
- 40 Björkman Nyqvist M, Svensson J, Yanagizawa-Drott D. Can good products drive out bad? A randomized intervention in the antimalarial medicine market in Uganda. *J Eur Econ Assoc* 2022;20:957–1000.
- 41 Silverman R, Keller J, Glassman A, et al. Tackling the triple transition in global health procurement. Center for Global Development, 2019. Available: https://www.cgdev.org/sites/default/files/better-healthprocurement-tackling-triple-transition.pdf
- 42 Carland C, Goentzel J, Montibeller G. Modeling the values of private sector agents in multi-echelon humanitarian supply chains. Eur J Oper Res 2018;269:532–43.

Appendix 1: Multivariate analysis of outlet location and stocking decision of antimalarials

			Professional				
	Financial reasons		recommended	Expe	erience/reputa	reputation	
X7 · 11	D. C. 11	T	Often prescribed by	Personal positive	Customers previous	Positive brand	
Variables	OR [95%	OR [95%	doctors	OR [95%	OR [95%	OR [95%	
	CI]	CI]	OR [95% CI]	CI]	CI]	CI]	
Place of location							
	0.43 [0.27.			0.54 [0.30.			
Urban	0.69]**			0.94]*			
Rural	1			1			
Ecological zone							
						2.44	
					0.47 [0.27,	[1.38,	
Southern					0.83]**	4.30]** 1.46	
					0.81 [0.49,	[0.83,	
Middle					1.35]	2.57]	
Northern					1	1	
Border location							
		1.73				0.58	
		[1.13,		2.01 [1.12,		[0.37,	
Border		2.65]*		3.58]*		0.92]*	
Non-border		1		1		1	
Has qualified							
pharmacist							
			0.41 [.021,				
Yes			0.80]*				
No			1				

Appendix S1 – Reflexivity Statement

1. How does this study address local research and policy priorities?

The findings of this study have implications for the management of anti-malarial quality across supply chains in Ghana, with potentially important consequences for malaria control, particular in lower-income areas where people rely on low-cost medication.

2. How were local researchers involved in study design?

The first category of local researchers involved were those with extensive experience of conducting, leading, or organising international research collaborations (SM, DAS and SAO in Ghana, Tanzania, and Malawi), FAJ in Ghana and UK, and ME in Ghana and Nigeria. The second category were local researchers for whom this was their first international collaboration (AO). In addition, there were high-income country researchers with extensive experience of conducting, leading, or organising international research collaborations involving low- and middle-income countries (KH and HH). Most of the authors originate from, and reside, in Ghana. The local researchers (AO, FAJ, SM, DAS, SAO, and EM) were involved in the design of the project, and they led the data collection and manuscript preparation.

3. How has funding been used to support the local research team?

This project started with a workshop attended by all authors in which the research was codesigned. Funding has also been used to support the local research team to employ and train research assistants in data collection and collation and to train colleagues in the University of Cape Coast (UCC), Ghana in several data analysis techniques such as GIS. These capacity building activities will continue to be supported by the senior authorship team to develop.

4. How are research staff who conducted data collection acknowledged?

All local research staff who attended the research design workshop were included as authors. Each team member was designated and delivered specific role(s) during the writing process (see acknowledgements).

5. Do all members of the research partnership have access to study data?

All members of the partnership have access to data.

6. How was data used to develop analytical skills within the partnership?

Attendees of the research workshop contributed directly to the research design. Fortnightly meetings where subsequently held via MS Teams over a 2-year period with the full research team to discuss progress and co-develop solutions for any problems that arose in the conduct of the research. The analytical skills of the local research team have been enhanced through constant feedback on project reports and draft manuscripts.

7. How have research partners collaborated in interpreting study data?

During the fortnightly MS Teams meetings, hypotheses and data analysis strategies were discussed in a structured format. Short papers and presentations were given by members of the team presenting the findings, which were discussed by the whole team. These frequent meetings have led to the preparation of this manuscript.

8. How were research partners supported to develop writing skills?

The authors are composed of senior and early career academics. The early career researchers (AO) on the authorship team were supported by senior colleagues to develop and refine their writing skills. This was achieved through the constant feedback on draft manuscripts by senior colleagues.

9. How will research products be shared to address local needs?

This article will be published as open access and findings will be discussed with research and policy leaders in global health in Ghana.

10. How is the leadership, contribution, and ownership of this work by LMIC researchers recognised within the authorship?

Authors AO, FAJ, SM, DAS, SAO, and ME worked as part of the senior authorship team in developing this manuscript, and their contribution has been recognised by the author order.

11. How have early career researchers across the partnership been included within the authorship team?

AO is an early career researcher and is first author.

12. How has gender balance been addressed within the authorship?

Six authors are male (AM, FAM, SM, SAS, SAO, and ME) and two authors female (HH and KH)

13. How has the project contributed to training of LMIC researchers?

The authorship team is primarily composed of senior researchers apart from AO who has been supported in his writing and analytical skills as first author.

14. What safeguarding procedures were used to protect local study participants and researchers?

Safeguarding issues were directly addressed through the ethics approval given by the IRBs at UCC Ghana and Durham University UK.