



Every Newborn Action Plan Metrics

Linked with

Ending Preventable Maternal Mortality

Data Collector Training Programme Session 1.1:

Overview of Training Syllabus







Session Planning

- Purpose: Outline overall course details and plans
- Target Audience: All data collectors & supervisors
- Length of session: 45 mins + 30 mins for quiz
- Type of session: (classroom / practical)
- Resources / tools required:
 Projector, slide deck, training handbook, additional materials/ resource pack.

Structure of this session

- Course Objectives
- Maintaining Data Quality
- Assessment Criteria
- Overview of sessions
- Timetable
- Pre-training knowledge quiz



Primary Training Objective

Participants should be able to competently conduct facility-based observations and data collection using the study tools (both in simulation and in a clinical setting).

Specifics Training Objectives

- 1. The training will address the following key steps to becoming a competent data collector:
- 2. Understand the purpose, objectives, plans for this study and the value of your role of data collector.
- 3. Recognise the value of reliable and high-quality data.
- 4. Review of data collection tools.
- 5. Learn how to complete the data collection tools completely and accurately
- 6. Gain skills in performing clinical observations/ maternal interviews/ data extraction or verification
- 7. Obtain competency by practising data collection and clinical observation skills.

Data Quality Management

By the end of this training, participants should be able to:

- Explain the importance of objective, standardised observation of clinical service delivery.
- Describe informed consent and why it is important.
- Describe how competency in observation is determined.
- Describe how criteria are used to determine skills performance.
- Describe the process used to develop adequate interrater reliability

Assessment Criteria

Demonstrate the ability to conduct clinical observation and use of data collection tools according to protocol

 Course participants should demonstrate the ability to follow study protocols in their use and administration of all tools and checklists, and achieve at least 80% (preferably higher) when evaluated for inter-rater reliability.

Knowledge of the study, observation methods, and related concepts

 Course participant should pass the final knowledge quizz with score of 80% or higher.

Take Pre- course knowledge quiz

Objective: To test participants baseline knowledge

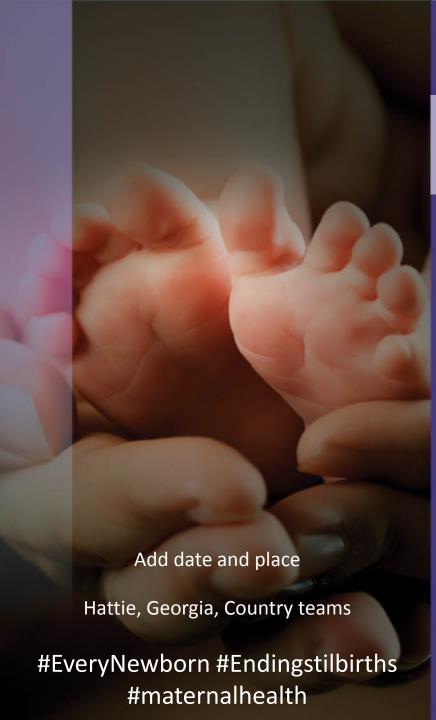
Time: 30 mins

Handouts / Materials: Knowledge quiz question paper; clock; pens

Instructions:

- All course participants should take the quiz as an individual without conferring with others. Participants should complete the quiz in silence and answers should be all their own work. The facilitator should mark and score the quiz results, saving the scores to compare with the post quiz knowledge test.
- No question papers or answer sheets should be removed from the room.







Every Newborn Action Plan Metrics

Linked with

Ending Preventable Maternal Mortality

Data Collector Training Programme Session 1.2:

Introduction to the Study







Session Planning

- Purpose: Outline background to the study
- Target Audience: All data collectors & supervisors
- Length of session: 30 mins
- Type of session: (classroom / practical)
- Resources / tools required:
 Projector, slide deck, training handbook, additional materials/ resource pack.

Structure of this session

- Background to the study
- What is the Every Newborn Action Plan?
- What is the purpose of the study?
- How long will it take
- How will the findings be used and disseminated?







At the end of the Millennium Development Goals era...



No woman should die while giving life

303,000 die



No baby stillborn

2.6 million die



No newborn born to die

2.7 million die



No child dying or stunted

3.2 million die

Progress slower than for child or maternal mortality

Almost 9 million deaths, two-thirds related to birth Marker of equity, and especially quality of care

From 8 MDGs to ...17 SDGs





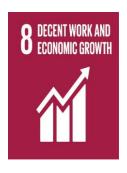








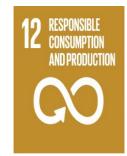














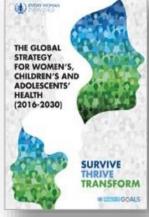












Making the case for health will have to be smarter – economic, environmental

Global Strategy for women, children & ADOLESCENTS (2016-2030)

1. SURVIVE

End preventable deaths for women, newborns, children and stillbirths



3. TRANSFORM

Achieve transformative and sustainable change



2. THRIVE

Realize highest attainable standard of health



Local context

NATIONAL	BANGLADESH (2015)	NEPAL (2015)	TANZANIA (2015)
POPULATION (000)	160,996	28,514	53,470
TOTAL MATERNAL DEATHS	5,200 ('13)	1,100 ('13)	7,900 ('13)
NMR /1000 LIVE BIRTHS	23	22	19
STILLBIRTH RATE /1000 LIVE BIRTHS	36 ('09)	23 ('09)	26 ('09)

What is the Every Newborn Action Plan?

- Based on evidence published in The Lancet Every Newborn series
- It is supported by 197 countries through a World Health Assembly resolution.
- It aims to help countries in reaching the Sustainable Development Goal (SDG) target of fewer than 12 newborn deaths per 1000 live births,
- And the ENAP target of fewer than 12 stillbirths per 1,000 total births by 2030.
- It is also closely linked to the Ending Preventable Maternal Mortality plan
- It aims to encourage government leaders, policymakers and program managers to improve quality of care, and end preventable maternal and newborn deaths, and stillbirths.

REACHING THE EVERY NEWBORN NATIONAL 2020 MILESTONES

COUNTRY PROGRESS, PLANS AND MOVING FORWARD



MAY 2017

What is the Purpose of the Study?

 To assess the quality of maternal and newborn health care in order to provide recommendations to national and global health facility monitoring systems.

Without this, investors (both governments and partners), program managers and advocates lack the reliable information needed to monitor the quality of care, prioritize use of resources, and hold the health system to account.

How will the results be used?

We expect this study to generate a better understanding of the current practices of care at birth, and for small or sick babies.

- To help reduce maternal and newborn mortality, and preventable stillbirths in Tanzania, Bangladesh and Nepal.
- To improve clinical outcomes for mothers and babies, and reduce preventable deaths and stillbirths.
- To improve and prioritise quality of care

These results will also be presented to the World Health Assembly as part of the ENAP, and will inform recommendations to improve the quality of maternal and newborn healthcare across the globe.

Implementation and Communication

How long will it take?

- Facility-based observation data collection completed by March 2018
- Overall research programme including feasibility testing completed by December 2018

How will the results be shared?

Local

Research team: internal webpage, information seminars, team meetings, workshops Participants: hospital notice board, simple language summary, hospital periodic journal, local news paper etc.

National

Internal webpage, national and regional conferences, webinars, meetings and workshops, policy briefs, national advisory committee.

International

Peer reviewed journals, blogs, news updates, consortium websites, reports, briefing summaries, conferences, webinars, UN events and key global meetings including the World Health Assembly.

Where to get more information



Lancet Every Newborn series: http://www.thelancet.com/series/everynewborn

Every Newborn Action Plan (ENAP):

http://www.who.int/maternal child adolescent/topics/newborn/enap consultation/en/



BMC Pregnancy and Childbirth series:

http://www.biomedcentral.com/bmcpregnancychildbirth/supplements/15/s2

ENAP WHO meeting report:

http://www.who.int/maternal_child_adolescent/documents/newborn-health-indicators/en/



MARCH MOOC:

http://www.lshtm.ac.uk/study/freeonlinecourses/women-children-health/index.html

UNICEF: www.childmortality.org

MARCH http://march.lshtm.ac.uk/



Healthy Newborn Network:

http://www.healthynewbornnetwork.org/page/newborn-numbers



INDEPTH: http://www.indepth-network.org/

Thanks to



This work would not have been possible without more than

80 partners involved in the Every Newborn

Action Plan and particularly those on the ENAP metrics coordination group

 Technical inputs from the Coverage Task teams, participants of the WHO meeting and 33 authors on the paper

Multi-partner plan and will take multi-country, multi partner action to work!









Every Newborn Action Plan Metrics

Linked with

Ending Preventable Maternal Mortality

Data Collector Training Programme Session 1.3

The Data Collection Team







Session Planning

Purpose:

Ensure all data collectors understand their roles and responsibilities within the team and how they relate to other team members

- Target Audience: Data collectors & supervisors
- Length of session:Presentation 45 minsActivity B (45 mins)
- Type of session: (classroom / practical)
- Resources / tools required:
 Projector, slide deck, training handbook, supporting materials/ resource pack, marker pens and paper.

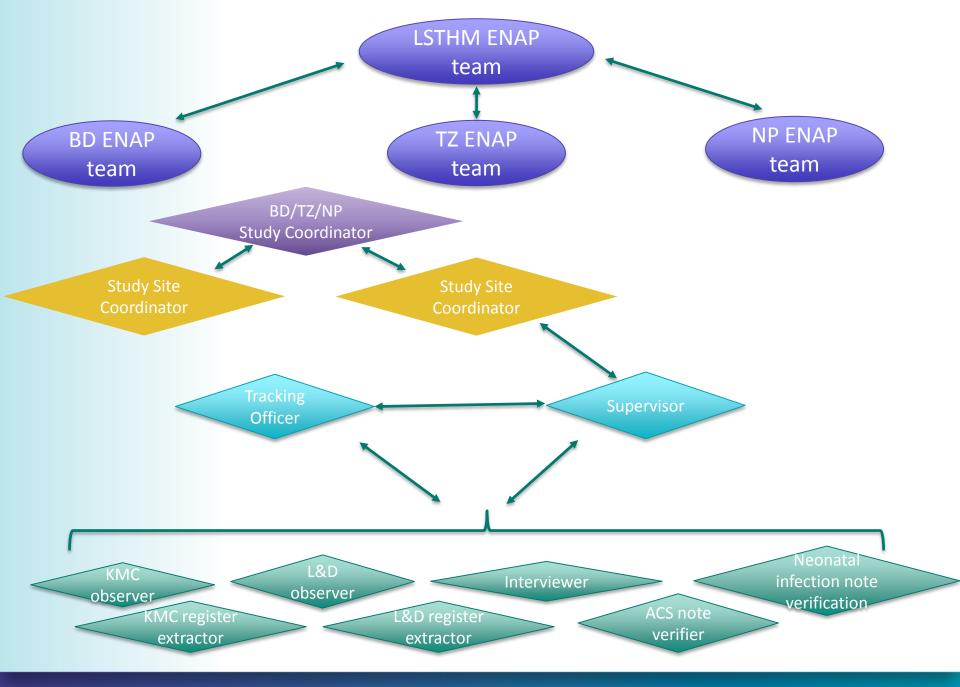
Aims & Learning Outcomes

- To have a clear understanding of the structure and make-up of the team
- 2. To have a clear understanding of the roles and responsibilities within the team
- To understand the flow of mothers and babies through the health facility and how this relates to data collection

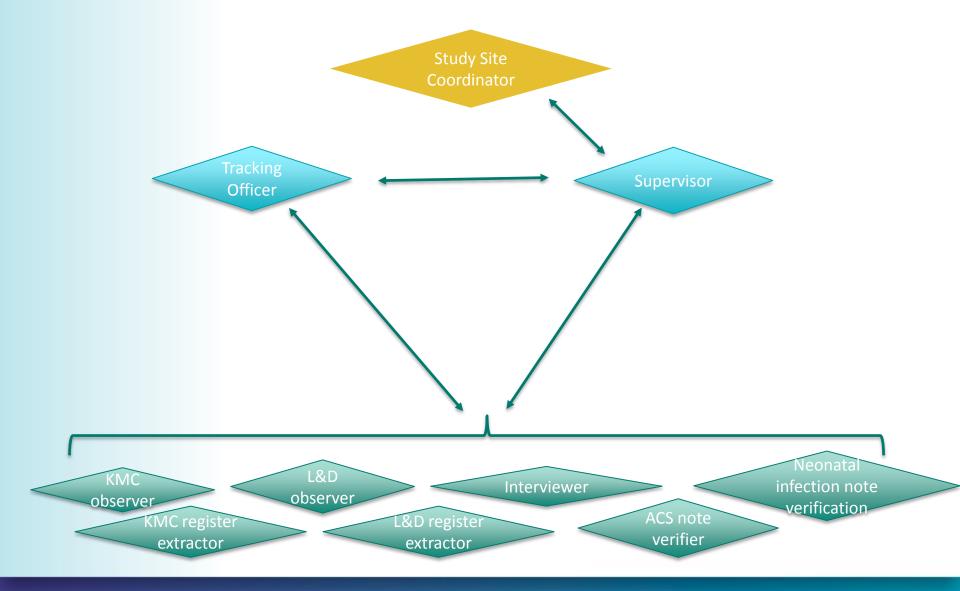


Data collection team

- Study site coordinator
- Supervisors
- Tracking officers (TO)
- L&D ward
- KMC ward observers
- Register data extractors
- Case note verifiers
- Interviewers
- Video data extractors (Nepal)



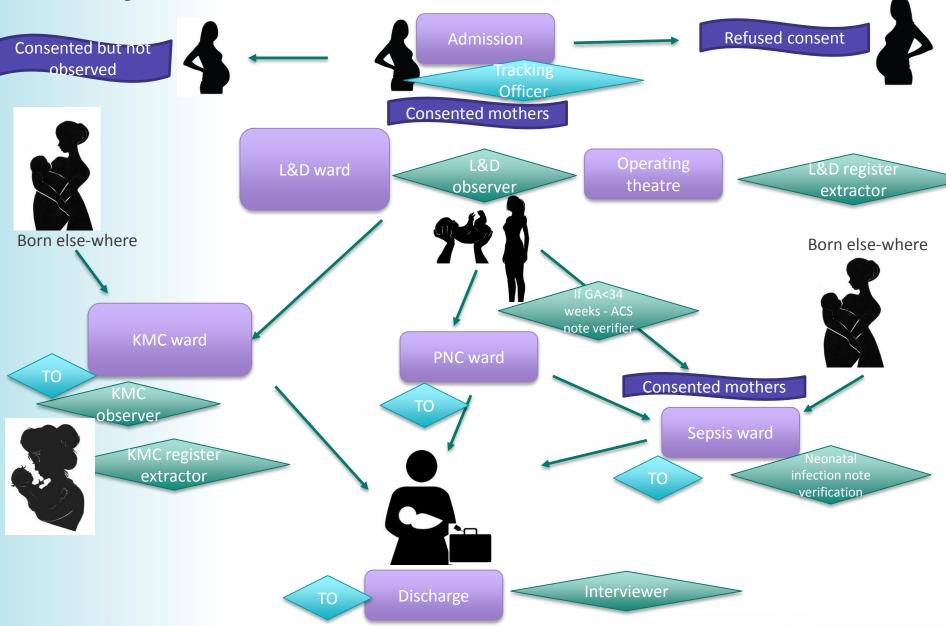
Structure of data collection team



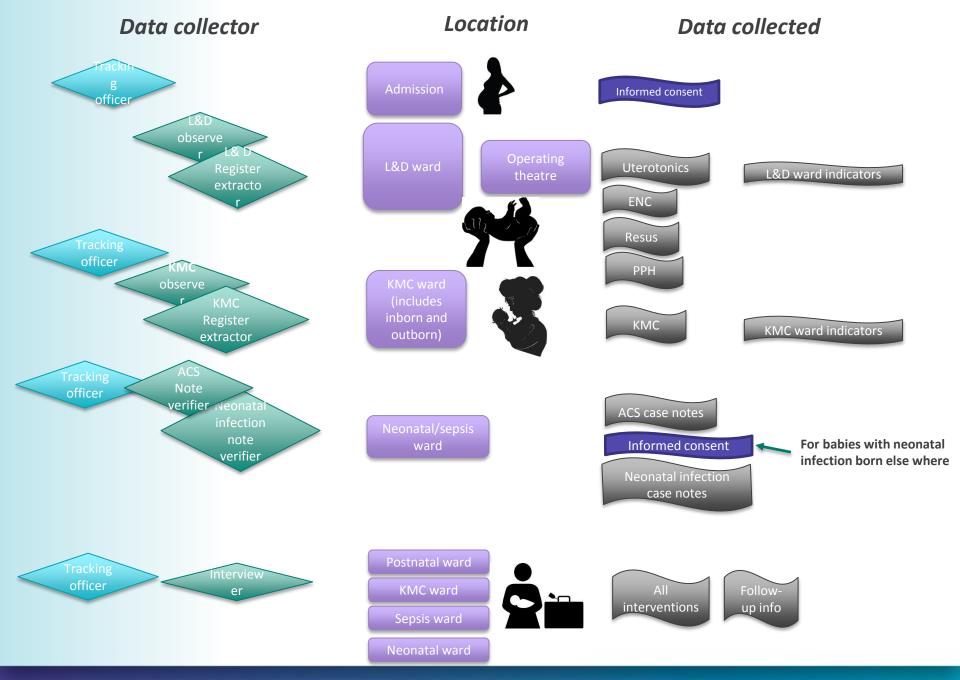
Roles and Responsibilities

Role	Key responsibilities	Location
Supervisors	Ensure all stages of data collection are conducted smoothly and to high quality	Everywhere
Tracking officers (TO)	Enrol mother/babies, collect informed consent, assign to observers, ensure smooth data collection and flow	Admission to L&D ward/OT and admission to KMC ward
L&D ward observers	Observe mothers on L&D ward/OT and collect all relevant data	L&D ward/operating theatre
KMC ward observers	Observe mothers on KMC ward and collect all relevant data	KMC ward
Register data extractors	Complete the L&D/KMC data extraction forms	L&D and/or KMC ward
Case note verifiers	Complete the ACS/neonatal infection verification forms	PNC and sepsis ward
Interviewers	Interview consented mothers pre-discharge	PNC ward
Video data extractors (Nepal)	Extract data from videos	Office

Data flow and location



Data flow and location Refused consent Admission Consented but not observed Tracking Officer **Consented mothers** L&D register extractor L&D **Operating** L&D ward L&D ward indicators theatre observer Uterotonics Resus Born else-where Born else-where ENC PPH ACS weeks - ACS note verifier KMC ward KMC PNC ward TO TO Consented mothers KMC observer Sepsis ward KMC register Neonatal infection note extractor verification KMC ward indicators Treatment of neonatal infection All interventions TO Interviewer Discharge



Activity B: Overview and Structure of data collection team (45 mins)

The trainers will now lead you through a group work activity to consolidate what you have learnt in this section 3 of the training.

Objective: To give you the opportunity to understand the flow of mothers/babies and location of data collectors through the health facility.

Handouts:

Title badges for each data collector you have in your team e.g. L&D observer, data extractor, tracking office. NB. Tablets are not needed.

Place markers- printed sign for each location of data collection, e.g. "Admission", "L&D ward", "KMC ward", "PNC ward", "Sepsis ward"

Time: 45 minute (roughly 30 minutes walking through different scenarios and 15 minutes for questions and discussion)

THE LANCET







Every Newborn Action Plan Metrics

Linked with

Ending Preventable Maternal Mortality

Data Collector Training Programme Session 1.4

The Data Collection Team







Session Planning

Purpose:

Ensure all data collectors understand what high quality data is and how to collect it. Ensure data collectors are familiar with the steps needed to maintain confidentiality.

- Target Audience: Data collectors & supervisors
- Length of session:
 Presentation 45 mins
- Type of session: (classroom / practical)
- Resources / tools required:
 Projector, slide deck, training handbook, supporting materials/ resource pack, marker pens and paper.

Aims & Learning Outcomes

- Data collectors will appreciate the value of high quality data and know how to improve the quality of their data collection
- Data collectors will understand how to ensure confidentiality while carrying out data collection

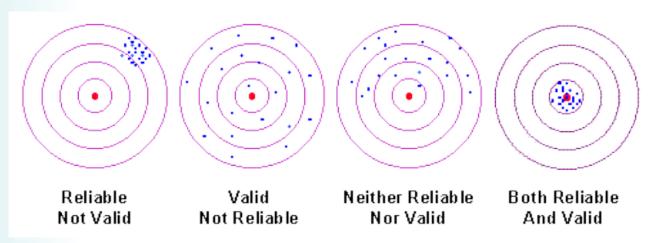


1 Goal:

Observations/interviews/data extraction/data verification is valid and reliable!

Data Quality

- Data are only valuable if they are of high quality
- The quality of the data determines the usefulness of the results
- Reliability: in research reliability means "repeatability" or "consistency"
 - A measure is reliable if it would give us the same result over and dover again.
- Validity: refers to the degree to which data reflects what actually happened



Common Data Collection Errors

- Misunderstanding the difference between "Didn't happen" or "Don't know"—
 - "Didn't happen" is to be used when you observed that an action was <u>not</u> done
 - e.g. you observe that bag and mask ventilation was not used on a baby who was not breathing
 - "Don't know" is to be used when you do not know the answer e.g. you could not see whether the baby was dried or not
- Missing/unreadable data
- Data entered incorrectly
- Delay in data entry

Can you think of any other reasons for errors in data?

Inter-rater reliability

- Inter-rater reliability is the extent to which two or more data collectors agree
- In an ideal world two data collectors would be present for each data entry so that their data could be compared
- For logistical reasons this isn't possible
- Instead: your supervisor will conduct double entry for 10% of cases

Inter-rater reliability

Goal for observers & interviewers: min. agreement of 80%

Goal for extractors & verifiers: min. agreement of 95%

Confidentiality

- Confidentiality refers to the treatment of information that an individual has disclosed in a relationship of trust and with the expectation that it will not be divulged to others without permission.
- You will be collecting personal and sensitive information on the mother and baby
- Everything you observe, read, hear during data collection should be kept confidential and not shared or discussed with others

Confidentiality

What actions can you think of to ensure confidentiality?

- Do not discuss with other data collectors information about a mother/baby unless you need to for the study.
- If you need to discuss a mother/baby with another data collector, use their study ID not their name
- Find a quiet place with some privacy to discuss a mother/baby with another data collector so that you are not over heard
- If a relative, friend, or someone you know comes to the health facility, please alert the supervisor or tracking officer. You should not collect data on someone you know.
- Do not discuss health workers or their behaviour with other data collectors or outside of the health facility.
- If there has been an event which you are required to report (as per clinical incidence guidelines) you should make sure this is only shared with your supervisor
- Never take pictures/videos or collect any other information unrelated to the study!

Quick Quiz

- What is the difference between reliability and validity?
- What things can commonly cause data errors?
- How will supervisors assess the accuracy of the data you are collecting?
- In the context of this study, why is confidentiality important?
- What will you do if you are asked to observe a mother on KMC and you realise you have met her before at your sister's birthday party?
- You witnessed an clinical event with out appropriate response while on the L&D ward- who will you tell and how will you go about telling them?







Every Newborn Action Plan Metrics

Linked with

Ending Preventable Maternal Mortality

Data Collector Training Programme Module 1 Section 1.5 & 1.6

Tablet and App use







Session Planning

1. Purpose:

Ensure data collectors know and understand how to use the tablet and are confident in using ENAP app

2. Target Audience: Everyone

3. Length of session:

Presentation: part i (60 mins)

Presentation: part ii (90 mins)

Activity C (90 mins)

Type of session: (classroom / practical)

Resources / tools required:

Projector, training manual, tablets with app installed

Aims & Learning Outcomes

- Understand how to operate the tablet and use the tablet's basic functions
- 2. Learn how to take care of a tablet on a daily basis
- 3. Understand how the ENAP data collection system works
- 4. Learn the basic elements of the ENAP app
- 5. Practice data collection using ENAP app





(60 MINUTES)

Tablets overview - introduction

- You will use a 10" screen tablet in L&D and 7" in all other locations
- All tablets have some functions that are not necessary for data collection
- You should not use functions of the tablet that are not necessary for data collection. Using extra features will severely reduce the battery life and can interact with the ENAP app functioning, leading to problems
- To respect the privacy of the patients do not use camera or listen to music or record any sounds



Tablets overview

- (A) Front Camera takes self-portraits/videos of yourself (will not be used during this project).
- **(B)** Power/Lock Button turns the device on/off, restarts it, or locks/wakes up the screen.
- **(C) Volume Buttons** adjusts the volume of your device (*will not be used during this project*).
- **(D) Memory Card Slot** optional memory card for removable file storage (*will not be used during this project*).
- **(E) Back Key** Returns to the previous screen, or closes a dialog box, menu, or keyboard.
- (F) Home Button Returns to the Home screen.
- **(G)** Recent Key Displays recent apps (will not be used during this project).

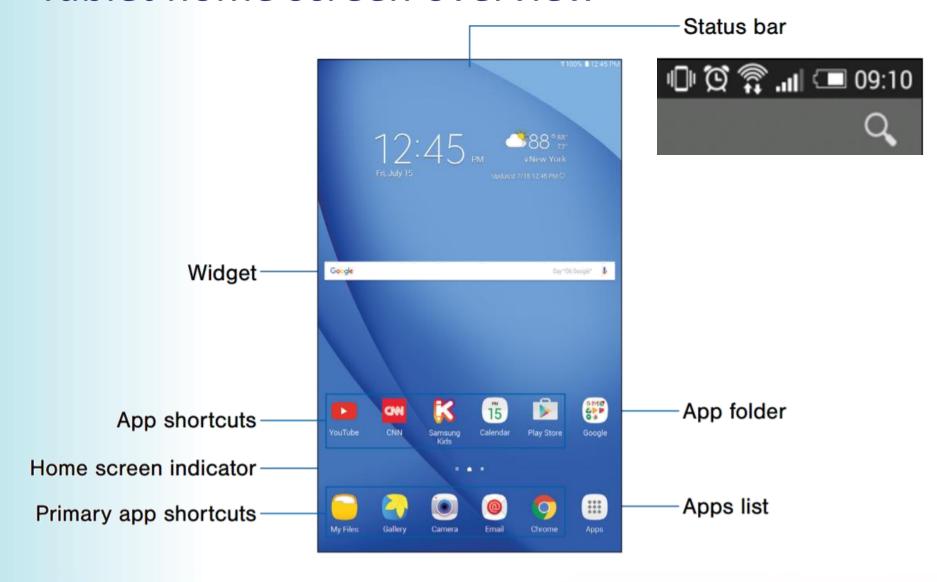


Tablets overview

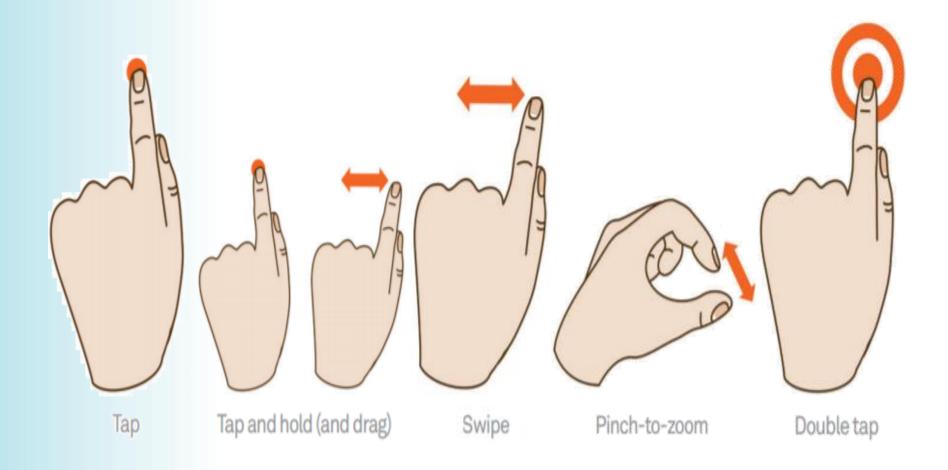
- (A) Microphone records audio and detects voice commands (will not be used during this project).
- (B) USB Charger/Accessory Port connects the USB charger.
- (C) Headset Jack connects a headset (will not be used during this project).
- (D) Rear Camera takes pictures and record videos (will not be used during this project).
- **(E) Flash** illuminates subjects in low-light environments when taking a photo or recording video (will not be used during this project).
- **(F) Speaker** plays music and other sounds (will not be used during this project).
- **(G) Speaker** -play music and other sounds (will not be used during this project).



Tablet home screen overview



Tablet – basic gestures

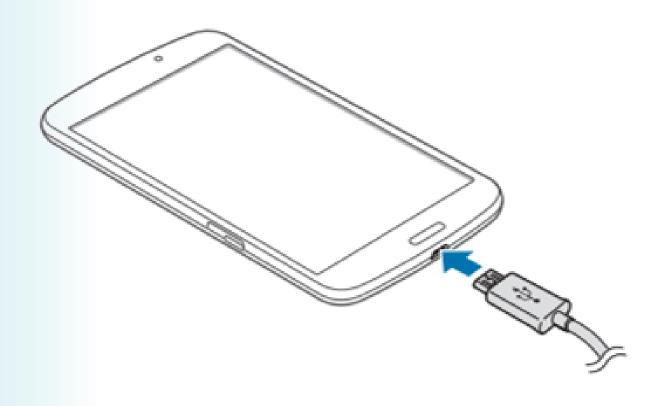


Tablet – text and numbers





Maintenance - charging



Maintenance – basic rules

Storage

• Keep it in the dry place, if possible in wooden cabinet. Always store the device in the lockable storage to protect it against theft.

Cleaning

- Clean the screen with a dry soft cloth only. A clean piece of cotton fabric works well.
- Never use any chemicals to clean the screen especially alcohol, ammonia, or solvents.

Physical protection

- Use a protective case. The case will protect from breaking by fall.
- Dropping your tablet on the floor or hitting it against sharp and dull edges will cause physical damage (break or crack the screen or body of the tablet, battery damage), leading to partial or complete malfunctioning of the device, requiring its full replacement.

Maintenance – basic rules

Digital protection

- Do not attempt to install any additional software.
- As instructed, do not give your log in details (user name and password) to anyone, including your colleagues.
- Your login details are person specific and are assigned on individual basis.
- If you have forgotten your login details, please contact your supervisor.

Device lost or malfunctioning

- If the tablet is lost or stolen, it is your responsibility to notify your manager as soon as possible.
- If your device is malfunctioning, please notify your manager as soon as possible. Your device might require complete reset and reconfiguration, or replacement.
- You can only attempt to reboot device, by turning it OFF and ON. If possible, do not forget to synchronise the data, otherwise, your data might be partially or completely lost.

Basic functions - practice

Turning your device ON and OFF

- To turn on your device, press and hold the Power key. To turn off your device, press and hold the Power key and select Power off → OK.
- When the tablet is ON, press the power button once it is locked and it will be closed.
 To unlock, first press the power / home button, then the lock / unlock screen will appear on the tablet.
- Unlocking any side of the screen with the finger pressure will be unlocked. If the screen does not work for some time (5 minutes), the screen will close automatically.

Unlocking the device

- When you do not use the device, your device turns off the touch screen and automatically locks the touch screen to prevent any unwanted device operations.
- To manually lock the touch screen, press the Power key. To unlock, turn on the screen by pressing the Power key, tap anywhere on the screen, and then flick your finger in any direction.

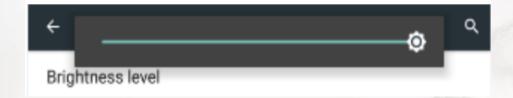
Basic functions - practice

Connecting to Wi-Fi – ON and OFF

- Your device will be configured to use the Wi-Fi or 3/4G connection. In case you
 have lost connectivity, to connect to Wi-Fi, follow the steps:
- 1. Tap the Apps icon in the top right corner of the tablet.
- 2. In the list locate the Settings icon (if you can't see it try scrolling left/right).
- 3. Tap on Wi-Fi to see the local wireless networks (make sure Wi-Fi is turned on, by sliding the switch at the top on the screen).
- 4. Tap the network you want to connect to and enter the wireless key.
- 5. Tap on Wi-Fi and turn it OFF

Basic functions - practice

Screen brightness adjustment



Turning off the sound

- To make sure your device is in the silent mode, adjust the device's volume by pressing the Volume key down.
- Alternatively, select and drag the sliders to adjust the volume level to the minimum.

Troubleshooting



Flight mode ON and OFF

- In Flight mode, your connection to all wireless networks is disabled.
- The Flight Mode is usually being activated in areas where wireless devices are prohibited, such as aeroplanes.
- To synchronise and keep the data up-to-date, your tablet needs to be connected to the network.
- If your network connection is disabled, your device might be in the Flight mode.
- To activate or deactivate Flight mode, go to Settings → more settings, and then check the check box next to Flight mode.

Troubleshooting



Device doesn't turn ON

- If the battery is completely discharged, you cannot turn on the device, even with the USB power adapter connected.
- Allow a depleted battery to charge for a few minutes before you try to turn on the device.
- Always charge the battery only with the original charger provided with the tablet to exclude the possibility of physical damage.

Network or service error message

- When you are in areas with weak signals or poor reception, you may lose reception.
- Move to another area and try again.

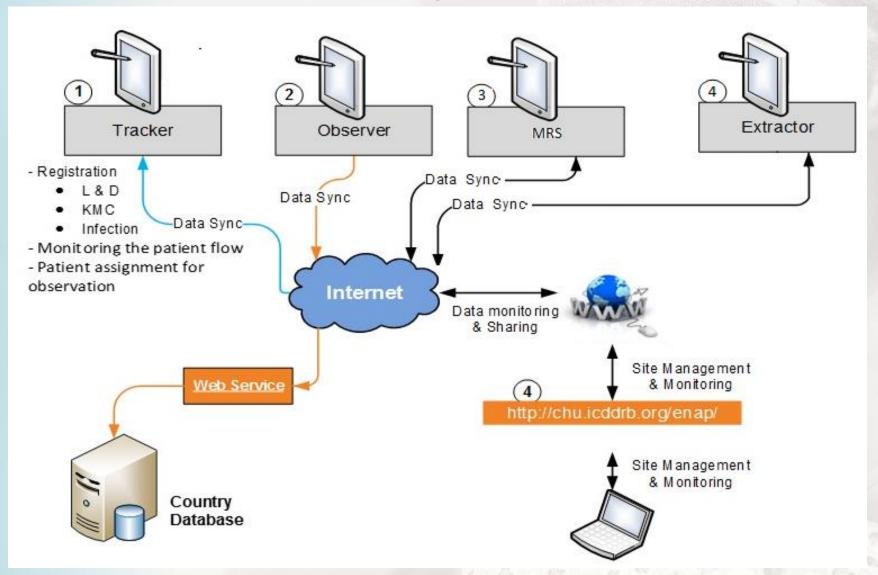
Concluding remarks

If you still experience problems...

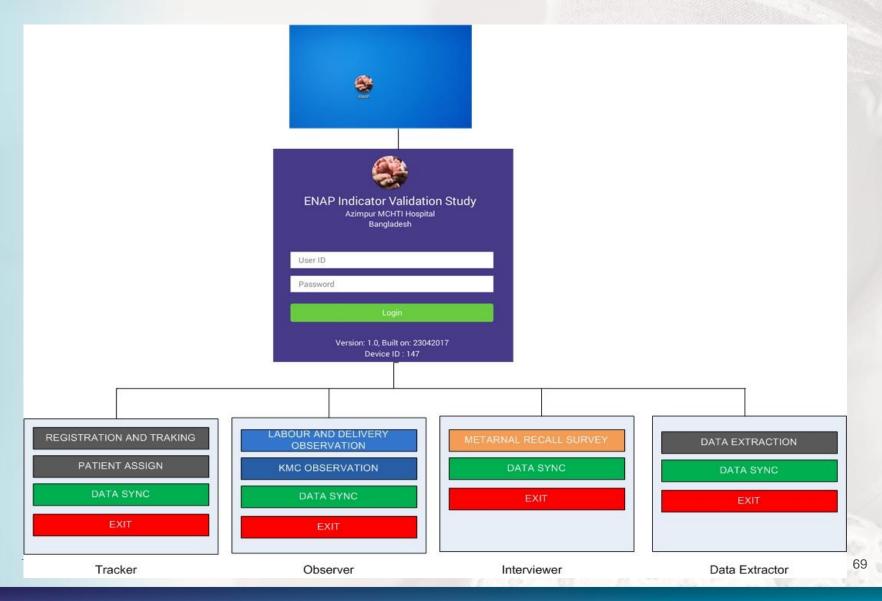
- Turn your device OFF and ON
- Make sure your tablet is charged
- If that doesn't help, please contact your supervisor



ENAP APP overview - flow

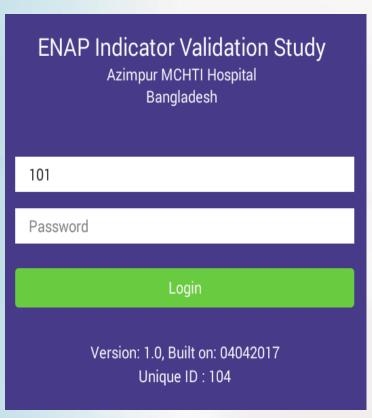


ENAP APP overview - roles



ENAP APP overview - logging in

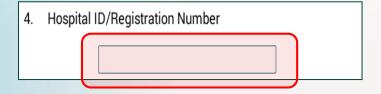
Logging in



- -Tracker
- -Observer
- -Interviewer
- -Data Extractor and Verifier

ENAP APP overview – data entry

Text box



Radio button

- Was the data collection team present in the selected cluster as per plan?
- Yes No
- Did you meet with the supervisor,
 O Yes
 O No inquired about the progress and identified any difficulties in data collection as per plan?

ENAP APP overview – data entry

Check box

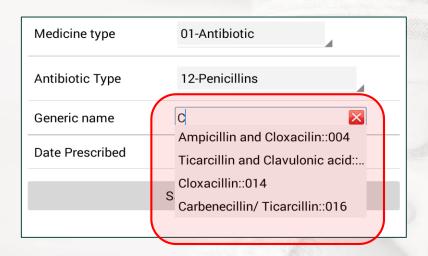
37. Maternal complications diagnosed or known at time of admission □ None/Not Recorded □ Hypertensive disease of pregnancy □ Hypertension □ Pre-eclampsia □ Other Hypertensive disease of pregnancy not specified □ Ante partum haemorrhage □ Not known □ Other complication specify

Drop-down list

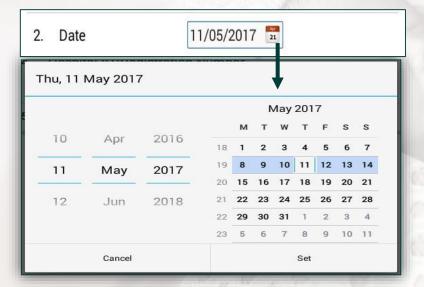


ENAP APP overview – data entry

Auto-complete field

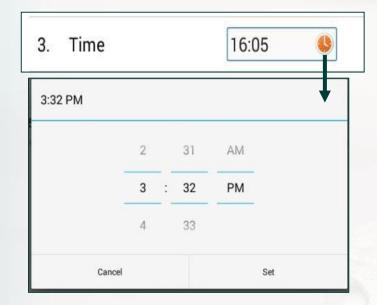


Date field



ENAP APP overview – data entry

Time



Keypad



74

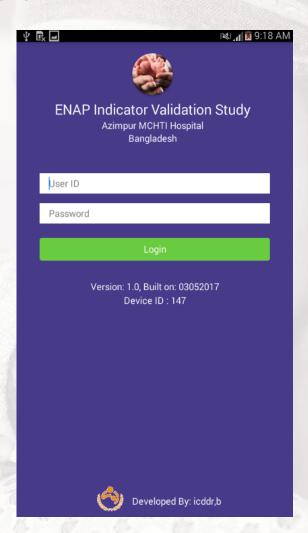
ENAP APP – colour coding

Color	Used in	Description		
Green	OB,MRS,DE,ACS,VER	When required action is completed / Observed-Done		
Red	OB,MRS,DE,ACS,VER	When action is incomplete / Observed-Not Done		
Orange	Patient	Patient switching button		
White	Observation Module	Don't know		
Yellow	Observation Module	Selected Tab		

ENAP APP – opening the app

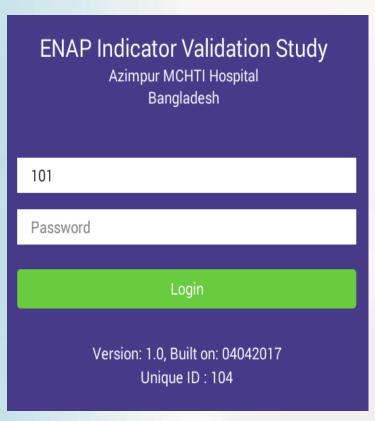






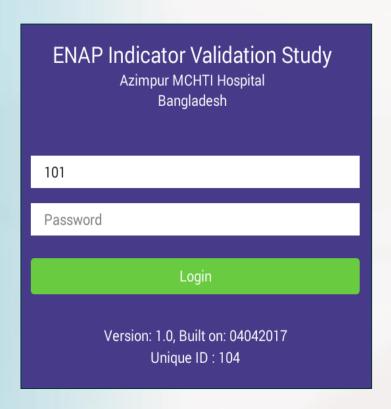
ENAP APP – logging in

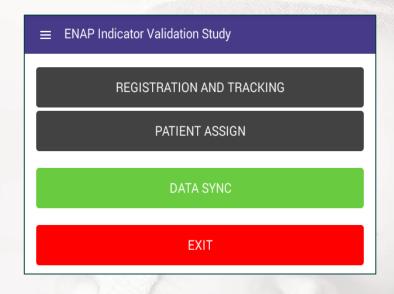
Logging in



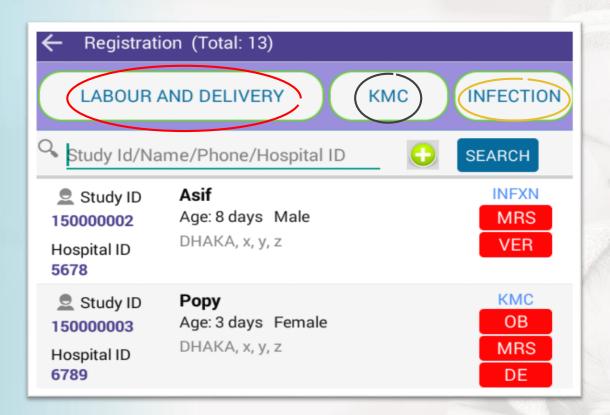
- -Tracker
- -Observer
- -Interviewer
- -Data Extractor and Verifier

Tracker: Logging in

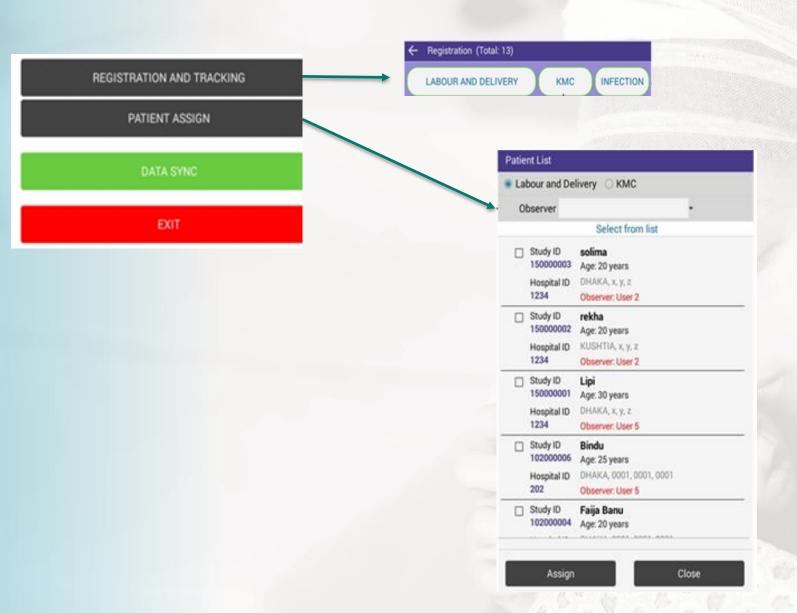


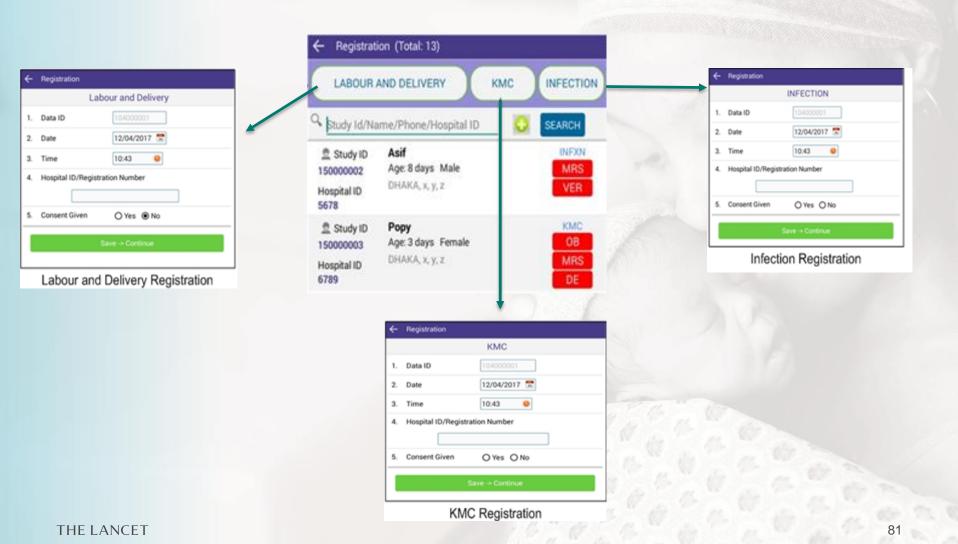


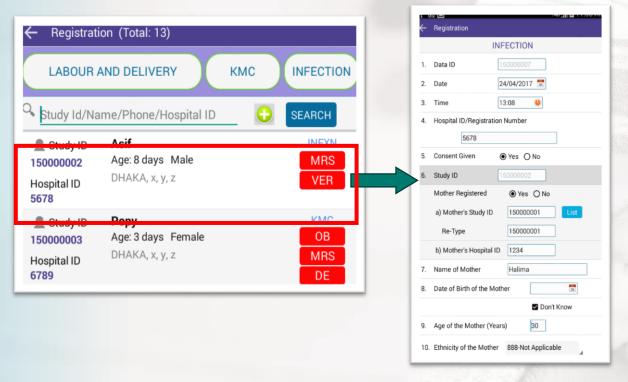
Tracker: registering and tracking



- Labour and Delivery Registration
- 2. KMC Registration
- 3. Infection Registration



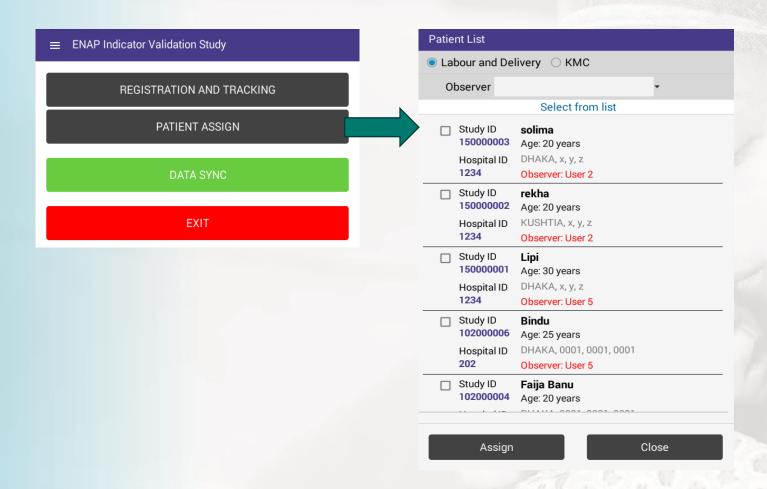




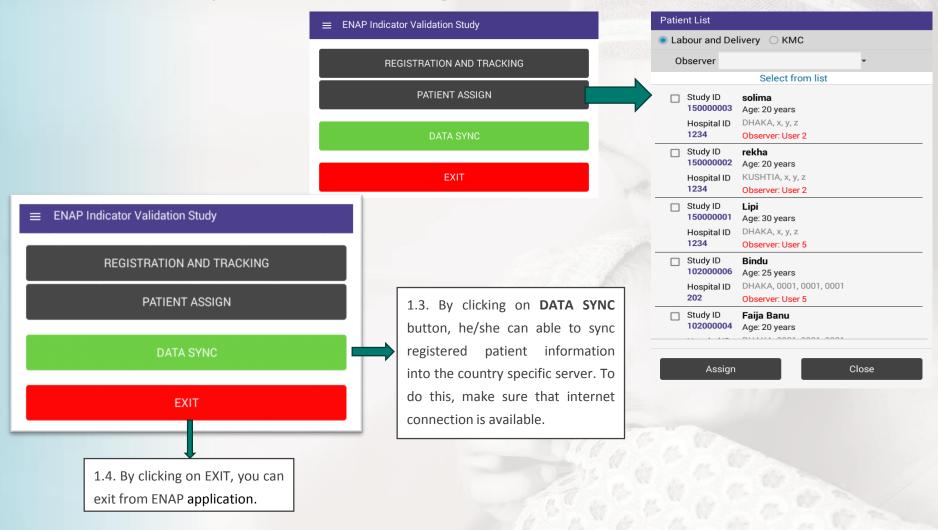
Registrati	on (Total: 13)		
LABOUR	INFECTION		
Study Id/Na	me/Phone/Hospital ID	_ (SEARCH
Study ID 150000002 Hospital ID 5678	Asif Age: 8 days Male DHAKA, x, y, z		MRS VER
Study ID 150000003 Hospital ID 6789	Popy Age: 3 days Female DHAKA, x, y, z		OB MRS

← Regist	ration (Total: 14)				
LABOU	R AND DELIVERY KMC INFECTION				
Study Id/	Name/Phone/Hospital ID SEARCH				
Date From	12/04/2017 To 12/04/2017 To				
Options	-				
Study II 102000006					
Hospital ID 202	3-Incomplete 4-Labour and Delivery Patient				
Study ID 102000005 Hospital ID	5-KMC Patient 6-Infection Patient				

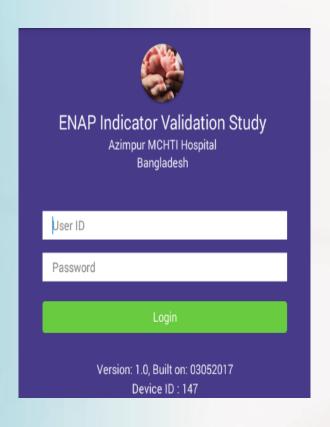
Tracker: patient assign

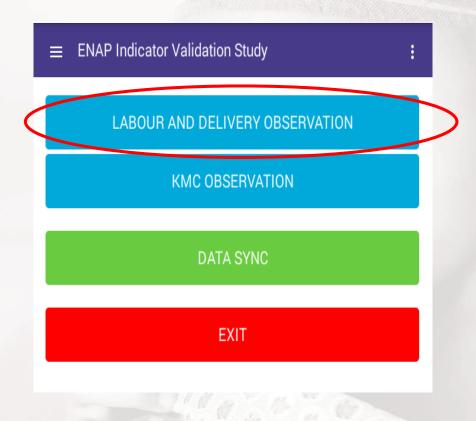


Tracker: patient assign

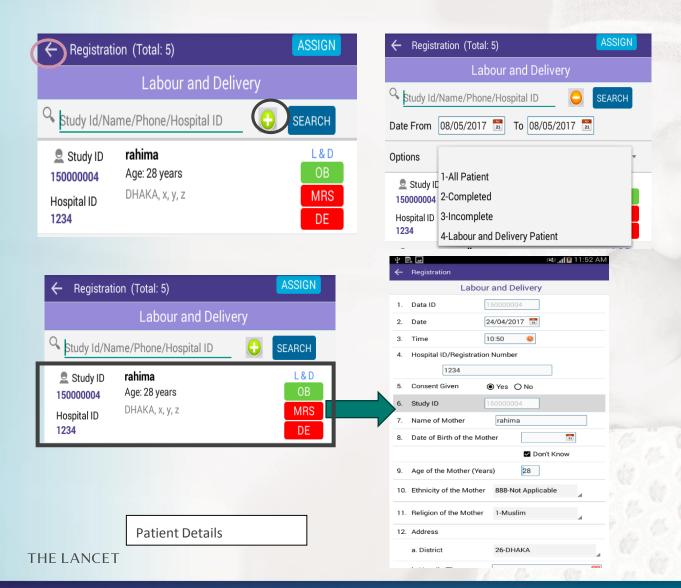


Observer: L&D observation



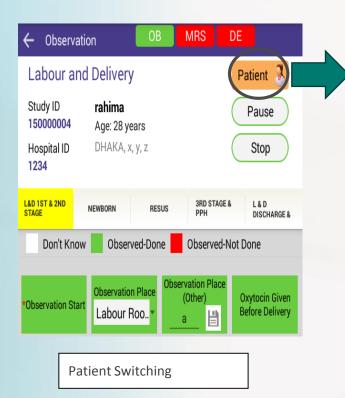


Observer: L&D observation



86

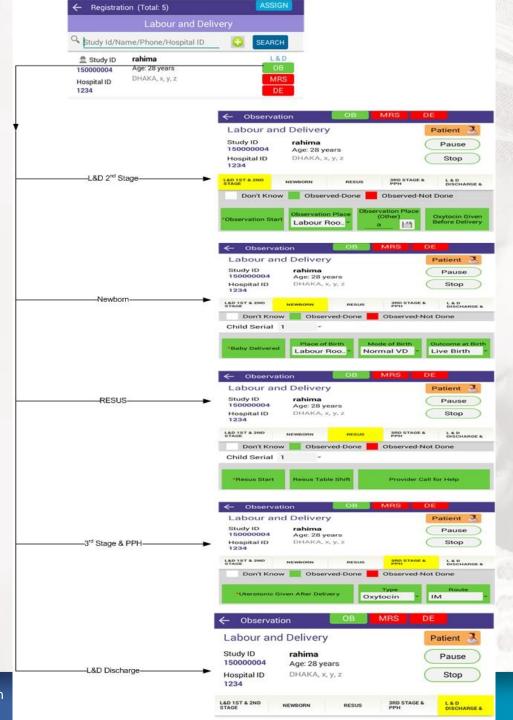
Observer: L&D observation



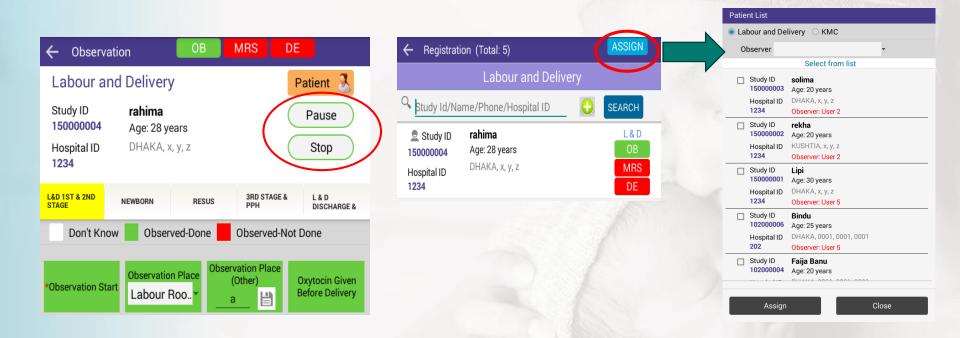




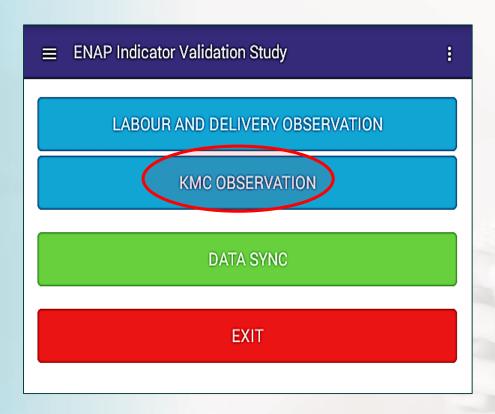
Observer



Observer: pause and stop and reassign



Observer: KMC

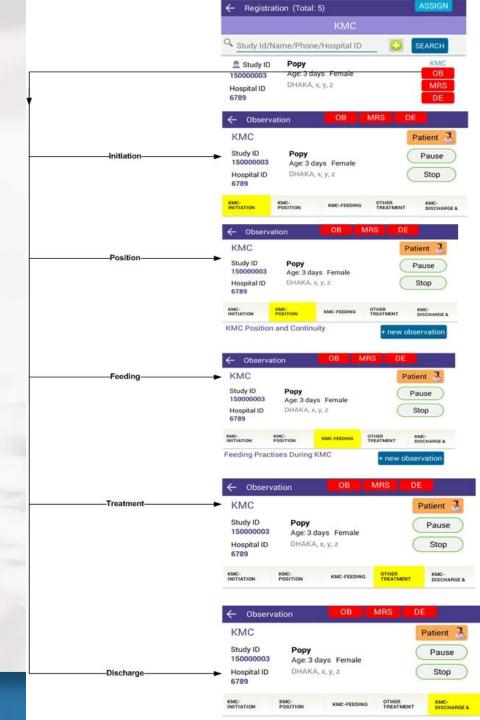


Observer: KMC observation

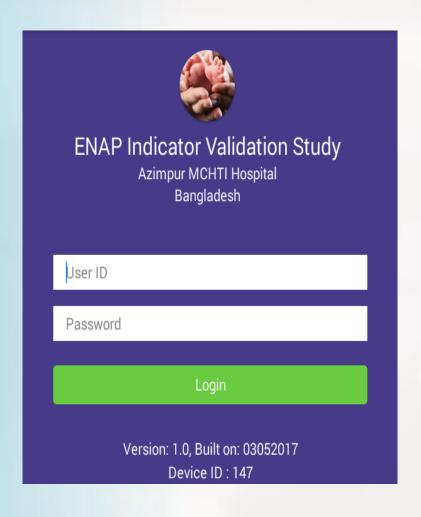


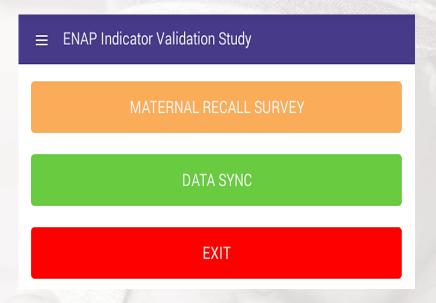


Observer: KMC



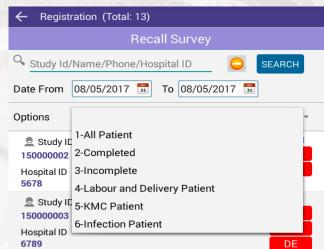
Interviewer





Interviewer





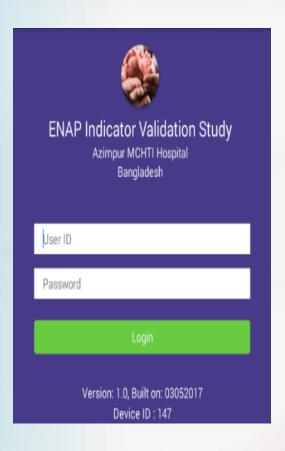


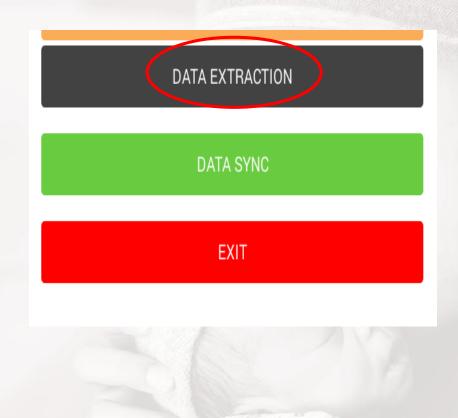
10. Ethnicity of the Mother 888-Not Applicable

Interviewer

Registration (Total: 13) ← Recall Survey Labour and Delivery Study ID Q Study Id/Name/Phone/Hospital ID 150000004 Age: 28 years Hospital ID DHAKA, x, y, z INFXN Study ID 1234 Age: 8 days Male MRS 150000002 DHAKA, x, y, z VER Hospital ID 5678 ← Recall Survey Labour and Delivery Study ID 150000004 Age: 28 years DHAKA, x, y, z Hospital ID 1234 LAD NEONATAL INFECTION SOCIO-ECONOMIC ← Recall Survey Labour and Delivery Study ID 150000004 Age: 28 years DHAKA, x, y, z Hospital ID 1234 ← Recall Survey Labour and Delivery Study ID 150000004 Age: 28 years DHAKA, x, y, z Hospital ID SOCIO-ECONOMIC L&D ← Recall Survey Labour and Delivery Study ID 150000004 Age: 28 years DHAKA, x, y, z Hospital ID 1234 ← Recall Survey Labour and Delivery Study ID 150000004 Age: 28 years Hospital ID DHAKA, x, y, z 1234 LAD

Data extractor





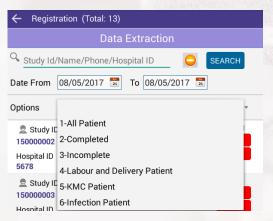
Data extractor

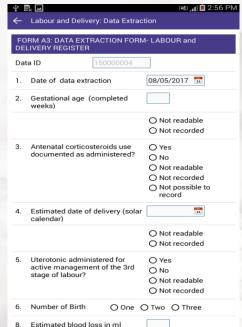






L&D





Activity C: Initial practise using the tablet and app

Instructions:

The trainer will now lead you through exercises that allow you to practice using the tablets and app, these exercises may not reflect the roles you will go on to do but they provide practice in app use.

- Objective: to ensure data collectors understand how to use the tablet and app for data collection.
- Time: 90 mins total
- Handouts / Materials: fully charged tablets with the ENAP app installed
- **Exercise 1 (30 minutes):** practicing tablet and app basics
- **Exercise 2 (20 minutes):** role playing as a tracking officer and a mother
- Exercise 2 (20 minutes): data entry as an observer on the LD ward
- Questions and discussion session (20 minutes)







Every Newborn Action Plan Metrics

Linked with

Ending Preventable Maternal Mortality

Data Collector Training Programme Session 1.8:

Procedure for Life Threatening Events where no appropriate action is being taken







Session Planning

- Purpose: Outline background to the study
- Target Audience: All data collectors & supervisors
- Length of session: 45 mins
- Type of session: classroom
- Resources / tools required:
 Projector, slide deck, training handbook, additional materials/ resource pack, critical incident forms, copy of the algorithm (last slide)

Aims & Structure of this session

- Define what is a life threatening event & what is meant by 'no appropriate action being taken' in this context
- Outline the procedure:
 - -when to intervene
 - -how to intervene
 - -what next within the clinical setting & for patient safety
 - -follow up & next steps required for data quality & research agenda
- Discussion: ethical, legal, clinical & data quality issues
- Role play activity: some example events to run through
- Identify and address any concerns, challenges or questions from the data collection team



What is a Clinical Alert?

- You will find clinical alert integrated into the KMC and, Labour and Delivery observer checklist app.
- They are automatically triggered on input of related clinical observations and will pop up on your screen.
- The flag will remind you of your responsibility during life threatening events directly associated with your observation.
- Once you are sure patient safety is not compromised, you can clear or resolve a clinical alert and continue your observation by.....

What is a life threatening event in this context?

This is any life threatening clinical incident that occurs in the absence of direct and appropriate care from facility healthcare providers.

- To who do you have a duty of care?
 - -research study participants
 - -other patients admitted to the study facilities
- What is direct and appropriate care?
 Care should be given by qualified registered healthcare providers and in accordance with local clinical guidelines

(these will be agreed at facility level ahead of commencing data collection)

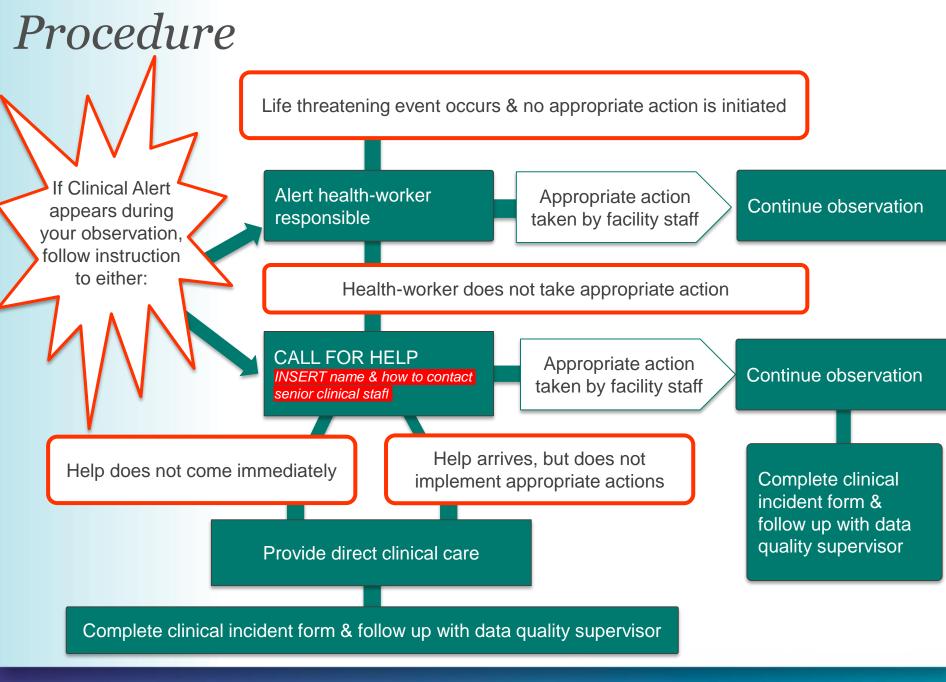
Events with no Clinical Alert

Discussion / group activity (10 mins):

Please discuss and provide examples of when and why you may wish to intervene in the absence of a clinical alert flag?

Example answers to be revealed following discussion:

- All cases of clinical shock not receiving appropriate care:
 obstructive (e.g. PE),
 cardiogenic,
 disruptive (anaphylaxis, septicemia),
 hypovolemic (hemorrhage, severe dehydration)
- Any failure to initiate resuscitation actions (adult or newborn) where required.
- Any undiagnosed or untreated obstetric or newborn emergency.
 (e.g APH, placenta previa, eclampsia / pre-eclampsia, hellp syndrome, cord prolapse, obstructed labor, fetal bradycardia, shoulder dystocia, mal-presentation, PPH, retained placenta, tears / trauma, etc.)
- Where clinical action may cause serious harm (e.g. wrong drug or overdose)



Clinical incident forms

CLINICAL INCIDENT FORM

Please complete all fields noting that these details are <u>not part of your observation</u>. They will be collected after any life-threatening incident where SOP was initiated in the failure of facility staff to act in accordanc with agreed procedures. You may use notes from your observation or the patient records (inpatient notes and drug charts gg_f) if required.

- What is it and where can I find it? This is a mandatory form that should be completed with support from your supervisor and in all cases indicated by the procedures.
- How do you complete it?
 Paper-based form (from your supervisor)
- What next?
 Contact the named responsible from hospital management team and ensure they have a copy of the clinical incident form
- What happens with this information? This information should be escalated to country and LSHTM level and will form the basis of monthly case reviews

Section I	Patient's details	Write or circle when applicable		
	Tracker/data extractor	Name	_	
	Facility	Name ID		
	Participant ID			
	Mother observed during labour & delivery	L&D	1	
	or KMC?	KMC	2	
	Patient's name			
	Patients' age	years		
	Patient's Date of birth	DD/MM/YY		
Section II	Time of Incident Events			
	Date of clinical incident	DD/MM/YY		
	Time health worker responsible alerted to problem	нн:мм		
	Time incident escalated to senior clinical staff as identified in the local SOP	нн:мм		
	Time observation stopped	HH:MM		
	If relevant, time observer initiated provision of direct clinical care	нн:мм		
Section ()	Facility staff in attendance			
		Name		
	Health worker Responsible	ID Time of control		
		Time of arrival		
	Role	ID		
		Time of arrival		
		Name		
	Role	ID		
		Time of arrival		
	Role	Name		
- 1		ID	I I	

Activity E: Role Play

Objective: To give students the opportunity to practice using the procedure algorithm / clinical incident forms.

Instructions (30 mins):

- 1. Break into teams of three.
- 2. Identify one person to be the data collector, one to be the research subject, and one person to watch. Each participant will have the opportunity to play each of the three roles: data collector, research subject, watching.
- Conduct a role play using the algorithm and testing different clinical scenarios you have experienced in your own practise.
- 4. Work together to identify any clinical or ethical challenges you may face and potential solutions.

Discussion (30 minutes)

- Ethical issues (ie. Patient safety)
- Legal issues (ie. permit to practice)
- Data quality issues (ie. Introducing bias)
- Etc.

Summary: Life-Threatening Event where no appropriate action is being taken

When should I intervene?

During any life threatening scenario where local staff are not taking the appropriate action, or when instructed by a clinical alert.

What should I do?

You should follow the procedure algorithm for these events, instructions provided by a clinical alert will direct you to the correct part of the algorithm.

What is a clinical incident form?

This is a mandatory form that should be completed with support from your supervisor and in all cases indicated by the procedure algorithm.

What next?

Your supervisor will contact the named responsible from hospital management team and ensure they have a copy of the clinical incident form. This form will also be used as part of incident review management at facility, country and international level.

