## **SnapShot Reporting practices for publishing results with human PSCs and tissue stem cells**

**ISSCR Task Force for Basic Research Standards<sup>1</sup>** 

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| Metadata  |                      |                                | Characterization of pluripotency and the undifferent  | ntiated state (PS            | Cs only)                   |
|---|----------------------|--------------------------------|---|------------------------------|----------------------------|
| Describe the source of the cells/cell line including:   | Reference<br>section | Page reported<br>in manuscript | Describe the following:   | Reference<br>section         | Page report<br>in manuscri |
| Name (or names)/alias of line   | 1.4; 5.1.2           |                                | Assay methodology   | 2.1; 2.2; 5.2;               |                            |
| Unique ID/registry # (name of registry)   | 1.4                  |                                |   | Appendix 4                   |                            |
| Source (vendor and catalog number if obtained commercially); biopsy site and derivation details (if derived)  | 4.1.1; 5.1           |                                | Quantitative results along with statistical analysis  | 2.1; 2.2; 5.2;<br>Appendix 4 |                            |
|   |                      |                                | Timing of analysis in relation to key experiments reported  | 2.1; 2.2; 5.2                |                            |
| Additional metadata as applicable (e.g., sex, ethnicity, disease information, known mutations, etc.)  | 4.1.2; 5.4.1         |                                | Confirmation of cell type (TSCs only)   |                              |                            |
| Culture details   |                      |                                | Describe the characterization of the following:   | Reference<br>section         | Page report<br>in manuscri |
| Describe methods used for isolation, maintenance, and   | Reference            | Page reported                  | Starting population(s) with recognized markers and methods  | 4.1; 4.3.1; 5.4.1            |                            |
| preservation of the cells including:  | section              | in manuscript                  | Phenotype of expanded cells   | 4.1; 4.3.1; 5.4.1            |                            |
| Passaging/dissociation/split ratio  | 3.2; 4.2.2; 5.1.1    |                                | Demonstration of lineage potential  | 4.1; 4.3.1                   |                            |
| Freezing and thawing  | 5.1.1                |                                | Molecular characterization  |                              |                            |
| Culture reagents used (e.g., media, matrices, growth factors, etc.) with vendor and catalog number  | 4.2.2; 5.1.1         |                                |   | Reference                    | Page report                |
| The passage number of the cryopreserved/characterized<br>Master Cell Bank or Working Cell Bank stocks used, and<br>the number of subsequent passages prior to and during<br>experimentation | 1.2; 3.2.2; 5.1.1    |                                | Describe the following:   | section                      | in manuscri                |
|   |                      |                                | Confirmation of disease mutation (if applicable)  | 4.3.4                        |                            |
|   |                      |                                | Confirmation of genetic modification (if applicable)  | 4.4.3; 4.4.4                 |                            |
| Basic characterization  |                      |                                | Experimental details  |                              |                            |
| Describe the assessment of the following including<br>when they were performed relative to the experiments:   | Reference<br>section | Page reported<br>in manuscript | Describe the following:   | Reference<br>section         | Page report                |
| Authentication  | 1.3; Appendix 1      |                                | Information regarding the experimental unit or sample type for each experiment (e.g. individuals, cell lines,   | 4.4.4; 5.4.2                 |                            |
| Mycoplasma  | 1.6; Appendix 1      |                                | clones, tissues, organoids, devices, batches, cells, etc.)  |                              |                            |
| Sterility (bacteriostasis/fungistasis)  | 1.6; Appendix 3      |                                | Number of replicates (biological/technical)   | 4.2.2; 5.4.2                 |                            |
| Genomic characterization  |                      |                                | Data practices  |                              |                            |
| Describe the genomic characterization including:  | Reference<br>section | Page reported<br>in manuscript | Information on:   | Reference<br>section         | Page report<br>in manuscri |
| Methodology used including sufficient detail to allow an  | 3.1; 5.3;            |                                | Statistical methods used  | 4.4.1; 5.4.2                 |                            |
| assessment of sensitivity (e.g. the number of cells<br>analyzed/resolution/depth of analysis)   | Appendix 5           |                                | Inclusion of the data and annotation code/software used for<br>phenotype classification for computationally derived<br>classifiers (if applicable)          | 5.4.4                        |                            |
| Timing of analysis in relation to key experiments reported  | 3.2                  |                                | Verification that FAIR (https://www.go-fair.org/fair-principles)<br>and CARE (https://www.gida-global.org/care) data<br>management principles were followed | 5.4.4                        |                            |

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STEMCELL Technologies, a company of Scientists Helping Scientists, is a passionate advocate for standardizing human pluripotent stem cell (hPSC) data reporting and quality control measures, limiting experimental variability, and ensuring that relevant and reproducible findings are shared.

The ISSCR's Standards for Human Stem Cell Use in Research document and this companion checklist represent critical steps toward these goals. By adhering to the reporting practices outlined in this wallchart and the principles that underlie them, you will help to improve quality and reproducibility for the field as a whole. Learn more about important cell quality attributes and find out how you can assess and maintain high-quality hPSCs by exploring the resources below.

## **Understanding the Standards**

Learn more about the principles discussed within the standards document and how STEMCELL can help you apply them to your research with this curated collection of resources. From webinars, to interviews, to articles, you will find the information you need to achieve higher quality results and more standardized practices. www.stemcell.com/ISSCR-Standards

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