The translation of cell-based therapies: clinical landscape and manufacturing challenges

Cell-based therapies have the potential to make a large contribution toward currently unmet patient need and thus effective manufacture of these products is essential. Many challenges must be overcome before this can become a reality and a better definition of the manufacturing requirements for cell-based products must be obtained. The aim of this study is to inform industry and academia of current cell-based therapy clinical development and to identify gaps in their manufacturing requirements. A total of 1342 active cell-based therapy clinical trials have been identified and characterized based on cell type, target indication and trial phase. Multiple technologies have been assessed for the manufacture of these cell types in order to facilitate product translation and future process development.

Keywords: bioreactor • cell-based therapy • clinical trials • manufacture • MSC • regenerative medicine • stem cell • translation

Regenerative medicine offers a transformative approach to healthcare, with the potential to not only treat, but cure disease. The therapeutic application of cells, or cell-based therapy, forms a fundamental part of this landscape and already has global revenues of over US$1 billion [1]. With an ever-increasing number of cell-based therapies in the pipeline this industry, if properly developed, has the potential to provide significant health and wealth benefits throughout the world.

The use of live cell-based therapies in medicine is not a new concept. The first successful allogeneic human hematopoietic stem cell transplant took place in 1968 [2] and is now a routine clinical procedure for bone marrow regeneration. The true value of stem cell-based therapies was not explored until some two decades later when the therapeutically relevant cells were considered for the regeneration of skeletal tissue [3] and later for broader therapeutic utility [4]. This led to the commercial venture into cellular therapy by Osiris Therapeutics (NASDAQ:OSIR), who have since received market approval in Canada and New Zealand for Prochymal™, a mesenchymal stem cell (MSC) therapy for juvenile graft versus host disease.

Since the turn of the millennium there has been a steady increase in the number of cell-based therapy clinical trials, with an increasing spread of phases and number of target indications [5]. This increased diversity of target clinical indications has been largely driven by growing evidence of the trophic effects of infused cells, whereby the putative mechanism of action is by secretion of proteins and paracrine factors rather than terminal engraftment [6]. Alongside adult stem cells, pluripotent stem cells have attracted significant interest for cell-based therapy applications due to their high proliferative potential and ability to form any tissue in the body. Both embryonic stem cell and induced pluripotent stem cell-derived products are currently entering clinical development [7] but face many technical and ethical hurdles particularly around product safety [8].

Clinical activity in cell-based therapies is clearly converging upon a critical mass,
with over 300,000 patients treated with regulatory-approved products since 1997 [9]. This does mean, however, that patient expectations are increasing and must be managed delicately to avoid a second wave of disillusionment in the sector, following the drop in commercial activity in the early 2000s [10]. In conjunction with this, great emphasis must be placed on patient safety by avoiding the exploitation of patients engaged in ‘stem cell tourism’ and the off-label or unlicensed use of stem cell-based therapies [11–13]. This is all part of a wider regulatory balance that on one hand ensures patient safety and on the other hand does not prevent much needed treatments reaching the clinic.

For the majority of clinical indications, the transplantation of primary donor cells may not be sufficient to meet the clinical need and the expansion of cells in culture will be required to address the shortage of therapeutically active cells available in the body. To ensure that these cell-based therapies can meet the needs of the patient, cells must be produced in sufficient numbers, with reproducible quality and at relatively low cost. For this to happen, suitable scalable manufacturing processes must be developed either before or during the clinical assessment phase, with the final optimal process used to produce batches for Phase III clinical studies typically carried forward to commercial production. The aim of these manufacturing processes is to significantly increase cell numbers without negatively affecting the therapeutic potential of the cell, which is known to deteriorate with time in culture [14]. It is possible that the manufacturing lot size required to meet this need will be in the order of trillions of cells, though this is dependent on the dose requirements per patient and market size of each indication [15]. Current manual planar culture technology is not sufficient to meet this requirement and new manufacturing methods must be developed to drive down the cost of goods to produce cost effective therapies and achieve product reimbursement.

There are currently many challenges facing the cell-based therapy industry particularly in the regulated manufacture of these products under GMP [16]. Conventional bioprocesses use cells to produce therapeutic agents, which can then be isolated and purified without the need to recover the cell. For the manufacture of cell-based therapies, retention of cell function and quality is of primary importance in order to preserve product efficacy [17]. This consideration is particularly pertinent, considering that a number of these cell types are adherent, meaning that they must be detached or harvested from a culture substrate, prior to downstream processing. To quantify this, effective manufacture is reliant on measuring these critical-to-quality attributes and establishing the allowable deviation of these product characteristics from a predetermined set point. The use of potency assays can provide a batch release test that validates whether the cell product is consistent, stable and of sufficient quality for therapeutic use, as well as providing a comparability metric to validate process changes [18]. Stringent safety assays must also be considered for cell-based therapies as many current processes use animal-derived products during culture, which could lead to pathogen transfer or immune complications once infused into the recipient [19]. Additionally, the use of pluripotent stem cells brings a new set of safety challenges within downstream purification, ensuring that the unlimited growth potential of these cells is not transferred to the patient [20]. The requirement for high levels of process and product characterization will result in significant direct costs in all process stages, from establishment of a master cell bank (for allogeneic products), to final product testing.

Cell-based therapies are complex biological products that are sensitive to their environment and display intrinsic variability within a tightly regulated industry. Variation in the product can come from two sources: process input material and process conditions [21]. For allogeneic therapies, where cells from one patient can be given to many patients, this input variability can be reduced by careful selection of comparable input material to the process. Alongside this, the variability in process conditions can be managed in part by automating the manufacturing process allowing for increased control and process capability [22–25]. This allogeneic or ‘off-the-shelf’ business model for cell-based therapies is far more akin to current biopharmaceuticals [26], where the product maintains long-term stability.

Conversely, autologous cell-based therapies, where cells are taken from a patient and returned to the same patient, represents a different manufacturing challenge as patient populations must be stratified to control this input variation. Issues surrounding the quality test burden and logistics of personalized (services based) medicine add to the complexity for large scale production and delivery of a cost effective autologous cell-based therapy [27]. Delivery of autologous cell-based products can be complex due to the short term preservation methods used for transport and delivery, for example the shelf life of Provenge, an autologous peripheral blood mononuclear cell-based therapy suspended in Lactated Ringer’s Injection solution is only 18 h at 2–8°C [28]. Several allogeneic therapies in development are also shipped to clinical delivery sites in a noncryopreserved format. This results in the need to maintain a constant level of production, even when clinical demand may vary greatly, as noncryopreserved products typically have a shelf life on the order of
days and significant product wastage can occur. This increases overall product cost and requires larger batch sizes to be produced. Considering the range of potential cell-based therapies it is clear that multiple manufacturing models must be evaluated in order to fill the need of both universal (scale up) and personalized (scale out) therapies and control the associated biological variation to achieve a consistent product.

Given the diversity of cell types, disease indications and delivery pathways it is unlikely that there will be a ‘one-size-fits-all’ manufacturing platform for cell-based therapies and therefore careful consideration must be given to process selection for commercial production during process definition, development and optimization, prior to clinical development. This study provides a snapshot of current cell-based therapy clinical trials by cell type, target clinical indication and trial phase and a comparative assessment of the process technology available to manufacture these cell-based therapies. The aim is twofold, firstly to inform industry and academia of current clinical activity involving their cell type and secondly to identify the challenges associated with the manufacture of these cell-based therapies. This analysis will provide a comparison tool for manufacturing technology and facilitate the development clinical indication specific quality assays by providing an overview of the current target clinical indications for cell-based therapies.

Methodology
To initiate this study, we searched for the term ‘cell’ on the ClinicalTrials.gov database [29] to include all clinical trials involving the use of cells up to January 1, 2014. This produced 29,467 results, which were further sorted on the database to show only open trials reducing the number to 9700 which were extracted for analysis, providing an overview of the current cell-based therapy clinical trials landscape.

As this study is evaluating cell-based therapy clinical trials, the following British Standard Institute definition was used to classify these as a ‘therapy in which cells are administered to the body to benefit the recipient’ [30]. This excluded:

- Trials where the application of cell was not the main therapeutic goal of the study (i.e., studying the effect of a drug infused as part of a cell-based therapy);
- Trials that were observational (i.e., no direct application of cells to the patient);
- Duplicate trials (i.e., the same trial taking place in more than one country).

The 9700 trials were sorted manually one at a time by one individual to identify the true cell-based therapies using the definition above and the listed exclusion criteria. This action left a database of 1342 cell-based therapy clinical trials that were checked to ensure the major industry clinical trials had been encompassed and further categorized as detailed below.

Cell type
Each of the 1342 clinical trials was assigned a cell type based on the clinical trial listing and the cell used for therapeutic benefit but where no cell type was indicated, the cell type was listed as unknown. Where a clinical trial was assessing a combination of more than one cell type, both cell types were included in the study.

Cell group
The cell types in use for the clinical trials have been divided into multiple groups namely, hematopoietic cells (CD34+, mononuclear cells and whole marrow), MSCs, lymphocytes, dendritic cells, hepatocytes, endothelial cells, fibroblasts, myocytes, epithelial cells, islet cells, chondrocytes, renal cells, embryonic stem cells and other cells. The purpose of these groups is to show the spread of cell types in current active cell-based therapy clinical trials.

Clinical indication
Categorized based on clinical speciality and disease mechanism of action or where an indication falls into more than one category, by the greatest need. An example of this is stroke, which is a vascular disease that manifests as a neurological disorder and therefore is categorized in neurology as this is where the cell-based therapy treatment would be targeted.

Trial phase
The clinical trial phase was taken from ClinicalTrials.gov and divided into Phases I, II, III and IV. Where trials were in the transition between phases, only the completed phase was used (i.e., a listing of Phase I/II would be classified as Phase I).

This process yielded a thorough database of categorized cell-based therapy clinical trials that are currently open as of January 1, 2014. Although this provides a comprehensive database of cell-based therapy clinical trials on ClinicalTrials.gov, it does have a limitation. ClinicalTrials.gov was initiated as a result of the US FDA Modernization Act of 1997 as a registry of clinical trial information for federally and privately funded trials conducted under investigational new drug applications (later expanded under FDA Amendments Act of 2007). This means that clinical trials in the USA...
must be registered on ClinicalTrials.gov but clinical trials initiated in other countries are only included on a voluntary basis. Therefore, this method does not include all cell-based therapy clinical trials, just those registered through the National Institutes of Health. Limitations of this database have been previously discussed [10]. The methodology presented here will, however, contain the vast majority of cell-based therapy clinical trials and certainly a large enough sample to draw meaningful conclusions.

Results
Clinical trial landscape for cell-based therapy
Our search of the clinical trial database yielded a total of 1342 active clinical trials that were judged to be true cell-based therapies from the British Standard Institute definition as of January 1, 2014. The vast majority of these cell-based therapies are using hematopoietic cells (n = 444), MSCs (n = 382), lymphocytes (n = 253) and dendritic cells (n = 91) (Figure 1). The use of tissue specific cells is less prevalent, representing 114 of current cell-based therapy activity with a mixture of cell types such as chondrocytes (n = 12), endothelial cells (n = 15), fibroblasts (n = 14), hepatocytes (n = 16), islet cells (n = 13) and neural cells (n = 13). Figure 2 shows the diversity of cell isolation sources represented from hematopoietic and MSC trials. The rapid increase in MSC activity is clear, with 382 clinical trials in progress from a number of isolation sources most notably 237 of which are from the bone marrow, 61 from umbilical cord and 59 from adipose tissue. The use of pluripotent stem cell-derived products is rare, though there are six clinical trials involving embryonic stem cells. Interestingly, all six involve differentiation into retinal pigment epithelium cells and are initiated by industry: CHA Bio & Diostech in Korea (two), Advanced Cell Technology (three) and Pfizer in the UK (in collaboration with University College London).

Disease indications targeted by cell-based therapy clinical trials
It is clear that the most common target for cell-based therapy clinical trials is oncology, which represents 46% of all cell-based therapy clinical trials identified (Figure 1), due to the use of traditional blood cell and immune cell-based therapies for the treatment of various cancers. Aside from these traditional cell-based therapies, cardiology is the second largest clinical target with 88 clinical trials (Figure 1), 51 of which are using MSCs (Figure 3). Immunological disorders remain a key target for cell-based therapies (Figure 4) with disorders such as graft versus host disease (n = 33), immune modulation following transplantation (n = 17) and Crohn’s disease (n = 16), as well as rheumatic disorders osteoarthritis (n = 15), lupus erythematosus (n = 9) and rheumatoid arthritis (n = 6) currently in Phase I.

Figure 1. Number of active clinical trials by cell type and target clinical indication. Displaying broader cell type categories of hematopoietic (blue), mesenchymal stem cells (red), immune cells (green), tissue-specific cells (orange), embryonic stem cells (purple) and other (aqua).
and above. Neurological indications also represent a significant target for novel cell-based therapies for indications such as multiple sclerosis, stroke and motor neuron disease (Figure 4). Figure 3 shows the broad therapeutic utility adopted for MSCs, with clinical indications covering all 18 clinical categories with the majority within cardiovascular, neurological and autoimmune indications predominantly in Phase I. Likewise, tissue specific cells cover the majority of clinical indication categories, owing to the diversity of cell populations within this category.

**Cell-based therapy clinical trial phase**

It is unsurprising that the majority of cell-based therapy clinical trials are in Phase I, with 232 MSC, 146 lymphocyte and 118 hematopoietic cell clinical trials (Figure 5). There is however, an increase in hematopoietic cell clinical trials moving to Phase II with 187 compared with 116 in Phase I. There are also four tissue specific cell-based clinical trials that have reached postmarket surveillance, or Phase IV (Figure 5). Most notable of these are the use of dermal fibroblasts for wound repair from Arita Medical (ReCell™) and Organogenesis (Apligraf™) as well as cell-based therapies to treat Type 2 diabetes and for cornea replacement. Figure 4 shows the major clinical categories by specific disease targets, with the vast majority targeting hematological malignancies within oncology, with equal weighting in Phases I and II. Cardiovascular indications also have a similar number of clinical trials in Phases I and II, with ten clinical trials in Phase III and above. MSC clinical trials are also well represented in Phase II (n = 88) and are even seeing progression to Phase III (Figure 5). Neurology
and rheumatology based indications that involve MSCs are mostly in Phase I (Figure 6), however clinical development using bone marrow-derived MSCs to treat acute coronary syndrome, Crohn’s disease and graft versus host disease are seeing progress into Phase II and above, with clinical trials taking place all over the world.

Discussion
Clinical translation of mesenchymal stem cell-based therapies
It is clear that outside of the traditional cell-based therapies using blood and immune cells to treat cancer, the majority of novel clinical trial activity is involving MSCs. The commercial pathway for MSC therapies was initiated after they were branded in 1991 by Arnold Caplan [3], leading to the generation of intellectual property by Osiris Therapeutics. The rebranding of MSCs has been used to generate intellectual property by defining a specific set of characterization criteria for the cell, for example, Athersys Inc. are developing a product termed MultiStem™, a Multipotent Adult Progenitor Cell therapy which has been patented under this name [32,33].

As the first clinical trials involving MSCs developed, the need for nomenclature and characterization standards led to the formation of the International Society for Cellular Therapy working group in 1992, who eventually published minimal criteria for MSCs in research [34,35]. The broad therapeutic utility currently adopted for MSC clinical indications was a product of work by Osiris Therapeutics [36] and led to a lot of publicity around the potential of stem cell-based therapies. The green shoots of industry that began to emerge in the 1990s soon dried up as the failure rate of clinical trials, the huge cash-burn and increased bankruptcy of cell-based therapy companies caused investors to lose confidence in the industry and as a
result, the capital value of publicly traded companies plummeted from US$2.5 billion in 2000 to US$300 million by the end of 2002 [10]. This decline was predominantly driven by a lack of fundamental understanding of the therapeutic mechanism of action of MSCs in vivo which is still under discussion [37]. This understanding will be imperative, as relevant potency assays must be used to optimize manufacturing processes based on a single clinical indication, even if the same master cell line is to be used for multiple therapeutic targets.

Clinical work in the mid-2000s showed that the immunomodulatory properties of MSCs played a key role in their therapeutic potential [38,39], suppressing tissue rejection by inhibiting the response of the patient’s lymphatic cells. This is reflected in Figure 6, as the use of MSCs for immunology and rheumatology based indications are common and are progressing through clinical development with five trials progressing to Phase III, one targeting osteoarthritis (Medipost), one targeting graft versus host disease (Mesoblast) and three targeting Crohn’s disease (Tigenix and two by Mesoblast). This demonstrates the increased emphasis that industry has placed on the immunomodulatory mechanism of action that MSCs have demonstrated. The mounting evidence that the therapeutic effect of MSCs is not mediated by engraftment and terminal differentiation but by cell secreted trophic factors is gaining increasing momentum with studies showing that donor DNA is only retained in the lungs, lymph nodes and intestine following MSC infusion [40]. There is limited clinical evidence supporting the optimal method for therapeutic delivery of cell-based products [41] and it is likely that, as with manufacturing processes, delivery systems will be developed specifically for each therapeutic indication [42].

**Case study: product manufacture during Phase I clinical trial**

Since the discovery of embryonic stem cells in 1998 [43], pluripotent stem cells have been considered a promising source of allogeneic stem cells for regenerative therapies. In 2010 Geron Co. went into Phase I clinical trials with an allogeneic human embryonic stem cell (hESC)-derived oligodendrocyte progenitor product for the treatment of spinal cord injury. Despite no adverse safety issues with this hESC-derived therapy, Geron Co. halted the clinical trial citing financial and regulatory issues [44]. Currently, all of the clinical trials involving hESCs are targeting ophthalmology related indications, in Stargardt’s macular dystrophy.
and dry age-related macular degeneration. Advanced Cell Technologies are sponsoring three of these clinical trials in the UK and USA, treating these disorders with hESC-derived retinal pigment epithelium (RPE) cells. One of the key challenges in the manufacture of a pluripotent product is in the differentiation of the hESC into the therapeutically active cell, which can dramatically impact product yield. The Advanced Cell Technologies RPE product is manufactured from the GMP-grade MA09 hESC line, cultured on mitomycin-inactivated murine embryonic fibroblasts for three passages. Following the formation of embryoid bodies, pigmented RPE patches are isolated and purified [45], before further expansion and cryopreservation for clinical use. Product characterization takes place in-process and postcryopreservation, in the form of pathogen testing, karyotype analysis and purity [46]. Much of this manufacturing process is labor-intensive and requires operator intervention, which may not be amenable to automated and scalable manufacture. Therefore, despite the positive safety data from these ESC clinical trials [44], it is clear that there are many manufacturing challenges that must be overcome before these therapies can be translated through clinical trial and into commercial production.

Case study: manufacture for commercial production
It is clear from the clinical trial results in Figure 1, that the most prevalent clinical category for cell based therapies is in oncology, with immune cells contributing to around half of these cell-based therapies. Sipuleucel-T is an autologous active cell-based immunotherapy product designed to stimulate an immune response against prostate cancer from Dendreon Corporation (WA, USA) [47]. In 2010, Sipuleucel-T (Provenge®) became the first FDA-approved autologous cell-based therapy, providing a scale-out service based approach to product manufacture and delivery. Patient cells are collected via leukapheresis [48] and cold shipped to a manufacturing site, where the cells are manipulated under GMP conditions to isolate and activate the target immune cells. The activated cell-based immune therapy is then cold shipped back to the patient to be reinfused, which is repeated three times to deliver the full therapeutic dose. Dendreon Corporation operates patient logistics from a central location, with distributed manufacturing taking place at multiple sites across the USA. This product handling and manipulation is largely manual and has led to high product operating costs, increasing the product cost of goods and therefore creating a high
reimbursement price, which is US$93,000 per patient. Efforts have been made to reduce the cost of goods by implementing automated process steps, with the intention of reducing these high operating costs. This case study highlights the importance of considering functionally closed and automated scale out processes early in clinical development as this will reduce the overall cost of goods during commercial production [49].

A proposed clinical development pathway would be to produce Phase I material in a manual, semiclosed process which could then be transferred to a scalable, closed and automated manufacturing process following the success of the initial clinical trials. This model would allow for reduced capital investment to evaluate the product at Phase I but then would transfer the process in a timely manner, ensuring a scalable and cost effective product supply for late stage clinical development and commercial supply. Considering the high cost and increased risk of validating sterilization cycles, it is likely that these closed-processes will utilize disposable technology, mimicking current therapeutic protein process development [50].

Progress from clinical development to product manufacture

Industry progression towards the successful manufacture of cell-based therapies is evident by international cooperation on the formation of global reference ‘ruler’ standards and protocols to facilitate manufacturing comparability of MSCs [51]. This will facilitate the development of consistent manufacturing processes across multiple sites and allow for a method to define each cell line. Defining desirable product characteristics is critical and will form the basis of release tests as well as setting the tolerances on the process, allowing for systematic product development and optimization.

A further sign that the commercial market for MSC application is increasing in competitiveness is the drive by companies such as Athersys and Osiris to clearly differentiate their product and its production process from ‘generic’ hMSCs. This phenomenon parallels the clinical development of MACITM (by Genzyme) and CCI™ (by TiGenix) where companies actively sought to distance their products from legacy clinical data on the use of Autologous Chondrocyte Implantation, which was inconclusive in demonstrating cost–effectiveness [14]. This approach to cell-based product development is largely driven by a fundamental lack of understanding of the product mechanism of action, as companies must differentiate their product to avoid a race to the bottom, whereby a lower cost therapy has an increased chance of reimbursement. Without the knowledge of the how these cell-based therapies elicit therapeutic benefit, manufacturing processes cannot be optimized to maximize this function and will inevitably be surpassed by lower cost therapies. Given the importance of regulation on the production of cell-based therapies, it is perhaps unsurprising that in the race to successfully commercialize cell products, companies have begun to raise the regulatory bar as a tactic to disrupt competitor companies. This action has the potential to hinder smaller companies who do not have the financial resource to comply with this increasing regulatory burden.

A further hallmark of the commercial progress of MSC therapies is the move towards developing automated processes as well as the implementation of functionally closed manufacturing systems and consideration of the associated logistics, supply chain and cost of goods [52]. Significant UK government investment in the form of the Cell Therapy Catapult has been tasked with ‘derisking’ cell-based therapy development for industry by providing a ‘center of excellence’ to bridge the current translation gap in the industry [53]. Outside of the UK, additional institutes have been established to provide resources in Canada (Centre for Commercialization of Regenerative Medicine) and the USA (California Institute for Regenerative Medicine and the National Institutes of Health) to facilitate this translational process. Despite this progress, there remains a requirement for a better understanding of potential manufacturing platforms and how they can be best utilized for cell-based therapy production.

Expansion technologies to achieve clinical scale manufacture

With cell-based therapies moving towards commercialization and multiple clinical trials in late stage development, it is clear that selecting suitable manufacturing technology is becoming increasingly important. It is imperative that potential pitfalls in developing scalable manufacturing methods are identified at an early stage and strategies are implemented that can streamline this development pathway (Table 1). A key consideration for streamlining this pathway is the tradeoff between clean room space and ongoing commercial supply as patient numbers increase from clinical development to commercial production. For an autologous therapy if you were treating, for example, 100 patients in a typical 2-week process for a Phase III clinical trial, you would likely require four clean room facilities in order to separate each patient lot. This is assuming that the process is not entirely closed and therefore each lot must be segregated to avoid cross contamination. The issues then arise if the product is successful and commercial production is carried out using the same process for say 1000 or even 10,000 patients per year, where the clean room and personnel requirements increase...
10-fold or 100-fold respectively, making the product cost-prohibitive at this scale. This will drive the development of expansion platforms that are fully closed, so that multiple patient lots can be manufactured in the same facility, greatly reducing fixed and operating costs as the product moves toward commercial production. Table 2 shows the number doses per lot achievable for multiple expansion technologies currently available based on an allogeneic MSC treatment for myocardial infarction [54] requiring 35–350 million cells per dose. This demonstrates the challenge of manufacturing an MSC-based product for 10,000 patients per year, given the number of doses per lot achievable using current expansion technology.

A manufacturing process that reduces biological divergence will inevitably yield a more consistent and higher quality product. This biological divergence is typical of cell expansion processes, whereby small changes to the cell environment at the start of culture will lead to large changes by the time the product is ready for clinical use. This divergence is exacerbated by the open and non-closed processes currently used in production, leading to high fixed costs and variable costs in the manufacturing process.

Table 1. Strategies for streamlining the path to scaled clinical manufacturing.

| Process requirement | Description | Benefits from implementation |
|---------------------|-------------|------------------------------|
| Implement end to end closed processing | Utilize upstream and downstream processes that can be interconnected in a functionally closed manner | While manufacturing systems are becoming functionally closed, the reliance on open or non-closed inter process steps and manipulations necessitates the use of a clean room facility – massively increasing fixed costs of manufacture |
| Use processing steps and processes with the highest practicable expansion ratio† | Maximize product output for a given master cell bank population or donor cell population | Any reduction in the number of unit processes that must be validated to pass through process qualification significantly reduces the cost of pre- and post-market validation |
| Increase process capability | Increase stability and consistency of individual process outputs | Reduce process output variation will provide a more consistent utilization of a manufacturer's fixed cost base. This in turn improves operational costs and reduces overall COGS |
| Increase inter-process pooling tolerances | All manufacturing platforms have a limit in the amount of surface area that can be manipulated (seeded or harvested) per unit time. This limit, in combination with cell pooling time tolerances between processing stages limits the effective batch size that may be cultured | Increase in overall batch size that may be achieved. For scale out systems such as closed single use bioreactors or the manual/automated manipulation of planar vessels an increase in pooling time tolerance could yield cost reductions in the form of reduced manufacturing labor, increased capacity utilization |
| Keep COGS low in short term while maintaining cost headroom | Develop a process that has a (realistic) reimbursable price point at all stages of scale, as market adoption rates are typically slow | Cell therapy manufacturing companies with approved products have struggled to achieve routine reimbursement at a level that would enable investment in facilities and process improvements that would reduce their costs. This effectively traps companies in a difficult financial position. Companies that maintain low COGS during clinical development can reinvest in process improvement and scale as they enter the market |
| Implement intermediate cryopreservation stages | Autologous – implement a cryopreservation step between manufacturing and product delivery Allogeneic – implement multiple in-process banking stages between cell sourcing, expansion and product delivery | In allogeneic manufacturing intermediate cryopreservation stages allow a more efficient use of master bank cell populations at low production capacities. Its also allows different stages of the process to be scaled up independently, allowing a staged validation program. In both autologous and allogeneic manufacturing strategies this reduces the need to release products at risk before end stage quality and safety testing is complete |

†Expansion ratio defined as the output of cells divided by the input population.

COGS: Cost of goods.
harvested. This has been demonstrated in the culture of embryonic stem cells whereby changes to dissolved oxygen levels in the culture medium can lead to changes in cell growth characteristics [55] and differentiation potential [56]. This potential divergence can be limited by reducing the heterogeneity of the culture conditions via mixing, so that the cell microenvironment remains consistent throughout the culture. Technology that does not induce mixing such as T-flasks will potentially suffer from heterogeneity in physical, chemical and hence physiological conditions, although it can also exist in other forms. Packed/fluidized-bed bioreactors, due to their high density will invariably suffer from axial concentration gradients as medium flows through the bed, particularly as the scale increases. Hollow fiber reactors have the potential to introduce heterogeneity via longitudinal concentration gradients as medium or dissociation reagent flowing down the bioreactor changes with distance through the fiber (under plug flow). Although these factors could impact the consistency of the product, there is a lack of evidence to back this up and far more work will be required to fully understand and develop the various manufacturing processes for cell-based therapies.

A lack of online process control is a key barrier for the consistent manufacture of cell-based therapies. Expansion technologies such as rotating flasks and multilayer flasks lack the capability for online cell visualization which could form the basis for a non-invasive control strategy based on cell coverage of a surface [57]. These technologies as well as T-flask automation currently lack the ability for online medium sampling which prohibits the control of key nutrient and metabolite concentrations, a staple of process control in the bioprocessing industry. Without an effective process control metric, expansion technology will suffer when attempting to maintain product consistency and will incur a higher cost for product validation. Process control strategies are poorly understood for novel cell expansion technologies, however are routine in stirred tank bioreactors for current bioprocesses [58,59]. In addition to this, the physical characterization of the stirred reactor system is well understood and can also be directly translated from traditional bioprocesses [60–62]. This development is analogous to process analytical technology (PAT) [63] in the current biopharmaceutical industry [64] and could be used as a model for the cell-based therapy process development.

The efficient harvest of cell-based therapies represents one of the few deviations from traditional bioprocessing and must be designed based on cell sensitivity and the pooling time limitations of the product. Scale-out processes such as rotating flasks [65,66], T-flasks [22] and multilayer flasks [67] will require high cell pooling times which will limit the effective scale to which these therapies can be manufactured as product quality will likely reduce during the pooling process (Table 2). For cell-based therapies that are surface-adherent, the dissociation of the cell from the surface is required during the expansion process. This unique constraint of cell-based therapy manufacture is often overlooked but has been considered for packed/fluidized bed [68], hollow fiber [69] and stirred tank bioreactors [70]. Rocking-motion [71,72] and pneumatically driven bioreactors [62,73] have been previously employed for suspension cell culture. However, transferring these systems to surface-adherent cell types will likely require an additional process step to transfer the culture to a

### Table 2. Potential scale per lot and harvest constraints for various cell therapy manufacturing platforms.

| Expansion technology | Estimated scale per lot | Estimated doses per lot | Adherent cell harvest |
|----------------------|-------------------------|-------------------------|-----------------------|
|                      | Surface area, cm² | Harvest volume, l | Number of cells (billion) | 35 million cells per dose | 350 million cells per dose | Detachment efficiency | Pooling time |
| Adherent             | Rotating flasks      | 850,000                | 100                     | 21.30                   | 609                          | 61                  | Low | High |
|                      | Packed/fluidized     | 440,000                | 5                       | 11.00                   | 314                          | 31                  | Medium | Medium |
|                      | Hollow fiber         | 210,000                | 5                       | 0.50                    | 14                           | 1                   | Medium | Medium |
|                      | T-flask (automated)  | 225,000                | 50                      | 0.56                    | 16                           | 2                   | High  | High  |
|                      | Multilayer flask     | 400,000                | 30                      | 10.00                   | 286                          | 29                  | High  | High  |
| Adherent and suspension | Stirred tank       | >4,000,000             | >1000                   | >100.00                 | >2857                         | >286               | High  | Low  |
|                      | Rocking-motion       | 2,000,000              | 500                     | 50.00                   | 1429                          | 143                | Low   | Low   |
|                      | Pneumatically driven | 2,000,000              | 500                     | 50.00                   | 1429                          | 143                | Low   | Low   |

In situ detachment efficiency: low <50%, medium 50–90% and high >90%.
Pool time: low <1 h, medium 1–5 h, high >5 h.
harvest, allowing for increased agitation to promote efficient detachment and harvest of cells from microcarriers, which may not be sufficient in these systems [74]. A further challenge will emerge if the cells are required to be dissociated during the process (i.e., if bead-to-bead transfer does not occur at a high enough rate to support one step expansion). This will add additional steps to the process and will likely influence the design and optimization of other aspects of the process, such as agitation strategy and culture medium.

Although much work is yet to be done on calculating the optimum volume for expanding cell-based products in stirred tank reactors, current estimates put the maximum value at around 1000 l [75]. It should be noted however, that industrial scale mammalian cell culture, once considered to be limited to a similar scale is now routinely operated at greater than 10,000 l [62,76]. Similar gains in effective cells per batch can also be made by increasing the microcarrier concentration, which is currently far from optimal for suspension based systems [77].

One feature that is a fundamental aspect of all culture processes is the need to supply oxygen to the cells. Due to its limited solubility, this supply is required to be provided continuously. As the cell density increases, it becomes necessary to provide that oxygen by sparging: typically with air, which also strips out the carbon dioxide produced [78]. For free suspension animal cells, sparging has required the inclusion in the media of protective agents, almost always the surfactant, PluronicF68, as bursting bubbles are detrimental to cell viability [62]. Because the cell densities reached to date and the specific oxygen uptake rate of MSCs [77] and embryonic stem cells [79], for example, are both low, such issues have not yet been addressed but will need to be as the density increases, especially since the presence of these protective agents in final product formulations may be an issue. For pneumatically driven bioreactors [62,73], bubbling gas is inherent to the way they function and therefore potential cell damage must be considered during the appraisal of expansion technology.

It is clear that other than stirred bioreactors, there is a distinct lack of comparable and peer reviewed data for the majority of manufacturing platforms discussed in this study. This could potentially be to the detriment of the field, as without reliable information to inform technology appraisal, process development is likely to default to stirred tank bioreactors due to the strong legacy data that exists for large scale mammalian cell culture. On the other hand, there is nothing to be gained by innovating unnecessarily. Many of the potential manufacturing platforms mentioned here were considered in the 1980s for culturing animal cells, when it was considered that because they lacked a cell wall, it would be impossible to do so in the presence of rotating stirrers. Yet today, whether in single use systems, at bench scale or the largest commercial scale, stirred bioreactors are commonplace in the biotechnology industry [80]. Indeed, they are now being preferred industrially even for clone selection in robotically controlled microbioreactors (ambrTM) at the 15 ml scale [81].

**Conclusion**

Providing an overview of the current cell-based therapy landscape has shown the breadth and diversity of the current cell-based therapies in clinical development. With an increasing number of cell types and clinical indications being assessed, it is clear that consideration must be given to how these products will be manufactured and subsequently delivered to patients on a clinically relevant scale. Although, several manufacturing systems are already available for the expansion and manipulation of therapeutically relevant cells, an optimal and universal manufacturing platform does not yet exist and may not be attainable due to the variety of cell types and clinical applications.

With the exception of MSCs, most therapeutically relevant cell types have only been demonstrated on a subset of the broad range of available platforms. Developers considering the range of available manufacturing technologies need to balance the competing pressures discussed. Nevertheless, the fact that both MSC culture and harvest have now been effectively demonstrated in stirred bioreactors based on sound physical principles is encouraging, especially given the inherent flexibility and controllability of this technology; and its success with free suspension and adherent animal cells from the 15 ml to the 25,000 l scale.

As an industry, cell-based therapies are still in the early stages of clinical development. It is clear that we must better understand the quality of our cell-based products in order to form a stable base for process evaluation and development. Investment in cell-based therapy manufacturing in the UK has created an urgent need to better understand these cell-based therapy products to facilitate successful manufacture based on strong clinical data demonstrating product efficacy. It is likely that there will be bespoke manufacturing processes for each of these cell-based products as well as ‘generic’ cell-based therapy products driven by this fundamental lack of product understanding, with a move toward functionally closed and single-use technology.

**Future perspective**

Moving forward, we must better characterize cell-based therapy clinical trials with accessible information for a host of variables, including cell dose, patient numbers and cell providence. This will allow for effi-
cient and accurate data collection on cell-based therapy clinical trials, facilitating decision making across the cell-based therapy sector. As understanding of the cell-based products increases, we will likely experience step change improvements in manufacturing capability. However, despite the current progress, we must recognize that cell-based therapy is still a relatively green industry with great potential yet to be realized.

Open Access
This work is licensed under the Creative Commons Attribution-NonCommercial 3.0 Unported License. To view a copy of this license, visit http://creativecommons.org/licenses/by-nc-nd/3.0/

Author contribution
CJ Hewitt, B Kara and AW Nienow designed the research questions; TRJ Heathman developed the clinical trial selection framework and completed the data analysis; TRJ Heathman, B Kara and MJ McCall compiled the manufacturing data; and TRJ Heathman drafted the manuscript for publication.

Acknowledgements
The authors would like to thank Dr Deyna Cardosa, Specialist Registrar at Guy’s and St Thomas’ NHS Trust for assistance with disease categorization within the clinical trial framework.

Financial & competing interests disclosure
This study has been funded by the Engineering and Physical Sciences Research Council and FUJIFILM Diosynth Biotechnologies. B Kara is a director of science and technology at FUJIFILM Diosynth Biotechnologies. The authors have no other relevant affiliations or financial involvement with any organization or entity with a financial interest in or financial conflict with the subject matter or materials discussed in the manuscript apart from those disclosed.

No writing assistance was utilized in the production of this manuscript.

Executive summary

Aims
• Inform industry and academia of current clinical activity involving the cell type with which they work.
• Identify gaps in the manufacturing requirements for cell-based therapies.

Materials & methods
• Searched for the term ‘cell’ on ClinicalTrials.gov to include all clinical trials involving the use of cells up to 1 January 2014.
• Results were further sorted on the database to show only open trials that conformed to the definition of a cell-based therapy.

Clinical translation of mesenchymal stem cell-based therapies
• Lack of fundamental understanding of the therapeutic mechanism of action of MSCs in vivo must be overcome to facilitate process development and optimization.
• There is limited clinical evidence supporting the optimal method for delivery of cell-based products and it is likely that they will be developed specifically for each therapeutic indication.

Progress from clinical development to product manufacture
• Progress reflected by development of automated processes, reference standards and implementation of functionally closed manufacturing systems.
• Companies must differentiate their product to avoid a race to the bottom, whereby a lower cost therapy has an increased chance of reimbursement.

Expansion technologies to achieve clinical scale manufacture
• Imperative that potential pitfalls in developing scalable manufacturing methods are identified at an early stage and strategies are implemented that can streamline this development pathway.
• Consideration must be given to cell harvest and pooling time limits as they represent key deviations from traditional bioprocesses.
• Other than stirred reactors with cells attached to microcarriers, there is a lack of comparable data for the majority of the cell-based manufacturing technologies.

Conclusion
• Multiple manufacturing platforms will likely be required to cover the diversity of cell types, clinical indications and business models.
• A greater understanding of cell quality metrics are required for bioprocess development.

Future perspective
• Need to better characterize cell-based therapy clinical trials and data should be immediately accessible.
• As understanding of the cell-based products increases, we will likely experience step change improvements in manufacturing capability.
• Despite the current progress, we must recognize that cell-based therapy is still a relatively green industry with great potential yet to be realized.
• First study surveying the clinical trial landscape for cell-based therapies, which has led to subsequent clinical trial analysis.

References
Papers of special note have been highlighted as:
• of interest; •• of considerable interest

1 Mason C, McCall MJ, Culme-Seymour EJ et al. The global cell therapy industry continues to rise during the second and third quarters of 2012. Cell Stem Cell 11(6), 735–739 (2012).

2 Tavassoli M, Crosby WH. Transplantation of marrow to extramedullary sites. Science 161(3836), 54–56 (1968).

3 Caplan AI. Mesenchymal stem cells. J. Orthop. Res. 9(5), 641–650 (1991).

4 Pittenger MF, Mackay AM, Beck SC et al. Multilineage potential of adult human mesenchymal stem cells. Science 284(5411), 143–147 (1999).

5 Culme-Seymour EJ, Davie NL, Brindley DA, Edwards-Paton S, Mason C. A decade of cell therapy clinical trials (2000–2010). Regen. Med. 7(4), 455–462 (2012).

• Review of the requirements for defining and measuring the quality of cell-based therapies, which is a key step in their successful manufacture.

19 Moll G, Hult A, Von Bahr L et al. Do ABO blood group antigens hamper the therapeutic efficacy of mesenchymal stem cells? PLoS ONE 9(1), e85040 (2014).

20 Rayment EA, Williams DJ. Concise review: mind the gap: challenges in characterizing and quantifying cell- and tissue-based therapies for clinical translation. Stem Cells 28(5), 996–1004 (2010).

21 Williams DJ, Thomas RJ, Hourd PC et al. Precision manufacturing for clinical-quality regenerative medicines. Philos. Trans. A Math. Phys. Eng. Sci. 370(1973), 3924–3949 (2012).

•• Outlines the importance of precision manufacturing in regenerative medicine and the requirements for the manufacturing technology.

22 Thomas RJ, Chandra A, Liu Y, Hourd PC, Conway PP, Williams DJ. Manufacture of a human mesenchymal stem cell population using an automated cell culture platform. Cytotechnology 55(1), 31–39 (2007).

23 Thomas RJ, Anderson D, Chandra A et al. Automated, scalable culture of human embryonic stem cells in feeder-free conditions. Biotechnol. Bioeng. 102(6), 1636–1644 (2009).

24 Thomas RJ, Hope AD, Hourd P et al. Automated, serum-free production of CTX0E03: a therapeutic clinical grade human neural stem cell line. Biotechnol. Lett. 31(8), 1167–1172 (2009).

25 Liu Y, Hourd P, Chandra A, Williams DJ. Human cell culture process capability: a comparison of manual and automated production. J. Tissue Eng. Regen. Med. 4(1), 45–54 (2010).

26 Mason C, Hoare M. Regenerative medicine bioprocessing: the need to learn from the experience of other fields. Regen. Med. 1(5), 615–623 (2006).

27 Mason C, Dunnill P. Assessing the value of autologous and allogeneic cells for regenerative medicine. Regen. Med. 4(6), 835–853 (2009).

28 European Medicines Agency. European Public Assessment Report: Provenge (2013). www.ema.europa.eu/ema/index.jsp?curl=pages/medicines/human/medicines/002513/human_med_001680.jsp&mid=WCM0b01ac058001d124

29 Clinical Trials.gov. http://clinicaltrials.gov

30 Cell therapy and regenerative medicine glossary. Regen. Med. 7(Suppl. 3), S14–S116 (2012).

31 Zarin DA, Tse T, Williams RJ, Califf RM, Ide NC. The clinicaltrials.gov results database – update and key issues. N. Engl. J. Med. 364(9), 852–860 (2011).

32 Breyer A, Estharabadi N, Oki M et al. Multipotent adult progenitor cell isolation and culture procedures. Exp. Hematol. 34(11), 1596–1601 (2006).

33 Jiang Y, Vaessen B, Lenvik T, Blackstad M, Reyes M, Verfaillie CM. Multipotent progenitor cells can be isolated from postnatal murine bone marrow, muscle, and brain. Exp. Hematol. 30(8), 896–904 (2002).
Perspective on the importance of understanding the identity in vivo

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Le Blanc K, Rasmusson I, Sundberg B et al. Treatment of severe acute graft-versus-host disease with third party haploidentical mesenchymal stem cells. *Lancet* 363(9419), 1439–1441 (2004).

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Le Blanc K, Frassoni F, Ball I et al. Mesenchymal stem cells for treatment of steroid-resistant, severe, acute graft-versus-host disease: a Phase II study. *Lancet* 371(9624), 1579–1586 (2008).

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Von Bahr L, Batsis I, Moll G et al. Multiple radial or radial and axial up-pumping impellers: a novel mixing concept for mammalian cell cultures. *Chem. Eng. Sci.* 68(3), 283–291 (2005).

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Joeris K, Frerichs JG, Konstantinov K, Scheper T. In-situ microscopy: online process monitoring of mammalian cell cultures. *Cytotechnology* 58(1–3), 129–134 (2002).

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Trummer E, Fauland K, Seidinger S et al. Process parameter shifting: part I. Effect of DOT, pH, and temperature on the performance of Epo-Fc expressing CHO cells cultivated in controlled batch bioreactors. *Biotechnol. Bioeng.* 94(6), 1033–1044 (2006).

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Butler M. Animal cell cultures: recent achievements and perspectives in the production of biopharmaceuticals. *Appl. Microbiol. Biotechnol.* 68(3), 283–291 (2005).

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Vråbel P, Van Der Lans RJJM, Luyben KCaM, Boon L, Nienow AW. Mixing in large-scale vessels stirred with multiple radial or radial and axial up-pumping impellers: modelling and measurements. *Chem. Eng. Sci.* 55(23), 5881–5896 (2000).

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Langheinrich C, Nienow AW, Eddleston T et al. Oxygen transfer in stirred bioreactors under animal cell culture conditions. *Food Bioproc. Procs.* 80(1), 39–44 (2002).

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Nienow AW. Reactor engineering in large scale animal cell culture. *Cytotechnology* 50(1–3), 9–33 (2006).

Clinical study demonstration the in vivo immunomodulatory properties of MSCs leading to shift in clinical use for MSCs.

Key paper addressing the issues associated with large scale engineering of suspension bioreactors.

Guidance for Industry. PAT - a framework for innovative pharmaceutical development, manufacturing and quality assurance (2004). www.fda.gov/downloads/Drugs/Guidances/ucm070305.pdf
First paper to show the use of intense agitation for cell detachment as a scale up technique with a sound theoretical base.

First published report of large scale microcarrier expansion and harvest of hMSCs, demonstrating the feasibility of stirred reactors for cell-based therapy production.