Cellular and Molecular Implications of Mature Adipocyte Dedifferentiation

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Abstract

There is a voluminous amount of scientific literature dealing with the involvement of adipocytes in molecular regulation of carcass composition, obesity, metabolic syndrome, or diabetes. To form adipocytes (process termed adipogenesis) nearly all scientific papers refer to the use of preadipocytes, adipofibroblasts, stromal vascular cells or adipogenic cell lines, and their differentiation to form lipid-assimilating cells containing storage triacylglyceride. However, mature adipocytes, themselves, possess ability to undergo dedifferentiation, form proliferative-competent progeny cells (the exact plasticity is unknown) and reinitiate formation of cells capable of lipid metabolism and storage. The progeny cells would make a viable (and alternative) cell system for the evaluation of cell ability to reestablish lipid assimilation, ability to differentially express genes (as compared to other adipogenic cells), and to form other types of cells (multi-lineage potential). Understanding the dedifferentiation process itself and/or dedifferentiated fat cells could contribute to our knowledge of normal growth processes, or to disease function. Indeed, the ability of progeny cells to form other cell types could turn-out to be important for processes of tissue reconstruction/engineering and may have implications in clinical, biochemical or molecular processes.

Key words: mature adipocytes, dedifferentiation, review.

Introduction

As a consequence of cellular adipogenesis, lipid metabolism, gene and protein expression and adipokine secretion, mature adipocytes containing a single large lipid droplet were generally considered to be in the terminal stage of differentiation [1-4]. Recent studies, however, have indicated that mature adipocytes possess the ability to dedifferentiate into a population of proliferation-competent progeny cells known as adipofibroblasts or dedifferentiated fat (DFAT) cells [5-13]. This ill-defined phenomenon challenges the traditional view of the terminality of mature adipocytes [1, 14].

Mature adipocyte dedifferentiation

Adipocyte dedifferentiation is readily seen in vitro [5-10, 15-33]. The key to obtaining knowledge
with progeny of mature adipocytes is to insure that initial cell cultures are pure, and do not possess contamination with spindle-shaped preadipocytes. In order to accomplish this, during the dedifferentiation process mature adipocytes attach to the surface of a culture flask (Figure 1) and change morphology from a round cell into an irregular shaped cell. Subsequently, the lipid droplet is sectioned into smaller droplets and the cell divides into two progeny cells processing different amounts of lipid [7, 15-17, 34]. Chen et al. (2009 a,b) [15,18] indicated that cultured mature pig-derived adipocytes expel lipid directly into the medium prior to becoming capable of cell division, rather than retaining lipids and transferring the lipid into progeny cells symmetrically or asymmetrically as do mature adipocytes from other animals [5,7,19,20,35]. These results suggest that lipid-processing by individual cells may be different among species, while the cellular and molecular regulation of the dedifferentiation process remains unclear. Although dedifferentiation may result in cellular/molecular markers being expressed in the diametrically-opposite direction of traditional adipogenesis [21], these checkpoints have not been defined, even though the phenomenon of mature adipocytes forming proliferative-competent progeny cells was first reported over 30 years ago by Adebonojo (1975a,b) while performing research on human adipocytes [22, 23].

Are there other explanations about these cells? Could DFAT cells that have been described as originating from mature adipocytes actually be cells like fibroblasts that possess the potential for extreme levels of lipid metabolism [5,7,36]? Alternatively, could these cells be a completely new type of cell which presents both differentiation and proliferation abilities [5,7,36]?

While our studies with mature adipocyte dedifferentiation involved pure cultures of cells derived from large meat animals, similar cell systems from other animals have been established. To clarify the molecular mechanisms of mature adipocyte dedifferentiation, Ono et al., [24] using porcine cells and microarray analysis, identified differentially expressed genes during dedifferentiation.

![Figure 1. Sequence of pig-derived (mature) adipocyte dedifferentiation in vitro. A. Initial ceiling culture, B. Mature adipocyte following purification by differential plating methods. C-F. Sequence whereby the mature cell (in B) undergoes lipid reorganization/extrusion and divides with asymmetric cell division. All photomicrographs were taken with a Sony RGB digital camera married to a Nikon Diaphot phase contrast microscope and Image Pro Plus image analysis software [20 x magnification].](http://www.jgenomics.com)
Figure 2: Stages of differentiation and dedifferentiation of fat cells. (1) adipofibroblasts, preadipocytes or stromal vascular cells proliferate until the cell population is expanded to the correct number. Under intrinsic and extrinsic signals, these cells (2) initiate the metabolism/accumulation of lipids where some of them can become a mature adipocyte (3) while some of them retain the capacity to lose lipid and return to a proliferative stage. Other types of cells (4) may also display both proliferative and lipid accumulative ability contributing for an increase in the mature adipocyte population.

Ono et al. [24] reported that dedifferentiation resulted in 308 genes being down-regulated [many of which were involved in metabolism of lipid such as ADIPOQ, LIPE, PDK4, LPL, FASN, PPARG and FABP4]. Concomitantly, 368 genes were clearly up regulated during dedifferentiation [up regulated genes were those related to proliferation and differentiation, such as SERPINE1, TIMP1, PLA2, SFRP2, AEBP1, PRRX2, PEG10, IGFBP5 and ID2]. A variety of other studies are available to indicate that during de-differentiation mature adipocytes lose their functional phenotype, reenter the cell cycle, gain a fibroblast-like appearance and eventually acquire a multipotent capacity for lineage differentiation [8-10,12,13,19] (Figure 2).

DFAT cells as a model for adipogenesis

DFAT cells apparently possess a strong ability to proliferate (up to 30 passages) without detectable lipid deposition [25], which suggests the availability of DFAT cells could be used as a novel model for adipocyte differentiation and tissue formation. Whereas a variety of different cell culture systems have been used to address questions dealing with adipogenesis, including the use of lines of Swiss mouse embryo 3T3-L1 and stromal vascular (SV) cells [5], some significant differences between these cells and DFAT cells have been reported. Markers such as ANPEP, ITGB1, CD34, CD44 and THY1, ENG, MCAM, HLA-A, HLA-B, HLA-C were found on both SV and DFAT cells [28, 37-39]. However, the surface immuno-phenotype partially changes in different passages of adipose-derived SV cells. At the early passages (primary to 4th) of adipose-derived SV cells, the hematopoietic-associated markers (ITGAL, CD14, PTPRC, CD86 and HLA-DR) decreased and the MSCs(Mesenchymal stem cells)-associated markers (ANPEP, ITGB1, CD34, CD44, CD63, NT5E, THY1 and ALCAM) increased significantly [28,39]. Conversely, the phenotype of DFAT cells remained considerably stable from the early 2nd to as long as the 30th passage [25].

Cellular origin, homogeneity of cultures, whether the cells were previously transformed, repeatability of obtaining the same proportions of cells
for culture, markers of adipogenesis and lipid metabolism, and viability of cells are all physiological changes that must be considered when evaluating SV cells, cell lines and progeny from DFAT adipocytes [3,5-7,11-14,19,32,40]. For example, the doubling time of DFAT cells was approximately 18 h whereas the time for the 3T3-L1 cells was 22 h [26], DFAT cells apparently possess higher GPDH (Glycerol-3-phosphate dehydrogenase) activity than 3T3-L1 cells, and expression of SREBF1 and LPL mRNA, which is related to lipid metabolism, has been detected during the growth phase in DFAT cells but not in 3T3-L1 pre-adipocytes [5]. The expression of PPARGC1A and CEBPA mRNA shows an earlier peak during differentiation in DFAT cells than in 3T3-L1 cells [26, 27]. Moreover, gene expression of re-differentiating DFAT progeny cells showed some differences to published reports using beef-derived SV cells [14], and unlike SV cells confluence is not required for DFAT cells to undergo differentiation in vitro [31]. Data from in vitro studies also suggest that DFAT cells possess greater ability to differentiate, not only into mature adipocytes, but also into vascular endothelial cells, forming a normal adipose tissue [27]. From these results, it could be inferred that DFAT cells could serve as an alternative model for studies of the mechanism of adipose tissue differentiation and formation for both in vitro and in vivo studies.

In meat animals, understanding variables regarding regulation, cellularity, growth and/or development of adipose tissue in different depots are important since fat deposition has an economic meaning in livestock production [19]. Different markers interact functionally to direct several regulatory processes involved in adipogenesis or lipid metabolism, such as PPAR and SREBP [41, 42]. However, the regulatory compounds of adipogenesis are not necessarily similar between ruminant and monogastric animals [19]. It has been reported that fat deposition follows the order in which adipocyte number and volume is achieved firstly in the visceral adipose depot and lastly in the intramuscular depot [36,43]. Chen et al [16] have indicated that PPARG and CEBPA mRNA were higher in DFAT cells from intramuscular than visceral adipose tissue derived from finishing pigs after differentiation, which means intramuscular DFAT cells may be more active in adipogenesis than visceral DFAT cells. This may imply that even if DFAT cells originated from the same animal, different depots present different rates of redifferentiation and molecular markers. Additional evidence to support this concept includes: high RNA abundance of FABP4 found in adipofibroblasts originated from dedifferentiation of bovine perimyscular mature adipocytes, suggesting that progeny preadipocytes may potentially have an ability to redifferentiate at early time point due to inherited characteristics of molecular components in the cytoplasm of progeny cells derived from the mature adipocytes, although shape of the cells remains fibroblast-like form [14]. Studies that provide knowledge of the dynamic of accumulation and metabolism of lipids may lead to the adoption of strategies in the livestock production in order to enhance fat deposition in specific (desired) depots [34, 35].

Multilineage potential of DFAT cells

A slightly different focus of DFAT cells studies is to understand the mechanisms that allow these cells to transdifferentiate in other types of cells [28-30]. Under specific in vitro conditions, beef-derived DFAT cells redifferentiate in a protracted manner, of which some cells contain lipid and some did not (only possessing the storage vesicles; [31]). DFAT cells are capable of transdifferentiating to other types of cells such as osteoblasts, chondrocytes, skeletal myocytes, smooth muscle cells, cardiomyocytes, vascular endothelial cells, and neural cells under appropriate culture conditions in vitro or in vivo [24]. A particularly elegant study found that RUNX2, SPP1, SP7, BGLAP, PTH1R and SOX9 were expressed in DFAT cells originating from mature adipocytes suggesting the osteogenic and chondrogenic potentials of these cells are enhanced [28]. Moreover, the transdifferentiation of DFAT cells into osteoblasts was observed in vivo by the implantation of collagen-based scaffolds [32]. Osteogenic differentiation of DFAT cells could also be induced by an analog of retinol which interacts with bone morphogenetic proteins to inhibit adipogenesis and enhance osteogenesis [32].

Although DFAT and SV cells share similarity with regard to multipotential lineage, a distinct aspect of the differentiation capacities between these cells exist in their angiogenesis behavior [25,44]. Previous studies have shown that adipose tissue-derived SV cells can differentiate into endothelial lineage and express endothelial markers and incorporate acetylated low density lipoprotein (Dil-ac-LDL) when agents such as fibroblast growth factor (FGF), epidermal growth factor (EGF) and vascular endothelial growth factor (VEGF) were applied [45-49]. On the other hand, DFAT cells with or without induction have negative expression of endothelial cell or progenitor markers such as such VWF, PECAM1 and CD34 [28].

So, do DFAT cells possess multipotent potential [11], or might the manipulation of the daughter cells

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simply be something manifested through cell culture manipulation [44, 50, 51]? Recently, it was demonstrated that the regulatory region of MYOD1, a master regulator for skeletal myogenesis, in DFAT cells shows a high degree of methylation [29]. Treatment of 5-azacytidine, a demethylating agent, led to the expression of MYOD1 and MYOG in DFAT cells, as well as the formation of multinucleated cells expressing MUTYH, although MYF5 was still absent after induction [29]. Another study found expression of specific cardiomyocyte-specific markers was observed in DFAT cells derived from mature adipocytes when co-cultured with neonatal cardiomyocytes, suggesting that DFAT cells can convert into cardiomyocyte phenotype under appropriate microenvironment [33]. Moreover, it was found that DFAT cells expressed cardiac phenotypic markers when grown on semi-solid methylcellulose medium in the absence of cardiomyocytes, indicating that DFAT cells have the potential to spontaneously convert to cardiomyocyte phenotype [33]. Sakuma et al., [30] using immunohistochemical analysis revealed that more than 50% of the DFAT cells were successfully transformed into α-smooth muscle actin-positive cells under specific culture condition. The expressions of TAGLN2, α-smooth muscle actin and smooth muscle-myosin heavy chain were increased in DFAT cells during the first week of differentiation culture.

Collectively, these studies suggest that the DFAT cells have potential to differentiate into several lineages, including adipose cells, osteoblasts, chondrocytes, skeletal myocytes, smooth muscle cells, cardiomyocytes, endothelial cells and neuronal cells [44, 50]. Moreover, although much of the observed multilineage potential was dependent on environmental conditions, such as the type of culture media used [51] once cells begin to express a phenotype for a different tissue form they can be utilized for regimens of tissue reconstruction or tissue regeneration/engineering requiring those specific cells. Further studies with DFAT cells should yield a number of uses for clinical, biochemical, metabolism and molecular application.

Summary

Dedifferentiation of mature adipocytes, and proliferation/transdifferentiation abilities of their daughter progeny cells provide us with a cellular system for exploring adipocyte involvement in fat-related diseases or improving animal productivity. Moreover, DFAT cells could be used as a unique adipogenesis model, and new information may be gained on comparative differences among these cells and SV cells, and cell lines such as 3T3-L1 cells. Mature adipocytes that possess the ability to lose lipid and form proliferative-competent daughter cells may be (in fact) a different type of cell than has ever been categorized. Continued studies may uncover this potentially exciting possibility. Finally, the multipotent lineage of DFAT cells may lead to the use of these cells in tissue reconstruction/engineering. Collectively, there are cellular and molecular implications of mature adipocyte dedifferentiation, including the potential uses in clinical, biochemical and molecular applications.

Conflict of Interest

The authors have declared that no conflict of interest exists.

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