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Minimum Information about T Regulatory Cells: A Step toward Reproducibility and Standardization

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Cellular therapies with CD4+ T regulatory cells (Tregs) hold promise of efficacious treatment for the variety of autoimmune and allergic diseases as well as posttransplant complications. Nevertheless, current manufacturing of Tregs as a cellular medicinal product varies between different laboratories, which in turn hampers precise comparisons of the results between the studies performed. While the number of clinical trials testing Tregs is already substantial, it seems to be crucial to provide some standardized characteristics of Treg products in order to minimize the problem. We have previously developed reporting guidelines called minimum information about tolerogenic antigen-presenting cells, which allows the comparison between different preparations of tolerance-inducing antigen-presenting cells. Having this experience, here we describe another minimum information about Tregs (MITREG). It is important to note that MITREG does not dictate how investigators should generate or characterize Tregs, but it does require investigators to report their Treg data in a consistent and transparent manner. We hope this will, therefore, be a useful tool facilitating standardized reporting on the manufacturing of Tregs, either for research purposes or for clinical application. This way MITREG might also be an important step toward more standardized and reproducible testing of the Tregs preparations in clinical applications.

Keywords: minimum information model, T regulatory cells, immunotherapy, good manufacturing practice, cell therapy, immune tolerance

INTRODUCTION

T regulatory cells (Tregs) are dominant cellular compounds of the immune system protecting the body from autoimmune reactions. These cells are also involved in imposing tolerance to alloantigens such as transplanted allogeneic cells and tissues

(1–5). For all these reasons, several Treg-based therapeutics are being tested in clinical trials as a prophylaxis or treatment of autoimmune diseases, graft-versus-host disease after hematopoietic stem cell transplants or rejections after solid organ transplants (6). The list of potential applications in the future is even wider. At the same time, manufacturing of Tregs

for preclinical and clinical experiments varies considerably between different centers, which significantly diminishes possible comparisons between the trials. For this reason, future development of these therapies is hampered as it happens that the available results from different trials are contradictory. The specificity of cellular products makes it difficult to verify the results in huge multicentre trials and therefore better standardization of early-phase trials as well as cellular products themselves might facilitate the progress in this promising branch of medicine.

We propose here a tool for standardization of Tregs studies designed on the basis of so-called minimum information models (MIMs). These models have gained increasing popularity among scientists as they enable the interpretation of reported data, comparison between data from different studies and facilitate experimental reproducibility (7, 8). MIMs provide mechanisms that all laboratories report at least the key facts about their analysis in a clear and consistent manner, allowing a comparison across the whole field. Our consortium has already designed the MIM called minimum information about tolerogenic antigen-presenting cells (MITAP). This is a reporting framework that makes transparent differences and similarities of different tolerogenic antigen-presenting cells (tolAPC) (9). It provides minimum reporting guidelines for the production process of tolAPC used in preclinical and/or clinical studies. We have followed the MITAP experience and designed a MIM for the manufacture of Tregs. We call it minimum information about T regulatory cells (MITREG). MITREG will be a useful resource for investigators reporting their data on the use of *in vitro* expanded natural Tregs or induced Tregs in preclinical models or clinical trials.

METHODS

Setting Up MITREG: Community Building and Initial Analysis

The community was mainly built on the experience of our completed MITAP initiative. For several years now, we have been working together in the field of tolerogenic cellular therapies under the umbrella of the consortium AFACTT (action to focus and accelerate cell-based tolerance-inducing therapies—<http://www.afactt.eu/>). It brings together European scientists and clinicians with the aim of jointly addressing issues related to the translation and clinical application of these new treatments. Having the experience of MITAP, we used this document as a template to describe Treg therapies. For MITREG, we also tried to extend the initiative beyond Europe and invited scientists working on tolerogenic cellular therapies from around the world. This way we ensured a broadly reflective discussion taking into account various opinions and current practices of many laboratories within the discipline.

The work on this MITREG document covered a series of “exercises” that provided some initial data. Like for MITAP, the exercises aimed at gathering “terms” in order to acquire basic vocabulary in use within the community. The first, so-called “sticky-note” exercise performed at several AFACTT meetings

assumed that each participant wrote a term on a sticky-note; these were then collated and clustered on a wall by the whole group, identifying synonyms and related terms. Second, we used the MITAP template to incorporate the collected terms and created an initial version of MITREG. This document underwent several rounds of face-to-face and online consultations with AFACTT members to improve its clarity. Internally agreed version was circulated to external specialists in the field. This external feedback was collected and implemented in the final version of the MITREG document. Finally, we used the existing literature to obtain a picture of how well the required information has been described in published articles.

RESULTS

Overview of the MITREG Document

The design of the MITREG document followed the concept of MITAP, which facilitated the whole process. It describes the manufacturing of Treg products in a chronological way. The document is divided into four sections highlighting critical points of the process and regulatory issues. The document describes the details that should be provided by investigators, which would allow other researchers to repeat the process. It also advises on the use of existing taxonomies and databases to provide the information in a uniform manner, and it suggests the use of other MIMs where appropriate. The full MITREG document can be found on archive.org (<http://w3id.org/ontolink/mitreg>) and it is also included in the Appendix A (MITREG document).

Section 1: Cells at the Start of the Procedure

This section describes characteristics of the biological material *before* it undergoes any manipulation. There are five subparts asking for (a) essential information about the donor, (b) source of the cells, (c) the methods used to separate Tregs, (d) the phenotype after separation, and (e) the number of Tregs after separation.

Section 2: Expansion/Differentiation

This section describes the protocol that has been used to expand or differentiate Tregs. The specificity of Tregs was a challenge here as different subsets can be obtained with a wide range of methods. Tregs can be either isolated and optionally expanded or can be induced from naive precursors. There are five subsections giving details on (a) preculture conditions, (b) culture conditions, (c) the protocol used to expand or differentiate cultured Tregs, (d) stimuli used during the process, and (e) the way Tregs are stored immediately after expansion/differentiation.

Section 3: Cells after Expansion/Differentiation

This section describes the characteristics of Tregs *after* the expansion or differentiation. It is mainly focused on the phenotype of the final Treg product as well as its suppressive activity verified in any form of functional assay. It also documents the cell yield from the entire process and, if the product is for clinical use or testing

of adoptive transfer in animals, the details on administration of the cells to the recipient.

Section 4: About the Protocol

This final section describes remaining details of the experimental or clinical protocol such as primary or secondary goals as well as regulatory issues such as adherence to particular acts or directives including compliance with good practice requirements (GCP, GLP, or GMP guidelines). Finally, the name and contact details of the corresponding author(s) must be provided.

The MITREG document is accompanied by a handy checklist to assist investigators in ensuring that all the relevant detail is provided before submitting their manuscripts for publication. The checklist can be found at archive.org (<http://w3id.org/ontolink/mitreg>) and is also included in the Appendix B (MITREG checklist).

Prevalence of MITREG Data in Extant Published Articles

The purpose of the MITREG document is to ensure that authors provide sufficient basic information about their production protocol. An implicit assumption is that currently some or all of this information is not being routinely described. To test this assumption, we reviewed a number of articles about Treg products and for each we determined whether it included data described in the MITREG document.

In detail, 19 Treg articles were selected (predominantly from members of AFACTT or from researchers well known in the field) and read in detail. The articles are given chronologically in the references but the order in **Figure 1** is different and anonymized (10–28). For each section of MITREG, we determined whether the information required was directly stated in the article (or referenced) (**Figure 1**: green squares), partly stated in the article (**Figure 1**: yellow triangles), not present at all (**Figure 1**: red circles), or whether information was not present due to lack of relevance for the publication (**Figure 1**: gray circles). For example, section 1-ai of MITREG describes the species used in the experimental setup. An article with the phrase “human” or “*Homo sapiens*” would fall into the first category (*included in the publication*). However, when mice are used and only the species is mentioned: “mouse” or “*Mus musculus*,” but not the strain, it would fall into the second category (*included but details missing*). Many articles do not describe their experimental methodology, but instead refer to another article (“as described previously”); in this case, we checked the article up to two references deep and if found, the information was considered as “present” (**Figure 1**: green squares), if not it was considered as “not present” (**Figure 1**: red circles). This work was performed by four independent scientists with experience in the field.

Results are shown in **Figure 1**. This figure shows that in some sections like the species, characteristics, ethics, and cell dose transferred sections, reporting is good with almost all revised articles describing these. However, other sections are often very poorly reported. For example, storage of cells, anticoagulant used and the number/viability of cells after each separate step are not

described in most articles. Moreover, important information (container type, concentration of cells) to repeat the performed experiments is missing in almost all articles.

Sustainability

We have taken particular care to consider the issues of digital sustainability for MITREG. A well-known problem with resources linked with URLs given in articles is that URLs are often lost over time: around a 25% loss 3 years after publication (29). We have, therefore, specifically addressed this issue by use of a stable identifier space; the MITREG document and checklist are hosted by archive.org, an organization committed to long-term digital preservation. In addition, we have used a permanent identifier (<http://w3id.org/ontolink/mitreg>) thereby providing a redirection-step.

Resources are available in a number of formats: both PDF and Word for manipulability, but also a simple HTML representation, ensuring vendor-neutrality and future-proofing, in so far as this is possible.

DISCUSSION

Minimum information models aid investigators by providing a specific guideline of what is required to interpret and compare experimental findings. Furthermore, reporting guidelines will facilitate independent validation of published results, a fundamental precept of scientific research. This is to our knowledge the first proposal of a minimum information standard on the description of experimental as well as clinical manufacturing and application of Tregs. The generation of MITREG was initiated by members of the European AFACTT consortium to fill a recognized gap in data reporting standards in the Treg community. MITREG was realized with the help of key international players in the Treg field.

Nine years after the first-in-man report, there are currently close to 30 recruiting or ongoing clinical trials administering Tregs in autoimmune settings, inflammatory diseases, transplantation and graft-versus-host disease (6). Clinical grade reagents for Treg isolation by magnetic activated cell sorting have become available to the growing community and off the shelf products and GMP-compatible fluorescence-based cell sorting is currently being available from multiple manufacturers of novel closed system devices, further increasing the diversity of isolation techniques (30). Given the low frequency of Tregs in the periphery, most clinical applications require an *in vitro* cell expansion culturing step classifying them as advanced therapy medicinal products. A growing number of culturing methods are being developed and published aiming at Treg induction, enhanced *ex vivo* expansion, alloreactivity and more recently, the implementation of specific T cell receptors or chimeric antigen receptors (17, 18, 25, 31–39). We are thus at a point where protocol diversity is growing exponentially, emphasizing the necessity to harmonize reporting regimens as a prerequisite of reproducibility and quality assurance. By analyzing extant articles according to the MITREG document (**Figure 1**), it also becomes clear that there is a big gap in what is currently being reported and what the community considers important and wants to receive in a Treg production/expansion

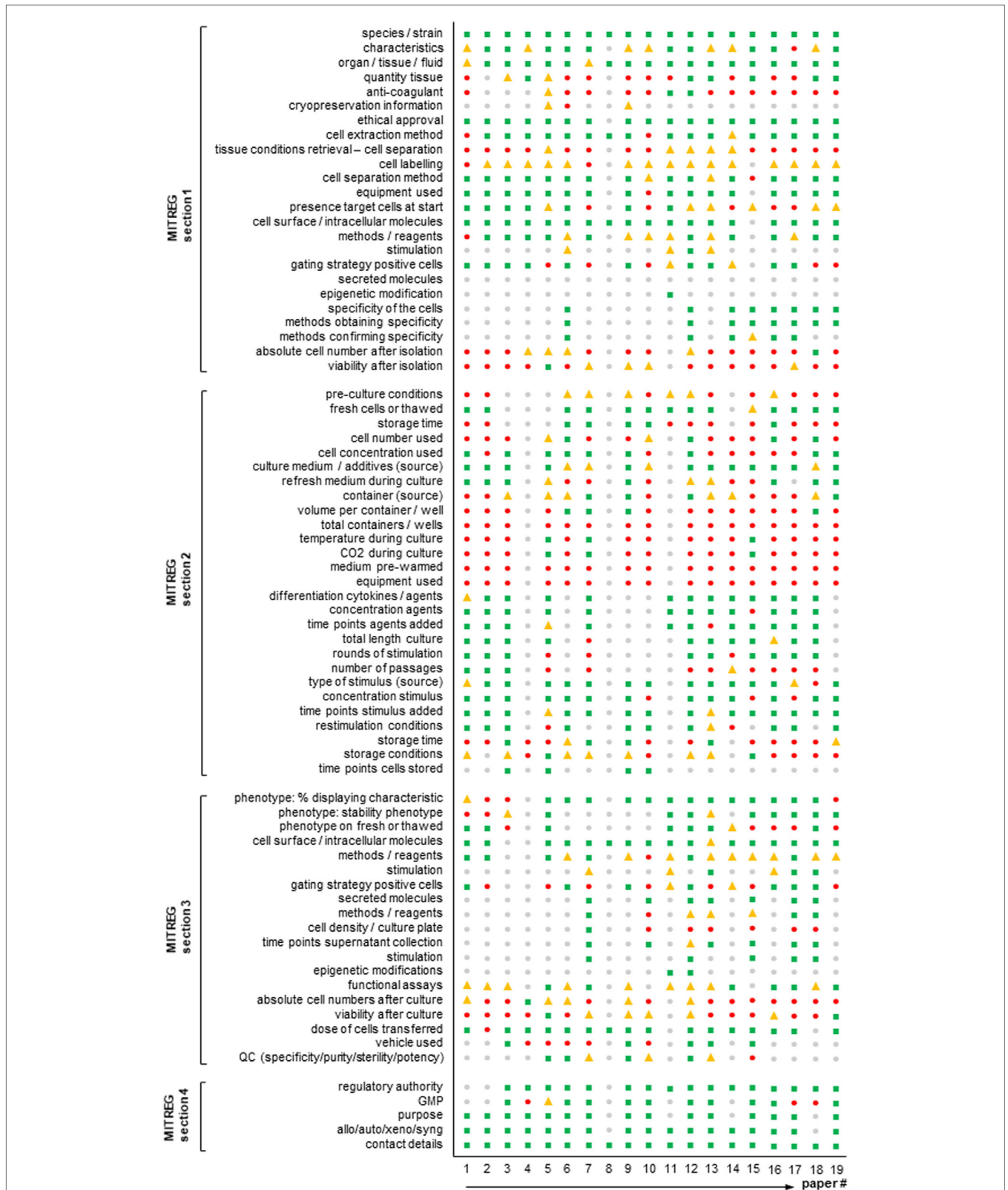


FIGURE 1 | Agreement of published T regulatory cell (Treg) articles with the minimum information about T regulatory cell (MITREG) document. Graph showing the results of a total of 19 Treg articles (10–28). The order in the figure is anonymized and different from that in the references. MITREG data directly stated in the article (■ green squares), partly stated in the article (▲ yellow triangles), not present at all (● red circles), or not present as it was not relevant for the publication (● gray circles).

protocol. For example, storage conditions, cell numbers and viability and anticoagulant used are almost never reported, but are most likely measured or known by the researcher. Moreover, essential information to allow experiments to be repeated is often missing.

Together with MITREG we provide a checklist that was designed with maximal flexibility to incorporate newly developed methodologies. While MITREG does not aim at uniform protocols or dictating quality checks, it is expected to enable a mere description of the growing diversity in production procedures. We expect it to mature as novel technologies arise and become a consensus guideline within the Treg community. Only by exact reporting we will be able to identify differences in Treg preparations that may help to understand results from clinical studies. We anticipate that MITREG will be a starting point for further joint efforts of the Treg community that will ultimately lead to optimized cellular therapy.

REFERENCES

- Sakaguchi S, Sakaguchi N, Asano M, Itoh M, Toda M. Immunologic self-tolerance maintained by activated T cells expressing IL-2 receptor alpha-chains (CD25). Breakdown of a single mechanism of self-tolerance causes various autoimmune diseases. *J Immunol* (1995) 155(3):1151–64. doi:10.1002/1521-4141(200104)31:4<1247::AID-IMMU1247>3.0.CO;2-M
- Stephens LA, Mottet C, Mason D, Powrie F. Human CD4(+) CD25(+) thymocytes and peripheral T cells have immune suppressive activity in vitro. *Eur J Immunol* (2001) 31(4):1247–54. doi:10.1002/1521-4141(200104)31:4<1247::AID-IMMU1247>3.0.CO;2-M
- Baecher-Allan C, Brown JA, Freeman GJ, Hafler DA. CD4+CD25 high regulatory cells in human peripheral blood. *J Immunol* (2001) 167(3):1245–53. doi:10.4049/jimmunol.167.3.1245
- Sakaguchi S, Vignali DA, Rudensky AY, Niec RE, Waldmann H. The plasticity and stability of regulatory T cells. *Nat Rev Immunol* (2013) 13(6):461–7. doi:10.1038/nri3464
- Abbas AK, Benoist C, Bluestone JA, Campbell DJ, Ghosh S, Hori S, et al. Regulatory T cells: recommendations to simplify the nomenclature. *Nat Immunol* (2013) 14(4):307–8. doi:10.1038/ni.2554
- Gliwiński M, Iwaszkiewicz-Grześ D, Trzonkowski P. Cell-based therapies with T regulatory cells. *BioDrugs* (2017) 31(4):335–47. doi:10.1007/s40259-017-0228-3
- Knudsen TB, Daston GP. Teratology society. MIAME guidelines. *Reprod Toxicol* (2005) 19:263. doi:10.1016/j.reprotox.2004.10.004
- Janetzki S, Britten CM, Kalos M, Levitsky HI, Maecker HT, Melief CJ, et al. “MIATA”-minimal information about T cell assays. *Immunity* (2009) 31:527–8. doi:10.1016/j.immuni.2009.09.007
- Lord P, Spiering R, Aguillon JC, Anderson AE, Appel S, Benitez-Ribas D, et al. Minimum information about tolerogenic antigen-presenting cells (MITAP): a first step towards reproducibility and standardisation of cellular therapies. *PeerJ* (2016) 4:e2300. doi:10.7717/peerj.2300
- Trzonkowski P, Bieniaszewska M, Juścińska J, Dobyszek A, Krzystyniak A, Marek N, et al. First-in-man clinical results of the treatment of patients with graft versus host disease with human ex vivo expanded CD4+CD25+CD127-T regulatory cells. *Clin Immunol* (2009) 133(1):22–6. doi:10.1016/j.clim.2009.06.001
- Di Ianni M, Falzetti F, Carotti A, Terenzi A, Castellino F, Bonifacio E, et al. Tregs prevent GVHD and promote immune reconstitution in HLA-haploidentical transplantation. *Blood* (2011) 117(14):3921–8. doi:10.1182/blood-2010-10-311894
- Brunstein CG, Miller JS, Cao Q, McKenna DH, Hippen KL, Curtsinger J, et al. Infusion of ex vivo expanded T regulatory cells in adults transplanted with umbilical cord blood: safety profile and detection kinetics. *Blood* (2011) 117(3):1061–70. doi:10.1182/blood-2010-07-293795

AUTHOR CONTRIBUTIONS

As described in the Section “Methods,” these recommendations are the common effort of all the authors, who were involved in the design, acquisition, and interpretation of available data on Tregs as well as revised critically and approved final version of the MITREG document. In addition, AF, MG, NG, and RS were involved in collecting and analysis of the data sent by the contributors and SG, CH, GL, PL, EC, and PT supervised the work and edited the article.

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- Edinger M, Hoffmann P. Regulatory T cells in stem cell transplantation: strategies and first clinical experiences. *Curr Opin Immunol* (2011) 23(5):679–84. doi:10.1016/j.coi.2011.06.006
- Desreumaux P, Foussat A, Allez M, Beaugerie L, Hébuterne X, Bouhnik Y, et al. Safety and efficacy of antigen-specific regulatory T-cell therapy for patients with refractory Crohn's disease. *Gastroenterology* (2012) 143(5):e1–2. doi:10.1053/j.gastro.2012.07.116
- Wu DC, Hester J, Nadig SN, Zhang W, Trzonkowski P, Gray D, et al. Ex vivo expanded human regulatory T cells can prolong survival of a human islet allograft in a humanized mouse model. *Transplantation* (2013) 96(8):707–16. doi:10.1097/TP.0b013e31829fa271
- Brunstein CG, Blazar BR, Miller JS, Cao Q, Hippen KL, McKenna DH, et al. Adoptive transfer of umbilical cord blood-derived regulatory T cells and early viral reactivation. *Biol Blood Marrow Transplant* (2013) 19(8):1271–3. doi:10.1016/j.bbmt.2013.06.004
- Putnam AL, Safinia N, Medvec A, Laszkowska M, Wray M, Mintz MA, et al. Clinical grade manufacturing of human alloantigen-reactive regulatory T cells for use in transplantation. *Am J Transplant* (2013) 13(11):3010–20. doi:10.1111/ajt.12433
- Landwehr-Kenzel S, Issa F, Luu SH, Schmück M, Lei H, Zobel A, et al. Novel GMP-compatible protocol employing an allogeneic B cell bank for clonal expansion of allospecific natural regulatory T cells. *Am J Transplant* (2014) 14(3):594–606. doi:10.1111/ajt.12629
- Martelli MF, Di Ianni M, Ruggeri L, Falzetti F, Carotti A, Terenzi A. HLA-haploidentical transplantation with regulatory and conventional T-cell adoptive immunotherapy prevents acute leukemia relapse. *Blood* (2014) 124(4):638–44. doi:10.1182/blood-2014-03-564401
- Bacchetta R, Lucarelli B, Sartirana C, Gregori S, Lupo Stanghellini MT, Miquieu P, et al. Immunological outcome in haploidentical-HSC transplanted patients treated with IL-10-nergized donor T cells. *Front Immunol* (2014) 5:16. doi:10.3389/fimmu.2014.00016
- Bluestone JA, Buckner JH, Fitch M, Gitelman SE, Gupta S, Hellerstein MK, et al. Type 1 diabetes immunotherapy using polyclonal regulatory T cells. *Science Transl Med* (2015) 7(315):315ra189. doi:10.1126/scitranslmed.aad4134
- Theil A, Tuve S, Oelschlägel U, Maiwald A, Döhler D, Oßmann D, et al. Adoptive transfer of allogeneic regulatory T cells into patients with chronic graft-versus-host disease. *Cytotherapy* (2015) 17(4):473–86. doi:10.1016/j.jcyt.2014.11.005
- Brunstein CG, Miller JS, McKenna DH, Hippen KL, DeFor TE, Sumstad D, et al. Umbilical cord blood-derived T regulatory cells to prevent GVHD: kinetics, toxicity profile, and clinical effect. *Blood* (2016) 127(8):1044–51. doi:10.1182/blood-2015-06-653667
- Marek-Trzonkowska N, Myśliwiec M, Iwaszkiewicz-Grześ D, Gliwiński M, Derkowska I, Żalińska M, et al. Factors affecting long-term efficacy

- of T regulatory cell-based therapy in type 1 diabetes. *J Transl Med* (2016) 14(1):332. doi:10.1186/s12967-016-1090-7
25. MacDonald KG, Hoeppli RE, Huang Q, Gillies J, Luciani DS, Orban PC, et al. Alloantigen-specific regulatory T cells generated with a chimeric antigen receptor. *J Clin Invest* (2016) 126(4):1413–24. doi:10.1172/JCI82771
 26. Duran-Struuck R, Sondermeijer HP, Bühler L, Alonso-Guallart P, Zitsman J, Kato Y, et al. Effect of ex vivo-expanded recipient regulatory T cells on hematopoietic chimerism and kidney allograft tolerance across MHC barriers in cynomolgus macaques. *Transplantation* (2017) 101(2):274–83. doi:10.1097/TP.0000000000001559
 27. Wolf D, Barreras H, Bader CS, Copsel S, Lightbourn CO, Pfeiffer BJ, et al. Marked in vivo donor regulatory T cell expansion via interleukin-2 and TLLA-Ig stimulation ameliorates graft-versus-host disease but preserves graft-versus-leukemia in recipients after hematopoietic stem cell transplantation. *Biol Blood Marrow Transplant* (2017) 23(5):757–66. doi:10.1016/j.bbmt.2017.02.013
 28. Theil A, Wilhelm C, Kuhn M, Petzold A, Tuve S, Oelschlägel U, et al. T cell receptor repertoires after adoptive transfer of expanded allogeneic regulatory T cells. *Clin Exp Immunol* (2017) 187(2):316–24. doi:10.1111/cei.12887
 29. Wren JD. URL decay in MEDLINE—a 4-year follow-up study. *Bioinformatics* (2008) 24(11):1381–5. doi:10.1093/bioinformatics/btn127
 30. Trzonkowski P, Bacchetta R, Battaglia M, Berglund D, Bohnenkamp HR, ten Brinke A, et al. Hurdles in therapy with regulatory T cells. *Sci Transl Med* (2015) 7(304):304s18. doi:10.1126/scitranslmed.aaa7721
 31. Hoffmann P, Eder R, Kunz-Schughart LA, Andreesen R, Edinger M. Large-scale in vitro expansion of polyclonal human CD4(+)CD25 high regulatory T cells. *Blood* (2004) 104(3):895–903. doi:10.1182/blood-2004-01-0086
 32. Brusko TM, Koya RC, Zhu S, Lee MR, Putnam AL, McClymont SA, et al. Human antigen-specific regulatory T cells generated by T cell receptor gene transfer. *PLoS One* (2010) 5(7):e11726. doi:10.1371/journal.pone.0011726
 33. Marek N, Bieniaszewska M, Krzystyniak A, Juścińska J, Myśliwska J, Witkowski P, et al. The time is crucial for ex vivo expansion of T regulatory cells for therapy. *Cell Transplant* (2011) 20(11–12):1747–58. doi:10.3727/096368911X566217
 34. Bin Dhuban K, d’Hennezel E, Nashi E, Bar-Or A, Rieder S, Shevach EM, et al. Coexpression of TIGIT and FCRL3 identifies Helios+ human memory regulatory T cells. *J Immunol* (2015) 194(8):3687–96. doi:10.4049/jimmunol.1401803
 35. Boardman DA, Philippeos C, Fruhwirth GO, Ibrahim MA, Hannen RF, Cooper D, et al. Expression of a chimeric antigen receptor specific for donor HLA class I enhances the potency of human regulatory T cells in preventing human skin transplant rejection. *Am J Transplant* (2017) 17(4):931–43. doi:10.1111/ajt.14185
 36. Noyan F, Zimmermann K, Hardtke-Wolenski M, Knoefel A, Schulde E, Geffers R, et al. Prevention of allograft rejection by use of regulatory T cells with an MHC-specific chimeric antigen receptor. *Am J Transplant* (2017) 17(4):917–30. doi:10.1111/ajt.14175
 37. McKenna DH Jr, Sumstad D, Kadidlo DM, Batdorf B, Lord CJ, Merkel SC, et al. Optimization of cGMP purification and expansion of umbilical cord blood-derived T-regulatory cells in support of first-in-human clinical trials. *Cytotherapy* (2017) 19(2):250–62. doi:10.1016/j.jcyt.2016.10.011
 38. Chandran S, Tang Q, Sarwal M, Laszik ZG, Putnam AL, Lee K, et al. Polyclonal regulatory T cell therapy for control of inflammation in kidney transplants. *Am J Transplant* (2017) 17(11):2945–54. doi:10.1111/ajt.14415
 39. Marek-Trzonkowska N, Piekarska K, Filipowicz N, Piotrowski A, Gućwa M, Vogt K, et al. Mild hypothermia provides Treg stability. *Sci Rep* (2017) 7(1):11915. doi:10.1038/s41598-017-10151-1

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The reviewer LH declared a shared affiliation, with no collaboration, with several of the authors RS, JI, CH, and PL to the handling editor.

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APPENDIX A

MINIMUM INFORMATION FOR T REGULATORY CELLS (MITREG)

Introduction

The purpose of this document is to enable the description of the generation of T regulatory cell (Treg) products for therapeutic application or experimental usage. It was designed to suit reports using endogenous, induced, antigen-specific, and polyclonal freshly isolated and expanded Tregs.

This document is split into four sections, each describing a different aspect of the process. Not all sections will be relevant to all Treg products.

Information in some sections of this document may be covered by other Minimum Information documents, or defined vocabularies. For example, flow cytometry is described in MIFlowCyt,¹ microarray data by MIAME,² T-cell assays by MIATA,³ and production of standardized tolerogenic antigen-presenting cells by MITAP;⁴ Authors are encouraged to use these resources as appropriate.

Use of Terminology

The key words “**must**,” “**should**,” and “**may**” in this document are to be interpreted as follows:

must: this word means that the information is an absolute requirement. Failure to provide this information is in strict violation of the specification.

EXAMPLE: the species and the source of the cell material are required for all experiments.

should: this word means that there may exist valid reasons for particular protocols to not provide these data, but that these data need to be provided if it is relevant to the protocol.

EXAMPLE: if the Tregs were generated or enriched using an antigen then this must be described, although there may be protocols where polyclonal Tregs are applied.

may: this word means that the data are optional and do not need to be included, but can be provided.

EXAMPLE: the health or age of the organism can be provided, but there may be protocols where this is not assessed, even though it could be.

These definitions are modified from RFC 2119 (<https://tools.ietf.org/html/rfc2119>).

¹<http://flowcyt.sourceforge.net/miflowcyt/>

²<http://fged.org/projects/miame/>

³<http://miatapproject.org>

⁴<https://doi.org/10.7717/peerj.2300>

(1) Cells at the start of procedure

This section describes the characteristics and state of the cells used in the procedure prior to any form of cell manipulation processes such as cell expansion and/or differentiation.

(a) Essential information about the donor

(i) Species and strain

The taxonomy of the organism from which the cells originated. You **must** use names according to the NCBI Taxonomy.⁵ If the strain of the species is known, you **should** indicate this.

EXAMPLE: Homo sapiens/human; Mus musculus, Rag^{-/-}γ_c⁻ (B6, H-2b)

(ii) Characteristics of the organism

Include information about the organism from which the cells originated that is not adequately described by the species/strain information. This **may** include details of their health, age, sex, or any treatments or environmental conditions to which they have been exposed to (e.g., medication). You **may** also include information that is specific to your laboratory, such as an individual identifier number. If you have purchased experimental animals (e.g., BALB/c mice) or tissues (e.g., human bone marrow) you **should** indicate the source of purchase.

EXAMPLE: healthy/volunteer/male/6-weeks-old/male/BALB/c mice/purchased from Charles River (Margate England)

(b) Source of cell material

The organ, tissue, or fluid from which the cells have been isolated **must** be stated. If you use a blood product you **should** state the product and the source (e.g., hospital department, blood bank) from where it was obtained. You **should** use terminology from Uberon,⁶ or the Foundational Model of Anatomy.⁷ You **should** also indicate the quantity of the sample by mass or volume, and, if applicable, which anti-coagulant was used. Additional details **must** be included if the source material was derived from cryopreserved samples (e.g., umbilical cord blood). This would include the methods and duration of storage and initial cell counts. The statement on use/ethics committee approval/written informed consent **MUST** be included.

EXAMPLE: apheresis/buffy coat/bone marrow aspirate/peripheral blood, Sanquin blood supply; 250 ml; EDTA

(c) Cell separation process

(i) Cell handling and labeling

The methodology used to extract the cells from the source material **must** be stated. You **should** also indicate the time between cell material retrieval and start of the isolation process. You **should** indicate how the tissue was kept during this time, including the

⁵<http://www.ncbi.nlm.nih.gov/taxonomy/>

⁶<http://www.uberon.org>

⁷<http://fme.biostr.washington.edu/FME>

temperature and you **may** indicate the container and fluid. You **must** indicate cell labeling procedures, including characteristics and source of labeling buffers and reagents. Other details, such as cell suspension volume and concentration, incubation temperature and washing steps **should** be included.

EXAMPLE: apheresis products were stored overnight at 4°C; Tregs were enriched by magnetic-activated cell sorting (MACS® Technology); Cells were labeled with anti-CD8-coated magnetic beads (CliniMACS® CD8 Reagent, Miltenyi Biotec) in 95 ml of PBS containing 1 mmol/l EDTA and 0.5% human albumin (PBS/EDTA buffer, Miltenyi Biotec) for 30 min at room temperature on an orbital shaker.

(ii) Cell separation equipment and process

The equipment (e.g., AutoMACS®, CliniMACS®, Aria III™ Fluorescence Activated Cell Sorter) and process used to enrich for the cells of interest **should** be stated. The presence of the target population in the starting material should be described.

EXAMPLE: anti-CD8 bead-labeled cells were resuspended in 100 ml of PBS/EDTA/0.5% HA. CD8⁺ cells were depleted with the use of the 2.1 depletion program on the CliniMACS® Cell Separation Device (Miltenyi Biotec).

(d) Phenotype

Characteristics of the cells that have been isolated **should** be described and how this has been determined. Where only a proportion of cells in the population display a characteristic, you **should** indicate the percentage.

(i) Cell surface and intracellular markers

Identifying molecules that are, or are not, expressed by the cells on their surface or intracellularly is useful. You **should** describe: (1) what you measured, (2) the methodology used for the measurement (including information on reagents; if using mAbs, information on clonotype, conjugate, and manufacturer **must** be provided), (3) whether the cells received a stimulus and for how long before the measurement was carried out, and (4) the method used to set marker or population positivity (e.g., fluorescence minus one method). You **should** use cluster of differentiation (CD) names when available (e.g., use CD62L instead of the alternative name L-selectin)—a full list of regularly updated CD numbers can be found on the website run by the HCDM⁸ (human cell differentiation molecules). Otherwise, you **may** use databases, e.g., Uniprot⁹ for proteins and ChEBI¹⁰ for non-protein organic molecules.

EXAMPLE: FOXP3 (PE-Cy7, clone PCH101, eBioscience) expression was measured directly after cell isolation by intracellular staining using the Foxp3/Transcription Factor Staining Buffer Set from eBioscience. Percentage of CD4⁺CD25^{high}CD127^{-/low}FOXP3⁺lin⁻do ublet⁻ Treg cells was determined by flow cytometry (FACS Canto

II™, Becton Dickinson). After the isolation, 98.0% (median, range 97–99.5%) of the cells presented this phenotype.

(ii) Secreted molecules

Molecules that are, or are not, secreted by the cells are useful to identify. These include cytokines (e.g., IL-10) and other soluble mediators. You **should** describe: (1) what you measured, (2) If using Abs, clone, conjugate and source of all antibodies and reagents used **must** be provided, (3) the methodology used for measurement, (4) cell density/milliliter of medium and plastic ware (e.g., 96 w round/flat bottom), (5) when supernatant was collected for cytokine concentration measurement, and (6) whether the cells received a stimulus and for how long before the measurement was carried out.

EXAMPLE: IFN-γ; ELISA; supernatant after 24 h of unstimulated cell culture.

(iii) Epigenetic modifications

Epigenetic modification relevant to the characteristics **should** be described if determined. Method of detection DNA demethylation **should** be clearly described.

EXAMPLE: the mean percentage of demethylated TSDR of the foxp3 gene in the Treg population was 7% (Epiontis, Berlin, Germany).

(iv) Specificity

Polyclonal or antigen-specific, especially genetic modifications to manipulate specificity **should** be described. You **should** describe: (1) what is the specificity of the cells, (2) the methodology used to obtain the specificity, and (3) the methodology used to confirm the specificity. To describe the specificity of your cells, you should use CD names when available (e.g., use CD19 instead of the alternative name B4)—a full list of regularly updated CD numbers can be found on the website run by the HCDM8 (human cell differentiation molecules). Otherwise, you may use databases, e.g., <http://hla.alleles.org>, for HLA alleles, Uniprot⁹ for proteins and ChEBI¹⁰ for non-protein organic molecules describing the targets for your cells.

EXAMPLE: HLA-A2-specific CAR (A2-CAR) Tregs were generated with lentiviral vectors encoding an HLA-A2-specific CAR by cloning and sequencing the heavy- and light-chain variable regions of the mAb and fusing the resulting scFv to portions of CD8, CD28, and CD3ζ in a second-generation CAR structure. Tetramers made from HLA-A2 were used to confirm the specificity of binding the cells to HLA-A2.

(e) Cell numbers

(i) Absolute cell number

You **should** indicate the total number of cells present after extraction, and how they have been counted.

EXAMPLE: 980 × 10⁶ cells as determined by Coulter counting.

(ii) Viability

You **should** indicate the percentage of cells that are alive, and how this has been determined. The percentage of apoptotic cells

⁸<http://www.hcdm.org/>

⁹<http://www.uniprot.org/>

¹⁰<https://www.ebi.ac.uk/chebi/>

should be stated if determined (indicate whether the starting material is fresh or frozen).

EXAMPLE: 95% viability as determined by trypan blue exclusion. 5% of CD3⁺ T-cells had a phenotype indicating early apoptosis (7-AAD⁻, AnnexinV⁺) as measured by flow cytometry.

(2) Expansion/differentiation

The section describes the protocol that has been used for expansion/differentiation of the isolated cells described in the previous section (Section 1). This process will hereafter be referred to as the expansion/differentiation process.

(a) Pre-culture conditions

The conditions under which the cells are kept after isolation but before starting the expansion/differentiation process (the fluid and type of container they are kept in, and at what temperature) **should** be described. The indication whether the starting material is fresh or thawed **must** be provided. You **should** also indicate the length of time between cell extraction and start of the expansion/differentiation process.

EXAMPLE: isolated cells were placed in PBS with 1% human serum albumin in a Falcon tube and kept at room temperature for up to 30 min before starting the culture.

(b) Culture conditions

The conditions under which the cells are kept during the expansion/differentiation process **should** be stated.

(i) Cell number

The number of cells used for the expansion/differentiation process **should** be stated, if different from numbers stated in Section 1*e*.

EXAMPLE: in total 5×10^6 cells were put into culture

(ii) Cell concentration

The concentration of cells in the medium at the start of and throughout the expansion/differentiation process **should** be stated as cells/milliliter.

EXAMPLE: cells were put into culture at a concentration of 1×10^6 cells/ml

(iii) Culture medium

The medium in which the cells are grown **must** be described, including its source, and whether it has any additives (e.g., antibiotics, inactivated serum), excluding the stimuli that are described later. If you use more than one type of medium, or refresh the medium during the culture, then you **should** describe that here.

EXAMPLE: X-VIVO15 (Lonza) supplemented with 5% human male type AB-serum (Sigma)

(iv) Culture container

The physical container in which the culture is carried out. This can include tissue culture plates, tissue culture bags or flasks. You **should** state the type of container, size and manufacturer. You

should also indicate the total cell culture volume per container or well, as well as the total number of containers used.

EXAMPLE: 20 ml of medium in a 100 ml MACS Good Manufacturing Practice (GMP) Cell Differentiation bag (Miltenyi Biotec); 1 bag

(v) Culture environment

Describe the physical environment in which the cells are kept during the expansion/differentiation process. This **should** include the temperature and CO₂ concentration. You **should** note whether medium has been pre-warmed. You **may** describe the equipment used to maintain the culture environment.

EXAMPLE: 37°C, 5% CO₂; Medium was pre-warmed to 37°C; Sanyo CO₂ incubator

(c) Expansion/Differentiation protocol

The protocol that is used to expand/differentiate the cells **should** be described. This **must** include the type and source of cytokine(s) or other agent(s) added into the medium, and at what time point and concentration **should** be included. You **should** also state the total length of the culture period as well as the rounds of stimulation, rounds of culture change, and the number of cell passages.

EXAMPLE: rapamycin (final concentration of 100 nM; Rapamune[®], Pfizer) was added on day 0, 2, 5, 7, and 9. IL-2 (final concentration of 500 IU/ml; Proleukin[®], Novartis) was added on day 2, 5, 7, and 9. Cells were harvested on day 12.

(d) Stimulus

It **should** be stated whether the cells are expanded/differentiated polyclonally or in an antigen-specific manner or against an alloantigen. The protein(s), antibody(ies), accessory cells or other preparation(s) (e.g., antigen-presenting cells; APCs) with which the cells are stimulated **must** be named. You **must** describe the source of the preparation, concentration, and time point(s) at which it/they are added to the cell culture. Restimulation conditions, if any, should also be stated.

EXAMPLE: cells were stimulated with CD3/CD28 MACS GMP ExpAct Treg Beads (Miltenyi Biotec) at a 4:1 bead:cell ratio. Cells were stimulated with CD40-activated allogeneic B cells (30 Gy-irradiated) at a ratio of 10 B cells per nTreg cell.

(e) Storage

The conditions in which the cells are kept after completion of the expansion/differentiation process, but before being used in any subsequent experimental assay or treatment **should** be described. You **should** indicate the fluid and temperature in/at what the cells are being kept, as well as the length of time. You **should** indicate if cells are being frozen, and give details on the freezing and thawing procedures, including cell recovery and viability after thawing. You **should** also indicate if cells are taken out of their culture environment for any length of time during the expansion/differentiation process (e.g., if cells are frozen

before completion of this process, with the aim to resume it at a later date).

EXAMPLE: cells were kept in PBS 1% human serum albumin (Sigma) in a 50 ml Falcon tube at room temperature for a maximum of 2 h; Cells were frozen in FCS/10% DMSO.

(3) Cells after expansion/differentiation

This section describes the characteristics and state of the cells at the end of the expansion/differentiation process described in the previous section (Section 2).

(a) Phenotype

Characteristics of the cells at the end of their expansion/differentiation, including their specificity and purity (e.g., as% of target cells) **must** be described. Where only a proportion of cells in the population display a characteristic, you **should** indicate the percentage. You **should** report on the stability of the phenotype and how you determined this. It **should** be indicated if the phenotype of the cells post-expansion was determined using fresh viable cells, or rather after a freeze–thaw cycle in a batched analysis.

(i) Cell surface and intracellular markers

A number of phenotypic markers help to define the Treg cellular phenotype and specificity and are associated with distinct expression levels of surface and intracellular proteins. These markers are often characteristic of the transcriptional program of a cellular lineage and provide important information regarding the phenotypic stability and function of resulting cell products. You **should** describe: (1) what you measured, (2) the methodology used for measurement (including information on reagents; if using mAbs, information on clonotype, conjugate and manufacturer) **must** be provided, (3) whether the cells received a stimulus and for how long before the measurement was carried out, and (4) the method used to set marker or population positivity (e.g., fluorescence minus one method). You **should** use CD names when available (e.g., use CD127 instead of the alternative name IL-7R α)—a full list of regularly updated CD numbers can be found on the website run by the HCDM (see footnote 8) (human cell differentiation molecules). Otherwise, you **may** use databases, e.g., <http://hla.alleles.org>, for HLA alleles, Uniprot (see footnote 9) for proteins and ChEBI (see footnote 10) for non-protein organic molecules.

EXAMPLE: intracellular IFN- γ and IL-17 expression was measured by flow cytometry after 4 h incubation with 20 ng/ml PMA and 1 μ g/ml Ionomycin in the presence of 1 μ l/ml GolgiPlugTM using the BD Cytotfix/CytopermTM buffer set.

(ii) Secreted molecules

Indicate molecules that are, or are not, secreted by the cells. These include cytokines (e.g., IL-10) and other soluble mediators. You **should** describe: (1) what you measured, (2) if using mAbs, clone, conjugate, and source of all antibodies and reagents used **must** be provided, (3) the methodology used for the measurement, (4)

cell density/ml of medium and plastic ware (e.g., 96 w round/flat bottom), (5) when supernatant was collected for cytokine concentration measurement, and (6) whether the cells received a stimulus and for how long before the measurement was carried out.

EXAMPLE: soluble IFN- γ , TNF- α , IL-17, and IL-10 were measured in the cell culture supernatant at a cell density of 1×10^6 cells/ml by ELISA according to the manufacturers' instruction.

(iii) Epigenetic modifications

Epigenetic modification relevant to the characteristics **should** be described if determined. Method of detection DNA demethylation **should** be clearly described.

EXAMPLE: the mean percentage of demethylated TSDR of the foxp3 gene in the Treg population was 97% (Epiontis, Berlin, Germany).

(b) Functional assay

You **should** describe any characteristic of the cells that has been measured by a functional assay (type of assays). This could either be the response of the cells to some stimulus or the behavior of other biological entities after exposure to the cells. There should be a clear indication of how the percentage of suppression was calculated (i.e., include formula). Whenever accessory cells such as responder cells are included in the assay, source and phenotype should be described. Behavior such as expression/production of molecules (described in Section 3a) does not need to be included.

EXAMPLE: proliferation-based suppression assay using CFSE labeled autologous CD4⁺CD25⁻ responder cells; IFN- γ based suppression assay

(c) Cell numbers

(i) Absolute cell number

You **must** indicate the total number of cells present at the end of the expansion/differentiation process, and how they have been counted and fold expansion **should** be included.

EXAMPLE: cell numbers were microscopically determined using C-Chip disposable counting chambers from NanoEnTek and fold expansion to day 0 was calculated.

(ii) Viability

You **must** indicate the percentage of cells that are alive and how this has been determined **should** be included.

EXAMPLE: 83% viability as determined by trypan blue exclusion

(d) Dosing

Whenever cells are transferred into an organism, details about dosing **must** be given. For clinical applications, information on the vehicle (solvent/medium) as well as intermediate components (trace amounts possible) **must** be given.

EXAMPLE: a single dose of 1×10^7 total nucleated cells per kilogram of body weight in 50 ml 0.9% NaCl was transfused i.v.

(e) Quality control

If the cells were produced for a clinical trial, you **must** describe release criteria and any methods used to determine sterility, specificity, purity, and quality of the product.

(4) About the protocol

In this section, we describe the general features about the protocol as a whole.

(a) Regulatory authority

Information about whether the protocol being used has been validated or quality-controlled to standards agreed to by an external regulatory authority **must** be stated. You **should** state the name of this authority. Also you **should** state whether the protocol follows GMP.

EXAMPLE: Medicines and Health Regulatory Authority

(b) Purpose

You **must** describe the overall purpose of the production of the cells.

EXAMPLE: prevention of transplant rejection; Treatment of patients affected by Crohns' disease.

(c) The relationship between the organism of origin of the cells and the target organism

You **must** state if the cell product is autologous/allogeneic/xenogeneic/syngeneic to the recipient.

EXAMPLE: patients receiving allogeneic kidney transplants and autologous Tregs. B6 mice receiving allogeneic (BALB/c xB6) heart transplants and syngeneic (B6) Tregs.

(d) Contact details

You **must** provide the name and contact information of the corresponding author(s).



(e) Citation

You **should** add information that your paper was written in accordance with the Minimum Information for T Regulatory Cells reporting guidelines.

APPENDIX B



			Must	Should	May	
(MITREG) Checklist						
Must	Should	May				
(1) Cells at the start of procedure						
(a) Essential information about the donor						
(i) Species and strain						
						Molecules measured [using cluster of differentiation (CD) names]
						Details of reagents used and source (incl. mAb clone, fluorochrome)
						Methodology
						Stimulus and time of stimulation (if applicable)
						Gating strategy to determine positive cells
						(ii) Secreted molecules
						Molecules measured
						Details of reagents used (incl. mAb clone, conjugate) and source
						Methodology
						Cell density/ml of medium and type of tissue culture plate
						Time point of supernatant collection
						Stimulus and time of stimulation (if applicable)
						(iii) Epigenetic modifications
						Epigenetic modification relevant to the characteristics
						(iv) Specificity
						Specificity of the cells (polyclonal or antigen-specific)
						Methodology used to obtain specificity
						Methodology used to confirm specificity
						(e) Cell numbers
						(i) Absolute cell number
						Total number of cells at the end of the isolation process
						Methodology
						(ii) Viability
						Percentage of viable cells
						Methodology
						(2) Expansion/differentiation
						(a) Pre-culture conditions
						Storage conditions
						Fluid
						Type of container
						Temperature
						Fresh or thawed
						Storage time
						(b) Culture conditions
						(i) Cell number
						The total number of cells put into culture
						(ii) Cell concentration
						The number of cells per ml of medium at start of culture
						(iii) Culture medium
						Type(s) of medium
						Source(s)
						(b) Source of cell material
						Organ, tissue, fluid, or blood product
						Source (if applicable)
						Quantity (volume, size, or weight)
						Anti-coagulant (if applicable)
						If using cryopreserved sample
						Method and duration of storage
						Initial cell counts
						Ethical committee approval/written informed consent
						(c) Cell separation process
						(i) Cell handling and labeling
						Cell extraction method
						Tissue conditions between tissue retrieval and cell separation
						Duration
						Temperature
						Container
						Fluid
						Cell labeling
						Buffers and reagents (incl. source)
						Cell suspension volume and concentration
						Incubation temperature and duration
						Washing steps
						(ii) Cell separation equipment and process
						Methodology
						Equipment
						Presence of target cells in starting material described
						(d) Phenotype
						For any of the below, indicate the percentage of cells displaying the characteristic (if known)
						(i) Cell surface and intracellular markers

Must	Should	May		Must	Should	May	
Must			Additives (excluding agents to maintain/induce T regulatory cells)		Should		Stability of the phenotype (if tested)
	Should		Refreshment of the medium		Should		Phenotype tested on fresh or thawed cells
			(iv) Culture container				(i) Cell surface and intracellular markers
	Should		Type of container		Should		Molecules measured (using CD names)
	Should		Size	Must			Details of reagents used and source
	Should		Manufacturer		Should		Methodology
	Should		Cell culture volume per container or well		Should		Stimulus and time of stimulation (if applicable)
	Should		Total number of containers or wells		Should		Gating strategy to determine positive cells
			(v) Culture environment				(ii) Secreted molecules
	Should		Temperature and CO ₂ concentration		Should		Molecules measured
	Should		Use of pre-warmed medium	Must			Details of reagents used and source
		May	Equipment		Should		Methodology
			(c) Differentiation/tolerization protocol				Cell density/milliliter of medium and type of tissue culture plate
Must			Name of cytokine(s) or other agent(s) used				Time point of supernatant collection
	Should		Concentrations				Stimulus and time of stimulation (if applicable)
	Should		Time point(s) added to cell culture				(iii) Epigenetic modifications
	Should		Total length of the culture period		Should		Epigenetic modification relevant to the characteristics
	Should		Rounds of stimulation				(b) Functional assay
	Should		Number of cell splitting		Should		Response of the cells to a defined stimulus
			(d) Stimulus				Behaviour of other biological entities after exposure to the cells
	Should		Polyclonal/antigen-specific/alloantigen				If using accessory cells, describe phenotype and source
Must			Stimulus (agent and/or accessory cell)				(c) Cell numbers
Must			Source				(i) Absolute cell number
	Should		Concentration	Must			Total number of cells at the end of the expansion process
	Should		Time point(s) added to culture		Should		Methodology
	Should		Restimulation conditions (if applicable)				(ii) Viability
			(e) Storage	Must			Percentage of viable cells
	Should		Storage time		Should		Methodology
			Storage conditions				(d) Dosing
	Should		If fresh	Must			Dose of cells transferred into organism (if applicable)
	Should		Fluid	Must			Vehicle (solvent/medium) and intermediate components (for clinical trials only)
	Should		Container				(e) Quality control (for clinical trial only)
	Should		Temperature				Specificity
			If cryopreserved				Purity
	Should		Freezing/thawing process				Sterility
	Should		Freezing medium				Potency
	Should		Cell recovery and viability after thawing				(4) About the protocol
			Time point at which cells are stored if different to the end of the culture process				(a) Regulatory authority
			(3) Cells after expansion/differentiation	Must			External authority that approved the protocol
			(a) Phenotype		Should		Does protocol follow Good Manufacturing Practice?
Must			For any of the below, indicate the percentage of cells displaying the characteristic (if known)				

Must	Should	May
		
		

(b) Purpose
The disorder for which the cell treatment has been manufactured

(c) Relationship between the source organism for the cells and the target organism
Allogeneic/autologous/ xenogeneic/syngeneic

Must	Should	May
		
		

(d) Contact details
Name and contact information of the corresponding author(s)

(e) Citation
Acknowledge the MITREG reporting guidelines