

Serosurveillance to Assess Measles and Rubella Prevalence

Country X, 20YY-YY

Standard Operating Procedures

Specimen Labeling

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1. Definitions:

- 1.1 Cryolabels: Synthetic labels designed for bio-medical research tubes/vials. Cryolabels used for this study must exhibit permanent adhesion to polypropylene tubes and be able to withstand temperatures as low as -196° C. They must also be resistant to multiple freeze-thaw cycles.
- 1.2 Participant ID: A unique ID associated with any child or woman who is enumerated and randomized to be a study participant. The ID is maintained after enrollment and used to link survey data to specimens.
- 1.3 Study Participant: Any child or woman enrolled into this study.

2. Purpose / Background:

- 2.1 Participant IDs must be standardized according to the scheme described in this SOP.

3. Scope / Applicability:

- 3.1 This SOP is applicable to all survey sites.

4. Roles / Responsibilities

- 4.1 *Site specific*

5. Prerequisites / Supplies Needed:

- 5.1 Pre-printed cryovial labels for specimen collection, specimen storage/freezer tubes, consent forms, Lab Register
- 5.2 Extra blank labels (as needed)
- 5.4 Black permanent marker for writing on blank labels

6. Safety/Risk Assessment:

- 6.1 N/A

7. Procedural Steps

Overview of IDs:

	Participant ID
Assigned to	Specimens collected from randomized individuals
Length	7 [8 for aliquots]
Format	CCC-A-RI-S-[#]
Description	CCC: Cluster-specific ID A: Age category A = 9 mo. - < 5 y B = 5 - < 15 y C = 15 – 49 y (female) RI: Randomized ID (1-13 per age group) S: Specimen indicator S = Sera by fingerprick V=Sera by venipuncture D = DBS #: Aliquot number (1-n)
Example	104-A-11-S2 2nd aliquot of serum specimen of the 11th child in age group 9m - 4y in cluster 104

7.1. Specimen IDs:

Specimen IDs are used to identify any specimen collected or stored in the laboratory (sera or DBS). The specimen ID is a **7-digit ID** (CCC - (A/B/C) - (XX) - (S/V/D)). The first 6 digits are participant ID (cluster ID number [CCC], age group category [A/B/C], and randomization ID [XX]). The last letter specifies specimen type: S (sera by fingerprick), V (sera by venipuncture) or D (DBS).

If participant has more than 1 aliquot, aliquot number (1, 2, 3..) will be added at the end of specimen ID.

Examples:

104A11S2: This is the 2nd aliquot of serum specimen (by fingerprick) of the 11th child in age group 9 months - 4 years selected for the survey in cluster 104).

112C02D: This is the DBS card (only 1 card collected) of the 2nd woman in the age 15-49 years category selected for the survey in cluster 112.

Cluster IDs have been assigned as follows (same as Cluster ID for Enumeration ID):

Survey Site	Cluster ID
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Site 1	101 – 130
Site 2	201 – 230
Site 3	301 – 330
Site 4	401 – 430
Site 5	501 – 530
Site 6	601 – 630
Site 7	701 – 730
Site 8	801 – 830
Site 9	901 – 930

Age group letter IDs have been assigned as follows:

Age Group	Letter ID
9 months – < 5 years	A
5 years – < 15 years	B
15 years – 49 years (female)	C

8. Creating labels

8.1. Cryolabels for each cluster will be pre-printed by central investigators or site coordinators using the standard participant ID scheme (CCC-A-RI). Labels will be printed for the following purposes:

- 8.1.1. To label the participant’s consent/parental permission form (in the field)
- 8.1.2. To label the participant’s specimen collection tubes (in the field)
- 8.1.3. To label aliquot tubes after processing
- 8.1.4. To label the laboratory register at the time of processing

8.2. For specimen collection tubes/DBS cards, the specimen indicator (S for sera by fingerprick; V for sera by venipuncture, or D for DBS) will be appended to the label using a waterproof marker prior to affixing to the specimen tube as appropriate.

8.3. For aliquots, the aliquot indicator (1-n, n=aliquot #) will be appended to the label using a waterproof marker prior to affixing to the aliquot tube. Aliquot number may also be marked on top of tube cap with waterproof marker.

9. Quality Assurance / Quality Control

9.1. Each site should have a system in place to ensure that each participant ID is used only once.

9.2. When the cluster is finished, the central study team will check tablet and laboratory data to ensure there are no errors or duplicates.