Pluripotent stem cell based medicinal products: A case study of process transfer related technical and manufacturing issues

[Abstract]

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staining and endotoxin were negative in all cases. Validation included the assessment of CD34+ cells stability after a 48h-storage.

**Conclusion**: Herein we demonstrated the establishment of a feasible, consistent and reproducible bioprocess for safe manufacturing of expanded CD34+ cells for clinical use, opening the way for a Phase II/III clinical trial which is currently recruiting.

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**THE PERFECT LOGISTIC TRANSPORT METHOD CONCERNING ALL KIND OF CELL THERAPY PRODUCTS**
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During the last few years different T-cell therapy products like CTLP (CMV-specific T-lymphocyte precursor cells) or CAR (chimeric antigen receptor) T-cells were further developed, it became increasingly important to find the most suitable transport method for starting, intermediate as well as for final products.

During clinical phase II and III studies GMP facilities as well as hospitals all over the world have to be connected. Beside a perfect logistic supply chain the right transport method and transport material is even more essential.

Ahead of a phase III study we have tested different transport methods in terms of safety, reliability and speed.

**On-board transport**
- Cargo transport
- Transport of cryopreserved products
  - First we put an eye on different transport material—we compared different types of LN2 shippers and on-board transport containers. In a second step we shipped mock products to test the transport under real conditions. Finally we performed during a phase III study on-board transports as well as transports of starting material, CD1 enriched and final products.
  - The test results showed clearly that the transport material has to be chosen in accordance to the product that has to be transported to avoid foreseeable product damage.
  - During the mock transports material differences turned up especially concerning the LN2 shippers. It was shown that some shippers are not well protected against shocks, in consequence the overwrap or even the product bag sometimes broke. Material used for on-board transports showed fewer differences.
  - The chosen transport method that is very decisive. Often unattended cargo transports bear the risk that shipments are handled roughly; but it could be proved that shipping starting material unattended within an appropriate cooler works perfectly.

**The right combination of transport material and transport method supplemented by project-related SOPs finally guarantees the safest and fastest result.**

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**THE OPTIMIZED ORGANIZATION OF CELL THERAPY RUNS**
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The development during the last few years shows clearly that T-cell therapy products such as CTLP (CMV-specific T-lymphocyte precursor cells) or CAR (chimeric antigen receptor) T-cells come to the market.

During clinical phase II and III studies different partners such as GMP facilities and hospitals worldwide have to work together to produce the best outcome within the shortest possible time frame. This is only feasible with a trained medical coordination team combined with a process that schedules and controls all steps and is connected 24/7 with all partners.

Ahead of a phase III study we have set up a semi-automatic workflow which is processed according to GMP guidelines combined with a computer program and an Annex 11 validated web based system.

- We put an eye on the essential steps:
  - Confirmation request/dates
  - Scheduling (apheresis site, GMP facility, manufacturing site, clinical site)
  - Organization of necessary transport legs
  - Documentation flow
  - Dunning system
  - Emergency management
  - Controlled compliance with Chain of Identity

We analyzed the workflow concerning efficiency, safety and control and we identified the phases as well as the steps that have to be optimized.

**Conclusion**: A recent process transfer of the human embryonic stem cell line MasterShef10 from the originator site to a manufacturing-focused site emphasized several technical challenges. This was due to the complex nature of the cells and the requirement to further optimise the maintenance methods. An initial analysis to identify the sources of variability in the cell expansion process of different vials from a working cell bank indicated that no substantial variability arose from independently cultured vials. Further it was also determined that no significant variability arise by a single passage when the cells were cultured for 12 passages and characterized at different stages. However a major variable identified were the steps involved with passages such as erratic post-seeding density.
and confluence during harvest. As a consequence of this, a cyclic variability in yield upon subsequent passages has occurred. Further analysis showed that the cell counting devise resulted in counting bias and resulting confluence-dependent cell aggregate size. This counting bias augmented variability in seeding density and quality of post pasaging survival of seeded cell colonies which caused further variability in yield.

This study points to the casual introduction of variability due to bias exists in measurement systems and the requirement for extensive process knowledge and characterization of the process steps to determine the underlying cause of the variation. Consequently currently available cell counting methods need to be improved in order to count cells that present as aggregates. Quantitative measurements of process stages and adequately detailed Standard Operating Procedures (SOP’s) will enable efficient process transfer of cell based medicinal products.

203 VALIDATED TRANSPORTATION SYSTEMS FOR CELL BASED PRODUCTS
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Background: With a growing interest in the development of cell based products, the need for validated transport systems suitable for temperature sensitive products grows. Conventional drug products are often designed to be stable at broad temperature ranges over a longer period of time and have been stress-tested at sub-optimal conditions. Cell based products are very sensitive, so transportation should take place within a short time frame and at strictly defined temperature ranges that are suitable for the kind of cellular product to assure their functionality.

Methods: We tested and validated systems consisting of durable, isolated transport boxes and temperature shells optimized for different temperature conditions. Our approach included systems suitable for the transportation at ranges of -160°C, 2–8°C, 18–25°C and 34–39°C.

For temperature ranges of 2–8°C, 18–25°C and 14–19°C, we relied on preconditioned temperature shells. As for frozen cells, their transportation is reportedly often performed using dry ice, reaching temperatures between -80 to -60°C. However, these transport systems are unsuitable for highly sensitive cells and hold a risk for the courier due to CO2 leakage. For this reason, we validated a LN2 devar system, containing the LN2 in a porous matrix, allowing the cells to reside in the vapour phase and minimizing the risk for the courier.

Results: With the preconditioned temperature shells we achieve reproducible stable temperatures for at least 64 hours in the 34–19°C system and 24 hours in the 2–8°C and 18–25°C systems in moderate external temperatures. With extreme external temperatures under 10°C and over 30°C, the capacity of the 2–8°C and 18–25°C systems reduces to at least 12 hours, the 34–19°C System has not been validated for extreme temperature ranges due to extensive courier qualification. The LN2 System is stable for at least 5 days in extreme conditions, which allows the shipment to more remote places.

Conclusions: Transport systems are to be validated considering different outdoor temperatures, reproducible and narrow temperature ranges and practicality for the courier. Highly detailed packaging instructions are to be developed, to meet temperature, reproducible and narrow temperature ranges and practicability for transport even in remote places.

204 QUALITY ASSURANCE OF FLOW CYTOMETRY ASSAYS: RESULTS OF THREE-YEAR SEMI-ANNUAL IMMUNE MONITORING INTER-SITE COMPARISONS OF THE TCRab-HAPLO2010 TRIAL
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Since cellular, immune and gene therapy are evolving and promising medical fields in research and daily routine, immune monitoring has become an essential tool for determining treatment-induced effects on immune cells. Flow cytometry offers the opportunity to monitor immunological parameters in clinical trials on low amounts of cellular material (e.g. whole blood) with a reasonable amount of time. However, reproducibility of flow cytometry-based immune monitoring is a matter of high importance, especially when multiple sites are involved.

Here we present results of the semi-annual inter-site comparisons conducted over 3 years as a quality assurance measure of flow cytometry-based immune monitoring for the TCRab-Haplo2010 trial. To assess the reproducibility of the immune monitoring in different flow cytometry core labs, blood from a healthy donor was centrally collected and distributed to the participating labs. Samples were assayed within 25 to 30 hours after blood collection by four operators using four different MACSQuant Analyzer devices at three different sites.

Immune cell subsets, which are relevant to evaluate immune reconstitution after haploidentical hematopoietic stem cell transplantation with TCRαβ-CD19 depleted stem cell grafts (T cells, T helper cells, cytotoxic T cells, B cells, NK cells, TCRαβ- and TCRγδ- T cells) showed an average coefficient of variation (CV) of 6.28 % (±3.89 %) for percentage of subsets among CD45 cells of 9.13 % (±5.48 %) for cells/jul (two platform approach). These inter-site comparisons over a period of 3 years demonstrated low variability of immune monitoring results and thereby confirm reliability and comparability of patient immune monitoring data obtained in the TCRab-Haplo2010 flow cytometry core labs.

205 RAPID AND ROBUST LARGE-SCALE EXPANSION OF HIGH QUALITY HUMAN MESENCHYMAL STEM CELLS FOR USE IN REGENERATIVE MEDICINE APPLICATIONS
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Human mesenchymal stem cells (hMSC) isolated from various tissues represent the most highly utilized cell type for cell therapy and regenerative medicine applications. However, a major impediment to progress in the field is availability of sufficient numbers of high quality hMSC for process development studies. Given that large numbers of high quality cells are required for many applications and that large surface area requirement for cell expansion is necessary, a search for a practical manufacturing platform for hMSC expansion has been ongoing for the last decade. Feasibility studies have established the utility of using microcarriers in dynamic cultures for expansion of hMSC and the results have positioned this technology as the platform of choice for the cost-effective generation of hMSC. Microcarrier cultures performed in fully closed single-use systems provide a platform for cell expansion and are a logical extension of traditional manufacturing technologies. We have developed a rapid and robust manufacturing platform that enables reproducible expansion of large numbers of high quality hMSC from fed-batch microcarrier cultures without the need for media exchanges or extensive process development. Studies with bone marrow-derived hMSC using sterile microcarriers in single-use bioreactors reproducibly generated over twenty-fold expansion in 6 L PadReactor® Mini bioreactors in 4 days. Cryopreserved Allergro™ Unison hMSC seeded directly onto SoloHill microcarriers in the PadReactor Mini bioreactor followed by transfer of cells to a larger volume PadReactor system showed >200-fold expansion in only 8 days. Cells expanded using this platform and processes maintain critical quality and functional attributes. This single-use system along with the processes developed in these studies is ready for use by others to generate billions of cells without the need for further development.

206 SIMPLE AND EFFICIENT CELL HARVEST METHODS FOR MICROcarrier CULTURES IN BIOreactors
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The utilization of cell therapies and regenerative medicine applications is dependent upon development of cost-effective, robust and reproducible technologies for generation of high quality cells. Microcarriers employed in single-use bioreactors provide a robust and reproducible system for cell expansion; however, efficient use of microcarriers for this purpose requires a scalable cell harvest process. Pall SoloHill microcarriers are rigid spheres that support rapid and efficient cell attachment and subsequent cell growth in bioreactors. These physical characteristics also allow users to employ standard cell culture harvest methods for detachment and separation of cells from microcarriers but techniques used