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## Predictive Value of Clinician Impression for Readmission and Post-Discharge Mortality among Neonates and Young Children in Dar es Salaam, Tanzania and Monrovia, Liberia

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## Predictive Value of Clinician Impression for Readmission and Post-Discharge Mortality among Neonates and Young Children in Dar es Salaam, Tanzania and Monrovia, Liberia

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**Key Words:** neonates; infants; young children; readmission; post-discharge mortality; Tanzania; Liberia

**Word Count:** 2,500

**Abbreviations:** AUC: area under the receiver operating characteristic curve; 95% CI: 95% confidence interval

### **Data Sharing Agreement**

Data may be made available upon reasonable request to the corresponding author.

### **Ethics Approval**

The study received ethical clearance from the Tanzania National Institute of Medical Research (#NIMR/HQ/R8a/Vol.IX/3494), the Muhimbili University of Health and Allied Sciences Research and Ethics Committee (#307/323/01), the John F. Kennedy Medical Center Institutional Review Board (#08062019), the Boston Children's Hospital Institutional Review Board (#P00033242), and the use of de-identified data was exempted from review by the Emory University Institutional Review Board (no number provided for exempted studies).

### **Transparency Declaration**

This manuscript is an honest and accurate account of the study being reported. No aspects of this study have been omitted or withheld.

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## Abstract

*Background:* There are no validated clinical decision aids to identify neonates and young children at risk of hospital readmission or post-discharge mortality in sub-Saharan Africa, leaving the decision to discharge a child to a clinician's impression. Our objective was to determine the accuracy of clinician impression to identify neonates and young children at risk for readmission and post-discharge mortality.

*Methods:* We conducted a survey study nested in a prospective observational cohort of neonates and children aged 1-59 months who were followed up to 60 days after hospital discharge from Muhimbili National Hospital in Dar es Salaam, Tanzania or John F. Kennedy Medical Center in Monrovia, Liberia. Clinicians who discharged each enrolled patient were surveyed at the time of discharge to determine their perceived probability of the patient's risk of 60-day hospital readmission or post-discharge mortality. We calculated the area under the receiver operating characteristic curve (AUC) to determine the accuracy of clinician impression for both outcomes.

*Results:* Of 4,247 discharged patients, 3,896 (91.7%) had available clinician surveys and 3,847 (98.7%) had 60-day outcomes available: 187 (4.8%) were readmitted and 120 (3.1%) died within 60 days of hospital discharge. Clinician impression had poor discriminatory value in identifying neonates and young children at risk of hospital readmission (AUC 0.46, 95% CI 0.43 to 0.49) and post-discharge mortality (AUC 0.53, 95% CI 0.50 to 0.55). Patients for whom clinicians attributed inability to pay for future medical treatment as the reason for risk for unplanned hospital readmission had 4.76 times the odds hospital readmission (95% CI 1.31 to 17.25,  $P=0.02$ ).

*Conclusions:* Given the poor discriminatory value of clinician impression alone to identify neonates and young children at risk of hospital readmission and post-discharge mortality, validated clinical decision aids are needed to aid in the identification of young children at risk for these outcomes.

### What is already known on this topic?

- In parts of sub-Saharan Africa, hospital readmissions and post-discharge mortality rates are estimated to be as high as 18% within months of hospital discharge
- There are no validated clinical decision aids to accurately identify neonates, infants, and young children at risk of hospital readmission or post-discharge mortality in sub-Saharan Africa

### What this study adds?

- In a survey study nested in a prospective observational cohort study conducted at referral hospitals in Tanzania and Liberia that included clinician surveys on 3,896 neonates, clinician impression alone had poor discriminatory value in identifying neonates, infants, and children at risk of hospital readmission within 60 days of hospital discharge
- Clinician impression also did not accurately identify neonates, infants, and children at risk of 60-day post-discharge mortality

### How this study might affect practice?

- Clinician impression alone is not sufficient to accurately identify neonates, infants, and children at risk of hospital readmission or post-discharge mortality
- Validated and objective clinical decision aids are urgently needed to better identify neonates, infants, and children at risk of hospital readmission and post-discharge mortality

## Introduction

The time after an inpatient hospital admission is particularly vulnerable in the life of a child in sub-Saharan Africa, where readmissions are common<sup>1,2</sup> and where more than half of the estimated 5 million annual deaths among children aged <5 years occurred in 2020.<sup>3,4</sup> Childhood mortality rates in the period immediately after hospitalization for an illness (i.e., the post-discharge period) may outpace rates of mortality during hospitalization.<sup>5,6</sup> In parts of sub-Saharan Africa, readmission and post-discharge mortality rates are estimated to be as high as 18% within six months after hospital discharge.<sup>5,7</sup>

Although clinical prediction rules for all-cause hospital readmissions among children in high-income settings have been developed,<sup>8</sup> to our knowledge, there are currently no clinical prediction rules for hospital readmissions among children in resource-limited settings. There are clinical prediction rules that have been developed to identify children at risk of post-discharge mortality in some settings in sub-Saharan Africa.<sup>6,9,10</sup> However, these clinical prediction rules lack external validation and thus are not widely used in clinical practice. Given the absence of validated risk assessment tools to identify young children at risk of readmission and post-discharge mortality in sub-Saharan Africa, the decision to safely discharge a young child from a hospital is often driven by clinical judgement.

Clinician impression relies on the clinician's ability to recognize patterns that may be associated with severe disease or an adverse outcome.<sup>11</sup> However, the accuracy of clinician impression to predict outcomes, such as severe disease from infections, among children has varied in previous studies conducted in high-income settings.<sup>12-14</sup> In a survey of 39 providers in Kenya, clinicians under-estimated the overall incidence of post-discharge mortality among children.<sup>15</sup> However, that study did not assess clinician impression of post-discharge mortality for individual patients and, to our knowledge, that has not been studied previously.

Given the absence of validated prognostic tools for hospital readmission and post-discharge mortality among children in sub-Saharan Africa, we aimed to determine the accuracy of treating clinicians' clinical impression to identify neonates and young children at risk for hospital readmission and post-discharge mortality in Dar es Salaam, Tanzania and Monrovia, Liberia.

## Methods

### *Study Design*

We conducted a survey nested in a prospective observational cohort study of pediatric patients discharged from Muhimbili National Hospital in Dar es Salaam, Tanzania and John F. Kennedy Medical Center in Monrovia, Liberia from October 2019 to January 2022. Details of our study protocol have been published previously.<sup>16</sup> Neonates and young children aged 1-59 months were enrolled at discharge from the neonatal or pediatric wards at each facility. Follow-up consisted of caregivers receiving phone calls up to 60 days after hospital discharge. Caregivers provided written consent for participation in Tanzania and oral consent in Liberia because of cultural preference and low rates of caregiver literacy.



### *Patient and public involvement statement*

The development of the research question was informed by the disease burden of readmission and post-discharge mortality among children in sub-Saharan Africa. Patients were not involved in the design, recruitment, or conduct of the study, nor were they advisers in this study. Results of this study will be made publicly available through publication.

### *Study Setting*

This study was conducted at two large, national referral hospitals supported by each country's Ministry of Health. They are in urban areas in their respective countries. Muhimbili National Hospital has a catchment of approximately 6 million people and John F. Kennedy Medical Center has a catchment of approximately 1.5 million people. Both hospitals are training hospitals for pediatric residents who are completing their specialty training.

### *Study Populations*

Neonates and young children discharged from the wards were consecutively enrolled. Neonates and young children who died during initial hospitalization were excluded. Neonates and young children whose caregivers did not have telephones for follow up or those who declined enrollment were excluded.

Surveyed clinicians included consultants/specialists, interns/residents, or medical officers. Consultants/specialists were certified pediatricians or pediatric specialists who completed medical school, residency, and subspecialty training (for specialists). Interns/residents had completed medical school and were completing residency training in pediatrics. Medical officers received three years of clinical training prior to providing clinical care to patients.

### *Study Procedures*

After obtaining informed consent from caregivers, trained research coordinators at each site approached the clinician who discharged each enrolled patient, obtained consent, and asked them to complete a survey near the time of the patient's hospital discharge. This survey modeled previous surveys that assessed clinician impression<sup>13</sup> and was developed through an iterative process by the research team with multiple opportunities to each investigator to refine the content. The survey was also reviewed by an expert in survey design (i.e., a survey methodologist) to ensure question clarity and appropriate response types (Appendix Survey). This survey was designed to allow clinician respondents to describe their perceived probability that each patient would experience both outcomes. Responses were recorded on standardized, electronic case report forms in electronic tablets using SQL (Tanzania) and KoboToolbox (Liberia).

### *Measurement and Outcomes*

The exposure variable was the impression of the discharging clinician of the patient's risk of: 1) unplanned hospital readmission within 60 days of hospital discharge or 2) all-cause, 60-day post-

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3 discharge mortality. Aligned with previous studies,<sup>13</sup> probabilities of perceived risk of  
4 readmission or post-discharge mortality included categorical options of 0%, 1-5%, 6-20%, 21-  
5 40%, 41-60%, 61-80%, 81-99%, and 100%. This survey also assessed discharging clinicians'  
6 perceptions of why readmission or post-discharge mortality were possible for those who were  
7 identified as at-risk for each outcome. Surveyed clinicians were familiar with the patients'  
8 clinical history and laboratory results during hospital admission. To assess for outcomes, phone  
9 calls to patients' caregivers were made by research staff at 7, 14, 30, 45, and 60 days after  
10 hospital discharge. Outcomes were determined as reported by caregivers to research staff.  
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### 13 *Statistical Analyses*

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16 The association of the discharging clinicians' predicted probability of readmission or post-  
17 discharge mortality and proportion of patients at each clinician-estimated risk threshold (e.g.,  
18 0%, 1-5%, etc.) who were readmitted or died was compared using Chi-square or Fisher's exact  
19 testing ( $P < 0.05$  for significance).<sup>13</sup> We calculated sensitivity, specificity, positive predictive  
20 value, and negative predictive value of treating clinician's impression at each percent risk  
21 threshold using caregiver-reported readmission or post-discharge mortality as the reference  
22 standards. We determined the accuracy of clinician impression for identifying patients at risk of  
23 readmission or post-discharge mortality by calculating the area under the receiver operating  
24 characteristic curve (AUC). 95% confidence intervals (95% CI) for the AUC were calculated  
25 through 2,000 bootstrap replicates. We conducted sub-analyses by the discharged patient's age  
26 group (i.e., neonate or young child), clinician experience level (i.e., consultant/specialist,  
27 intern/resident, or medical officer), site, and time to outcome. We conducted binary logistic  
28 regression analyses to assess whether the perceived reason for risk for each outcome was  
29 associated with the patient's likelihood of each outcome. All tests were two-sided tests and used  
30 a 0.05 significance level. AUC analyses were conducted through the pROC package in R.<sup>17</sup> All  
31 analyses were performed in R Version 4.1.3 (R Foundation for Statistical Computing, Vienna,  
32 Austria) and SAS 9.4 (SAS Institute Inc., Cary, NC).  
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### 36 **Results**

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39 There were 4,460 discharged patients, 4,247 (95.2%) enrolled, and 3,896 (91.7%) had complete  
40 clinician surveys (Figure 1). Enrollment was approximately equal between the two sites  
41 (Tanzania  $n=1,997$ , 51.3%, Liberia  $n=1,899$ , 48.7%) (Table 1). There were 2,173 (55.8%)  
42 neonates and 1,723 (44.2%) young children who had clinician surveys available.  
43

44 Sixty-day outcomes were available for 3,847 (98.7%) enrolled patients. The median age of  
45 enrolled neonates was 2 days (interquartile range [IQR] 1-7) and 12 months (IQR 5-24) for  
46 infants and children. The most common discharge diagnoses among neonates were sepsis  
47 (29.7%,  $n=609$ ), prematurity (28.8%,  $n=591$ ), and birth asphyxia (15.8%,  $n=323$ ). Among infants  
48 and children, pneumonia (12.1%,  $n=223$ ), diarrheal disease (10.1%,  $n=186$ ), and malaria (7.2%,  
49  $n=133$ ) were the most common discharge diagnoses.  
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52 There were 187 (4.8%) patients readmitted and 120 (3.1%) died within 60 days of discharge.  
53 There were 80 (3.6%) neonates who were readmitted and 61 (2.8%) died within 60 days of  
54 hospital discharge. Among infants and children, 107 (6.2%) were readmitted and 59 (3.4%) died  
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3 after hospital discharge. The median time from hospital discharge to readmission was 30 days  
4 (IQR 7-45). The median time from hospital discharge to mortality was 30 days (IQR 14-45).  
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### 6 *Clinician Impression and Hospital Readmission*

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9 Nearly three quarters of patients were perceived to have 0% risk of readmission within 60 days  
10 (Table 2). Clinician assigned probability of readmission was associated with readmission  
11 ( $P<0.001$ ; Table 2). However, among the 187 neonates and young children who were readmitted,  
12 80.7% (n=151) were perceived to have 0% risk of readmission.  
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15 Overall, clinician impression had poor discriminatory value in identifying neonates and young  
16 children at risk of readmission (AUC 0.46, 95% CI 0.43 to 0.49) (Table 3). Among medical  
17 officers, clinician impression had fair discriminatory value in identifying children at risk of  
18 readmission (AUC 0.67, 95% CI 0.55 to 0.79); this group was better at identifying patients at  
19 risk of readmission than interns/residents and consultants/specialists.  
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22 By clinician type, medical officer clinician impression had poor discriminatory value in  
23 identifying neonates (Supplemental Table 1) but fair discriminatory value in identifying infants  
24 and children at risk for readmission (Supplemental Table 2). In site-specific analyses, clinician  
25 impression was poor in identifying neonates and young children at risk of readmission at both  
26 sites (Supplemental Table 3). Regardless of the time from hospital discharge to readmission,  
27 clinician impression had poor discrimination in identifying neonates and young children at risk  
28 for readmission (Supplemental Table 4).  
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### 30 *Clinician Impression and Post-Discharge Mortality*

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33 Most (97.2%, n=3,746) patients were assigned 0% risk of post-discharge mortality (Table 4).  
34 Clinician assigned probability of post-discharge mortality was associated with the outcome  
35 ( $P=0.002$ ; Table 4). Among the 120 neonates and young children who died within 60 days of  
36 hospital discharge, 90.8% (n=109) were estimated to have a 0% probability of post-discharge  
37 mortality.  
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40 Overall, clinician impression had poor discriminatory value in identifying neonates and young  
41 children at risk post-discharge mortality (AUC 0.53, 95% CI 0.50 to 0.55) and did not vary  
42 substantially among interns/residents, specialist/consultants, or medical officers (Table 5).  
43 Clinician impression had poor discriminatory value in identifying post-discharge mortality  
44 among neonates (Supplemental Table 5) and infants and children (Supplemental Table 6). When  
45 analyzed by site, clinician impression in both Tanzania and Liberia had poor discriminatory  
46 value in identifying neonates and young children at risk of post-discharge mortality  
47 (Supplemental Table 7). Clinician impression had poor discriminatory value regardless of the  
48 time to post-discharge mortality (Supplemental Table 8).  
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### 50 *Reason for Perceived Risk of Hospital Readmission and Post-Discharge Mortality*

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54 Patients for whom clinicians attributed inability to pay for treatment as the reason for  
55 readmission had 4.76 times the odds readmission (95% CI 1.31 to 17.25,  $P=0.02$ ) compared to  
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those with no perceived risk (Table 6). Patients whose clinician cited “other” reasons to be at risk had lower odds of readmission compared to those whose clinician did not believe they were at risk for readmission (OR 0.24, 95% CI 0.09 to 0.66,  $P=0.005$ ). Patients for whom clinicians attributed inability to pay for treatment as the reason for potential post-discharge mortality had 5.53 times the odds of post-discharge mortality (95% CI 1.22 to 25.10,  $P=0.03$ ).

## Discussion

Among nearly 3,900 neonates and young children discharged from referral hospitals in Dar es Salaam, Tanzania and Monrovia, Liberia, clinician impression had poor discriminatory value for identifying those at risk of unplanned hospital readmissions and post-discharge mortality. Medical officer clinician impression at both sites had fair discriminatory value in identifying young children at risk of readmission. Clinician perception of inability to pay for treatment was associated with readmission and post-discharge mortality.

The poor discriminatory value among clinicians in identifying neonates and young children at risk of readmission and post-discharge mortality differs from findings in studies in high-income settings that assessed the diagnosis of acute coronary syndrome and sinusitis in adults,<sup>18,19</sup> the presence of pneumonia<sup>20</sup> or the development of severe pneumonia among children,<sup>13</sup> or the presence of appendicitis among children.<sup>21</sup> This difference is likely multifactorial in nature and may consist of differences in available diagnostic and prognostic resources. Prior studies suggest that laboratory capabilities in resource-limited settings are inadequate,<sup>22–24</sup> leading to dependence on clinical exam findings to make diagnoses and determine prognosis,<sup>25</sup> which may hinder the accuracy of clinician impression in identifying neonates and young children at risk of untoward post-discharge outcomes. Moreover, clinicians may not consider key factors in the home (e.g., access to healthcare facilities and maternal health) that may contribute to post-discharge outcomes.

Clinician impression had poor discriminatory value in identifying neonates and young children at risk for readmission or post-discharge mortality regardless of the time from discharge to either event. Prior studies of clinician impression assessed outcomes within hours or days<sup>13,18,19</sup> and not up to 60 days. Several clinical prediction rules for post-discharge mortality have been developed but none have been validated or implemented in sub-Saharan Africa.<sup>6,9,10</sup> Thus, the current state of prognostic determination after discharge depends on clinician impression. Clinical decision aids to identify at-risk neonates and young children that perform well outside the population of derivation are urgently needed. Such clinical decision aids should include commonly collected variables and may include biomarkers to add precision to risk stratification to identify neonates and young children at risk of post-discharge morbidity and mortality.<sup>26–28</sup>

Clinician impression among medical officers had fair discriminatory value in identifying young children at risk of readmission. This may be due to the combination of more clinical experience than interns/residents and more time spent with patients than consultants/specialists who often spend less time directly with patients and more time supervising clinical care. Prior studies conducted in high-income settings demonstrate that clinician impression of less experienced clinicians may have less discriminatory value than that of more experienced clinicians.<sup>13</sup>

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3 Our examination of reasons for estimated outcomes suggested that clinician perception of  
4 inability to pay for future treatment was associated with higher risk of readmission and post-  
5 discharge mortality. Young children from lower socioeconomic status have poorer overall health  
6 outcomes compared to young children from higher socioeconomic households in sub-Saharan  
7 Africa.<sup>29-31</sup> This is particularly relevant in the post-discharge period during which the financial  
8 burden of care seeking may influence the ability for a family to seek additional clinical care after  
9 a potentially costly hospital admission.  
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### 12 *Limitations*

14 Clinician impression is multifactorial and depends on clinical training as well as available  
15 laboratory, radiological, and clinical data that may not have been available to all discharging  
16 clinicians. We did not assess the availability of these in our analysis. We did not include nurses  
17 or clinical officers in our study, which is a limitation as these groups may have good insight into  
18 potential adverse outcomes after discharge. We could not account for the potential role that  
19 variations in the quality of clinical care provided to patients may have had in readmissions or  
20 post-discharge mortality.  
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### 23 **Conclusions**

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26 Clinician impression had poor discriminatory value in identifying neonates and young children at  
27 risk of unplanned hospital readmission and post-discharge mortality at two referral hospitals in  
28 Dar es Salaam, Tanzania and Monrovia, Liberia. Validated and objective clinical decision aids to  
29 assist clinicians in the identification of young children at risk of readmission and post-discharge  
30 mortality may facilitate the identification of those at greatest risk.  
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3 **Author Contributions:** CAR, RK, RCI, AS, EG, HKM, CRS, MN, KPM, and CD  
4 conceptualized and designed the study. CAR, RCI, JK, AS, EG, CRS, KPM, and CD oversaw  
5 data collection and verified the underlying data. CAR and AM verified the underlying data. AM  
6 conducted the statistical analyses. CAR wrote the first draft of the manuscript. CAR, RK, RCI,  
7 JK, Y-JC, AS, EG, HKM, CRS, AM, MN, CRM, CGW, RFB, KPM, and CD interpreted the  
8 data, reviewed, and provided input to the final draft. CAR had final responsibility for the  
9 decision to submit for publication.

10  
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**Figure 1.** Flow diagram for included neonates and children aged 1-59 months discharged from the neonatal wards and pediatric wards in Dar es Salaam, Tanzania and Monrovia, Liberia

Confidential: For Review Only



**Table 1.** Characteristics of neonates and young children included in the evaluation of clinician impression on predicting 60-day hospital readmission or post-discharge mortality

Patient Characteristics	Overall Population, (N=3,896) n (%)	Readmitted to Hospital 60 Days After Hospital Discharge, (N=187) n (%)	Died 60 Days After Hospital Discharge, (N=120) n (%)
<b>Discharged from Neonatal Ward<sup>1</sup></b>	2,173 (55.8)	80 (3.6)	61 (2.8)
Age in days at discharge, median (IQR)	2 (1, 7)	4 (2, 13)	2 (0, 10)
<b>Discharged from Pediatrics Ward</b>	1,723 (44.2)	107 (6.2)	59 (3.4)
Age in months for pediatric patients aged 1-59 months, median (IQR) <sup>2</sup>	12 (5, 24)	8 (4, 18)	8 (4, 21)
<b>Sex<sup>3</sup></b>			
Male	2,198 (56.4)	109 (5.0)	74 (3.4)
Female	1,691 (43.4)	78 (4.6)	46 (2.7)
<b>Country</b>			
Tanzania	1,997 (51.3)	140 (7.0)	63 (3.2)
Liberia	1,899 (48.7)	47 (2.5)	57 (3.0)
<b>Disposition from the Hospital</b>			
Discharge	3,775 (96.8)	186 (4.9)	106 (2.8)
Left against medical advice	119 (3.1)	1 (0.8)	14 (11.8)
Transfer to another facility	2 (0.05)	0 (0.0)	0 (0.0)

<sup>1</sup>24 neonates had missing age.

<sup>2</sup>31 young children had missing age.

<sup>3</sup>7 participants did not have a documented sex.



**Table 2.** Association of discharging clinicians' predicted probability and unplanned 60-day hospital readmission among neonates and young children

	Total, n (%) (N=3,896)	Readmitted 60 Days After Hospital Discharge, n (%) (N=187)	P Value <sup>1</sup>
<b>Clinician predicted probability</b>			<0.001
0%	2,838 (94.1)	151 (5.1)	
1-5%	9 (100)	0 (0.0)	
6-20%	26 (89.7)	3 (10.3)	
21-40%	736 (97.8)	16 (2.2)	
41-60%	38 (92.7)	3 (7.3)	
61-80%	237 (95.9)	10 (4.1)	
81-99%	11 (73.3)	4 (26.7)	

<sup>1</sup>By Chi square test to assess independence of clinician predicted probability and likelihood of hospital readmission.

Note: There was 1 respondent who estimated that the discharged child was at risk of hospital readmission but did not assign a proportion.

**Table 3.** Test characteristics for clinician predicted probability of unplanned,60-day hospital readmission among all enrolled neonates and young children aged 1-59 months overall and by clinician type

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>All Clinicians (N=3,895)<sup>1</sup></b>					<b>0.46 (0.43, 0.49)</b>
≤5	19.3 (13.9, 25.1)	72.5 (71.0, 73.8)	3.4 (2.5, 4.4)	94.7 (94.3, 95.1)	
≤20	19.3 (13.9, 25.1)	72.7 (71.3, 74.1)	3.4 (2.5, 4.5)	94.7 (94.4, 95.1)	
≤40	17.7 (12.3, 23.5)	73.7 (72.3, 75.1)	3.3 (2.4, 4.3)	94.7 (94.3, 95.0)	
≤60	16.0 (11.2, 21.9)	74.3 (72.9, 75.7)	3.0 (2.1, 4.1)	94.6 (94.3, 94.9)	
≤80	7.5 (4.2, 11.8)	93.7 (92.9, 94.4)	5.6 (3.1, 8.7)	95.3 (95.1, 95.5)	
≤99	2.1 (0.5, 4.3)	99.8 (99.7, 99.9)	36.4 (9.1, 66.7)	95.3 (95.2, 95.4)	
<b>Consultant/Specialist (N=175)</b>					<b>0.41 (0.28, 0.55)</b>
≤5	16.7 (0, 50.0)	69.2 (62.1, 75.8)	1.9 (0.0, 5.7)	95.8 (94.7, 97.6)	
≤20	16.7 (0, 50.0)	69.8 (62.7, 76.3)	1.9 (0, 5.9)	95.9 (94.7, 97.6)	
≤40	16.7 (0, 50.0)	73.4 (66.3, 79.9)	2.1 (0, 6.7)	96.1 (95.0, 97.7)	
≤60	0.0 (0.0, 0.0)	75.2 (68.6, 81.1)	0.0 (0.0, 0.0)	95.5 (95.1, 95.8)	
≤80	0.0 (0.0, 0.0)	85.2 (79.9, 90.5)	0.0 (0.0, 0.0)	96.0 (95.7, 96.2)	
≤99	0.0 (0.0, 0.0)	100 (100, 100)	NA	96.6 (96.6, 96.6)	
<b>Intern/Resident (N=3,502)</b>					<b>0.44 (0.41, 0.47)</b>
≤5	14.9 (9.9, 20.5)	72.4 (70.8, 73.9)	2.5 (1.7, 3.5)	94.6 (94.3, 95.0)	
≤20	14.9 (9.9, 20.5)	72.5 (70.9, 74.0)	2.5 (1.7, 3.5)	94.6 (94.3, 95.0)	
≤40	14.9 (9.9, 20.5)	73.2 (71.6, 74.7)	2.6 (1.7, 3.6)	94.7 (94.4, 95.1)	
≤60	13.6 (8.7, 19.3)	73.5 (71.9, 75.0)	2.4 (1.6, 3.4)	94.6 (94.3, 94.9)	
≤80	5.6 (2.5, 9.3)	94.1 (93.3, 94.9)	4.3 (1.9, 7.3)	95.4 (95.2, 95.6)	
≤99	1.2 (0.0, 3.1)	99.9 (99.8, 100)	40.0 (0.0, 100)	95.5 (95.4, 95.5)	
<b>Medical Officer (N=217)</b>					<b>0.67 (0.55, 0.79)</b>
≤5	55.0 (30.0, 75.0)	76.1 (70.1, 81.7)	18.9 (11.6, 26.5)	94.3 (91.6, 96.9)	
≤20	55.0 (30.0, 75.0)	78.7 (72.6, 84.3)	20.7 (12.7, 29.4)	94.5 (91.9, 96.9)	
≤40	40.0 (20.0, 60.0)	82.2 (76.6, 87.3)	18.4 (9.4, 28.9)	93.1 (90.8, 95.5)	
≤60	40.0 (20.0, 60.0)	86.8 (81.7, 91.4)	23.5 (12.1, 36.4)	93.4 (91.3, 95.7)	
≤80	25.0 (9.8, 45.0)	93.9 (90.4, 96.9)	29.4 (10.5, 50.0)	92.5 (90.9, 94.4)	
≤99	10.0 (0.0, 25.0)	97.9 (95.9, 99.5)	33.3 (0.0, 80.0)	91.5 (90.6, 92.8)	

<sup>1</sup>There was 1 respondent who estimated that the discharged child was at risk of hospital readmission but did not assign a proportion.

**Table 4.** Association of discharging clinicians' predicted probability and all-cause,60-day post-discharge mortality among neonates and young children

	<b>Total, n (%) (N=3,896)</b>	<b>Died 60 Days After Hospital Discharge, n (%) (N=120)</b>	<b>P Value<sup>1</sup></b>
<b>Clinician predicted probability</b>			0.002
0%	3,746 (97.2)	109 (2.8)	
1-5%	10 (100)	0 (0.0)	
6-20%	9 (81.8)	2 (18.2)	
21-40%	13 (86.7)	2 (13.3)	
41-60%	95 (95.0)	5 (5.0)	
61-80%	21 (95.5)	1 (4.5)	
81-99%	2 (66.7)	1 (33.3)	

<sup>1</sup>By Chi square test to assess independence of clinician predicted probability and likelihood of hospital readmission.

**Table 5.** Test characteristics for clinician predicted probability of all-cause, 60-day post-discharge mortality among all enrolled neonates and young children aged 1-59 months by clinician type

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>All Clinicians (N=3,895)<sup>1</sup></b>					<b>0.53 (0.50, 0.55)</b>
≤5	9.2 (4.2, 15.0)	96.3 (95.7, 96.9)	7.3 (3.5, 11.9)	97.1 (96.9, 97.3)	
≤20	9.2 (4.2, 15.0)	96.6 (95.9, 97.2)	7.7 (3.8, 12.7)	97.1 (96.9, 97.3)	
≤40	7.5 (3.3, 12.5)	96.7 (96.2, 97.3)	6.8 (2.9, 11.6)	97.1 (96.9, 97.2)	
≤60	5.8 (1.7, 10.8)	97.1 (96.5, 97.6)	5.8 (1.9, 10.5)	97.0 (96.9, 97.2)	
≤80	1.7 (0.0, 4.2)	99.4 (99.2, 99.7)	8.3 (0.0, 22.7)	96.9 (96.9, 97.0)	
≤99	0.8 (0.0, 2.5)	99.9 (99.9, 100)	50.0 (0.0, 100)	96.9 (96.9, 96.9)	
<b>Specialist or Consultant (N=175)</b>					<b>0.49 (0.48, 0.50)</b>
≤5	0.0 (0.0, 0.0)	97.6 (95.2, 99.4)	0.0 (0.0, 0.0)	95.9 (95.8, 95.9)	
≤20	0.0 (0.0, 0.0)	97.6 (95.2, 99.4)	0.0 (0.0, 0.0)	95.9 (95.8, 95.9)	
≤40	0.0 (0.0, 0.0)	97.6 (95.2, 99.4)	0.0 (0.0, 0.0)	95.9 (95.8, 95.9)	
≤60	0.0 (0.0, 0.0)	97.6 (95.2, 99.4)	0.0 (0.0, 0.0)	95.9 (95.8, 95.9)	
≤80	0.0 (0.0, 0.0)	98.8 (97.0, 100)	0.0 (0.0, 0.0)	95.9 (95.9, 96.0)	
≤99	0.0 (0.0, 0.0)	100 (100, 100)	-	96.0 (96.0, 96.0)	
<b>Intern/Resident (N=3,502)</b>					<b>0.53 (0.50, 0.56)</b>
≤5	8.8 (3.9, 14.7)	97.1 (96.5, 97.6)	8.3 (3.7, 13.8)	97.3 (97.1, 97.4)	
≤20	8.8 (3.9, 14.7)	97.2 (96.6, 97.7)	8.5 (3.8, 14.2)	97.3 (97.1, 97.4)	
≤40	6.9 (2.9, 12.7)	97.2 (96.6, 97.7)	6.8 (2.7, 12.0)	97.2 (97.1, 97.4)	
≤60	5.8 (1.9, 10.8)	97.2 (96.7, 97.8)	5.9 (1.9, 11.0)	97.2 (97.1, 97.3)	
≤80	1.9 (0.0, 4.9)	99.6 (99.4, 99.8)	11.8 (0.0, 30.8)	97.1 (97.1, 97.2)	
≤99	0.9 (0.0, 2.9)	99.9 (99.9, 100.0)	50.0 (0.0, 100.8)	97.1 (97.1, 97.2)	
<b>Medical Officer (N=218)</b>					<b>0.51 (0.38, 0.64)</b>
≤5	18.2 (0.0, 45.5)	82.6 (77.8, 87.9)	5.0 (0, 12.2)	94.9 (93.8, 96.5)	
≤20	18.2 (0.0, 45.5)	85.9 (81.2, 90.3)	6.1 (0, 14.8)	95.1 (94.1, 96.7)	
≤40	18.2 (0.0, 45.5)	88.9 (84.5, 92.8)	7.7 (0, 18.7)	95.3 (94.2, 96.8)	
≤60	9.1 (0.0, 27.3)	93.7 (90.3, 96.6)	6.7 (0, 23.1)	95.1 (94.5, 96.1)	
≤80	0.0 (0.0, 0.0)	97.6 (95.2, 99.5)	0.0 (0.0, 0.0)	94.8 (94.7, 94.9)	
≤99	0.0 (0.0, 0.0)	100 (100, 100)	-	94.9 (94.9, 94.9)	

<sup>1</sup>There was 1 respondent who estimated that the discharged child was at risk of hospital readmission but did not assign a proportion.

**Table 6.** Reasons for perceived risk of hospital readmission and post-discharge mortality

<b>Clinician Cited Reason for Outcome</b>	<b>Hospital Readmission, n (%)</b>	<b>No Hospital Readmission, n (%)</b>	<b>Odds Ratio (95% Confidence Interval)</b>	<b>P value<sup>1</sup></b>
No risk	148 (79.1)	2584 (69.7)	<i>Referent</i>	--
Clinician perceived inability to pay for treatment	3 (1.6)	11 (0.3)	4.76 (1.31, 17.25)	0.02
Clinician perceived social concerns	3 (1.6)	70 (1.9)	0.75 (0.23, 2.41)	0.63
Clinician perceived progression of illness	29 (15.5)	754 (20.3)	0.67 (0.45, 1.01)	0.05
Other <sup>2</sup>	4 (2.1)	290 (7.8)	0.24 (0.09, 0.66)	0.005
<b>Clinician Cited Reason for Outcome</b>	<b>Died Within 60 Days, n (%)</b>	<b>Did not Die Within 60 Days, n (%)</b>	<b>Odds Ratio (95% Confidence Interval)</b>	<b>P value<sup>1</sup></b>
No risk	80 (66.7)	2652 (70.2)	<i>Referent</i>	--
Clinician perceived inability to pay for treatment	2 (1.7)	12 (0.3)	5.53 (1.22, 25.10)	0.03
Clinician perceived social concerns	1 (0.8)	72 (1.9)	0.46 (0.06, 3.36)	0.44
Clinician perceived progression of illness	29 (24.2)	754 (20.0)	1.28 (0.83, 1.97)	0.27
Other <sup>2</sup>	8 (6.7)	286 (7.6)	0.93 (0.44, 1.94)	0.84

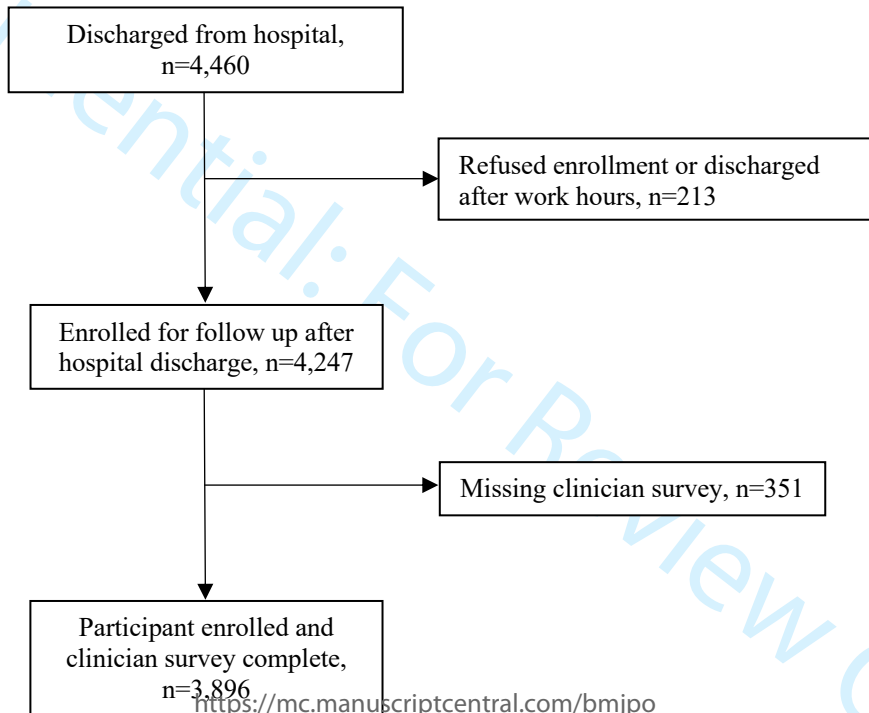
<sup>1</sup>Calculated using ordinal logistic regression analyses to assess whether the perceived reason for risk was associated with the participant's perceived likelihood of the outcomes.

<sup>2</sup>Included concerns about caregiver understanding, general clinician impression, and patient with history of recurrent illness.

## References

1. Kedebe F. Time to readmission and associated factors after post treatment discharge of severe acute malnourished under-five children in Pawe General Hospital. *J Heal Popul Nutr* 2022;41:29.
2. Pavlinac PB, Singa BO, Tickell KD, et al. Azithromycin for the prevention of rehospitalisation and death among Kenyan children being discharged from hospital: a double-blind, placebo-controlled, randomised controlled trial. *Lancet Glob Heal* 2021;9:e1569-e1578.
3. You D, Hug L, Ejdemyr S, et al. Global, regional, and national levels and trends in under-5 mortality between 1990 and 2015, with scenario-based projections to 2030: A systematic analysis by the un Inter-Agency Group for Child Mortality Estimation. *Lancet* 2015;386:2275-2286.
4. Abbafati C, Machado DB, Cislighi B, et al. Global age-sex-specific fertility, mortality, healthy life expectancy (HALE), and population estimates in 204 countries and territories, 1950–2019: a comprehensive demographic analysis for the Global Burden of Disease Study 2019. *Lancet* 2020;396:1160-1203.
5. Wiens MO, Pawluk S, Kissoon N, et al. Pediatric Post-Discharge Mortality in Resource Poor Countries: A Systematic Review. *PLoS One* 2013;8:e66698.
6. Diallo AH, Sayeem Bin Shahid ASM, Khan AF, et al. Childhood mortality during and after acute illness in Africa and south Asia: a prospective cohort study. *Lancet Glob Heal* 2022;10:e673-e684.
7. Connon R, George EC, Olupot-Olupot P, et al. Incidence and predictors of hospital readmission in children presenting with severe anaemia in Uganda and Malawi: a secondary analysis of TRACT trial data. *BMC Public Health* 2021;21:1480.
8. Goodman DM, Casale MT, Rychlik K, et al. Development and Validation of an Integrated Suite of Prediction Models for All-Cause 30-Day Readmissions of Children and Adolescents Aged 0 to 18 Years. 2022;5:e2241513.
9. Wiens MO, Kumbakumba E, Larson CP, et al. Postdischarge mortality in children with acute infectious diseases: Derivation of postdischarge mortality prediction models. *BMJ Open* 2015;5:e009449.
10. Madrid L, Casellas A, Sacoer C, et al. Postdischarge Mortality Prediction in Sub-Saharan Africa. *Pediatrics* 2019;143:e20180606.
11. Horwood J, Cabral C, Hay AD, Ingram J. Primary care clinician antibiotic prescribing decisions in consultations for children with RTIs: a qualitative interview study. *Br J Gen Pr* 2016;66:e207-e213.
12. Van den Bruel A, Thompson M, Buntinx F, Mant D. Clinicians' gut feeling about serious infections in children: observational study. *BMJ* 2012;345:e6144.
13. Gao HM, Ambroggio L, Shah SS, Ruddy RM, Florin TA. Predictive value of clinician "gestalt" in pediatric community-acquired pneumonia. *Pediatrics* 2021;147:e2020041582.
14. Turnbull S, Lucas PJ, Redmond NM, et al. What gives rise to clinician gut feeling, its influence on management decisions and its prognostic value for children with RTI in primary care: a prospective cohort study. *BMC Fam Pr* 2018;19:25.
15. Paul S, Tickell KD, Ojee E, et al. Knowledge, attitudes, and perceptions of Kenyan healthcare workers regarding pediatric discharge from hospital. *PLoS One*

- 2021;16:e0249569.
16. Rees CA, Kisenge R, Ideh RC, et al. A Prospective, observational cohort study to identify neonates and children at risk of postdischarge mortality in Dar es Salaam, Tanzania and Monrovia, Liberia: The PPDM study protocol. *BMJ Paediatr Open* 2022;6:e001379.
  17. Robin X, Turck N, Hainard A, et al. pROC: an open-source package for R and S+ to analyze and compare ROC curves. *BMC Bioinformatics* 2011;12:77.
  18. Williams JW Jr, Simel DL, Roberts L SG. Clinical evaluation for sinusitis. Making the diagnosis by history and physical examination. *Ann Intern Med* 1992;117:705-710.
  19. Oliver G, Reynard C, Morris N, Body R. Can Emergency Physician Gestalt “Rule In” or “Rule Out” Acute Coronary Syndrome: Validation in a Multicenter Prospective Diagnostic Cohort Study. *Acad Emerg Med* 2020;27:24-30.
  20. Neuman MI, Scully KJ, Kim D, Shah S, Bachur R. Physician assessment of the likelihood of pneumonia in a pediatric emergency department. *Pediatr Emerg Care* 2010;26:817-822.
  21. Lee WH, O’Brien S, Skarin D, et al. Accuracy of clinician gestalt in diagnosing appendicitis in children presenting to the emergency department. *Emerg Med Australas* 2019;31:612-618.
  22. Yadav H, Shah D, Sayed S, Horton S, Schroeder LF. Availability of essential diagnostics in ten low-income and middle-income countries: results from national health facility surveys. *Lancet Glob Heal* 2021;9:e1553-e1560.
  23. Peeling RW, Mabey D. Point-of-care tests for diagnosing infections in the developing world. *Clin Microbiol Infect* 2010;16:1062-1069.
  24. Singh H, Schiff GD, Graber ML, Onakpoya I TM. The global burden of diagnostic errors in primary care. *BMJ Qual Saf* 2017;26:484-494.
  25. World Health Organization. Integrated Management of Childhood Illness: management of the sick young infant aged up to 2 months: IMCI chart booklet. [https://www.who.int/maternal\\_child\\_adolescent/documents/management-sick-young-infant-0-2-months/en/](https://www.who.int/maternal_child_adolescent/documents/management-sick-young-infant-0-2-months/en/). Published 2019. Accessed September 24, 2021.
  26. Njunge JM, Tickell K, Diallo AH, et al. The Childhood Acute Illness and Nutrition (CHAIN) network nested case-cohort study protocol: a multi-omics approach to understanding mortality among children in sub-Saharan Africa and South Asia. *Gates Open Res* 2022;6:77.
  27. Njunge JM, Gwela A, Kibinge NK, et al. Biomarkers of post-discharge mortality among children with complicated severe acute malnutrition. *Sci Rep* 2019;9:5981.
  28. Sarangam ML, Namazzi R, Datta D, et al. Intestinal Injury Biomarkers Predict Mortality in Pediatric Severe Malaria. *MBio* 2022;13:e0132522.
  29. Ogbo FA, Ezech OK, Awosemo AO, et al. Determinants of trends in neonatal, post-neonatal, infant, child and under-five mortalities in Tanzania from 2004 to 2016. *BMC Public Health* 2019;19:1243.
  30. Burkey MD, Weiser SD, Fehmie D, et al. Socioeconomic determinants of mortality in HIV: evidence from a clinical cohort in Uganda. *J Acquir Immune Defic Syndr* 2014;66:41-47.
  31. Ezech OK, Agho KE, Dibley MJ, Hall JJ, Page AN. Risk factors for post neonatal, infant, child and under-5 mortality in Nigeria: a pooled cross-sectional analysis. *BMJ Open* 2015;5:e006779.





**Supplemental Table 1.** Test characteristics for clinician predicted probability of all-cause, 60-day hospital readmission for neonates at both sites

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>All Clinicians (N=2,173)</b>					<b>0.41 (0.38, 0.46)</b>
≤5	8.75 (3.75, 15)	73.77 (71.91, 75.68)	1.25 (0.51, 2.19)	95.48 (95.19, 95.81)	
≤20	8.75 (3.75, 15)	73.87 (72, 75.73)	1.25 (0.51, 2.19)	95.48 (95.19, 95.81)	
≤40	8.75 (3.75, 15)	74.34 (72.48, 76.21)	1.28 (0.52, 2.23)	95.51 (95.23, 95.84)	
≤60	7.5 (2.5, 13.75)	74.58 (72.72, 76.4)	1.1 (0.36, 2.04)	95.46 (95.2, 95.78)	
≤80	2.5 (0, 6.25)	94.84 (93.88, 95.8)	1.75 (0, 4.47)	96.21 (96.11, 96.36)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.32 (96.32, 96.32)	
<b>Interns/Residents (N=2,076)</b>					<b>0.41 (0.38, 0.44)</b>
≤5	7.79 (2.6, 14.29)	73.69 (71.68, 75.54)	1.12 (0.37, 2.03)	95.39 (95.11, 95.72)	
≤20	7.79 (2.6, 14.29)	73.74 (71.74, 75.59)	1.12 (0.37, 2.04)	95.4 (95.12, 95.73)	
≤40	7.79 (2.6, 14.29)	74.09 (72.09, 75.94)	1.14 (0.37, 2.07)	95.42 (95.14, 95.74)	
≤60	7.79 (2.6, 14.29)	74.29 (72.34, 76.14)	1.15 (0.38, 2.08)	95.43 (95.15, 95.76)	
≤80	2.6 (0, 6.49)	95.15 (94.15, 96.05)	1.96 (0, 4.85)	96.2 (96.09, 96.35)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.29 (96.29, 96.29)	
<b>Specialist or Consultant (N=66)</b>					<b>0.53 (0.13, 0.93)</b>
≤5	50 (0, 100)	65.62 (53.12, 76.56)	4.35 (0, 10.53)	97.67 (94.87, 100)	
≤20	50 (0, 100)	67.19 (54.69, 79.69)	4.55 (0, 11.11)	97.73 (95.12, 100)	
≤40	50 (0, 100)	71.88 (60.94, 82.81)	5.26 (0, 12.5)	97.87 (95.45, 100)	
≤60	0 (0, 0)	73.44 (62.5, 84.38)	0 (0, 0)	95.92 (95.24, 96.43)	
≤80	0 (0, 0)	82.81 (73.44, 92.19)	0 (0, 0)	96.36 (95.92, 96.72)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.97 (96.97, 96.97)	
<b>Medical Officer (N=30)</b>					<b>0.50 (NA, NA)</b>
≤5	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤20	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤40	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤60	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤80	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	

**Supplemental Table 2.** Test characteristics for clinician predicted probability of all-cause, 60-day hospital readmission for young children aged 1-59 months at both sites

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>All Clinicians (N=1,722)</b>					<b>0.49 (0.45, 0.54)</b>
≤5	27.1 (18.69, 35.51)	70.77 (68.54, 73)	5.77 (4.09, 7.57)	93.6 (92.9, 94.33)	
≤20	27.1 (18.69, 35.51)	71.21 (68.98, 73.44)	5.85 (4.14, 7.69)	93.64 (92.95, 94.36)	
≤40	24.3 (15.89, 32.71)	72.76 (70.65, 74.86)	5.51 (3.81, 7.4)	93.53 (92.9, 94.23)	
≤60	22.43 (14.95, 30.84)	73.87 (71.7, 75.98)	5.34 (3.59, 7.35)	93.48 (92.88, 94.19)	
≤80	11.21 (5.61, 17.76)	92.26 (90.84, 93.44)	8.63 (4.32, 13.71)	94 (93.63, 94.45)	
≤99	3.74 (0.93, 7.48)	99.57 (99.26, 99.88)	35.71 (10, 66.67)	93.98 (93.81, 94.21)	
<b>Interns/Residents (N=1,426)</b>					<b>0.46 (0.42, 0.51)</b>
≤5	21.43 (13.1, 30.95)	70.57 (68.18, 73.03)	4.32 (2.7, 6.11)	93.47 (92.81, 94.2)	
≤20	21.43 (13.1, 30.95)	70.72 (68.33, 73.1)	4.35 (2.71, 6.13)	93.48 (92.82, 94.22)	
≤40	21.43 (13.1, 30.95)	71.83 (69.45, 74.14)	4.51 (2.81, 6.35)	93.58 (92.93, 94.29)	
≤60	19.05 (10.71, 27.38)	72.35 (69.97, 74.67)	4.08 (2.45, 5.93)	93.43 (92.83, 94.12)	
≤80	8.33 (3.57, 14.29)	92.55 (91.06, 93.96)	6.25 (2.48, 11.2)	94.14 (93.83, 94.54)	
≤99	2.38 (0, 5.95)	99.78 (99.48, 100)	40 (0, 100)	94.23 (94.09, 94.43)	
<b>Specialist or Consultant (N=109)</b>					<b>0.36 (0.31, 0.40)</b>
≤5	0 (0, 0)	71.43 (62.86, 80)	0 (0, 0)	94.94 (94.29, 95.45)	
≤20	0 (0, 0)	71.43 (62.86, 80)	0 (0, 0)	94.94 (94.29, 95.45)	
≤40	0 (0, 0)	74.29 (65.71, 82.86)	0 (0, 0)	95.12 (94.52, 95.6)	
≤60	0 (0, 0)	76.19 (67.62, 83.81)	0 (0, 0)	95.24 (94.67, 95.65)	
≤80	0 (0, 0)	86.67 (79.98, 92.38)	0 (0, 0)	95.79 (95.45, 96.04)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.33 (96.33, 96.33)	
<b>Medical Officer (N=187)</b>					<b>0.67 (0.54, 0.80)</b>
≤5	57.89 (36.84, 78.95)	72.02 (64.88, 79.17)	19.2 (12.07, 26.32)	93.85 (90.76, 96.85)	
≤20	57.89 (36.84, 78.95)	75 (67.86, 81.55)	20.93 (13.46, 28.85)	94.07 (91.11, 96.95)	
≤40	42.11 (21.05, 63.16)	79.17 (72.62, 85.12)	18.75 (9.37, 28.57)	92.36 (89.61, 95.17)	
≤60	42.11 (21.05, 63.16)	84.52 (78.57, 89.88)	23.68 (12.12, 36)	92.86 (90.26, 95.42)	
≤80	26.32 (10.53, 47.37)	92.86 (88.69, 96.43)	30 (10.53, 50)	91.76 (89.88, 93.98)	
≤99	10.53 (0, 26.32)	97.62 (95.24, 99.4)	33.33 (0, 80)	90.61 (89.56, 92.18)	

**Supplemental Table 3.** Test characteristics for clinician predicted probability of all-cause, 60-day hospital readmission among all participants by study site

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>Tanzania</b>					
All Clinicians (N=1,996)					0.53 (0.51, 0.56)
≤5	10.71 (6.43, 15.71)	95.64 (94.67, 96.5)	15.56 (9.2, 22.64)	93.42 (93.1, 93.78)	
≤20	10.71 (6.43, 15.71)	96.07 (95.15, 96.93)	17.05 (10.23, 24.76)	93.44 (93.13, 93.81)	
≤40	8.57 (4.29, 13.57)	97.04 (96.23, 97.79)	17.81 (9.86, 27.59)	93.36 (93.08, 93.7)	
≤60	7.14 (3.57, 11.43)	97.79 (97.09, 98.38)	19.61 (9.76, 30.77)	93.32 (93.06, 93.62)	
≤80	4.29 (1.43, 7.86)	99.14 (98.71, 99.52)	26.92 (9.52, 47.62)	93.2 (93.01, 93.46)	
≤99	2.86 (0.71, 5.71)	99.68 (99.41, 99.89)	40 (10, 72.73)	93.15 (93, 93.35)	
<b>Liberia</b>					
All Clinicians (N=1,899)					0.52 (0.43, 0.60)
≤5	55.32 (40.43, 70.21)	50.76 (48.49, 53.08)	2.77 (2.04, 3.47)	97.81 (97.1, 98.5)	
≤20	55.32 (40.43, 70.21)	50.7 (48.49, 52.97)	2.77 (2.04, 3.47)	97.81 (97.1, 98.49)	
≤40	55.32 (40.43, 70.21)	49.78 (47.52, 52.11)	2.72 (2.01, 3.42)	97.77 (97.04, 98.46)	
≤60	57.45 (42.55, 70.21)	49.24 (47.03, 51.62)	2.8 (2.09, 3.47)	97.86 (97.13, 98.55)	
≤80	82.98 (72.34, 93.62)	11.77 (10.37, 13.23)	2.34 (2.02, 2.61)	96.51 (94.09, 98.6)	
≤99	100 (100, 100)	0.05 (0, 0.16)	2.48 (2.47, 2.48)	100 (100, 100)	

**Supplemental Table 4.** Test characteristics for clinician predicted probability of all-cause, 60-day hospital readmission among all participants by time to unplanned hospital readmission

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>Day 7</b>					0.49 (0.42, 0.56)
≤5	23.4 (10.64, 36.17)	72.77 (71.34, 74.22)	1.03 (0.49, 1.62)	98.72 (98.54, 98.95)	
≤20	23.4 (10.64, 36.17)	73 (71.57, 74.45)	1.04 (0.5, 1.64)	98.73 (98.54, 98.95)	
≤40	23.4 (10.64, 36.17)	74.01 (72.61, 75.39)	1.08 (0.51, 1.7)	98.75 (98.56, 98.96)	
≤60	23.4 (10.64, 36.17)	74.66 (73.28, 76.09)	1.1 (0.53, 1.74)	98.76 (98.57, 98.97)	
≤80	12.77 (4.26, 23.4)	93.69 (92.96, 94.49)	2.38 (0.78, 4.35)	98.87 (98.76, 99.01)	
≤99	4.26 (0, 10.64)	99.77 (99.58, 99.9)	16.67 (0, 46.15)	98.84 (98.79, 98.92)	
<b>Day 14</b>					0.45 (0.41, 0.50)
≤5	17.28 (9.88, 25.93)	72.68 (71.24, 74.07)	1.31 (0.75, 1.95)	97.63 (97.43, 97.87)	
≤20	17.28 (9.88, 25.93)	72.92 (71.47, 74.28)	1.32 (0.75, 1.97)	97.64 (97.43, 97.88)	
≤40	16.05 (8.64, 23.46)	73.89 (72.47, 75.25)	1.28 (0.69, 1.89)	97.64 (97.43, 97.86)	
≤60	16.05 (8.64, 23.46)	74.57 (73.13, 75.91)	1.31 (0.71, 1.94)	97.66 (97.46, 97.88)	
≤80	7.41 (2.47, 13.58)	93.68 (92.87, 94.47)	2.38 (0.8, 4.43)	97.94 (97.83, 98.08)	
≤99	2.47 (0, 6.17)	99.76 (99.61, 99.9)	16.67 (0, 44.44)	97.97 (97.91, 98.04)	
<b>Day 30</b>					0.46 (0.42, 0.49)
≤5	18.64 (11.86, 25.42)	72.57 (71.06, 73.95)	2.06 (1.34, 2.87)	96.61 (96.34, 96.91)	
≤20	18.64 (11.86, 25.42)	72.82 (71.33, 74.21)	2.08 (1.35, 2.9)	96.62 (96.35, 96.92)	
≤40	16.95 (10.17, 23.73)	73.79 (72.31, 75.17)	1.97 (1.23, 2.74)	96.6 (96.33, 96.87)	
≤60	15.25 (9.32, 22.03)	74.42 (72.97, 75.8)	1.81 (1.1, 2.62)	96.56 (96.31, 96.83)	
≤80	6.78 (2.54, 11.86)	93.65 (92.82, 94.39)	3.15 (1.24, 5.53)	96.98 (96.85, 97.14)	
≤99	1.69 (0, 4.24)	99.76 (99.6, 99.89)	16.67 (0, 44.44)	97.01 (96.96, 97.09)	
<b>Day 45</b>					0.46 (0.43, 0.49)
≤5	18.67 (12.67, 24.67)	72.52 (71.13, 73.91)	2.64 (1.84, 3.51)	95.7 (95.39, 96.02)	
≤20	18.67 (12.67, 24.67)	72.76 (71.35, 74.15)	2.66 (1.85, 3.54)	95.71 (95.41, 96.03)	
≤40	16.67 (11.33, 22.67)	73.72 (72.26, 75.09)	2.47 (1.69, 3.33)	95.66 (95.38, 95.97)	
≤60	15.33 (10, 21.33)	74.34 (72.92, 75.73)	2.33 (1.56, 3.19)	95.63 (95.37, 95.92)	
≤80	7.33 (3.33, 11.33)	93.67 (92.87, 94.45)	4.4 (2.21, 7.17)	96.19 (96.04, 96.37)	
≤99	2.67 (0.67, 5.35)	99.81 (99.68, 99.95)	35.71 (9.09, 66.67)	96.24 (96.16, 96.35)	

**Supplemental Table 5.** Test characteristics for clinician predicted probability of all-cause, 60-day post-discharge mortality for neonates at both sites

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>All Clinicians (N=2,173)</b>					<b>0.52 (0.49, 0.55)</b>
≤5	6.56 (1.64, 13.11)	96.97 (96.21, 97.73)	5.71 (1.41, 12.09)	97.29 (97.15, 97.48)	
≤20	6.56 (1.64, 13.11)	96.97 (96.21, 97.73)	5.71 (1.41, 12.09)	97.29 (97.15, 97.48)	
≤40	6.56 (1.64, 13.11)	96.97 (96.21, 97.73)	5.71 (1.41, 12.09)	97.29 (97.15, 97.48)	
≤60	4.92 (0, 11.48)	97.02 (96.21, 97.77)	4.35 (0, 9.86)	97.24 (97.11, 97.42)	
≤80	1.64 (0, 4.92)	99.62 (99.34, 99.86)	10 (0, 36.36)	97.23 (97.18, 97.32)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	97.19 (97.19, 97.19)	
<b>Interns/Residents (N=2,076)</b>					<b>0.52 (0.49, 0.55)</b>
≤5	7.14 (1.79, 14.29)	96.93 (96.14, 97.67)	5.95 (1.43, 11.94)	97.41 (97.26, 97.62)	
≤20	7.14 (1.79, 14.29)	96.93 (96.14, 97.67)	5.95 (1.43, 11.94)	97.41 (97.26, 97.62)	
≤40	7.14 (1.79, 14.29)	96.93 (96.14, 97.67)	5.95 (1.43, 11.94)	97.41 (97.26, 97.62)	
≤60	5.36 (0, 12.5)	96.98 (96.19, 97.72)	4.62 (0, 10.17)	97.36 (97.22, 97.55)	
≤80	1.79 (0, 5.36)	99.65 (99.36, 99.9)	11.11 (0, 42.86)	97.34 (97.29, 97.44)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	97.3 (97.3, 97.3)	
<b>Specialist or Consultant (N=66)</b>					<b>0.48 (0.46, 0.51)</b>
≤5	0 (0, 0)	96.77 (91.94, 100)	0 (0, 0)	93.75 (93.44, 93.94)	
≤20	0 (0, 0)	96.77 (91.94, 100)	0 (0, 0)	93.75 (93.44, 93.94)	
≤40	0 (0, 0)	96.77 (91.94, 100)	0 (0, 0)	93.75 (93.44, 93.94)	
≤60	0 (0, 0)	96.77 (91.94, 100)	0 (0, 0)	93.75 (93.44, 93.94)	
≤80	0 (0, 0)	98.39 (95.16, 100)	0 (0, 0)	93.85 (93.65, 93.94)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	93.94 (93.94, 93.94)	
<b>Medical Officer (N=30)<sup>1</sup></b>					<b>0.50 (NA, NA)</b>
≤5	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤20	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤40	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤60	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤80	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	

<sup>1</sup>All medical officers reported a perceived probability of 0% for neonates.

**Supplemental Table 6.** Test characteristics for clinician predicted probability of all-cause, 60-day post-discharge mortality for young children aged 1-59 months at both sites

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>All Clinicians (N=1,723)</b>					<b>0.54 (0.49, 0.58)</b>
≤5	11.86 (5.08, 20.34)	95.49 (94.53, 96.45)	8.43 (3.23, 14.77)	96.83 (96.56, 97.14)	
≤20	11.86 (5.08, 20.34)	96.09 (95.19, 97.06)	9.59 (3.66, 16.67)	96.85 (96.59, 97.16)	
≤40	8.47 (1.69, 16.95)	96.51 (95.61, 97.42)	7.84 (1.75, 14.82)	96.75 (96.52, 97.03)	
≤60	6.78 (1.69, 13.56)	97.12 (96.27, 97.96)	7.55 (1.79, 15.69)	96.7 (96.52, 96.95)	
≤80	1.69 (0, 5.08)	99.22 (98.74, 99.58)	6.25 (0, 25)	96.6 (96.54, 96.73)	
≤99	1.69 (0, 5.08)	99.94 (99.82, 100)	50 (0, 100)	96.63 (96.57, 96.74)	
<b>Interns/Residents (N=1,426)</b>					<b>0.54 (0.50, 0.59)</b>
≤5	10.87 (2.17, 19.57)	97.32 (96.38, 98.19)	11.63 (2.78, 21.95)	97.03 (96.76, 97.33)	
≤20	10.87 (2.17, 19.57)	97.54 (96.67, 98.33)	12.5 (3.03, 23.68)	97.04 (96.77, 97.34)	
≤40	6.52 (0, 15.22)	97.61 (96.74, 98.41)	8.11 (0, 18.18)	96.9 (96.7, 97.17)	
≤60	6.52 (0, 15.22)	97.61 (96.74, 98.41)	8.11 (0, 18.18)	96.9 (96.7, 97.17)	
≤80	2.17 (0, 6.52)	99.49 (99.06, 99.86)	11.11 (0, 42.86)	96.82 (96.75, 96.97)	
≤99	2.17 (0, 6.52)	99.93 (99.78, 100)	50 (0, 100)	96.84 (96.77, 96.98)	
<b>Specialist or Consultant (N=109)</b>					<b>0.49 (0.48, 0.50)</b>
≤5	0 (0, 0)	98.11 (95.28, 100)	0 (0, 0)	97.2 (97.12, 97.25)	
≤20	0 (0, 0)	98.11 (95.28, 100)	0 (0, 0)	97.2 (97.12, 97.25)	
≤40	0 (0, 0)	98.11 (95.28, 100)	0 (0, 0)	97.2 (97.12, 97.25)	
≤60	0 (0, 0)	98.11 (95.28, 100)	0 (0, 0)	97.2 (97.12, 97.25)	
≤80	0 (0, 0)	99.06 (97.17, 100)	0 (0, 0)	97.22 (97.17, 97.25)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	97.25 (97.25, 97.25)	
<b>Medical Officer (N=188)</b>					<b>0.50 (0.36, 0.64)</b>
≤5	20 (0, 50)	79.78 (73.6, 85.39)	5.13 (0, 12.12)	94.63 (93.24, 96.55)	
≤20	20 (0, 50)	83.71 (78.09, 88.76)	6.25 (0, 15)	94.87 (93.59, 96.71)	
≤40	20 (0, 50)	87.08 (82.02, 91.57)	7.69 (0, 19.05)	95.06 (93.83, 96.84)	
≤60	10 (0, 30)	92.7 (88.76, 96.07)	6.67 (0, 23.08)	94.8 (94.12, 96.02)	
≤80	0 (0, 0)	97.19 (94.38, 99.44)	0 (0, 0)	94.54 (94.38, 94.65)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	94.68 (94.68, 94.68)	

**Supplemental Table 7.** Test characteristics for clinician predicted probability of all-cause, 60-day post-discharge mortality among all participants by study site

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>Tanzania</b>					
All Clinicians (N=1,997)					0.53 (0.50, 0.56)
≤5	7.94 (1.59, 15.87)	97.88 (97.21, 98.5)	10.62 (2.63, 20.46)	97.02 (96.83, 97.27)	
≤20	7.94 (1.59, 15.87)	98.4 (97.83, 98.91)	13.51 (3.57, 25.93)	97.04 (96.85, 97.28)	
≤40	6.35 (1.59, 12.7)	98.76 (98.24, 99.22)	13.64 (3.45, 28.12)	97 (96.85, 97.21)	
≤60	4.76 (0, 11.11)	99.28 (98.91, 99.64)	16.67 (0, 37.51)	96.97 (96.82, 97.16)	
≤80	1.59 (0, 4.76)	99.69 (99.43, 99.9)	12.5 (0, 50)	96.88 (96.83, 96.99)	
≤99	1.59 (0, 4.76)	99.95 (99.84, 100)	50 (0, 100)	96.89 (96.84, 96.99)	
<b>Liberia</b>					
All Clinicians (N=1,899)					0.53 (0.49, 0.57)
≤5	7.94 (1.59, 15.87)	97.88 (97.21, 98.5)	10.62 (2.63, 20.46)	97.02 (96.83, 97.27)	
≤20	7.94 (1.59, 15.87)	98.4 (97.83, 98.91)	13.51 (3.57, 25.93)	97.04 (96.85, 97.28)	
≤40	6.35 (1.59, 12.7)	98.76 (98.24, 99.22)	13.64 (3.45, 28.12)	97 (96.85, 97.21)	
≤60	4.76 (0, 11.11)	99.28 (98.91, 99.64)	16.67 (0, 37.51)	96.97 (96.82, 97.16)	
≤80	1.59 (0, 4.76)	99.69 (99.43, 99.9)	12.5 (0, 50)	96.88 (96.83, 96.99)	
≤99	1.59 (0, 4.76)	99.95 (99.84, 100)	50 (0, 100)	96.89 (96.84, 96.99)	

**Supplemental Table 8.** Test characteristics for clinician predicted probability of all-cause, 60-day post-discharge mortality among all patients by time to post-discharge mortality

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area Under the Curve (95% CI)
<b>Day 7</b>					0.57 (0.48, 0.67)
≤5	17.65 (0, 35.29)	96.21 (95.59, 96.78)	1.97 (0, 4.29)	99.63 (99.55, 99.71)	
≤20	17.65 (0, 35.29)	96.47 (95.88, 97.01)	2.11 (0, 4.58)	99.63 (99.55, 99.71)	
≤40	17.65 (0, 35.29)	96.7 (96.13, 97.24)	2.26 (0, 4.92)	99.63 (99.55, 99.71)	
≤60	17.65 (0, 35.29)	97.04 (96.49, 97.55)	2.52 (0, 5.5)	99.63 (99.55, 99.71)	
≤80	5.88 (0, 17.65)	99.43 (99.2, 99.66)	4.17 (0, 14.29)	99.59 (99.56, 99.64)	
≤99	0 (0, 0)	99.95 (99.87, 100)	0 (0, 0)	99.56 (99.56, 99.56)	
<b>Day 14</b>					0.55 (0.49, 0.60)
≤5	13.16 (2.63, 23.68)	96.24 (95.62, 96.81)	3.23 (0.7, 6.3)	99.12 (99.01, 99.23)	
≤20	13.16 (2.63, 23.68)	96.5 (95.93, 97.05)	3.45 (0.75, 6.78)	99.12 (99.02, 99.23)	
≤40	13.16 (2.63, 23.68)	96.73 (96.16, 97.25)	3.7 (0.81, 7.26)	99.12 (99.02, 99.23)	
≤60	7.89 (0, 18.42)	97.02 (96.47, 97.54)	2.42 (0, 5.51)	99.07 (99, 99.17)	
≤80	2.63 (0, 7.89)	99.43 (99.2, 99.66)	4 (0, 14.29)	99.04 (99.02, 99.1)	
≤99	0 (0, 0)	99.95 (99.87, 100)	0 (0, 0)	99.02 (99.02, 99.02)	
<b>Day 30</b>					0.53 (0.49, 0.57)
≤5	9.52 (3.17, 17.46)	96.24 (95.64, 96.82)	3.95 (1.29, 7.24)	98.48 (98.37, 98.61)	
≤20	9.52 (3.17, 17.46)	96.5 (95.9, 97.08)	4.23 (1.39, 7.76)	98.48 (98.37, 98.61)	
≤40	9.52 (3.17, 17.46)	96.74 (96.16, 97.29)	4.51 (1.48, 8.33)	98.49 (98.38, 98.62)	
≤60	6.35 (1.59, 12.7)	97.03 (96.45, 97.55)	3.28 (0.79, 6.96)	98.44 (98.36, 98.55)	
≤80	3.17 (0, 7.94)	99.45 (99.22, 99.69)	8.33 (0, 21.74)	98.42 (98.37, 98.5)	
≤99	1.59 (0, 4.76)	99.97 (99.92, 100)	50 (0, 100)	98.41 (98.38, 98.46)	
<b>Day 45</b>					0.52 (0.50, 0.55)
≤5	8.6 (3.23, 13.98)	96.27 (95.66, 96.87)	5.26 (2.17, 8.86)	97.73 (97.61, 97.87)	
≤20	8.6 (3.23, 13.98)	96.53 (95.9, 97.11)	5.63 (2.36, 9.45)	97.74 (97.61, 97.87)	
≤40	8.6 (3.23, 13.98)	96.77 (96.13, 97.32)	6.02 (2.5, 10.17)	97.74 (97.62, 97.88)	
≤60	6.45 (2.15, 11.83)	97.05 (96.48, 97.56)	4.96 (1.64, 9.26)	97.69 (97.59, 97.83)	
≤80	2.15 (0, 5.38)	99.45 (99.21, 99.66)	8.33 (0, 22.22)	97.65 (97.6, 97.73)	
≤99	1.08 (0, 3.23)	99.97 (99.92, 100)	50 (0, 100)	97.64 (97.61, 97.69)	



**Appendix Survey.** Survey that clinicians filled out near the time of discharge for each enrolled neonate or young child in Dar es Salaam, Tanzania and Monrovia, Liberia

### Provider Discharge Survey

Date of Data Collection \_\_\_\_\_  
DD/MM/YYYY

Patient's FIRST name \_\_\_\_\_

Patient's LAST name \_\_\_\_\_

Patient's SEX:

- Male
- Female

Participant's Hospital Identifier (Unique number assigned to each patient) \_\_\_\_\_

Discharging provider year of training:

- Intern
- First year resident
- Second year resident
- Third year resident
- Specialist
- Consultant
- Medical officer
- Other. Please describe. \_\_\_\_\_

In your estimation, is the patient who was discharged at risk of any of the following?

- Re-admission to the hospital within 60 days
- Death in the next 60 days
- None of the above

If yes to any of the above, how likely is that outcome?

- 0%
- 1-5%
- 6-20%
- 21-40%
- 41-60%
- 61-80%
- 81-99%
- 100%

If yes, why do you think the patient is at risk of any of the above outcomes?

- Progression of illness
- Social concerns
- Inability to pay for future medical
- Other. Please describe. \_\_\_\_\_

# BMJ Paediatrics Open

## Predictive Value of Clinician Impression for Readmission and Post-Discharge Mortality among Neonates and Young Children in Dar es Salaam, Tanzania and Monrovia, Liberia

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## Predictive Value of Clinician Impression for Readmission and Post-Discharge Mortality among Neonates and Young Children in Dar es Salaam, Tanzania and Monrovia, Liberia

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**Key Words:** neonates; infants; young children; readmission; post-discharge mortality; Tanzania; Liberia

**Word Count:** 2,967

**Abbreviations:** AUC: area under the receiver operating characteristic curve; 95% CI: 95% confidence interval

### **Data Sharing Agreement**

Data may be made available upon reasonable request to the corresponding author.

### **Ethics Approval**

The study received ethical clearance from the Tanzania National Institute of Medical Research (#NIMR/HQ/R8a/Vol.IX/3494), the Muhimbili University of Health and Allied Sciences Research and Ethics Committee (#307/323/01), the John F. Kennedy Medical Center Institutional Review Board (#08062019), the Boston Children's Hospital Institutional Review Board (#P00033242), and the use of de-identified data was exempted from review by the Emory University Institutional Review Board (no number provided for exempted studies).

### **Transparency Declaration**

This manuscript is an honest and accurate account of the study being reported. No aspects of this study have been omitted or withheld.

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## Abstract

*Background:* There are no validated clinical decision aids to identify neonates and young children at risk of hospital readmission or post-discharge mortality in sub-Saharan Africa, leaving the decision to discharge a child to a clinician's impression. Our objective was to determine the precision of clinician impression to identify neonates and young children at risk for readmission and post-discharge mortality.

*Methods:* We conducted a survey study nested in a prospective observational cohort of neonates and children aged 1-59 months followed 60 days after hospital discharge from Muhimbili National Hospital in Dar es Salaam, Tanzania or John F. Kennedy Medical Center in Monrovia, Liberia. Clinicians who discharged each enrolled patient were surveyed to determine their perceived probability of the patient's risk of 60-day hospital readmission or post-discharge mortality. We calculated the area under the precision-recall curve (AUPRC) to determine the precision of clinician impression for both outcomes.

*Results:* Of 4,247 discharged patients, 3,896 (91.7%) had available clinician surveys and 3,847 (98.7%) had 60-day outcomes available: 187 (4.8%) were readmitted and 120 (3.1%) died within 60 days of hospital discharge. Clinician impression had poor precision in identifying neonates and young children at risk of hospital readmission (AUPRC 0.06, 95% CI 0.04 to 0.08) and post-discharge mortality (AUC 0.05, 95% CI 0.03 to 0.08). Patients for whom clinicians attributed inability to pay for future medical treatment as the reason for risk for unplanned hospital readmission had 4.76 times the odds hospital readmission (95% CI 1.31 to 17.25,  $P=0.02$ ).

*Conclusions:* Given the poor precision of clinician impression alone to identify neonates and young children at risk of hospital readmission and post-discharge mortality, validated clinical decision aids are needed to aid in the identification of young children at risk for these outcomes.

### What is already known on this topic?

- In parts of sub-Saharan Africa, hospital readmissions and post-discharge mortality rates are estimated to range from 1 to 18% within months of hospital discharge
- There are no validated clinical decision aids to accurately identify neonates, infants, and young children at risk of hospital readmission or post-discharge mortality in sub-Saharan Africa

### What this study adds?

- Clinician impression alone had poor precision in identifying neonates, infants, and children at risk of hospital readmission within 60 days of hospital discharge
- Clinician impression also did not accurately identify neonates, infants, and children at risk of 60-day post-discharge mortality

### How this study might affect practice?

- Clinician impression alone is not sufficient to accurately identify neonates, infants, and children at risk of hospital readmission or post-discharge mortality
- Validated and objective clinical decision aids are urgently needed to better identify neonates, infants, and children at risk of hospital readmission and post-discharge mortality

## Introduction

Mortality rates among children aged <5 years in sub-Saharan Africa are 74 per 1,000 live births, which is the highest in the world and is 14 times higher than rates in Europe and North America.[1] The time after an inpatient hospital admission is particularly vulnerable in the life of a child in sub-Saharan Africa. Recent studies suggest that readmissions occur in 8-18% of young children and as much as 1-20% of young children die within six months after hospital discharge.[2-5] Childhood mortality rates in the period immediately after hospitalization for an illness (i.e., the post-discharge period) approximate and may even outpace rates of mortality during hospitalization.[5,6]

Although clinical prediction rules for all-cause hospital readmissions among children in settings like the United States have been developed that include variables such as prior healthcare utilization and markers of illness severity,[7] to our knowledge, there are currently no clinical prediction rules for hospital readmissions among children in sub-Saharan Africa, where readmissions are common. Differences in healthcare access and disease prevalence in the United States and sub-Saharan Africa necessitate the creation of clinical prediction rules catered to settings in sub-Saharan Africa. Clinical prediction rules have been developed to identify both young children and those aged <15 years at risk of post-discharge mortality in some settings in sub-Saharan Africa.[5,8,9] However, these clinical prediction rules lack external validation and thus are not widely used in clinical practice. Given the absence of validated risk assessment tools to identify young children at risk of readmission and post-discharge mortality in sub-Saharan Africa, the decision to safely discharge a young child from a hospital is often driven by clinical judgement.

Clinician impression relies on a clinician's ability to recognize patterns that may be associated with severe disease or an adverse outcome.[10] However, the accuracy of clinician impression to predict outcomes, such as severe disease from infections, among children has varied in previous studies conducted in high-income settings.[11-13] In a survey of 39 providers in Kenya, clinicians under-estimated the overall incidence of post-discharge mortality among children.[14] However, that study did not assess clinician impression of post-discharge mortality for individual patients and, to our knowledge, that has not been studied previously.

Given the absence of validated prognostic tools for hospital readmission and post-discharge mortality among children in sub-Saharan Africa, our primary objective was to determine the precision of treating clinicians' clinical impression to identify neonates and young children at risk for hospital readmission and post-discharge mortality in Dar es Salaam, Tanzania and Monrovia, Liberia. Our secondary objective was to evaluate factors associated with accuracy of treating clinicians' clinical impression to identify neonates and young children at risk for hospital readmission and post-discharge mortality.

## Methods

### *Study Design*



1  
2  
3 We conducted a survey nested in a prospective observational cohort study of pediatric patients  
4 discharged from Muhimbili National Hospital in Dar es Salaam, Tanzania and John F. Kennedy  
5 Medical Center in Monrovia, Liberia from October 2019 to January 2022. Details of our study  
6 protocol have been published previously.[15] Besides differences in estimated and actual  
7 enrollment, there were no deviations from that protocol. Neonates and young children aged 1-59  
8 months were enrolled at discharge from the neonatal or pediatric wards at each facility. Follow-  
9 up consisted of caregivers receiving phone calls up to 60 days after hospital discharge.  
10 Caregivers provided written consent for participation in Tanzania and oral consent in Liberia  
11 because of cultural preference and low rates of caregiver literacy.  
12  
13

#### 14 *Patient and public involvement statement*

15  
16  
17 The development of the research question was informed by the disease burden of readmission  
18 and post-discharge mortality among children in sub-Saharan Africa. Patients were not involved  
19 in the design, recruitment, or conduct of the study, nor were they advisers in this study. Results  
20 of this study will be made publicly available through publication.  
21  
22

#### 23 *Study Setting*

24  
25 This study was conducted at two large, national referral hospitals supported by each country's  
26 Ministry of Health. They are in urban areas in their respective countries. Muhimbili National  
27 Hospital has a catchment of approximately 6 million people and John F. Kennedy Medical  
28 Center has a catchment of approximately 1.5 million people. Both hospitals are training hospitals  
29 for pediatric residents who are completing their specialty training.  
30  
31

#### 32 *Study Populations*

33  
34 Neonates and young children discharged from the wards were consecutively enrolled. Neonates  
35 and young children who died during initial hospitalization were excluded. Neonates and young  
36 children whose caregivers did not have telephones for follow up or those who declined  
37 enrollment were excluded.  
38  
39

40 Surveyed clinicians included consultants/specialists, interns/residents, or medical officers.  
41 Consultants/specialists were certified pediatricians or pediatric specialists who completed  
42 medical school, residency, and subspecialty training (for specialists). Interns/residents had  
43 completed medical school and were completing residency training in pediatrics. Medical officers  
44 received three years of clinical training prior to providing clinical care to patients.  
45  
46

#### 47 *Study Procedures*

48  
49 After obtaining informed consent from caregivers, trained research coordinators at each site  
50 approached the clinician who discharged each enrolled patient, obtained consent, and asked them  
51 to complete a survey near the time of the patient's hospital discharge. This survey modeled  
52 previous surveys that assessed clinician impression[12] and was developed through an iterative  
53 process by the research team with multiple opportunities to each investigator to refine the  
54 content. The survey was also reviewed by an expert in survey design (i.e., a survey  
55  
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57  
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60

methodologist) to ensure question clarity and appropriate response types (Appendix Survey). This survey was designed to allow clinician respondents to describe their perceived probability that each patient would experience both outcomes. Responses were recorded on standardized, electronic case report forms in electronic tablets using SQL (Tanzania) and KoboToolbox (Liberia).

### *Measurement and Outcomes*

The exposure variable was the impression of the discharging clinician of the patient's risk of: 1) unplanned hospital readmission within 60 days of hospital discharge or 2) all-cause, 60-day post-discharge mortality. Aligned with previous studies,[12] probabilities of perceived risk of readmission or post-discharge mortality included categorical options of 0%, 1-5%, 6-20%, 21-40%, 41-60%, 61-80%, 81-99%, and 100%. This survey also assessed discharging clinicians' perceptions of why readmission or post-discharge mortality were possible for those who were identified as at-risk for each outcome. Surveyed clinicians were familiar with the patients' clinical history and laboratory results during hospital admission. To assess for outcomes, phone calls to patients' caregivers were made by research staff at 7, 14, 30, 45, and 60 days after hospital discharge. Outcomes were determined as reported by caregivers to research staff.

### *Statistical Analyses*

The association of the discharging clinicians' predicted probability of readmission or post-discharge mortality and proportion of patients at each clinician-estimated risk threshold (e.g., 0%, 1-5%, etc.) who were readmitted or died was compared using Chi-square or Fisher's exact testing ( $P < 0.05$  for significance).[12] We calculated sensitivity, specificity, positive predictive value, and negative predictive value of treating clinician's impression at each percent risk threshold using caregiver-reported readmission or post-discharge mortality as the reference standards. We determined the precision of clinician impression for identifying patients at risk of readmission or post-discharge mortality by calculating the area under the precision-recall curve (AUPRC) which is useful for evaluating binary classifiers in imbalanced datasets.[16] 95% confidence intervals (95% CI) for the AUPRC were calculated through 5-fold cross-validation. We compared AUPRC and corresponding 95% CIs to baseline chance of the outcome occurring in that group. A 95% CI higher than baseline chance indicated better precision than random chance.

We conducted sub-analyses by the discharged patient's age group (i.e., neonate or young child), clinician experience level (i.e., consultant/specialist, intern/resident, or medical officer), site, and time to outcome. We conducted binary logistic regression analyses to assess whether the perceived reason for risk for each outcome was associated with the patient's likelihood of each outcome. Additionally, we conducted binary logistic regression analyses to assess whether the clinician probability for each outcome was associated with the patient's likelihood of each outcome after adjusting for clinician type, patient age at discharge (months), whether the discharge diagnosis was infectious or not, and duration of hospitalization (days). Due to small sample sizes in the non-0% clinician probability categories, we reduced the categorization of perceived risk to 0%, 1-20%, 21-60%, and 61-99%. The clinician cited reason for the outcome was also considered in the model but was removed due to collinearity as assessed by the variance

inflation factor. All tests were two-sided tests and used a 0.05 significance level. AUPRC analyses were conducted through the precrec package in R.[17] All analyses were performed in R Version 4.1.3 (R Foundation for Statistical Computing, Vienna, Austria) and SAS 9.4 (SAS Institute Inc., Cary, NC).

## Results

There were 4,460 discharged patients, 4,247 (95.2%) enrolled, and 3,896 (91.7%) had complete clinician surveys (Figure 1). Enrollment was approximately equal between the two sites (Tanzania n=1,997, 51.3%, Liberia n=1,899, 48.7%) (Table 1). There were 2,173 (55.8%) neonates and 1,723 (44.2%) young children who had clinician surveys available.

Sixty-day outcomes were available for 3,847 (98.7%) enrolled patients. The median age of enrolled neonates was 2 days (interquartile range [IQR] 1-7) and 12 months (IQR 5-24) for infants and children. The most common discharge diagnoses among neonates were sepsis (29.7%, n=609), prematurity (28.8%, n=591), and birth asphyxia (15.8%, n=323). Among infants and children, pneumonia (12.1%, n=223), diarrheal disease (10.1%, n=186), and malaria (7.2%, n=133) were most common.

There were 187 (4.8%) patients readmitted and 120 (3.1%) died within 60 days of discharge. There were 80 (3.6%) neonates who were readmitted and 61 (2.8%) died within 60 days of hospital discharge. Among infants and children, 107 (6.2%) were readmitted and 59 (3.4%) died after hospital discharge. The median time from hospital discharge to readmission was 30 days (IQR 7-45). The median time from hospital discharge to mortality was 30 days (IQR 14-45).

### *Clinician Impression and Hospital Readmission*

Nearly three quarters of patients were perceived to have 0% risk of readmission within 60 days (Table 2). Patients who were readmitted were more likely to have a perceived risk of readmission of 0% than those who were not readmitted (81% vs. 72%,  $P<0.001$ ; Table 2). Among the 187 neonates and young children who were readmitted, 80.7% (n=151) were perceived to have 0% risk of readmission.

Overall, clinician impression had poor precision in identifying neonates and young children at risk of readmission (AUPRC 0.06, 95% CI 0.04 to 0.08) (Table 3). Among medical officers, clinician impression had greater precision in identifying children at risk of readmission (AUPRC 0.23, 95% CI 0.17 to 0.34); this group was marginally better at identifying patients at risk of readmission than interns/residents and consultants/specialists.

By clinician type, medical officer clinician impression had poor precision in identifying neonates (Supplemental Table 1) but greater precision when identifying infants and children at risk for readmission (Supplemental Table 2). In site-specific analyses, clinician impression was imprecise when identifying neonates and young children at risk of readmission in Tanzania (AUPRC 0.11, 95% CI 0.07 to 0.15, chance: 0.07) and Liberia (AUPRC 0.03, 95% CI 0.02 to 0.04, chance: 0.02). Regardless of the time from hospital discharge to readmission, clinician

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2  
3 impression had poor precision in identifying neonates and young children at risk for readmission  
4 (Supplemental Table 3).  
5

### 6 *Clinician Impression and Post-Discharge Mortality*

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8  
9 Most (96.1%, n=3,746) patients were assigned 0% risk of post-discharge mortality (Table 4).  
10 Patients who died within 60 days of discharge were more likely to have a perceived risk of 0%  
11 than patients who survived (96% vs. 90.8%,  $P=0.002$ ; Table 4). Among the 120 neonates and  
12 young children who died within 60 days of hospital discharge, 90.8% (n=109) were estimated to  
13 have a 0% probability of post-discharge mortality.  
14

15  
16 Overall, clinician impression had poor precision in identifying neonates and young children at  
17 risk post-discharge mortality (AUPRC 0.05, 95% CI 0.03 to 0.08) and did not vary substantially  
18 among interns/residents, specialist/consultants, or medical officers (Table 5). Clinician  
19 impression had poor precision in identifying post-discharge mortality among neonates (AUPRC  
20 0.04, 95% CI 0.03 to 0.06, chance: 0.03) and infants and children (AUPRC 0.06, 95 % CI 0.03 to  
21 0.10, chance: 0.03). When analyzed by site, clinician impression in both Tanzania (AUPRC 0.08,  
22 95% CI 0.03 to 0.13, chance: 0.03) and Liberia (AUPRC 0.04, 95% CI 0.03 to 0.06, chance:  
23 0.04) had poor precision in identifying neonates and young children at risk of post-discharge  
24 mortality. Clinician impression had poor precision regardless of the time to post-discharge  
25 mortality (Supplemental Table 4).  
26  
27

### 28 *Reason for Perceived Risk of Hospital Readmission and Post-Discharge Mortality*

29  
30  
31 Patients for whom clinicians attributed inability to pay for treatment as the reason for  
32 readmission had 4.76 times the odds readmission (95% CI 1.31 to 17.25,  $P=0.02$ ) compared to  
33 those with no perceived risk (Table 6). Patients whose clinician cited “other” reasons to be at risk  
34 had lower odds of readmission compared to those whose clinician did not believe they were at  
35 risk for readmission (OR 0.24, 95% CI 0.09 to 0.66,  $P=0.005$ ). Patients for whom clinicians  
36 attributed inability to pay for treatment as the reason for potential post-discharge mortality had  
37 5.53 times the odds of post-discharge mortality (95% CI 1.22 to 25.10,  $P=0.03$ ).  
38  
39

40 In multivariable analyses, patients whom clinicians estimated to be at moderate risk for hospital  
41 readmission (i.e., 21-60%) were at decreased odds of hospital readmission (aOR 0.45, 95% CI  
42 0.26 to 0.74,  $P=0.003$ ) (Table 7). No other factors were independently associated with either  
43 hospital readmission or post-discharge mortality.  
44

## 45 **Discussion**

46  
47  
48 Among nearly 3,900 neonates and young children discharged from referral hospitals in Dar es  
49 Salaam, Tanzania and Monrovia, Liberia, clinician impression had poor precision for identifying  
50 those at risk of unplanned hospital readmissions and post-discharge mortality. Medical officer  
51 clinician impression at both sites had slightly greater precision in identifying young children at  
52 risk of readmission. Clinician perception of inability to pay for treatment was associated with  
53 readmission and post-discharge mortality.  
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2  
3 The poor precision of clinician impression in identifying neonates and young children at risk of  
4 readmission and post-discharge mortality differs from findings in studies in high-income settings  
5 that assessed the diagnosis of acute coronary syndrome and sinusitis in adults,[18,19] the  
6 presence of pneumonia[20] or the development of severe pneumonia among children,[12] or the  
7 presence of appendicitis among children.[21] This difference is likely multifactorial in nature  
8 and may consist of differences in available diagnostic and prognostic resources. Prior studies  
9 suggest that laboratory capabilities in resource-limited settings are inadequate,[22-24] leading to  
10 dependence on clinical exam findings to make diagnoses and determine prognosis,[25] which  
11 may hinder the accuracy of clinician impression to identify neonates and young children at risk  
12 of untoward post-discharge outcomes. Moreover, clinicians may not consider key factors in the  
13 home (e.g., access to healthcare facilities and maternal health) that may contribute to post-  
14 discharge outcomes. Additionally, clinician impression had poor precision in identifying  
15 neonates and young children at risk for readmission or post-discharge mortality regardless of the  
16 time from discharge to either event. Prior studies of clinician impression assessed outcomes  
17 within hours or days[12,18,19] and not up to 60 days, which may contribute to the difference in  
18 our results compared to prior studies assessing clinician impression in prognostication of  
19 outcomes.  
20  
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22  
23

24 Clinician impression among medical officers had fair precision in identifying young children at  
25 risk of readmission. This may be due to the combination of more clinical experience than  
26 interns/residents and more time spent with patients than consultants/specialists who often spend  
27 less time directly with patients and more time supervising clinical care. Prior studies conducted  
28 in high-income settings demonstrate that clinician impression of less experienced clinicians may  
29 have less discriminatory value than that of more experienced clinicians.[12]  
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32 Several clinical prediction rules for post-discharge mortality have been developed among young  
33 children aged 2-23 months in several countries in sub-Saharan Africa and South Asia, 6 months-  
34 5 years in Uganda, and all children aged <15 years in Mozambique.[5,8,9] These clinical  
35 prediction rules include variables such as clinical diagnoses and anthropometry to assign  
36 weighted points to included variables to assess an individual patient's risk for post-discharge  
37 mortality up to six months after discharge. However, none of these have focused specifically on  
38 neonates and none have been externally validated, which is a necessary step to assess  
39 discriminatory value prior to clinical use. Thus, the current state of prognostic determination for  
40 young children after discharge in sub-Saharan Africa depends on clinician impression and, given  
41 its poor precision demonstrated in our study, validated clinical prediction rules to identify  
42 neonates and young children at risk of post-discharge mortality are urgently needed. Such  
43 clinical decision aids should include commonly collected variables and may include biomarkers  
44 to add precision to risk stratification to identify neonates and young children at risk of post-  
45 discharge morbidity and mortality.[26-28]  
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49 Our examination of reasons for estimated outcomes suggested that clinician perception of  
50 inability to pay for future treatment was associated with higher risk of readmission and post-  
51 discharge mortality. Young children from lower socioeconomic status have poorer overall health  
52 outcomes compared to young children from higher socioeconomic households in sub-Saharan  
53 Africa.[29-31] This is particularly relevant in the post-discharge period during which the  
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3 financial burden of care seeking may influence the ability for a family to seek additional clinical  
4 care after a potentially costly hospital admission.  
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### 6 *Limitations*

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9 Clinician impression is multifactorial and depends on clinical training as well as available  
10 laboratory, radiological, and clinical data that may not have been available to all discharging  
11 clinicians. We did not assess the availability of these in our analysis. We did not include nurses  
12 or clinical officers in our study, which is a limitation as these groups may have good insight into  
13 potential adverse outcomes after discharge. We could not account for the potential role that  
14 variations in the quality of clinical care provided to patients may have had in readmissions or  
15 post-discharge mortality.  
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### 17 **Conclusions**

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20 Clinician impression had poor precision in identifying neonates and young children at risk of  
21 unplanned hospital readmission and post-discharge mortality at two referral hospitals in Dar es  
22 Salaam, Tanzania and Monrovia, Liberia. Validated and objective clinical decision aids to assist  
23 clinicians in the identification of young children at risk of readmission and post-discharge  
24 mortality may facilitate the identification of those at greatest risk.  
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33 conceptualized and designed the study. CAR, RCI, JK, AS, EG, CRS, KPM, and CD oversaw  
34 data collection and verified the underlying data. CAR and AM verified the underlying data. AM  
35 conducted the statistical analyses. CAR wrote the first draft of the manuscript. CAR, RK, RCI,  
36 JK, Y-JC, AS, EG, HKM, CRS, AM, MN, CRM, CGW, RFB, KPM, and CD interpreted the  
37 data, reviewed, and provided input to the final draft. CAR had final responsibility for the  
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**Figure 1.** Flow diagram for included neonates and children aged 1-59 months discharged from the neonatal wards and pediatric wards in Dar es Salaam, Tanzania and Monrovia, Liberia

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**Table 1.** Characteristics of neonates and young children included in the evaluation of clinician impression on predicting 60-day hospital readmission or post-discharge mortality

Patient Characteristics	Overall Population, (N=3,896) n (%)	Readmitted to Hospital 60 Days After Hospital Discharge, (N=187) n (%)	Died 60 Days After Hospital Discharge, (N=120) n (%)
<b>Discharged from Neonatal Ward<sup>1</sup></b>	2,173 (55.8)	80 (3.6)	61 (2.8)
Age in days at discharge, median (IQR)	2 (1, 7)	4 (2, 13)	2 (0, 10)
<b>Discharged from Pediatrics Ward</b>	1,723 (44.2)	107 (6.2)	59 (3.4)
Age in months for pediatric patients aged 1-59 months, median (IQR) <sup>2</sup>	12 (5, 24)	8 (4, 18)	8 (4, 21)
<b>Sex<sup>3</sup></b>			
Male	2,198 (56.4)	109 (5.0)	74 (3.4)
Female	1,691 (43.4)	78 (4.6)	46 (2.7)
<b>Country</b>			
Tanzania	1,997 (51.3)	140 (7.0)	63 (3.2)
Liberia	1,899 (48.7)	47 (2.5)	57 (3.0)
<b>Disposition from the Hospital</b>			
Discharge	3,775 (96.8)	186 (4.9)	106 (2.8)
Left against medical advice	119 (3.1)	1 (0.8)	14 (11.8)
Transfer to another facility	2 (0.05)	0 (0.0)	0 (0.0)

<sup>1</sup>24 neonates had missing age.

<sup>2</sup>31 young children had missing age.

<sup>3</sup>7 participants did not have a documented sex.



**Table 2.** Association of discharging clinicians' predicted probability and unplanned 60-day hospital readmission among neonates and young children

	Total, n (%) (N=3,896)	Not Readmitted 60 Days After Hospital Discharge, n (%) (N=3,709)	Readmitted 60 Days After Hospital Discharge, n (%) (N=187)	P Value <sup>1</sup>
<b>Clinician predicted probability</b>				<0.001
0%	2,838 (72.8)	2,687 (72.4)	151 (80.7)	
1-5%	9 (0.2)	9 (0.2)	0 (0.0)	
6-20%	26 (0.7)	23 (0.6)	3 (1.6)	
21-40%	736 (18.9)	720 (19.4)	16 (8.6)	
41-60%	38 (1.0)	35 (0.9)	3 (1.6)	
61-80%	237 (6.1)	227 (6.1)	10 (5.3)	
81-99%	11 (0.3)	7 (0.2)	4 (2.1)	

<sup>1</sup>By Fisher's exact test to assess independence of clinician predicted probability and likelihood of hospital readmission.

Note: There was 1 respondent who estimated that the discharged child was at risk of hospital readmission but did not assign a proportion.

**Table 3.** Test characteristics for clinician predicted probability of unplanned,60-day hospital readmission among all enrolled neonates and young children aged 1-59 months overall and by clinician type

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area under Precision-Recall Curve (95% CI)
All Clinicians (N=3,895) <sup>1</sup>					0.06 (0.04, 0.08), chance=0.05
≤5	19.3 (13.9, 25.1)	72.5 (71.0, 73.8)	3.4 (2.5, 4.4)	94.7 (94.3, 95.1)	
≤20	19.3 (13.9, 25.1)	72.7 (71.3, 74.1)	3.4 (2.5, 4.5)	94.7 (94.4, 95.1)	
≤40	17.7 (12.3, 23.5)	73.7 (72.3, 75.1)	3.3 (2.4, 4.3)	94.7 (94.3, 95.0)	
≤60	16.0 (11.2, 21.9)	74.3 (72.9, 75.7)	3.0 (2.1, 4.1)	94.6 (94.3, 94.9)	
≤80	7.5 (4.2, 11.8)	93.7 (92.9, 94.4)	5.6 (3.1, 8.7)	95.3 (95.1, 95.5)	
≤99	2.1 (0.5, 4.3)	99.8 (99.7, 99.9)	36.4 (9.1, 66.7)	95.3 (95.2, 95.4)	
Consultant/Specialist (N=175)					0.05 (0.02, 0.08), chance=0.03
≤5	16.7 (0, 50.0)	69.2 (62.1, 75.8)	1.9 (0.0, 5.7)	95.8 (94.7, 97.6)	
≤20	16.7 (0, 50.0)	69.8 (62.7, 76.3)	1.9 (0, 5.9)	95.9 (94.7, 97.6)	
≤40	16.7 (0, 50.0)	73.4 (66.3, 79.9)	2.1 (0, 6.7)	96.1 (95.0, 97.7)	
≤60	0.0 (0.0, 0.0)	75.2 (68.6, 81.1)	0.0 (0.0, 0.0)	95.5 (95.1, 95.8)	
≤80	0.0 (0.0, 0.0)	85.2 (79.9, 90.5)	0.0 (0.0, 0.0)	96.0 (95.7, 96.2)	
≤99	0.0 (0.0, 0.0)	100 (100, 100)	NA	96.6 (96.6, 96.6)	
Intern/Resident (N=3,502)					0.06 (0.03, 0.08), chance=0.05
≤5	14.9 (9.9, 20.5)	72.4 (70.8, 73.9)	2.5 (1.7, 3.5)	94.6 (94.3, 95.0)	
≤20	14.9 (9.9, 20.5)	72.5 (70.9, 74.0)	2.5 (1.7, 3.5)	94.6 (94.3, 95.0)	
≤40	14.9 (9.9, 20.5)	73.2 (71.6, 74.7)	2.6 (1.7, 3.6)	94.7 (94.4, 95.1)	
≤60	13.6 (8.7, 19.3)	73.5 (71.9, 75.0)	2.4 (1.6, 3.4)	94.6 (94.3, 94.9)	
≤80	5.6 (2.5, 9.3)	94.1 (93.3, 94.9)	4.3 (1.9, 7.3)	95.4 (95.2, 95.6)	
≤99	1.2 (0.0, 3.1)	99.9 (99.8, 100)	40.0 (0.0, 100)	95.5 (95.4, 95.5)	
Medical Officer (N=217)					0.23 (0.17, 0.34), chance=0.09
≤5	55.0 (30.0, 75.0)	76.1 (70.1, 81.7)	18.9 (11.6, 26.5)	94.3 (91.6, 96.9)	
≤20	55.0 (30.0, 75.0)	78.7 (72.6, 84.3)	20.7 (12.7, 29.4)	94.5 (91.9, 96.9)	
≤40	40.0 (20.0, 60.0)	82.2 (76.6, 87.3)	18.4 (9.4, 28.9)	93.1 (90.8, 95.5)	
≤60	40.0 (20.0, 60.0)	86.8 (81.7, 91.4)	23.5 (12.1, 36.4)	93.4 (91.3, 95.7)	
≤80	25.0 (9.8, 45.0)	93.9 (90.4, 96.9)	29.4 (10.5, 50.0)	92.5 (90.9, 94.4)	
≤99	10.0 (0.0, 25.0)	97.9 (95.9, 99.5)	33.3 (0.0, 80.0)	91.5 (90.6, 92.8)	

<sup>1</sup>There was 1 respondent who estimated that the discharged child was at risk of hospital readmission but did not assign a proportion.

**Table 4.** Association of discharging clinicians' predicted probability and all-cause, 60-day post-discharge mortality among neonates and young children

	Total, n (%) (N=3,896)	Did not Die 60 Days After Hospital Discharge, n (%) (N=3,776)	Died 60 Days After Hospital Discharge, n (%) (N=120)	<i>P</i> Value <sup>1</sup>
<b>Clinician predicted probability</b>				0.002
0%	3,746 (96.1)	3,637 (96.3)	109 (90.8)	
1-5%	10 (0.3)	10 (0.3)	0 (0.0)	
6-20%	9 (0.2)	7 (0.2)	2 (1.7)	
21-40%	13 (0.3)	11 (0.3)	2 (1.7)	
41-60%	95 (2.4)	90 (2.4)	5 (4.2)	
61-80%	21 (0.5)	20 (0.5)	1 (0.8)	
81-99%	2 (0.1)	1 (0.03)	1 (0.8)	

<sup>1</sup>By Fisher's exact test to assess independence of clinician predicted probability and likelihood of hospital readmission.

**Table 5.** Test characteristics for clinician predicted probability of all-cause, 60-day post-discharge mortality among all enrolled neonates and young children aged 1-59 months by clinician type

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area under Precision-Recall Curve (95% CI)
All Clinicians (N=3,895) <sup>1</sup>					0.05 (0.03, 0.08), chance=0.03
≤5	9.2 (4.2, 15.0)	96.3 (95.7, 96.9)	7.3 (3.5, 11.9)	97.1 (96.9, 97.3)	
≤20	9.2 (4.2, 15.0)	96.6 (95.9, 97.2)	7.7 (3.8, 12.7)	97.1 (96.9, 97.3)	
≤40	7.5 (3.3, 12.5)	96.7 (96.2, 97.3)	6.8 (2.9, 11.6)	97.1 (96.9, 97.2)	
≤60	5.8 (1.7, 10.8)	97.1 (96.5, 97.6)	5.8 (1.9, 10.5)	97.0 (96.9, 97.2)	
≤80	1.7 (0.0, 4.2)	99.4 (99.2, 99.7)	8.3 (0.0, 22.7)	96.9 (96.9, 97.0)	
≤99	0.8 (0.0, 2.5)	99.9 (99.9, 100)	50.0 (0.0, 100)	96.9 (96.9, 96.9)	
Specialist or Consultant (N=175)					0.05 (0.02, 0.07), chance=0.04
≤5	0.0 (0.0, 0.0)	97.6 (95.2, 99.4)	0.0 (0.0, 0.0)	95.9 (95.8, 95.9)	
≤20	0.0 (0.0, 0.0)	97.6 (95.2, 99.4)	0.0 (0.0, 0.0)	95.9 (95.8, 95.9)	
≤40	0.0 (0.0, 0.0)	97.6 (95.2, 99.4)	0.0 (0.0, 0.0)	95.9 (95.8, 95.9)	
≤60	0.0 (0.0, 0.0)	97.6 (95.2, 99.4)	0.0 (0.0, 0.0)	95.9 (95.8, 95.9)	
≤80	0.0 (0.0, 0.0)	98.8 (97.0, 100)	0.0 (0.0, 0.0)	95.9 (95.9, 96.0)	
≤99	0.0 (0.0, 0.0)	100 (100, 100)	-	96.0 (96.0, 96.0)	
Intern/Resident (N=3,502)					0.05 (0.02, 0.08), chance=0.03
≤5	8.8 (3.9, 14.7)	97.1 (96.5, 97.6)	8.3 (3.7, 13.8)	97.3 (97.1, 97.4)	
≤20	8.8 (3.9, 14.7)	97.2 (96.6, 97.7)	8.5 (3.8, 14.2)	97.3 (97.1, 97.4)	
≤40	6.9 (2.9, 12.7)	97.2 (96.6, 97.7)	6.8 (2.7, 12.0)	97.2 (97.1, 97.4)	
≤60	5.8 (1.9, 10.8)	97.2 (96.7, 97.8)	5.9 (1.9, 11.0)	97.2 (97.1, 97.3)	
≤80	1.9 (0.0, 4.9)	99.6 (99.4, 99.8)	11.8 (0.0, 30.8)	97.1 (97.1, 97.2)	
≤99	0.9 (0.0, 2.9)	99.9 (99.9, 100.0)	50.0 (0.0, 100.8)	97.1 (97.1, 97.2)	
Medical Officer (N=218)					0.05 (0.03, 0.08), chance=0.05
≤5	18.2 (0.0, 45.5)	82.6 (77.8, 87.9)	5.0 (0, 12.2)	94.9 (93.8, 96.5)	
≤20	18.2 (0.0, 45.5)	85.9 (81.2, 90.3)	6.1 (0, 14.8)	95.1 (94.1, 96.7)	
≤40	18.2 (0.0, 45.5)	88.9 (84.5, 92.8)	7.7 (0, 18.7)	95.3 (94.2, 96.8)	
≤60	9.1 (0.0, 27.3)	93.7 (90.3, 96.6)	6.7 (0, 23.1)	95.1 (94.5, 96.1)	
≤80	0.0 (0.0, 0.0)	97.6 (95.2, 99.5)	0.0 (0.0, 0.0)	94.8 (94.7, 94.9)	
≤99	0.0 (0.0, 0.0)	100 (100, 100)	-	94.9 (94.9, 94.9)	

<sup>1</sup>There was 1 respondent who estimated that the discharged child was at risk of hospital readmission but did not assign a proportion.

**Table 6.** Reasons for perceived risk of hospital readmission and post-discharge mortality

<b>Clinician Cited Reason for Outcome</b>	<b>No Hospital Readmission, n (%)</b>	<b>Hospital Readmission, n (%)</b>	<b>Odds Ratio (95% Confidence Interval)</b>	<b>P value</b>
No risk	2,584 (69.7)	148 (79.1)	<i>Referent</i>	--
Clinician perceived inability to pay for treatment	11 (0.3)	3 (1.6)	4.76 (1.31, 17.25)	0.02
Clinician perceived social concerns	70 (1.9)	3 (1.6)	0.75 (0.23, 2.41)	0.63
Clinician perceived progression of illness	754 (20.3)	29 (15.5)	0.67 (0.45, 1.01)	0.05
Other <sup>1</sup>	290 (7.8)	4 (2.1)	0.24 (0.09, 0.66)	0.005
<b>Clinician Cited Reason for Outcome</b>	<b>Did not Die Within 60 Days, n (%)</b>	<b>Died Within 60 Days, n (%)</b>	<b>Odds Ratio (95% Confidence Interval)</b>	<b>P value</b>
No risk	2,652 (70.2)	80 (66.7)	<i>Referent</i>	--
Clinician perceived inability to pay for treatment	12 (0.3)	2 (1.7)	5.53 (1.22, 25.10)	0.03
Clinician perceived social concerns	72 (1.9)	1 (0.8)	0.46 (0.06, 3.36)	0.44
Clinician perceived progression of illness	754 (20.0)	29 (24.2)	1.28 (0.83, 1.97)	0.27
Other <sup>1</sup>	286 (7.6)	8 (6.7)	0.93 (0.44, 1.94)	0.84

<sup>1</sup>Included concerns about caregiver understanding, general clinician impression, and patient with history of recurrent illness.

**Table 7.** Multivariable regression model for all-cause 60-day hospital readmission and post-discharge mortality among young children discharged in Dar es Salaam, Tanzania and Monrovia, Liberia

Characteristics	No Hospital Readmission, n (%), N=3,709	Hospital Readmission, n (%), N=187	Adjusted Odds Ratio (95% CI), N=3,426	P value
<b>Perceived risk</b>				
0%	2,687 (72%)	151 (81%)	<i>Referent</i>	--
1-20%	44 (1.2%)	3 (1.6%)	0.96 (0.23, 2.77)	0.95
21-60%	743 (20%)	19 (10%)	0.45 (0.26, 0.74)	0.003
61-99%	234 (6.3%)	14 (7.5%)	1.08 (0.57, 1.90)	0.79
<b>Discharge Provider Type</b>				
Intern/Resident	3,341 (90%)	161 (86%)	<i>Referent</i>	--
Specialist or Consultant	169 (4.6%)	6 (3.2%)	0.78 (0.30, 1.65)	0.55
Medical Officer	198 (5.3%)	20 (11%)	2.07 (1.21, 3.39)	0.01
<b>Patient Age at Discharge, months</b>	1 (0, 10)	3 (1, 11)	1.00 (0.98, 1.01)	0.45
<b>Discharge Diagnosis</b>				
Non-infectious	1,492 (40%)	87 (47%)	<i>Referent</i>	--
Infectious	2,217 (60%)	100 (53%)	0.86 (0.62, 1.19)	0.35
<b>Duration of Hospitalization, days</b>	7 (3, 13)	8 (3, 16)	1.002 (1.0001, 1.005)	0.04
Characteristics	Did not Die Within 60 Days, n (%), N=3,776	Died Within 60 Days, n (%), N=120	Adjusted Odds Ratio (95% CI), N=3,426	P value
<b>Perceived risk</b>				
0%	3,637 (96%)	109 (91%)	<i>Referent</i>	--
1-20%	17 (0.5%)	2 (1.7%)	2.95 (0.44, 11.8)	0.18
21-60%	101 (2.7%)	7 (5.8%)	1.92 (0.73, 4.20)	0.14
61-99%	21 (0.6%)	2 (1.7%)	2.94 (0.46, 10.4)	0.15
<b>Discharge Provider Type</b>				
Intern/Resident	3,400 (90%)	102 (85%)	<i>Referent</i>	--
Specialist or Consultant	168 (4.5%)	7 (5.8%)	1.53 (0.63, 3.15)	0.29
Medical Officer	207 (5.5%)	11 (9.2%)	1.76 (0.82, 3.45)	0.12
<b>Patient Age at Discharge, months</b>	1 (0, 11)	1 (0, 7)	0.99 (0.97, 1.00)	0.13
<b>Discharge Diagnosis</b>				
Non-infectious	1,531 (41%)	48 (40%)	<i>Referent</i>	--
Infectious	2,245 (59%)	72 (60%)	0.95 (0.64, 1.42)	0.80
<b>Duration of Hospitalization, days</b>	7 (3, 13)	8 (3, 14)	1.00 (0.99, 1.00)	0.88

## References

1. World Health Organization. Child mortality (under 5 years). <https://www.who.int/news-room/fact-sheets/detail/levels-and-trends-in-child-under-5-mortality-in-2020>. Published 2022. Accessed May 2, 2023.
2. Pavlinac PB, Singa BO, Tickell KD, et al. Azithromycin for the prevention of rehospitalisation and death among Kenyan children being discharged from hospital: a double-blind, placebo-controlled, randomised controlled trial. *Lancet Glob Heal* 2021;9:e1569-e1578.
3. Connon R, George EC, Olupot-Olupot P, et al. Incidence and predictors of hospital readmission in children presenting with severe anaemia in Uganda and Malawi: a secondary analysis of TRACT trial data. *BMC Public Health* 2021;21:1480.
4. Nemetchek B, English L, Kissoon N, et al. Paediatric postdischarge mortality in developing countries: a systematic review. *BMJ Open*. 2018;8:e023445.
5. Diallo AH, Sayeem Bin Shahid ASM, Khan AF, et al. Childhood mortality during and after acute illness in Africa and south Asia: a prospective cohort study. *Lancet Glob Heal* 2022;10:e673-e684.
6. Wiens MO, Pawluk S, Kissoon N, et al. Pediatric Post-Discharge Mortality in Resource Poor Countries: A Systematic Review. *PLoS One* 2013;8:e66698.
7. Goodman DM, Casale MT, Rychlik K, et al. Development and Validation of an Integrated Suite of Prediction Models for All-Cause 30-Day Readmissions of Children and Adolescents Aged 0 to 18 Years. *JAMA Netw Open*. 2022;5:e2241513.
8. Wiens MO, Kumbakumba E, Larson CP, et al. Postdischarge mortality in children with acute infectious diseases: Derivation of postdischarge mortality prediction models. *BMJ Open* 2015;5:e009449.
9. Madrid L, Casellas A, Sacoor C, et al. Postdischarge Mortality Prediction in Sub-Saharan Africa. *Pediatrics* 2019;143:e20180606.
10. Horwood J, Cabral C, Hay AD, Ingram J. Primary care clinician antibiotic prescribing decisions in consultations for children with RTIs: a qualitative interview study. *Br J Gen Pr* 2016;66:e207-e213.
11. Van den Bruel A, Thompson M, Buntinx F, Mant D. Clinicians' gut feeling about serious infections in children: observational study. *BMJ* 2012;345:e6144.
12. Gao HM, Ambroggio L, Shah SS, Ruddy RM, Florin TA. Predictive value of clinician "gestalt" in pediatric community-acquired pneumonia. *Pediatrics* 2021;147:e2020041582.
13. Turnbull S, Lucas PJ, Redmond NM, et al. What gives rise to clinician gut feeling, its influence on management decisions and its prognostic value for children with RTI in primary care: a prospective cohort study. *BMC Fam Pr* 2018;19:25.
14. Paul S, Tickell KD, Ojee E, et al. Knowledge, attitudes, and perceptions of Kenyan healthcare workers regarding pediatric discharge from hospital. *PLoS One* 2021;16:e0249569.
15. Rees CA, Kisenge R, Ideh RC, et al. A Prospective, observational cohort study to identify neonates and children at risk of postdischarge mortality in Dar es Salaam, Tanzania and Monrovia, Liberia: The PPDM study protocol. *BMJ Paediatr Open* 2022;6:e001379.
16. Saito T, Rehmsmeier M. The precision-recall plot is more informative than the ROC plot when evaluating binary classifiers on imbalanced datasets. *PLoS One*. 2015;10:e0118432.

17. Saito T, Rehmsmeier M. Precrec: fast and accurate precision-recall and ROC curve calculations in R. *Bioinformatics*. 2017;33:145-147.
18. Williams JW Jr, Simel DL, Roberts L SG. Clinical evaluation for sinusitis. Making the diagnosis by history and physical examination. *Ann Intern Med* 1992;117:705-710.
19. Oliver G, Reynard C, Morris N, Body R. Can Emergency Physician Gestalt “Rule In” or “Rule Out” Acute Coronary Syndrome: Validation in a Multicenter Prospective Diagnostic Cohort Study. *Acad Emerg Med* 2020;27:24-30.
20. Neuman MI, Scully KJ, Kim D, Shah S, Bachur R. Physician assessment of the likelihood of pneumonia in a pediatric emergency department. *Pediatr Emerg Care* 2010;26:817-822.
21. Lee WH, O’Brien S, Skarin D, et al. Accuracy of clinician gestalt in diagnosing appendicitis in children presenting to the emergency department. *Emerg Med Australas* 2019;31:612-618.
22. Yadav H, Shah D, Sayed S, Horton S, Schroeder LF. Availability of essential diagnostics in ten low-income and middle-income countries: results from national health facility surveys. *Lancet Glob Heal* 2021;9:e1553-e1560.
23. Peeling RW, Mabey D. Point-of-care tests for diagnosing infections in the developing world. *Clin Microbiol Infect* 2010;16:1062-1069.
24. Singh H, Schiff GD, Graber ML, Onakpoya I TM. The global burden of diagnostic errors in primary care. *BMJ Qual Saf* 2017;26:484-494.
25. World Health Organization. Integrated Management of Childhood Illness: management of the sick young infant aged up to 2 months: IMCI chart booklet. [https://www.who.int/maternal\\_child\\_adolescent/documents/management-sick-young-infant-0-2-months/en/](https://www.who.int/maternal_child_adolescent/documents/management-sick-young-infant-0-2-months/en/). Published 2019. Accessed September 24, 2021.
26. Njunge JM, Tickell K, Diallo AH, et al. The Childhood Acute Illness and Nutrition (CHAIN) network nested case-cohort study protocol: a multi-omics approach to understanding mortality among children in sub-Saharan Africa and South Asia. *Gates Open Res* 2022;6:77.
27. Njunge JM, Gwela A, Kibinge NK, et al. Biomarkers of post-discharge mortality among children with complicated severe acute malnutrition. *Sci Rep* 2019;9:5981.
28. Sarangam ML, Namazzi R, Datta D, et al. Intestinal Injury Biomarkers Predict Mortality in Pediatric Severe Malaria. *MBio* 2022;13:e0132522.
29. Ogbo FA, Ezeh OK, Awosemo AO, et al. Determinants of trends in neonatal, post-neonatal, infant, child and under-five mortalities in Tanzania from 2004 to 2016. *BMC Public Health* 2019;19:1243.
30. Burkey MD, Weiser SD, Fehmie D, et al. Socioeconomic determinants of mortality in HIV: evidence from a clinical cohort in Uganda. *J Acquir Immune Defic Syndr* 2014;66:41-47.
31. Ezeh OK, Agho KE, Dibley MJ, Hall JJ, Page AN. Risk factors for post neonatal, infant, child and under-5 mortality in Nigeria: a pooled cross-sectional analysis. *BMJ Open* 2015;5:e006779.



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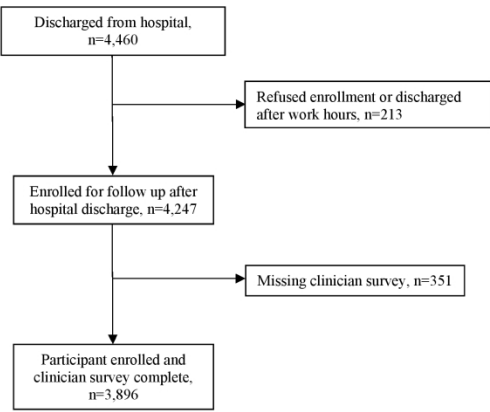


Figure 1. Flow diagram for included neonates and children aged 1-59 months discharged from the neonatal wards and pediatric wards in Dar es Salaam, Tanzania and Monrovia, Liberia

215x279mm (300 x 300 DPI)

**Supplemental Table 1.** Test characteristics for clinician predicted probability of all-cause, 60-day hospital readmission for neonates at both sites

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area under Precision-Recall Curve (95% CI)
All Clinicians (N=2,173)					0.03 (0.02, 0.04), chance=0.04
≤5	8.75 (3.75, 15)	73.77 (71.91, 75.68)	1.25 (0.51, 2.19)	95.48 (95.19, 95.81)	
≤20	8.75 (3.75, 15)	73.87 (72, 75.73)	1.25 (0.51, 2.19)	95.48 (95.19, 95.81)	
≤40	8.75 (3.75, 15)	74.34 (72.48, 76.21)	1.28 (0.52, 2.23)	95.51 (95.23, 95.84)	
≤60	7.5 (2.5, 13.75)	74.58 (72.72, 76.4)	1.1 (0.36, 2.04)	95.46 (95.2, 95.78)	
≤80	2.5 (0, 6.25)	94.84 (93.88, 95.8)	1.75 (0, 4.47)	96.21 (96.11, 96.36)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.32 (96.32, 96.32)	
Interns/Residents (N=2,076)					0.03 (0.02, 0.04), chance=0.04
≤5	7.79 (2.6, 14.29)	73.69 (71.68, 75.54)	1.12 (0.37, 2.03)	95.39 (95.11, 95.72)	
≤20	7.79 (2.6, 14.29)	73.74 (71.74, 75.59)	1.12 (0.37, 2.04)	95.4 (95.12, 95.73)	
≤40	7.79 (2.6, 14.29)	74.09 (72.09, 75.94)	1.14 (0.37, 2.07)	95.42 (95.14, 95.74)	
≤60	7.79 (2.6, 14.29)	74.29 (72.34, 76.14)	1.15 (0.38, 2.08)	95.43 (95.15, 95.76)	
≤80	2.6 (0, 6.49)	95.15 (94.15, 96.05)	1.96 (0, 4.85)	96.2 (96.09, 96.35)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.29 (96.29, 96.29)	
Specialist or Consultant (N=66)					0.09 (0.01, 0.18) , chance=0.07
≤5	50 (0, 100)	65.62 (53.12, 76.56)	4.35 (0, 10.53)	97.67 (94.87, 100)	
≤20	50 (0, 100)	67.19 (54.69, 79.69)	4.55 (0, 11.11)	97.73 (95.12, 100)	
≤40	50 (0, 100)	71.88 (60.94, 82.81)	5.26 (0, 12.5)	97.87 (95.45, 100)	
≤60	0 (0, 0)	73.44 (62.5, 84.38)	0 (0, 0)	95.92 (95.24, 96.43)	
≤80	0 (0, 0)	82.81 (73.44, 92.19)	0 (0, 0)	96.36 (95.92, 96.72)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.97 (96.97, 96.97)	
Medical Officer (N=30)					0.03 (NA, NA), chance=0.04
≤5	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤20	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤40	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤60	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤80	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.67 (96.67, 96.67)	

**Supplemental Table 2.** Test characteristics for clinician predicted probability of all-cause, 60-day hospital readmission for young children aged 1-59 months at both sites

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area under Precision-Recall Curve (95% CI)
All Clinicians (N=1,722)					0.09 (0.04, 0.14), chance=0.06
≤5	27.1 (18.69, 35.51)	70.77 (68.54, 73)	5.77 (4.09, 7.57)	93.6 (92.9, 94.33)	
≤20	27.1 (18.69, 35.51)	71.21 (68.98, 73.44)	5.85 (4.14, 7.69)	93.64 (92.95, 94.36)	
≤40	24.3 (15.89, 32.71)	72.76 (70.65, 74.86)	5.51 (3.81, 7.4)	93.53 (92.9, 94.23)	
≤60	22.43 (14.95, 30.84)	73.87 (71.7, 75.98)	5.34 (3.59, 7.35)	93.48 (92.88, 94.19)	
≤80	11.21 (5.61, 17.76)	92.26 (90.84, 93.44)	8.63 (4.32, 13.71)	94 (93.63, 94.45)	
≤99	3.74 (0.93, 7.48)	99.57 (99.26, 99.88)	35.71 (10, 66.67)	93.98 (93.81, 94.21)	
Interns/Residents (N=1,426)					0.08 (0.04, 0.11), chance=0.06
≤5	21.43 (13.1, 30.95)	70.57 (68.18, 73.03)	4.32 (2.7, 6.11)	93.47 (92.81, 94.2)	
≤20	21.43 (13.1, 30.95)	70.72 (68.33, 73.1)	4.35 (2.71, 6.13)	93.48 (92.82, 94.22)	
≤40	21.43 (13.1, 30.95)	71.83 (69.45, 74.14)	4.51 (2.81, 6.35)	93.58 (92.93, 94.29)	
≤60	19.05 (10.71, 27.38)	72.35 (69.97, 74.67)	4.08 (2.45, 5.93)	93.43 (92.83, 94.12)	
≤80	8.33 (3.57, 14.29)	92.55 (91.06, 93.96)	6.25 (2.48, 11.2)	94.14 (93.83, 94.54)	
≤99	2.38 (0, 5.95)	99.78 (99.48, 100)	40 (0, 100)	94.23 (94.09, 94.43)	
Specialist or Consultant (N=109)					0.05 (0.02, 0.08), chance=0.06
≤5	0 (0, 0)	71.43 (62.86, 80)	0 (0, 0)	94.94 (94.29, 95.45)	
≤20	0 (0, 0)	71.43 (62.86, 80)	0 (0, 0)	94.94 (94.29, 95.45)	
≤40	0 (0, 0)	74.29 (65.71, 82.86)	0 (0, 0)	95.12 (94.52, 95.6)	
≤60	0 (0, 0)	76.19 (67.62, 83.81)	0 (0, 0)	95.24 (94.67, 95.65)	
≤80	0 (0, 0)	86.67 (79.98, 92.38)	0 (0, 0)	95.79 (95.45, 96.04)	
≤99	0 (0, 0)	100 (100, 100)	NA (NA, NA)	96.33 (96.33, 96.33)	
Medical Officer (N=187)					0.30 (0.18, 0.41), chance=0.10
≤5	57.89 (36.84, 78.95)	72.02 (64.88, 79.17)	19.2 (12.07, 26.32)	93.85 (90.76, 96.85)	
≤20	57.89 (36.84, 78.95)	75 (67.86, 81.55)	20.93 (13.46, 28.85)	94.07 (91.11, 96.95)	
≤40	42.11 (21.05, 63.16)	79.17 (72.62, 85.12)	18.75 (9.37, 28.57)	92.36 (89.61, 95.17)	
≤60	42.11 (21.05, 63.16)	84.52 (78.57, 89.88)	23.68 (12.12, 36)	92.86 (90.26, 95.42)	
≤80	26.32 (10.53, 47.37)	92.86 (88.69, 96.43)	30 (10.53, 50)	91.76 (89.88, 93.98)	
≤99	10.53 (0, 26.32)	97.62 (95.24, 99.4)	33.33 (0, 80)	90.61 (89.56, 92.18)	

**Supplemental Table 3.** Test characteristics for clinician predicted probability of all-cause, 60-day hospital readmission among all participants by time to unplanned hospital readmission

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area under Precision-Recall Curve (95% CI)
Day 7					0.03 (0.01, 0.05), chance=0.01
≤5	23.4 (10.64, 36.17)	72.77 (71.34, 74.22)	1.03 (0.49, 1.62)	98.72 (98.54, 98.95)	
≤20	23.4 (10.64, 36.17)	73 (71.57, 74.45)	1.04 (0.5, 1.64)	98.73 (98.54, 98.95)	
≤40	23.4 (10.64, 36.17)	74.01 (72.61, 75.39)	1.08 (0.51, 1.7)	98.75 (98.56, 98.96)	
≤60	23.4 (10.64, 36.17)	74.66 (73.28, 76.09)	1.1 (0.53, 1.74)	98.76 (98.57, 98.97)	
≤80	12.77 (4.26, 23.4)	93.69 (92.96, 94.49)	2.38 (0.78, 4.35)	98.87 (98.76, 99.01)	
≤99	4.26 (0, 10.64)	99.77 (99.58, 99.9)	16.67 (0, 46.15)	98.84 (98.79, 98.92)	
Day 14					0.03 (0.02, 0.04), chance=0.02
≤5	17.28 (9.88, 25.93)	72.68 (71.24, 74.07)	1.31 (0.75, 1.95)	97.63 (97.43, 97.87)	
≤20	17.28 (9.88, 25.93)	72.92 (71.47, 74.28)	1.32 (0.75, 1.97)	97.64 (97.43, 97.88)	
≤40	16.05 (8.64, 23.46)	73.89 (72.47, 75.25)	1.28 (0.69, 1.89)	97.64 (97.43, 97.86)	
≤60	16.05 (8.64, 23.46)	74.57 (73.13, 75.91)	1.31 (0.71, 1.94)	97.66 (97.46, 97.88)	
≤80	7.41 (2.47, 13.58)	93.68 (92.87, 94.47)	2.38 (0.8, 4.43)	97.94 (97.83, 98.08)	
≤99	2.47 (0, 6.17)	99.76 (99.61, 99.9)	16.67 (0, 44.44)	97.97 (97.91, 98.04)	
Day 30					0.03 (0.01, 0.06), chance=0.03
≤5	18.64 (11.86, 25.42)	72.57 (71.06, 73.95)	2.06 (1.34, 2.87)	96.61 (96.34, 96.91)	
≤20	18.64 (11.86, 25.42)	72.82 (71.33, 74.21)	2.08 (1.35, 2.9)	96.62 (96.35, 96.92)	
≤40	16.95 (10.17, 23.73)	73.79 (72.31, 75.17)	1.97 (1.23, 2.74)	96.6 (96.33, 96.87)	
≤60	15.25 (9.32, 22.03)	74.42 (72.97, 75.8)	1.81 (1.1, 2.62)	96.56 (96.31, 96.83)	
≤80	6.78 (2.54, 11.86)	93.65 (92.82, 94.39)	3.15 (1.24, 5.53)	96.98 (96.85, 97.14)	
≤99	1.69 (0, 4.24)	99.76 (99.6, 99.89)	16.67 (0, 44.44)	97.01 (96.96, 97.09)	
Day 45					0.05 (0.04, 0.06), chance=0.04
≤5	18.67 (12.67, 24.67)	72.52 (71.13, 73.91)	2.64 (1.84, 3.51)	95.7 (95.39, 96.02)	
≤20	18.67 (12.67, 24.67)	72.76 (71.35, 74.15)	2.66 (1.85, 3.54)	95.71 (95.41, 96.03)	
≤40	16.67 (11.33, 22.67)	73.72 (72.26, 75.09)	2.47 (1.69, 3.33)	95.66 (95.38, 95.97)	
≤60	15.33 (10, 21.33)	74.34 (72.92, 75.73)	2.33 (1.56, 3.19)	95.63 (95.37, 95.92)	
≤80	7.33 (3.33, 11.33)	93.67 (92.87, 94.45)	4.4 (2.21, 7.17)	96.19 (96.04, 96.37)	
≤99	2.67 (0.67, 5.35)	99.81 (99.68, 99.95)	35.71 (9.09, 66.67)	96.24 (96.16, 96.35)	

**Supplemental Table 4.** Test characteristics for clinician predicted probability of all-cause, 60-day post-discharge mortality among all patients by time to post-discharge mortality

Threshold %	Sensitivity (95% CI)	Specificity (95% CI)	Positive Predictive Value (95% CI)	Negative Predictive Value (95% CI)	Area under Precision-Recall Curve (95% CI)
Day 7					0.02 (0.001, 0.03), chance=0.004
≤5	17.65 (0, 35.29)	96.21 (95.59, 96.78)	1.97 (0, 4.29)	99.63 (99.55, 99.71)	
≤20	17.65 (0, 35.29)	96.47 (95.88, 97.01)	2.11 (0, 4.58)	99.63 (99.55, 99.71)	
≤40	17.65 (0, 35.29)	96.7 (96.13, 97.24)	2.26 (0, 4.92)	99.63 (99.55, 99.71)	
≤60	17.65 (0, 35.29)	97.04 (96.49, 97.55)	2.52 (0, 5.5)	99.63 (99.55, 99.71)	
≤80	5.88 (0, 17.65)	99.43 (99.2, 99.66)	4.17 (0, 14.29)	99.59 (99.56, 99.64)	
≤99	0 (0, 0)	99.95 (99.87, 100)	0 (0, 0)	99.56 (99.56, 99.56)	
Day 14					0.02 (0.005, 0.03), chance=0.01
≤5	13.16 (2.63, 23.68)	96.24 (95.62, 96.81)	3.23 (0.7, 6.3)	99.12 (99.01, 99.23)	
≤20	13.16 (2.63, 23.68)	96.5 (95.93, 97.05)	3.45 (0.75, 6.78)	99.12 (99.02, 99.23)	
≤40	13.16 (2.63, 23.68)	96.73 (96.16, 97.25)	3.7 (0.81, 7.26)	99.12 (99.02, 99.23)	
≤60	7.89 (0, 18.42)	97.02 (96.47, 97.54)	2.42 (0, 5.51)	99.07 (99, 99.17)	
≤80	2.63 (0, 7.89)	99.43 (99.2, 99.66)	4 (0, 14.29)	99.04 (99.02, 99.1)	
≤99	0 (0, 0)	99.95 (99.87, 100)	0 (0, 0)	99.02 (99.02, 99.02)	
Day 30					0.04 (0.01, 0.07), chance=0.02
≤5	9.52 (3.17, 17.46)	96.24 (95.64, 96.82)	3.95 (1.29, 7.24)	98.48 (98.37, 98.61)	
≤20	9.52 (3.17, 17.46)	96.5 (95.9, 97.08)	4.23 (1.39, 7.76)	98.48 (98.37, 98.61)	
≤40	9.52 (3.17, 17.46)	96.74 (96.16, 97.29)	4.51 (1.48, 8.33)	98.49 (98.38, 98.62)	
≤60	6.35 (1.59, 12.7)	97.03 (96.45, 97.55)	3.28 (0.79, 6.96)	98.44 (98.36, 98.55)	
≤80	3.17 (0, 7.94)	99.45 (99.22, 99.69)	8.33 (0, 21.74)	98.42 (98.37, 98.5)	
≤99	1.59 (0, 4.76)	99.97 (99.92, 100)	50 (0, 100)	98.41 (98.38, 98.46)	
Day 45					0.04 (0.02, 0.06), chance=0.02
≤5	8.6 (3.23, 13.98)	96.27 (95.66, 96.87)	5.26 (2.17, 8.86)	97.73 (97.61, 97.87)	
≤20	8.6 (3.23, 13.98)	96.53 (95.9, 97.11)	5.63 (2.36, 9.45)	97.74 (97.61, 97.87)	
≤40	8.6 (3.23, 13.98)	96.77 (96.13, 97.32)	6.02 (2.5, 10.17)	97.74 (97.62, 97.88)	
≤60	6.45 (2.15, 11.83)	97.05 (96.48, 97.56)	4.96 (1.64, 9.26)	97.69 (97.59, 97.83)	
≤80	2.15 (0, 5.38)	99.45 (99.21, 99.66)	8.33 (0, 22.22)	97.65 (97.6, 97.73)	
≤99	1.08 (0, 3.23)	99.97 (99.92, 100)	50 (0, 100)	97.64 (97.61, 97.69)	

**Appendix Survey.** Survey that clinicians filled out near the time of discharge for each enrolled neonate or young child in Dar es Salaam, Tanzania and Monrovia, Liberia

### Provider Discharge Survey

Date of Data Collection \_\_\_\_\_  
DD/MM/YYYY

Patient's FIRST name \_\_\_\_\_

Patient's LAST name \_\_\_\_\_

Patient's SEX:

- Male
- Female

Participant's Hospital Identifier (Unique number assigned to each patient) \_\_\_\_\_

Discharging provider year of training:

- Intern
- First year resident
- Second year resident
- Third year resident
- Specialist
- Consultant
- Medical officer
- Other. Please describe. \_\_\_\_\_

In your estimation, is the patient who was discharged at risk of any of the following?

- Re-admission to the hospital within 60 days
- Death in the next 60 days
- None of the above

If yes to any of the above, how likely is that outcome?

- 0%
- 1-5%
- 6-20%
- 21-40%
- 41-60%
- 61-80%
- 81-99%
- 100%

If yes, why do you think the patient is at risk of any of the above outcomes?

- Progression of illness
- Social concerns
- Inability to pay for future medical
- Other. Please describe. \_\_\_\_\_