

## NIGERIAQUAL PMTCT AUDIT FORM

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A. FACILITY DETAILS	B. LEVEL (Check one)			
FACILITY NAME:	Primary Health Centre Secondary Hospital			
STATE: LGA:				
IMPLEMENTING PARTNER:	Tertiary Hospital Federal med. Centre Specialist Hospital			
NAME OF ASSESSOR:	Teaching Hospital			
Date of Assessment (dd/mm/yyyy):	Ownership 🔲 Public 🔄 Faith-based			
	Private			
C. PATIENT TYPE				
Booked Unbooked	If unbooked, fill Section D and then skip to			
☐ HIV-infected prior to booking ☐ Known HIV-infected prior to arriving in lab	or Section G			
Unknown status prior to booking				
D. PATIENT DEMOGRAPHICS				
	Patient ANC No.			
Last Name	Age Parity			
First Name				
EDD / / / Date of 1st booking / / Gestational age at 1st booking weeks				
E. ANC TESTING AND COUNSELING (Complete this section for only booked patients)				
Woman counseled for HCT 🔲 Yes 📄 No				
Woman accepted HCT $\Box$ Vas $\Box$ No (If VES complete section E if NO as to section C)				
HCT done       Yes       No       Date HCT done (dd/mm/yyyy)       //         Time HCT done       Before pregnancy (Past)       ANC       Labor/Delivery       After delivery				
HIV status  Positive  Negative  Date of HIV diagnosis (dd/mm/yyyy)	/			
Infant feeding counseling received Yes No				
Woman agreed to partner notification  Yes  No				
Date result received (dd/mm/yyyy):	]			
WHO Clinical Stage				
Patient found to be eligible for ART by Clinician  Yes No Not Recorded				
ARV received 🔲 Yes 🔲 No If yes, date regimen started (dd/mm/yyyy):	/			
Pre pregnancy HAART Yes No				
Ante partum ARV:  ZDV (only opt A) HAART for prophylaxis (opt B) HAART for treatment	None Unknown/Not Indicated Other			
Referred for ART: Yes No Not applicable Not Recorded				
F. PARTNER REGISTER				
Partner of HIV + woman tested and received result Yes No Not Recorded				
Partner HIV status:  Positive  Negative  Not Recorded				



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G. DELIVERY REGISTER (Fill this section for all booked and unbooked patients)					
Date of delivery (dd/mm/yyyy):					
Maternal intrapartum ARV regimen received SdNVP+3TC+ZDV (opt A) HAART for prophylaxis (opt B) HAART for treatment None Unknown/Not Indicated Other					
Mode of delivery: 🔲 Vaginal 🔲 Elective C section 📄 Emergency C section 📄 Other (specify):					
Gestational age at delivery weeks					
Episiotomy: 🗌 Yes 📄 No					
Infant feeding choice:  Exclusive breastfeeding Exclusive breast milk substitute Mixed feeding Other (specify): Maternal outcome: Alive Dead					
Child status:  Still birth  Neonatal death  Alive					
H. CHILD FOLLOW-UP REGISTER					
Was a Dried Blod Spot (DBS) sample collected? Yes No If Yes, date of collection:					
RESULT Date of Sample collection (dd/mm/yyyy) Date Caregiver received results (dd/m	m/yyyy)				
1st PCR (EID)       Done:       Yes       No       Positive         Image: Description of the second seco					
2nd PCR (EID)   Done:   Yes   No   Positive     Negative   Image: Construction of the second secon					
Rapid Test Done:       Yes       No       Positive         at <12 months)					
Rapid Test   Done:   Yes   No   Positive     at 18 months)   Negative   Image: Construction of the second seco					
Referred to ARV clinic: Yes No					
ARV prophylaxis given: 🗌 Yes 🔲 No					
Did the infant receive NVP within 72 hours of delivery?  Yes No					
Did the infant receive any of the following:					
Daily NVP until 1 week after cessation of breastfeeding Daily NVP for 6 weeks sdNVP + daily ZDV for 6 weeks Other Not Recorded					
Infant received CPT: Yes No If YES, age of cotrim initiation: days weeks months					
I. MATERNAL FOLLOW-UP REGISTER					
Mother accessed Family planning: 🔲 Yes 🔲 No					
If yes, method used:  Hormonal Condom IUD Abstinence Other (specify):					
Infant feeding method used: 🔲 Exclusive breastfeeding 🔄 Exclusive breast milk substitute 📄 Mixed feeding 📄 Other (specify):					
Maternal referral:  Family Planning  Support group  Pap smear  Other					
Infant referral: EID OVC ART TB Evaluation Other					