This basic results template is intended to help prepare and organize study-specific information to report results for a **3-arm study** in the Protocol Registration System (PRS) for ClinicalTrials.gov. Separate templates are available for 1- or 2-arm studies. For 4 or more arms, use Microsoft Word to insert table columns as needed or contact the UCLA PRS Administrator to request assistance.

* Shaded cells (Pink) represent areas to be customized or completed with study-specific information.
* **[Hover mouse over hyperlinked text to view Help tips](#Hover" \o "Hyperlinked text displays help tips)**

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| **PARTICIPANT FLOW** |
| **Recruitment details:** *Key information relevant to the recruitment process for the overall study, such as dates of the recruitment period and types of location (e.g., medical clinic), to provide context. Example:* At 1 U.S. clinical site (UCLA), 35 patients were enrolled to the study between 10/15/2007 and 12/03/2009 |
| **Pre-assignment details:** *Describe any significant events and approaches for the overall study (e.g., wash out, run-in, transition) following enrollment, but prior to group assignment, e.g., explain any exclusion of enrolled participants before assignment to groups.* |
| **Reporting Groups:** |
| Arm/Group 1 Title (Description):  |  |
| Arm/Group 2 Title (Description): |  |
| Arm/Group 3 Title (Description): |  |
| **Period:** *Overall Study* | **Arm 1** | **Arm 2** | **Arm 3** |
| **How many started study** |  |  |  |
| **Study-specific midpoint (if applicable)** |  |  |  |
| **Study-specific midpoint (if applicable)** |  |  |  |
| **How many completed study** |  |  |  |
| **How many did not complete study** |  |  |  |

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| **BASELINE CHARACTERISTICS** |
| Arm/Group 1 Title (Description):  | *Provide if different from Participant Flow Arm 1 Title(Description)* |
| Arm/Group 2 Title (Description): | *Provide if different from Participant Flow Arm 2 Title(Description)* |
| Arm/Group 3 Title (Description): | *Provide if different from Participant Flow Arm 3 Title(Description)* |
|  | **Arm 1** | **Arm 2** | **Arm 3** | **Overall (total)** |
| **N evaluated** |  |  |  |  |
| Measure of [**Central Tendency**](#CentralT) **(**[**Dispersion**](#Dispersion)**)\***↓ |  |
| **Age**  | mean(sd)\* | \_\_\_\_ (\_\_\_\_) | \_\_\_\_ (\_\_\_\_) | \_\_\_\_ (\_\_\_\_) | \_\_\_\_ (\_\_\_\_) |
| **Gender:** | **Arm 1** | **Arm 2** | **Arm 3** | **Overall (total)** |
|  **Female**  | **(N)\*** |  |  |  |  |
|  **Male**  | **(N)\*** |  |  |  |  |
| **Region of Enrollment:** | **Arm 1** | **Arm 2** | **Arm 3** | **Overall (total)** |
|  **U.S.**  | **(N)\*** |  |  |  |  |
| **Study-specific Baseline Measures, if applicable:** |  |
| Measure of [**Central Tendency**](#CentralT) **(**[**Dispersion**](#Dispersion)**)\***↓ | **Arm 1** | **Arm 2** | **Arm 3** | **Overall (total)** |
| *Name of measure* |  | \_\_\_\_\_\_ (\_\_\_\_ \_)\* | \_\_\_\_\_ (\_\_\_\_\_\_)\* | \_\_\_\_\_ (\_\_\_\_\_\_)\* | \_\_\_\_\_ (\_\_\_\_\_\_)\* |
| *Name of measure* |  | \_\_\_\_\_ (\_\_\_\_\_\_)\* | \_\_\_\_\_ (\_\_\_\_\_\_)\* | \_\_\_\_\_ (\_\_\_\_\_\_)\* | \_\_\_\_\_ (\_\_\_\_\_\_)\* |

\*Report measure of central tendency (e.g., mean, median) and measure of dispersion (e.g., std dev, range); Measures reported as “Number” (N) do not require a measure of dispersion.

**PRIMARY OUTCOME MEASURE(S) –** ClinicalTrials.gov requires results, and statistical analyses as appropriate, for ALL pre-specified Primary and Secondary Outcome Measures in an [Applicable Clinical Trial (ACT)](https://clinicaltrials.gov/ct2/manage-recs/fdaaa#WhichTrialsMustBeRegistered).

* **[Hover mouse over hyperlinked text to view Help tips](#Hover" \o "Hyperlinked text displays help tips)**

Copy and paste the table if reporting multiple Primary Outcomes

|  |
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| **PRIMARY OUTCOME MEASURE** |
| [**Measure Title:**](#Title) |  |
| [**Measure Description:**](#Description) |  |
| [**Analysis Population Description:**](#Analysis_population) |  |
| **Safety Issue?** | *Yes or No* |
| [**Time Frame:**](#Time_Frame) |  |
| Arm/Group 1 Title (Description):  | *Provide if different from Participant Flow Arm 1 Title(Description)* |
| Arm/Group 2 Title (Description): | *Provide if different from Participant Flow Arm 2 Title(Description)* |
| Arm/Group 3 Title (Description): | *Provide if different from Participant Flow Arm 3 Title(Description)* |
| **Reporting Groups:** | **Arm 1** | **Arm 2** | **Arm 3** | **Overall (total)** |
| **N evaluated** |  |  |  |  |
| **[Central Tendency](#CentralT" \o "Result may be reported as Number, Mean, Median, Least Squares Mean, Geometric Mean or Log mean) (**[**Dispersion**](#Dispersion)**)\*** | \_\_\_\_(\_\_\_\_\_) | \_\_\_\_\_\_(\_\_\_\_\_) | \_\_\_\_\_\_(\_\_\_\_\_) | \_\_\_\_\_\_(\_\_\_\_\_) |
| **Unit of measure:**  |
| *Click “Add Statistical Analysis” in the PRS as appropriate to describe within and between-group comparisons. Required fields will depend on how you report the outcomes in the table, and the type of analyses performed.*  |

\*Report measure of central tendency (e.g., mean, median) and measure of dispersion (e.g., std dev, range); Measures reported as “Number” (N) do not require a measure of dispersion.

**SECONDARY OUTCOME MEASURE(S) –** ClinicalTrials.gov requires results, and statistical analyses as appropriate, for ALL pre-specified Primary and Secondary Outcome Measures in an [Applicable Clinical Trial (ACT)](https://clinicaltrials.gov/ct2/manage-recs/fdaaa#WhichTrialsMustBeRegistered).

* **[Hover mouse over hyperlinked text to view Help tips](#Hover" \o "Hyperlinked text displays help tips)**

Copy and paste the table if reporting multiple Secondary Outcomes

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| **SECONDARY OUTCOME MEASURE** |
| [**Measure Title:**](#Title) |  |
| [**Measure Description:**](#Description) |  |
| [**Analysis Population Description:**](#Analysis_population) |  |
| **Safety Issue?** | *Yes or No* |
| [**Time Frame:**](#Time_Frame) |  |
| Arm/Group 1 Title (Description):  | *Provide if different from Participant Flow Arm 1 Title(Description)* |
| Arm/Group 2 Title (Description): | *Provide if different from Participant Flow Arm 2 Title(Description)* |
| Arm/Group 3 Title (Description): | *Provide if different from Participant Flow Arm 3 Title(Description)* |
| **Reporting Groups:** | **Arm 1** | **Arm 2** | **Arm 3** | **Overall (total)** |
| **N evaluated** |  |  |  |  |
| [**Central Tendency**](#CentralT) **(**[**Dispersion**](#Dispersion)**)\*** | \_\_\_\_(\_\_\_\_\_) | \_\_\_\_\_\_(\_\_\_\_\_) | \_\_\_\_\_\_(\_\_\_\_\_) | \_\_\_\_\_\_(\_\_\_\_\_) |
| **Unit of measure:**  |
| *Click “Add Statistical Analysis” in the PRS as appropriate to describe within and between-group comparisons. Required fields will depend on how you report the outcomes in the table, and the type of analyses performed.*  |

\*Report measure of central tendency (e.g., mean, median) and measure of dispersion (e.g., std dev, range); Measures reported as “Number” (N) do not require a measure of dispersion.

**OTHER PRE-SPECIFIED OUTCOME MEASURE(S) –** ClinicalTrials.gov requires results, and statistical analyses as appropriate, for ALL pre-specified Primary and Secondary Outcome Measures in an [Applicable Clinical Trial (ACT)](https://clinicaltrials.gov/ct2/manage-recs/fdaaa#WhichTrialsMustBeRegistered).

* **[Hover mouse over hyperlinked text to view Help tips](#Hover" \o "Hyperlinked text displays help tips)**

Copy and paste the table if reporting multiple Other Pre-Specified Outcomes

|  |
| --- |
| **OTHER PRE-SPECIFIED OUTCOME MEASURE** |
| [**Measure Title:**](#Title) |  |
| [**Measure Description:**](#Description) |  |
| [**Analysis Population Description:**](#Analysis_population) |  |
| **Safety Issue?** | *Yes or No* |
| [**Time Frame:**](#Time_Frame) |  |
| Arm/Group 1 Title (Description):  | *Provide if different from Participant Flow Arm 1 Title(Description)* |
| Arm/Group 2 Title (Description): | *Provide if different from Participant Flow Arm 2 Title(Description)* |
| Arm/Group 3 Title (Description): | *Provide if different from Participant Flow Arm 3 Title(Description)* |
| **Reporting Groups:** | **Arm 1** | **Arm 2** | **Arm 3** | **Overall (total)** |
| **N evaluated** |  |  |  |  |
| [**Central Tendency**](#CentralT) **(**[**Dispersion**](#Dispersion)**)\*** | \_\_\_\_(\_\_\_\_\_) | \_\_\_\_\_\_(\_\_\_\_\_) | \_\_\_\_\_\_(\_\_\_\_\_) | \_\_\_\_\_\_(\_\_\_\_\_) |
| **Unit of measure:**  |
| *Click “Add Statistical Analysis” in the PRS as appropriate to describe within and between-group comparisons. Required fields will depend on how you report the outcomes in the table, and the type of analyses performed.*  |

\*Report measure of central tendency (e.g., mean, median) and measure of dispersion (e.g., std dev, range); Measures reported as “Number” (N) do not require a measure of dispersion.

**ADVERSE EVENTS** – The PRS has separate modules for reporting Serious Adverse Events (SAEs) and Other (Non-Serious) Adverse Events (AEs)

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| **SERIOUS ADVERSE EVENTS\*** |
| Arm/Group 1 Title (Description):  | *Provide if different from Participant Flow Arm 1 Title(Description)* |
| Arm/Group 2 Title (Description): | *Provide if different from Participant Flow Arm 2 Title(Description)* |
| Arm/Group 3 Title (Description): | *Provide if different from Participant Flow Arm 3 Title(Description)* |
|  | **Arm 1** | **Arm 2** | **Total** |
| # participants affected by any SAE **/** # at risk: |  |  |  |
| **SAE description**  | **System Organ Class\*** | **# affected/at risk** | **# affected/at risk** | **# affected/at risk** |
|  |  |  |  |  |
|  |  |  |  |  |
|  |  |  |  |  |

Add table rows as needed to report additional SAEs

|  |
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| **OTHER (NON-SERIOUS) ADVERSE EVENTS\*** |
| Arm/Group 1 Title (Description):  | *Provide if different from Participant Flow Arm 1 Title(Description)* |
| Arm/Group 2 Title (Description): | *Provide if different from Participant Flow Arm 2 Title(Description)* |
| Arm/Group 3 Title (Description): | *Provide if different from Participant Flow Arm 3 Title(Description)* |
|  | **Arm 1** | **Arm 2** | **Total** |
| # participants affected by any AE **/** # at risk: |  |  |  |
| **AE description**  | **System Organ Class\*** | **# affected/at risk** | **# affected/at risk** | **# affected/at risk** |
|  |  |  |  |  |
|  |  |  |  |  |
|  |  |  |  |  |

Add table rows as needed to report additional AEs

**\*[MedDRA System Organ Classes (SOC):](http://www.meddra.org/how-to-use/basics/hierarchy)**

1. Blood and lymphatic system disorders
2. Cardiac disorders
3. Congenital, familial and genetic disorders
4. Ear and labyrinth disorders
5. Endocrine disorders
6. Eye disorders
7. Gastrointestinal disorders
8. General disorders and administration site conditions
9. Hepatobiliary disorders
10. Immune system disorders
11. Infections and infestations
12. Injury, poisoning and procedural complications
13. Investigations
14. Metabolism and nutrition disorders
15. Musculoskeletal and connective tissue disorders
16. Neoplasms benign, malignant and unspecified (incl cysts and polyps)
17. Nervous system disorders
18. Pregnancy, puerperium and perinatal conditions
19. Product issues
20. Psychiatric disorders
21. Renal and urinary disorders
22. Reproductive system and breast disorders
23. Respiratory, thoracic and mediastinal disorders
24. Skin and subcutaneous tissue disorders
25. Social circumstances
26. Surgical and medical procedures
27. Vascular disorders