REVIEW

Dietary interventions as regulators of stem cell behavior in homeostasis and disease

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Stem cells maintain tissues by balancing self-renewal with differentiation. A stem cell's local microenvironment, or niche, informs stem cell behavior and receives inputs at multiple levels. Increasingly, it is becoming clear that the overall metabolic status of an organism or metabolites themselves can function as integral members of the niche to alter stem cell fate. Macroscopic dietary interventions such as caloric restriction, the ketogenic diet, and a high-fat diet systemically alter an organism's metabolic state in different ways. Intriguingly, however, they all converge on a propensity to enhance self-renewal. Here, we highlight our current knowledge on how dietary changes feed into stem cell behavior across a wide variety of tissues and illuminate possible explanations for why diverse interventions can result in similar stem cell phenotypes. In so doing, we hope to inspire new avenues of inquiry into the importance of metabolism in stem cell homeostasis and disease.

Adult stem cells serve as the stewards of the body by maintaining homeostasis and repairing tissues in response to stress. For stem cells to carry out these tasks, they must balance self-renewal and commitment to differentiate. When exposed to insults such as inflammatory stimuli, pathogens, mechanical stress, or oncogenic stress, stem cells are faced with new hurdles to restore homeostasis. If they cannot cope, an imbalance can occur in the stem cell's equilibrium, which if left unchecked can place the tissue at risk for hyperproliferation, chronic wounding, and/or cancer.

During normal homeostasis, stem cells integrate cues from their local microenvironment or niche to maintain tissue equilibrium. These cues include signaling pathways (BMPs, WNTs, TGF β s, NOTCH, SHH, and growth factor/tyrosine kinase receptors), extracellular matrix (ECM) ligands (fibronectin, laminins, and collagens), mechanical forces, cell–cell interactions, and metabolites, all of which have been shown to impact stem cell behavior

(Gonzales and Fuchs 2017; Intlekofer and Finley 2019; Fuchs and Blau 2020). In recent years, intracellular metabolites and metabolic pathways have gained particular attention as it was learned that they not only support growth and proliferation, but they can also work in conjunction with signal transduction pathways (Mihaylova et al. 2014; Martínez-Reyes and Chandel 2020; Jewell et al. 2013) and chromatin modifying enzymes to regulate gene expression (Ward and Thompson 2012; Schvartzman et al. 2018). Accordingly, metabolic inputs at all levels, from macroscopic dietary interventions to intracellular pathways, have been implicated in the fate decisions that are made by tissue stem cells.

Understanding how metabolism regulates stem cell function in vivo has begun to shed light on the convergence of phenotypes along the spectrum of normal homeostasis and cancer and possible therapeutic avenues for interventions that target metabolic pathways. To date, several methods have been employed to alter organismal metabolism using dietary interventions. The most thoroughly studied interventions include caloric restriction, prolonged fasting, ketogenic diet, and the high-fat “Western” diet (Table 1).

Engaging in caloric restriction, prolonged fasting, or a ketogenic diet elicits similar systemic changes to metabolic homeostasis, albeit through slightly different mechanisms. Caloric restriction, defined as either reducing calorie intake by >20% over ad libitum feeding or merely lengthening the time between meals (Pugh et al. 1999), induces a state of ketogenesis that is recapitulated by a ketogenic diet. Prolonged fasting elicits a similar outcome, but here, the lengths of fasted versus fed periods is able to better control and distinguish it from a ketogenic diet.

During the fasted or nutrient-poor state, glucose levels decline and ketone biogenesis increases. The liver produces and releases into circulation the ketones acetone and β -hydroxybutyrate (β -OHB). Ketone bodies then feed into the tricarboxylic acid (TCA) cycle as acetyl-CoA, thereby bypassing glucose oxidation, typically a major source of

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Table 1. Comparison of properties and effects of dietary interventions

	Control chow diet	Pure caloric restriction	Prolonged fasting	Ketogenic diet	High-fat diet	Western diet
Caloric content	4 kcal/gram chow	≥20% decrease	Unchanged or decreased	6 kcal/gram chow	5 kcal/gram chow	4.5 kcal/gram chow
Dietary protein content	20%	Decreased	Unchanged or decreased	20%–25%	20%–25%	17%
Dietary carbohydrate content	53%	Decreased	Unchanged or decreased	8%	20%–25%	49%
Dietary lipid content	5%	Decreased	Decreased	55%	34%	21%
Induces ketogenesis	No	No	Yes	Yes	No	No
Effect on insulin sensitivity	N/A	Increased	Increased	Increased	Decreased	Decreased
General trend in stem cell fate	N/A	Increased self-renewal	Increased self-renewal	Increased self-renewal	Increased self-renewal, carcinogenesis	Increased self-renewal, carcinogenesis

The compositions and major effects of the macroscopic dietary interventions discussed are summarized by intervention. Note: the percent compositions are by mass and not energy composition.

acetyl-CoA in cells (Shi and Tu 2015). Ketone bodies can be generated through administration of the ketogenic diet, which is a high-fat, low-carbohydrate diet that was originally developed to treat pediatric seizure disorders (Peterman 1924; Helmholz 1927; Neal et al. 2008). This feeding paradigm again pushes an organism into a state of transient ketogenesis, wherein the cell switches from oxidizing glucose to fatty acids and ketone bodies (β -hydroxybutyrate [β -OHB] and acetone), which derive from acetyl-CoA. Thus, the major systemic changes derived from these dietary interventions are (1) a decrease in glucose availability and oxidation, (2) an increase in lipid utilization, and (3) the production of ketone bodies as alternate carbon sources.

In contrast to dietary interventions that cause a state of ketogenesis, the high-fat diet elevates circulating glucose despite increasing insulin levels (Winzell and Ahrén 2004). By definition, the high-fat diet reduces an organism's insulin sensitivity and the efficiency by which glucose enters tissues. The high-fat diet also alters the fatty acid composition within the circulation, elevating total polyunsaturated, total n-6 (ω -6), and cholesteryl ester fatty acids without detectable changes in triacylglycerol fatty acids (Raatz et al. 2001). Overall, the high-fat diet (1) increases glucose availability, but not necessarily utilization; (2) increases circulating levels of certain types of unsaturated fatty acids, which may act as ligands for various signaling pathways; and (3) results in insulin resistance.

Interestingly, while these different dietary paradigms exert unique effects on organismal-level metabolism, they all converge on enforcing stem cell self-renewal across several different tissue types. The molecular mechanisms that enable these disparate dietary interventions to enact similar changes in stem cell behavior are currently unknown. However, there are several factors that may contribute to these convergent phenotypes. First, the decline in glucose availability and increase in lipid utilization confers increased stemness to cells across different tissues. Second, these macroscopic dietary interventions feed into stem cell-intrinsic glucose and fatty acid metabolism. Third, dietary interventions that affect the fasted and fed states bias stem cells toward either a self-renewal or differentiation program by exacerbating changes already present when shifting between these two nutrient states. Last, the diets activate similar metabolic stress pathways and yield their common phenotypes through similar mechanisms of compensation (Fig. 1).

Systemic dietary interventions converge on glucose and lipid metabolism

Disparate dietary interventions promote stemness across tissue types

One of the most heavily studied systems with regard to organismal-level dietary interventions and their effects on stem cell behavior, normal homeostasis, and disease is

Figure 1. Disparate dietary interventions converge on stem cell self-renewal. Macroscopic dietary interventions have been shown to influence stem cell behavior across a wide array of tissue types. Here, we summarize the major findings across these tissue types and propose possible explanations for why disparate dietary interventions converge on similar phenotypes. (SC) Stem cell, (ISC) intestinal stem cell, (HSC) hematopoietic stem cell, (EpdSC) epidermal stem cells, (HFSC) hair follicle stem cell, (MuSC) muscle stem cell (satellite cell), (NSPC) neural stem/progenitor cell.

the small intestine. These intestinal stem cells (ISCs) reside at the base of the crypts of Lieberkühn, where they proliferate continuously to fuel the flux of differentiating intestinal epithelial populations in the contiguous villus. Caloric restriction in this context promotes mTORC1-dependent enhanced self-renewal predominantly through Paneth cell niche-induced changes (Yilmaz et al. 2012; Igarashi and Guarente 2016), whereas high-fat or cholesterol-rich diets and intermittent fasting directly alter ISCs and progenitor function (Beyaz et al. 2016; Wang et al. 2018). The effects of a high-fat diet extend past homeostasis. In a model of ionizing radiation-induced intestinal injury, administration of a high-fat diet increased the number of surviving, proliferating, and *Olfm4*/OLFM4-positive crypts without altering numbers of enteroendocrine or goblet cells (Beyaz et al. 2016).

Both a high-fat diet and intermittent fasting lead to ISC-autonomous activation of PPAR δ signaling (Fig. 2), which enhances self-renewal through multiple mechanisms, including partial activation of a Wnt signaling program and enhancing fatty acid oxidation (Beyaz et al. 2016; Mihaylova et al. 2018). A high-fat diet and intermittent fasting could also profoundly alter the serum composition of lipids, as a high-fat diet likely enriches for circulating lipids that differ from those that are released from adipose tissue breakdown during intermittent fasting. These changes would translate to different PPAR ligands that can engage the pathway. In this regard, even though caloric restriction, a high-fat diet, and fasting all converge on enhanced crypt proliferation and survival in both homeostasis and injury-based regenerative programs, they may do so through different underlying mechanisms.

An overall metabolic state of ketogenesis can be achieved both through caloric restriction and a ketogenic diet. Whereas caloric restriction increases both ISC and Paneth cell numbers (Yilmaz et al. 2012), month-long administration of a ketogenic diet enriched the small intestine for both ISCs (*Lgr5*^{high}) and *Lgr5*^{low} short-lived

proliferative intestinal progenitors, without altering differentiated Paneth cell numbers (Cheng et al. 2019). What regulates the sensitivity of different cell types to macroscopic dietary interventions remains an open question, but it may reflect the intrinsic metabolic programs of cells within the niche. For example, ISCs, but not Paneth cells, are highly enriched for the expression of the rate-limiting enzyme for ketone biosynthesis, 3-hydroxy-3-methylglutaryl-CoA synthase 2 (*Hmgcs2*), and accordingly engage in de novo ketogenesis even in the absence of a ketogenic diet. The product of ketogenesis, β -hydroxybutyrate (β -OHB), can alter gene expression by inhibiting histone deacetylases (HDACs) (Shimazu et al. 2013), which enhances ISC self-renewal (Cheng et al. 2019).

Although the paucity of *Hmgcs2* expression relative to ISCs makes it unlikely that Paneth cells biosynthesize their own β -OHB, indirect effects could arise from increased availability or utilization of β -OHB in circulation (Fig. 2). Indeed, systemic β -OHB is known to increase during fasting, caloric restriction and/or ketogenic diets to millimolar concentrations, largely due to hepatic production (Cahill et al. 1966; Shimazu et al. 2013). Whether β -OHB contributes to enhanced self-renewal of ISCs under caloric restriction, a high-fat diet, and fasting remains to be explored. Additionally, while it is currently unclear whether β -OHB can accumulate at sufficient concentrations in the nucleocytosolic compartment to inhibit HDACs, β -OHB can be taken up from circulation either through passive diffusion or by facilitated transport via monocarboxylate transporters (MCTs) (Halestrap 1978; Shimazu et al. 2013). Thus, free diffusion into the nucleus through the large nuclear pores after sufficient uptake would enable high concentrations of this molecule to accumulate and alter gene expression.

While the data in ISCs provide compelling evidence that ketogenesis can promote stemness in the small intestine,

Figure 2. Integration of diet-induced metabolic changes into stem cell fate decisions. Macroscopic dietary interventions such as caloric restriction, prolonged fasting, the ketogenic diet, and a high-fat diet systemically alter an organism's metabolic state and the availability of key metabolic substrates. Interestingly, these interventions converge on pro-self-renewal phenotypes across stem cells in several different tissues. Stem cells integrate locally available metabolites such as fatty acids, ketone bodies (β -OHB and acetone), and microbiota-derived butyrate to enact changes in stem cell fate, skewing them toward a greater propensity for stemness as well as carcinogenesis. (β -OHB) β -Hydroxybutyrate, (CD36) cluster of differentiation 36 (fatty acid translocase), (HDAC) histone deacetylase, (MCT1) monocarboxylate transporter 1, (PPAR δ) peroxisome proliferator-activated receptor δ .

it remains to be seen whether other tissue stem cell types respond similarly to enforcing organismal ketogenesis. If so, it could be a molecular path to increasing longevity. Indeed, with regard to caloric restriction more generally, the benefits to stem cell renewal are broadly applicable across diverse tissue types. Stem cells of the muscle (satellite cells or MuSCs), bone marrow (hematopoietic stem cells, HSCs), and hair follicle (hair follicle stem cells, HFSCs) all spend much of their time in a relatively quiescent state during homeostasis, which enables them to preserve their self-renewing, tissue-regenerative capacity (Fuchs and Blau 2020). In these cases, caloric restriction favors their preservation and/or function in both young and old animals (Cerletti et al. 2012; Tang et al. 2016; Forni et al. 2017). Exactly how is less clear. For the hair follicle, which undergoes cyclical bouts of hair regeneration, the HFSC pool appeared to increase while hair coat turnover decreased when mice were restricted in their food intake (Forni et al. 2017). For muscle, the repair ability was enhanced (Cerletti et al. 2012). For the bone marrow, caloric restriction improved repopulation but impaired lymphoid differentiation capacity of the HSCs (Tang et al. 2016).

One of the complexities involved in sifting through the many beneficial effects of caloric restriction on stem cell

behavior and tissue fitness is that secondary effects often surface. For instance, calorie restriction was recently shown to increase the fitness of circulating memory T lymphocytes, cells that allow mammals to rapidly and effectively control subsequent challenges to the immune system. Enhancing their fitness triggered the memory T cells to home to the bone marrow, thereby imparting enhanced protection to the HSCs and promoting their survival (Collins et al. 2019). In a similar but even more circuitous vein, short-term fasting resulted in the activation of low-energy sensor 5'-AMP-activated protein kinase (AMPK) in hepatocytes, suppressing systemic CCL2 production by PPAR α and in turn reducing the number of circulating monocytes. The outcome was dampening of unwanted chronic inflammation without compromising monocyte mobilization in tissue repair, two features that favorably impact stem cells (Jordan et al. 2019). This result is consistent with the large body of literature implicating the immunosuppressive effects of starvation, suggesting that these well-documented immunosuppressive effects may benefit the host organism (Lord et al. 1998; Freitag et al. 2000). While the hormone leptin may rescue starvation-induced abrogation of T lymphocyte function, it is unclear whether leptin

administration is also sufficient to rescue any defects in hematopoietic stem cell function or differentiation. Given the critical role of immune cells in regulating stem cell behavior, further dissecting how metabolic inputs influence this crosstalk will be an interesting avenue of future study.

A high-fat diet can profoundly impact stem cell behavior

In striking contrast to the metabolic effects of caloric restriction and a ketogenic diet, a high-fat diet induces obesity in mice within months. Paradoxically, even though a high-fat diet is typically associated with poor health outcome, it can in some contexts favor stem cell expansion. A case in point is embryonic development of the vertebrate hematopoietic system, where superactivating the master regulator of cholesterol biosynthesis, *Srebp2*, either genetically or through a high-cholesterol diet, accelerates HSC emergence (Gu et al. 2019).

Similarly, under conditions of normal intestinal homeostasis, a high-fat diet increases ISC number and proliferation. However, unlike caloric restriction and ketone and fasting diets, a high-fat diet in the intestine depletes Paneth cells via activation of PPAR δ signaling, which in turn induces Wnt target genes in ISCs (Beyaz et al. 2016). As Paneth cells are ISC progeny and ISCs in turn depend upon WNT signals from Paneth cells, this imbalance within the intestinal crypt could create a secondary response by the ISCs. Moreover, a high-fat diet endowed progenitors in the crypt with stem cell properties, which may explain the maladaptive effects of a high-fat diet such as enhanced propensity for tumorigenesis.

Further insights come from investigating the adult bone marrow niche, where a high-fat diet affects stem cell behavior, but mostly by altering stem cell fate rather than numbers per se. Thus, when hypercholesterolemia-prone mice are fed a “Western diet,” their hematopoietic stem cells skew their fate away from a lymphoid lineage and towards a myeloid lineage (Christ et al. 2018). Elevated fatty food intake can also elicit an inflammatory response, which becomes all the more deleterious, given recent evidence that tissue stem cells retain an epigenetic memory of their inflammatory experiences, irrespective of whether the source of these experiences is a topical irritant, pathogen infection, or fatty diet (Naik et al. 2017; Kaufmann et al. 2018; Mitroulis et al. 2018; Naik et al. 2018; Netea et al. 2020). Consistently, abnormal myelopoiesis can exacerbate metabolic disease: In both patients and mice, enhanced inflammatory signaling in clonally expanded HSCs and their myeloid daughter cells accelerates atherosclerosis under a Western diet (Fuster et al. 2017; Jaiswal et al. 2017). Thus, the interplay between a high-fat diet and stem cell behavior may form a positive feedback loop leading to organismal health decline.

Taken together, although metabolites emanating from a high-fat diet can confer a pro-self-renewal paradigm to tissue stem cells, the long-term effects of such food remain to be evaluated, and in a number of contexts, the

diet is likely to alter stem cells in ways that skew their fates and/or exacerbate their response to inflammation.

Changes to intracellular glucose and lipid metabolism inform stem cell behavior

An enticing hypothesis is that macroscopic dietary changes feed into the individual metabolism of stem cells and their niche partners to cell-intrinsically coordinate self-renewal and differentiation. These dietary changes induce critical changes with respect to glucose and fatty acid metabolism. Therefore, it is likely that these interventions shift the balance in glucose and fatty acid metabolism intracellularly.

Unifying effects of lipid metabolism on self-renewal

Availability of circulating lipids substantially increases under conditions of caloric restriction, the ketogenic diet, and Western diets, and the accompanying increased fatty acid oxidation provides insights into the possible roots of the convergence of phenotypes observed with these diets (Corton et al. 2004; Sikder et al. 2018; Steinhäuser et al. 2018; Rosenbaum et al. 2019). At the level of transcriptional regulation, fasting increased PPAR-dependent fatty acid oxidation in ISCs to maintain stemness (Mihaylova et al. 2018). Orthogonal methods to enhance ISC fatty acid oxidation, such as genetic inhibition of glucose oxidation, similarly enhance ISC self-renewal (Bensard et al. 2020). Notably, ablation of PPAR γ in HFSCs resulted in a phenotype similar to scarring alopecia with destruction of the pilosebaceous units, indicating the importance of this pathway to proper hair follicle homeostasis (Karnik et al. 2009). Furthermore, neural stem/progenitor cells increase fatty acid oxidation during periods of quiescence and increase fatty acid synthesis after stem cell activation and during proliferative periods (Knobloch et al. 2013, 2017). Similarly, in the *Drosophila* germline, lipid utilization supports, whereas lipid synthesis antagonizes, stem cell maintenance (Sênos Demarco et al. 2019).

Lipids are also likely at the roots of the epigenetic memory induced in stem cells by their inflammatory experiences. In response to inflammation, both epidermal stem cells and HSCs not only activate cholesterol biosynthesis genes but also maintain them in a chromatin-poised state where they are more rapidly activated upon the next inflammatory encounter (Naik et al. 2017; Mitroulis et al. 2018). Interestingly, although cholesterol levels per se are not required, the activation of the cholesterol synthesis pathway appears to be an essential facet of the memory in HSCs, as it is erased by treatment with atorvastatin (Bekker et al. 2018; Mitroulis et al. 2018). In HSCs, this was linked to a key role of mevalonate, an intermediate in cholesterol biosynthesis, in promoting inflammatory memory. Although these findings suggest a key role for cholesterol synthesis in epigenetically altering stem cell landscapes, the mechanisms await further investigation.

An interesting beneficial twist to increased cholesterol synthesis is the improvement of the response of certain

stem cells to functional challenges. For HSCs, it conferred enhanced myeloablative effects of chemotherapy and abrogated the DNA damage response upon induction of systemic inflammation (Bekkering et al. 2018; Mitroulis et al. 2018). For epidermal stem cells, although not necessarily rooted in cholesterol synthesis per se, inflammatory memory enhanced their ability to heal wounds (Naik et al. 2017). The ability of different cell types to gain fitness through lipid metabolism provides insights into why seemingly deleterious metabolic outcomes related to increased cholesterol levels may have been evolutionarily conserved.

Glucose utilization dictates stem cell behavior

Cells oxidize glucose to fuel energy production and anabolic pathways to support growth and proliferation. Under fasting conditions in vivo, glucose feeds into the tricarboxylic acid (TCA) cycle by circulating lactate in all tissues except the brain, and glucose primarily tends to feed into glycolysis and the pentose phosphate pathway versus other metabolic pathways (Hui et al. 2017). In the fed state, direct glucose oxidation increases, but lactate oxidation remains the primary mechanism by which glucose feeds the TCA cycle. It remains unknown whether these shifts in glucose metabolism at the tissue level also occur in stem cells during fasting and refeeding. Studying this through new technologies such as imaging mass spectrometry may provide new insights into dietary regulation of stem cell behavior.

Emerging evidence suggests that glucose utilization can regulate a stem cell's decision to remain quiescent or activate. For example, HSCs are thought to reside in hypoxic areas of the niche and elevate glycolysis during periods of quiescence (Parmar et al. 2007; Simsek et al. 2010; Takubo et al. 2010; Nombela-Arrieta et al. 2013). Notably, maintenance of HSC quiescence requires the action of pyruvate dehydrogenase kinase, which, by preventing the shunting of pyruvate for mitochondrial oxidation, enhances glycolysis (Takubo et al. 2013). Similarly, inhibition of the glycolytic enzyme hexokinase (HK2) and lactate dehydrogenase (LDH) during normoxia resulted in proliferation defects in neural progenitor cells (Zheng et al. 2016). Conversely, HFSCs and ISCs rely on glycolysis to sustain proliferation. HFSCs that cannot activate LDH activity, which sustains redox balance to promote glycolysis, fail to activate hair cycling (Flores et al. 2017). Similarly, ISCs that cannot oxidize glucose in the TCA cycle have enhanced proliferation and self-renewal. Finally, recent evidence also suggests that glucose fate regulates stem cell differentiation: Blocking glucose entry into the TCA cycle in epidermal stem cells promotes their differentiation (Goguet-Rubio et al. 2016; Baksh et al. 2020), whereas it enhances ISC self-renewal (Schell et al. 2017; Bensard et al. 2020). Altogether, these data suggest cell type-specific effects of glucose metabolism on stem cell proliferation and fate. Notably, blocking glucose oxidation in the TCA cycle triggers compensatory oxidation of other substrates, including FAs, which may provide a link between decreased glucose availability and enhanced fatty acid oxidation in stem cells upon dietary interventions (Corton et al.

2004; Sikder et al. 2018; Steinhauser et al. 2018; Rosenbaum et al. 2019).

Pathogens and microbiota can alter the metabolic milieu of tissue stem cell niches

Diet can confer profound changes to the composition of gastrointestinal microbiota, which can subsequently alter the local metabolic milieu to which stem cells are exposed (Schulz et al. 2014; Olson et al. 2018). One of the most fascinating examples of the effects of microbes and pathogens to impact the metabolic milieu of stem cell environments comes from helminths, common intestinal parasites that, if allowed to persist, can elicit an inflammatory response. Recently, it was shown that helminth-derived metabolites, specifically succinate, can stimulate resident innate immune cells, called ILC2s, that respond by producing IL-13. In turn, IL-13 skews the ISCs to produce an excess of Tuft cells, which in turn secrete IL-25, which then further fuels ILC2 proliferation to create a vicious inflammatory circuit (Schneider et al. 2018). Thus, metabolites derived from gastrointestinal helminths can skew cell fate of ISCs and bias differentiation toward a specific lineage.

While the identification of certain metabolites that dictate stem cell behavior is ongoing, the source of these metabolites continues to elude the field. However, in certain circumstances, the gastrointestinal microbiota is emerging as a key source of these metabolites. *Lactobacillus*-derived lactate promoted intestinal epithelial turnover in the colon, suggesting this microbiota-derived source of carbon can augment stem cell self-renewal (Okada et al. 2013). In an esophageal tumor initiation model, the high-fat diet-associated increase in Lgr5⁺ progenitors was dependent on the microbiota within the gastrointestinal tract, as germ-free mice did not display the same propensity for tumor formation, and fecal microbiota transfer from mice fed a high-fat diet to mice fed the control diet also displayed increased tumor formation (Münch et al. 2019). Although a molecular mechanism is still needed that would link metabolites produced by distinct species of microbiota, general dysbiosis caused by a high-fat diet has been associated with tumor progression through lack of secreted butyrate as a waste product by the microbiota (Schulz et al. 2014). Of note, butyrate can also serve as an HDAC inhibitor (Candido et al. 1978). Thus, both a high-fat diet and ketogenesis can enhance stemness through enforcing self-renewal via microbiota-derived butyrate and β -OHB accumulation, respectively (Fig. 2). Since butyrate is a more potent HDAC inhibitor than β -OHB (Shimazu et al. 2013), a high-fat diet may be enforcing gene expression to enable carcinogenesis and may explain why increased carcinogenesis is not observed under ketogenic conditions.

Pathogens additionally elicit profound changes in organismal metabolism that may influence stem cell behavior. For example, induction of anorexia or feeding behaviors in response to pathogens influences host survival (Wang et al. 2016; Rao et al. 2017). Glucose availability and utilization by the host in response to pathogens

has been linked to pathogen virulence and host survival (Wang et al. 2016; Sanchez et al. 2018). Given the known effects of diet on stem cell behavior and the critical role stem cells play in response to pathogens and tissue injury, it will be of interest to interrogate more deeply how microbes, organismal metabolism, and stem cell behavior intersect during infectious processes.

Fasting and refeed states

While many studies have demonstrated effects of fasting on stem cell behavior, these studies do not focus on how physiologic refeeding after fasting influences stem cell behavior and the tissue and lineage-dependent differences in stem cell fate due to overall metabolic status of an organism (fasting or fed). Furthermore, nonautonomous factors that integrate into the control of stem cells during fasting and refeeding have largely been unexplored. Studies in the *Drosophila* midgut have demonstrated refeeding after nutrient deprivation initiates signals for ISCs to proliferate. These responses were nonautonomous and dependent on insulin signaling (Choi et al. 2011; O'Brien et al. 2011). Furthermore, intrinsic insulin signaling regulates the stability of adherens junctions between ISCs and their daughter cells, as insulin receptor knockdown ISCs increase the stability of these junctions to prohibit ISC proliferation (Choi et al. 2011). Cycles of prolonged fasting increased survival in mice subjected to chemotherapy-induced bone marrow ablation and increased chimerism in noncompetitive transplant models (Cheng et al. 2014). Strikingly, fasting specifically depleted circulating white blood cells and increased HSC numbers, but these numbers were rescued by refeeding. In the Peyer's patches of the small intestine, fasting-refeeding markedly depleted tissue B-cells and caused some to migrate to the bone marrow and return to the Peyer's patches upon refeeding (Nagai et al. 2019). In the central nervous system, refeeding stimulates hypothalamic neurons that subsequently activate a subset of NSCs to proliferate, driving production of specific neuronal subtypes (Paul et al. 2017). These results imply that the transition between fasting and refeeding can dynamically change the tissue microenvironment, highlighting the importance of elucidating the underlying mechanisms during this metabolic transition. They also implicate heretofore unidentified growth factor or hormonal signals as potential mediators of these phenotypes and the dynamic balance of stem cell state during periods of fasting and refeeding, independent of diet-induced changes in nutrient availability. Thus, both nutrient availability and hormonal changes may exert discrete effects on stem cell function.

Effects of generalized metabolic stress

Macroscopic dietary interventions that prompt similar changes in stem cell behavior may engage the same metabolic signaling pathways to enable cells to mount a response to subsequent nutrient stresses. Caloric restriction/prolonged fasting, ketogenesis, and a high-fat

diet all diminish the levels of circulating glucose. Numerous intracellular signaling pathways sense systemic glucose availability. mTORC1 senses glucose through the glycolytic intermediate dihydroxyacetone phosphate in a Rag-GTPase-dependent mechanism (Efeyan et al. 2013; Orozco et al. 2020). Although its effects on stem cell behavior remain to be explored, carbohydrate response element binding protein (ChREBP) functions as a glucose sensor in the liver by binding to promoters of glucose-sensitive genes upon activation by glycolytic intermediates together with its obligate binding partner Max-like protein X (MLX) (Yamashita et al. 2001; Stoeckman et al. 2004; Ma et al. 2005, 2006). AMPK may play a role if these systemic dietary interventions result in a lack of biosynthetic materials for growth, as AMPK turns on in these circumstances (González et al. 2020). Additionally, many of these diets increase PPAR signaling due to their intimate ties with lipid metabolism. Insulin sensitivity can also change in response to dietary interventions (Kinzig et al. 2010; Rosenbaum et al. 2019). However, the effect of dietary changes on stem cell-intrinsic insulin signaling remains mostly unexplored. The sole exception is in ISCs where insulin receptor ablation prevents ISC proliferation by stabilizing adherens junctions (Choi et al. 2011). Overall, this area represents a relatively unexplored avenue in the field that will help to better understand the similarities and differences between these dietary interactions.

How diet can impact cancer

Stem cells are increasingly recognized as the root of malignancy, as oncogenic mutations lock stem cells in states of ongoing self-renewal that facilitate tumorigenesis (Bjerkvig et al. 2005; Kreso and Dick 2014; Ge and Fuchs 2018). Consistent with the effects of organismal metabolism on stem cell behavior, diet can confer profound effects on the course of tumor initiation and progression in vivo. In ad libitum-fed mice, intestinal stem cells but not progenitors represent the cell of origin of intestinal adenomas and adenocarcinomas (Barker et al. 2009). Strikingly, a high-fat diet was sufficient to bestow tumor-forming capacity in premalignant intestinal progenitors independent of the niche. Enforced PPAR δ signaling through pharmacologic administration of a PPAR δ agonist was able to phenocopy this finding, suggesting that a high-fat diet works at least in part through PPAR δ signaling (Beyaz et al. 2016).

Intiguously, a high-fat diet, and not an oncogenic mutation, served as the so-called second hit that enables tumor initiation in this one-hit model in ISCs. These data demonstrate that exogenous metabolic shifts in homeostasis imparted tumor-initiating capacity to a cell type that is not normally recognized as tumor-initiating. The fact that intermittent fasting has not yielded the same propensity for tumor formation suggests that a high-fat diet and fasting differ with regard to the quantity or strength of the PPAR ligands generated by the dietary intervention. In the same vein, administration of a Western diet or perturbation of cholesterol homeostasis altered stem cell fate

during homeostasis and during tumor initiation. A Western diet increased the number of polyps formed in premalignant mouse models of colorectal carcinoma (Baltgalvis et al. 2009). The increased tumorigenic features of a Western diet on premalignant ISCs suggests that a Western diet also promotes stem cell self-renewal. That said, in the hematopoietic system, a Western diet skewed hematopoietic stem cell fate from lymphoid to myeloid lineages (Christ et al. 2018), a feature that may also play out in the intestinal stem cell niche, where multiple ISC fates including those altering inflammation are also possible.

Concordant with the data in ISCs, a high-fat diet administered to mice during an esophageal carcinoma tumor initiation model increased *Lgr5*⁺ progenitor cells and promoted tumor growth. These mice had premalignant esophageal lesions, and a high-fat diet again acted as the second hit to initiate tumors *in situ* (Münch et al. 2019). Overall, it is becoming clear that the metabolic state of an organism serves as a key constituent of the tissue milieu that can govern stem cell fate during homeostasis and disease.

In a model of oral squamous cell carcinoma, a high-fat diet promoted metastasis in a CD36-dependent fashion (Pascual et al. 2017). Since the tumor-initiating cells in squamous cell carcinoma normally up-regulate the fatty acid transporter CD36, the findings suggest that uptake of fatty acids into tumor-initiating cells, and likely their oxidation, is at least partially responsible for the observed increase in metastasis after a high-fat diet. Similarly, CD36-mediated fatty acid uptake enhances leukemic stem cell resistance to chemotherapy, promoting disease progression (Ye et al. 2016). Overall, these phenotypes begin to explain the pro-stemness effects of the dietary manipulations discussed above (Fig. 2). Might therapeutically increasing fatty acid oxidation provide a tempting new avenue to safeguard stem cell populations across tissue types? Perhaps, but since prolonged bias toward self-renewal potentially confers increased cancer risk, more work must be done to tease apart which permissive factors enable increased self-renewal without promoting carcinogenesis.

Future directions

Metabolic pathways for future investigation

Although much is known about how these dietary interventions alter glucose and lipid metabolism, how the dietary interventions affect other central carbon pathways is relatively unexplored (Box 1). With regard to central carbon metabolism, more work must be done to understand exactly how these dietary interventions affect central carbon pathways such as glycolysis, the pentose phosphate pathway, TCA cycle, and amino acid biosynthesis, especially through organismal isotopic tracing studies as has been previously described during conditions of standard mouse chow (Hui et al. 2017). For example, does decreasing glucose availability lower the rate of glycolysis and limit intermediaries used for amino acid synthesis? Do these changes in amino acid synthesis exert downstream

effects on cell fate by altering the availability of molecules that govern the epigenome of a cell (Box 1)?

Amino acid availability and metabolism are known regulators of stemness (Shyh-Chang et al. 2013; Carey et al. 2015; Taya et al. 2016; Wang et al. 2019; Baksh et al. 2020). Similar to dietary interventions, such as a high-fat diet, that abrogate insulin sensitivity, increased circulating levels of branched-chain amino acids (BCAAs) have long been associated with insulin resistance and type 2 diabetes (Newgard 2012). Notably, BCAA metabolism can modulate stem cell function through the enzyme branched chain amino transaminase 1 (BCAT1), which converts BCAAs to branched-chain ketoacids through transamination of α -ketoglutarate (α KG) to glutamate.

α KG is a key TCA cycle intermediate in the mitochondrial matrix. However, α KG also acts as a cosubstrate for a class of enzymes called α KG-dependent dioxygenases, many of which serve as histone and DNA demethylases and can enact changes in nuclear gene expression. Furthermore, α KG's immediate downstream metabolite in the TCA cycle, succinate, can competitively inhibit α KG-dependent dioxygenases and antagonize the effects of α KG (Schvartzman et al. 2018).

Leukemia cells up-regulate BCAT1 and deplete α KG, which results in inhibition of ten-eleven translocation (Tet) DNA demethylases and subsequent DNA hypermethylation (Hattori et al. 2017; Raffel et al. 2017). While maintenance of increased total α KG:succinate pools maintains pluripotency in mouse embryonic stem cells, α KG accumulation has been implicated as a prodifferentiation factor across several tissue types, such as premalignant epidermal and intestinal stem cells and pancreatic cancer cells (Morris et al. 2019; Baksh and Finley 2020; Baksh et al. 2020; Tran et al. 2020). Notably, other transaminases such as phosphoserine aminotransferase (PSAT1), alanine aminotransferase (GPT1/2), and aspartate aminotransferase (GOT1/2) can also contribute to cytosolic accumulation or depletion of α KG. For example, activation or suppression of the serine biosynthesis pathway can alter α KG levels through modulation of the activity of PSAT1 (Possemato et al. 2011; Baksh et al. 2020). The fact that α KG can alter gene expression and stem cell fate also suggests that accumulation or reduction of this metabolite must eventually occur in the nucleocytoplasmic compartment, possibly through the action of these

Box 1. Future directions and outstanding questions

- How do dietary interventions affect pathways involved in central carbon metabolism?
- Can dietary interventions regulate the epigenome through modulation of intracellular metabolite levels?
- How do dietary interventions reshape the stem cell niche?
- How might nonautonomously produced growth factors signal to stem cells and prime them to adapt to tissue stress?

transaminases or through the mitochondrial α KG/malate carrier protein (SLC25A11) (Palmieri et al. 1972). Therefore, exploring the role of BCAAs and understanding whether any other amino acids that feed into these pathways are altered in the macroscopic dietary interventions described herein may be an interesting avenue of further research in understanding how systemic metabolic perturbations regulate stem cell fate in vivo.

Dietary interventions can fundamentally change the cellular composition of the niche

There is evidence that these macroscopic dietary interventions can substantially change the cellular census of stem cell niche (Box 1). For example, caloric restriction can reshape the local HFSC niche by decreasing white adipose depots in the dermis (Forni et al. 2017). Interestingly, local white adipocytes can maintain HFSC quiescence by providing inhibitory BMP2 (Plikus et al. 2008). Paradoxically however, contraction of the local white adipose tissue due to caloric restriction would disinhibit HFSCs and enable activation of the hair cycle, contrary to the observations of decreased hair turnover. Further work will be needed to reconcile these differences as well as elucidate the compositional changes across tissue types induced by macroscopic dietary perturbations. Taking cues from the ISC niche, further investigation is warranted to determine how sensitive or responsive the stem cells in these tissues are to caloric restriction versus any identified or unidentified niche players in their local microenvironments that regulate stem cell fate similar to Paneth cells.

Conclusion

Stem cells integrate cues from their local niche to inform their behavior and cell fate. While metabolites continue to emerge as key members of the niche, there has been little focus on how the macroscopic dietary status of an organism might contribute to stem cell behavior. Comparisons between caloric restriction/prolonged fasting, the ketogenic diet, and high-fat diets suggest the importance of the balance between glucose and fatty acid utilization as oxidative substrates and beg the question why these disparate dietary interactions converge on similar results. Thus, a closer investigation of how these major dietary interventions augment the balance of intracellular metabolites and their respective biosynthetic and catabolic pathways is warranted. Studies from fasting and refeeding implicate nonautonomous production of insulin as a mediator. Additionally, this finding implies that other types of growth factors can signal to stem cells how their behavior might change to adapt to local tissue stress, independent of differences in diet-induced nutrient availability (Box 1). Overall, more work must be done to better link how macroscopic dietary changes impact the overall metabolic status of an organism, how this status contributes to the stem cell niche, and finally how these changes translate into changes in stem cell fate and behavior. A better understanding of these processes will lead to a bet-

ter understanding of homeostatic and disease states as well as suggest new avenues of inquiry for therapeutic intervention in a wide array of diseases, especially those characterized by states of inflammation and cancer.

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We are grateful to the many friends and colleagues in the stem cell and metabolism fields whose work inspired us to write this review. We centered this review largely on three common dietary restrictions, drawing parallels and distinctions between them and the responses they elicit from tissue stem cells. While text constraints limited our focus, we tried to illustrate paradigms gained from examining the impact of pathogens, microbiota, and stress, including oncogenic stress, on metabolism in the tissue microenvironment and its resident stem cells. Similarly, due to reference constraints, we regret that we could not cite all of the worthy papers on this topic. From our treatise, we hope that the exciting questions in stem cell biology and future avenues for exploring how tissue stem cells cope with metabolic changes will shine through and inspire new work. E.F. is an Investigator of the Howard Hughes Medical Institute and the Rebecca C. Lancefield Professor. J.S.S.N. and S.C.B. are predoctoral fellows of the Weill Cornell/Rockefeller/Sloan Kettering Tri-Institutional Medical Scientist Training Program (T32-GM007739). S.C.B. is a Ruth Kirschstein National Institutes of Health (NIH) predoctoral fellow (F31-CA236465). This work was supported by grants from the NIH (R01-AR050452, R01-AR31737, R01-AR27883, E.F.), the Starr Foundation (E.F., J.S.S.N., and S.C.B.), and NYSYSTEM C32585GG (E.F.).

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