Unintentional excessive weight loss has been documented in association with chronic illness throughout medical history. Dating back to ancient Greece (460–377 BC), Hippocrates wrote “the flesh is consumed and becomes water … the abdomen fills with water; the feet and legs swell, the shoulders, clavicles, chest and thighs melt away”1. This description contains most of the main elements by which cachexia was defined in the early medical records2,3. Cachexia typically manifests itself alongside chronic inflammatory illnesses, various aggressive cancers (such as gastrointestinal and lung cancers) and chronic infections, including HIV infection and Mycobacterium tuberculosis infection4–6. The severe emaciation resulting from cachexia not only has significant mental health implications but can eventually reach a debilitating state that renders the patient incapable of fulfilling basic daily needs. Depleted muscle mass and strength may result in cardiac arrhythmias, respiratory weakness and other complications that lead to premature death7. The latest international consensus defined cancer-associated cachexia (CAC) as “a multifactorial syndrome characterized by an ongoing loss of skeletal muscle mass (with or without loss of fat mass) that cannot be fully reversed by conventional nutritional support and is driven by a variable combination of reduced food intake and abnormal metabolism”7. This definition is relevant for both CAC and infection-associated cachexia (IAC) and provides a clear framework representing the most consistent features of cachexia from a clinical and metabolic perspective. Notably, this definition does not address the immunological aspects of the syndrome, likely due to the limited data that are currently available in this area. However, while we still do not fully understand why cachexia occurs, it is clear that the onset of cachexia is driven by the inflammatory programme triggered by the underlying illness. In this Review, we focus on the immunological aspects of cachexia within cancer and infection to better understand the commonalities and distinctions within the elements involved in the cachexia programme. Additionally, we will discuss the immunometabolic consequences of cachexia by highlighting the pathways through which immune cell activation can influence and be influenced by altered metabolic environments.

Cytokines and cachexia
Much of the immunological data available from patients with cachexia are derived from measurements of circulating cytokines and acute-phase proteins. Tumour necrosis factor (TNF), IL-1β, IL-6 and interferon-γ (IFNγ) are among the most consistently upregulated cytokines in cachexia4. These cytokines are abundantly expressed by both immune and non-immune cell types downstream of pattern-recognition receptor activation by pathogen-associated molecular patterns and/or damage-associated molecular patterns8. This leads to activation of JAK–STAT and NF-κB signalling pathways and downstream transcriptional regulation that can induce various catabolic pathways in muscles and adipose tissue9,10. More studies are required to clarify the precise roles of pattern-recognition receptor activation in cachexia-inducing diseases; in the following sections...
we discuss our current understanding of cytokines in cachexia associated with cancer or infection.

**Classical cachexia-associated cytokines**

The link between immunology and cachexia first became a point of interest after the isolation of TNF\(^1\), which was known as ‘cachectin’ at that time\(^2\). In 1985, Beutler et al. showed that adipocytes exposed to TNF suppress their expression of lipoprotein lipase (LPL), thereby reducing lipid uptake\(^1\). This built on the observations made by Rouzer and Cerami 5 years earlier, which showed that rabbits infected with *Trypanosoma brucei brucei* exhibit hypertriglyceridaemia as a result of reduced triglyceride clearance from the circulation\(^1\). These animals were also emaciated, a phenotype that was later shown in murine models to be a manifestation of cachexia\(^1\). Subsequent studies in adipocyte cultures showed that TNF promotes adipose triglyceride lipase (ATGL)-mediated lipolysis by depleting the ATGL-inhibitory protein G0/G1 switch protein 2 (G0S2)\(^3\). In myoblast cultures, exposure to TNF, IL-1β, IL-6 and IFNγ inhibits myoblast differentiation and induces protein loss through STAT3-mediated NF-κB activation\(^4\–\(^8\). These results were further validated in mouse models, where mice into which tumour cells overexpressing TNF, IL-1α, IL-6 or IFNγ had been implanted exhibited severe weight loss, adipose tissue depletion and muscle atrophy\(^9\–\(^12\) (Fig. 1a). In models where tumour cell lines overexpressing IL-6 or IFNγ were injected into mice, blockade of IL-6 or IFNγ, respectively, ameliorated the cachectic phenotype. Notably, the overexpression of IL-1α and IL-6 in these models also reduced overall survival and led to more aggressive tumour progression\(^9\–\(^12\). In a different approach, mice into which pancreatic tumour cells lacking IL-6 expression had been injected showed reduced loss of fat tissue and protection against muscle atrophy\(^9\). Further analysis of adipose tissue and muscles, both in vivo and in vitro, revealed a catabolic feedforward loop of IL-6 signalling between the tumour, the adipose tissue and the muscles\(^9\).

In addition to promoting catabolism through NF-xB activation, cytokines can promote cachexia through NF-xB-independent mechanisms. For instance, IL-1β signalling in the hypothalamus can trigger the activation of the hypothalamic–pituitary–adrenal axis, which ramps up glucocorticoid production, resulting in catabolic effects on both muscle tissue and adipose tissue\(^13\–\(^16\) (Fig. 1a). Administration of IL-1β through intracerebroventricular injection was sufficient to elicit muscle atrophy. This was associated with increased expression of factors involved in muscle catabolism, such as muscle atrophy F-box protein (MAFbx; also known as FBXO32 and atrogin 1), MURF1 (also known as E3 ubiquitin–protein ligase TRIM63) and forkhead box protein O1 (FOXO1), reduced muscle mass and reduced muscle fibre cross-sectional area\(^17\) (Fig. 2a). Intraperitoneal administration of IL-1β did not induce muscle loss\(^17\), suggesting that IL-1β acts predominantly by directly stimulating the hypothalamic–pituitary–adrenal axis. Of note, tissue mass comparison with a pair-fed group demonstrated that the loss of fat mass subsequent to IL-1β administration was coupled to food intake, whereas the loss of muscle mass occurred independently of food intake\(^17\).

In summary, these data show that pro-inflammatory cytokines such as TNF, IL-1, IL-6 and IFNγ are able to activate diverse catabolic processes across multiple organs in experimental models of CAC (FIGS 1, 2). However, targeting these cytokines in a clinical setting has shown underwhelming results. Over the years, a number of cytokine inhibitors, mainly those targeting TNF, have been tested in patients with cancer who were undergoing cachexia. Most of these treatments showed no significant increase in body weight, strength or survival\(^18\–\(^20\). In two small randomized controlled trials, patients with CAC were treated with thalidomide for its wide immunomodulatory functions and its ability to inhibit cytokines, including TNF\(^20\,2^1\). In these trials, thalidomide treatment resulted in a small increase of body weight and muscle mass, with no increase in hand-grip strength or survival\(^20\,2^1\). This highlights the possibility that cytokines are not singularly sufficient to induce cachexia, but can be elements of a more complex coordinated immune and metabolic response leading to cachexia.

Mechanistic data examining the role of cytokines during IAC are scarce, and available experimental models come with their own intrinsic limitations (BOX 1). Mice chronically infected with *Toxoplasma gondii*, an obligatory intracellular parasite, develop cachexia as evident by their severe weight loss and depletion of fat and lean mass\(^22\). This occurs within the first week of infection and coincides with elevated serum levels of cytokines, including TNF, IL-1β, IL-6 and IFNγ\(^22\). In this context, TNF and IL-1β support the production of IFNγ, which is essential in controlling *T. gondii* infection and limiting parasite replication\(^23\,2^4\) (Fig. 1b). Notably, animals deficient in TNF or IFNγ show accelerated death and an increase in parasite burdens\(^25\–\(^27\). On the other hand, studies that inhibit IL-1 signalling through genetic disruption of the IL-1 receptor (IL-1R) pathway have reported conflicting results. For instance, one study showed that disruption of IL-1 signalling through the abrogation of inflammesome activation in mice (using animals that lacked caspase 1 and caspase 11, NLRRP3, NLRRP1 or IL-1R) led to increased parasite burdens and reduced survival\(^28\). Meanwhile, a more recent study found that, compared with wild-type mice, IL-1R-deficient mice showed increased survival, reduced weight loss and increased lean and fat mass following *T. gondii* infection\(^29\). These effects were independent of pathogen clearance, brain inflammation, immune cell infiltration and anorexia. Notably, IL-1β signalling through the gut–brain signalling axis has been associated with the induction of anorexia in other models of infection\(^30\) (BOX 2).

In a model of IAC that involves chronic lymphocytic choriomeningitis virus (LCMV) infection with LCMV strain clone 13 (LCMV-G13) (BOX 1), the depletion of TNF, IFNγ or both cytokines together had no effect on weight loss even though the levels of both cytokines were increased in the serum during infection\(^31\). By contrast, abrogation of type 1 interferon signalling ameliorated weight loss in this model\(^31\). Importantly,
Relevant cytokines in the development of cachexia. a | Tumourigenesis is associated with the release of a wide range of cytokines by tumour cells, by the surrounding tissue and from innate and adaptive immune cells downstream of pattern-recognition receptor (PRR) activation by tumour cell-associated damage-associated molecular patterns. Tumour necrosis factor (TNF), interferon-γ (IFNγ), IL-6 and IL-1β are able to induce tissue catabolism by modulating gene-expression profiles in both adipose tissue and muscle cells. IL-1β also contributes to cachexia through the central nervous system (CNS), where it modulates food intake and activates the hypothalamic–pituitary–adrenal (HPA) axis. The subsequent release of glucocorticoids contributes to tissue catabolism. GDF15 release induces weight loss through effects on the CNS that modulate food intake and by increasing adipose tissue lipolysis through sympathetic nervous system (SNS) signalling. IL-20 also contributes to adipose tissue depletion, while IL-4 shows protective effects on muscle cells. Transforming growth factor-β (TGFβ) released from tumour cells and/or bone (during bone metastasis-induced osteoclastic resorption) can induce adipose tissue fibrosis and compromise muscle strength.

b | During parasitic infection, TNF and IL-1 release downstream of PRRs supports the production of IFNγ, which is important for controlling the pathogen load. IL-1 is also involved in the catabolism of adipose tissues and muscles. During viral infection, type I interferon (IFN) signalling to CD8+ T cells during antigen recognition and T cell activation is an important step in triggering cachexia.
**Fig. 2 | Mechanisms of myocyte and adipocyte catabolism.**

**a | Muscle cells**

Cytokines induce muscle catabolism by regulating the transcription of various genes. NF-κB signalling suppresses Myod1 expression, which leads to inhibition of myoblast differentiation. Forkhead box protein O (FOXO) activation results in its translocation to the nucleus, where it upregulates the expression of genes involved in proteasomal degradation (for example, Murf1 and Mafbx) and autophagy (for example, Becn1 and Map1lc3b2, which encode beclin 1 and LC3B, respectively). Transforming growth factor-β (TGFβ)–SMAD signalling increases β is induced through the binding of catecholamines to NF-κB, which activates cAMP phosphorylation of protein kinase A (PKA) downstream of GNAS and adenylyl cyclase. PKA is then able to phosphorylate p-SMAD, phosphorylated SMAD; SR, sarcoplasmic reticulum; TG, triacylglycerol.

**b | Adipocytes**

Adipocyte triglyceride lipase (ATGL) and hormone-sensitive lipase (HSL), resulting in their translocation to the surface of lipid droplets, where lipolysis occurs. ATGL-mediated lipolysis is modulated through its interaction with G0/G1 switch protein 2 (G0S2) and CGI58. Cytokine signalling in the adipose tissue induces transcriptional changes downstream of NF-κB and STAT signalling, resulting in an increase in the levels of lipolytic enzymes and a suppression of genes involved in lipogenesis. Perilipin 1 (PLIN1) encoded by PLIN1 coats the surface of the lipid droplet and is also transcriptionally suppressed during lipolysis to expose a wider surface area to lipase activity. IFNγ, interferon-γ; IFN, interferon; IFNγR, interferon-γ receptor; IL-6R, IL-6 receptor; NEFA, non-esterified fatty acid; p-HSL, phosphorylated HSL; p-SMAD, phosphorylated SMAD; SR, sarcoplasmic reticulum; TG, triacylglycerol; TGFβR, TGFβ receptor; TNF, tumour necrosis factor; TNFR, tumour necrosis factor receptor.

**Reviews**

**Mechanisms of myocyte and adipocyte catabolism.**

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**Other cytokines associated with cachexia**

In the preceding section, we discussed some of the key inflammatory cytokines that have been associated with cachexia. However, in both cancer and infection the concentration levels of numerous other cytokines are increased in both the circulation and within local tissue environments. Many of these cytokines have not been systematically probed for their role in the pathophysiology of cachexia. In the following subsections we discuss a few additional examples of cytokines that have been implicated in the development of cachexia.

**II-20.** II-20 is a pro-inflammatory cytokine produced mainly by haematopoietic cells, facilitating their communication with epithelial cells. It has an important role in enhancing innate defence mechanisms and tissue repair processes at epithelial surfaces. II-20 is highly expressed in the tumours of patients with pancreatic cancer. In mouse models of pancreatic ductal adenocarcinoma (PDAC), levels of II-20 correlate with increased tumour fibrosis, PDL1 expression and an overall reduction in survival. In the cachectic mouse models of PDAC and Lewis lung carcinoma (LLC), anti-II-20 treatment ameliorated body weight loss and prevented loss of adipose tissue mass by decreasing ATGL and hormone-sensitive lipase (HSL) expression. However, this treatment had no effect on muscle atrophy, supporting that the pro-cachectic function of II-20 is linked to its influence on the adipose tissue.

**IL-4.** IL-4 is produced by both innate and adaptive immune cells during type 2 immune responses. A recent study demonstrated a protective role for IL-4 in the C26 colon cancer model. Daily administration of IL-4 to C26 tumour-bearing mice prevented excessive weight loss, improved physical performance and muscle protein synthesis and regeneration, and increased survival. Moreover, IL-4 administration increased tumour necrosis and promoted CD8+ T cell and macrophage infiltration into tumours. In the context of muscle injury...
induced by cardiotoxin administration, IL-4 secretion activates STAT6 signalling in fibroadipogenic progenitor (FAP) cells. This leads to their differentiation into fibroblast cells that are capable of phagocytosing cellular debris, thereby promoting muscle regeneration. However, FAPs may also differentiate into adipocytes and, in this context, can be detrimental to muscle growth. For instance, glycerol-mediated adipocytes and, in this context, can be detrimental for investigations both into the fate of FAPs during different models of CAC and for understanding the potential protective effect of IL-4 administration in CAC (FIG. 1a). Similarly to IL-4, other cytokines with anabolic effects on muscles warrant further study. This includes IL-15, which is known to induce myogenesis and mitigate the catabolic effect of TNF in cultured myotubes.

**TGFβ family cytokines.** Transforming growth factor-β (TGFβ) signalling is involved in multiple aspects of cancer, including tumour suppression, cell-cycle arrest, modulation of the tumour microenvironment and tumour metastasis. In the context of cachexia, biopsies of tumours and adipose tissue from patients with cancer with cachexia showed an association between high levels of TGFβ and tissue fibrosis that is not seen in weight-stable patients with cancer.

In mouse models of bone metastasis, bone-derived TGFβ directly affected muscle strength and mediated the phosphorylation of SMAD signalling factors. This resulted in increased

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**Box 1 | Experimental models of cachexia**

Experimental models are indispensable in studying cachexia due to the limited data that can be derived from patients with cachexia. Mechanistic studies relied on the use of cell lines and animal models to map out the cachectic pathways leading to muscle and adipose tissue depletion. Cell lines such as C2C12 myoblasts and 3T3-L1 adipocytes provided the most simplified models to closely examine the effects of stimulants such as cytokines and tumour-cell conditioned media. The repertoire of animal models of cachexia is heterogeneous and keeps evolving to better represent human cachexia. Here we discuss some of the important sources of variations across cancer-associated cachexia (CAC) models and their limitations, and provide a summary of available models for the study of infection-associated cachexia (IAC).

**CAC**

Most of the available mechanistic data on cachexia have been derived from CAC mouse models that rely either on tumour cell implantation or genetically engineered mice. For tumour cell implantation models, the cancer cell type used and the route of administration are important factors to consider. The genetic background of tumour cell lines depend on the mouse strain of origin and on whether they are derived from naturally occurring tumours (for example, the Lewis lung carcinoma lines established by Margaret Lewis in 1951) or are chemically induced (for example, C26 colon cancer cells isolated from mice exposed to the carcinogen N-nitroso-N-methylurethane). These and other tumour cells are injected either ectopically through subcutaneous, intraperitoneal or intramuscular injection or orthotopically, by injecting, for example, pancreatic tumour cells directly into the pancreas. Both routes have their limitations and result in phenotypical variations in tumour progression and the onset of cachexia. Orthotopic injections are invasive and require surgery, which introduces complications and variables that are difficult to control. Ectopically injected tumours grow at a very high rate, and manifest cachexia at advanced stages of tumour progression, leaving a narrow window to study cachexia before termination of the experiment. Even when ectopic injections are used, variability can occur depending on the site, affecting the pathophysiology and/or time frame of cachexia development, which also affects the rate of tumour metastasis. Human-derived tumours implanted in immune-deficient mice have also been used to allow a better representation of human cancer cachexia and its response to therapeutic agents; however, these models have a major caveat of lacking proper antitumour immune responses. Genetically engineered mouse models attempt to produce naturally occurring tumours with mutational signatures similar to those seen in human cancer. Among these models, certain considerations must be taken into account, including the age at which tumour development occurs and the effect of the genetic mutations on mouse development and tissue function.

**IAC**

Few models of infection have been studied in the context of cachexia. The parasites *Toxoplasma gondii* and *Trypanosoma cruzi* provide valuable experimental models of IAC with a gradual manifestation of cachexia sustained over a longer time frame, which better simulates the prolonged nature of the syndrome in humans and allows long-term investigations into therapeutic opportunities. When *Toxoplasma* cysts are ingested, mice lose up to 20% of their initial body weight within the first 10 days after infection. This weight loss is sustained for more than 90 days after infection, independently of the parasite load. During *Trypanosoma* infection, weight loss occurs within a time frame between 20 and 40 days after infection. Lymphocytic choriomeningitis virus (LCMV) is a negative-strand RNA virus from the family Arenaviridae that provides a viral infection model of IAC, whereby cachexia manifests itself within 1 week of infection and gradually resolves thereafter. Importantly, this noncytolytic virus elicits a strong antiviral immune response led by type I interferon signalling and followed by a robust CD8+ T cell response. As CD8+ T cells clear infected cells, their cytolytic activity drives the immunopathology associated with the infection. In the case of chronic infection with LCMV strain clone C113, CD8+ T cell are also the main triggers for IAC by a still unknown mechanism. Therefore, this model provides a unique opportunity to study the cachexia-related immune response with presumably little direct involvement of the pathogen and offers defined experimental variations (for example, viral strain, dose, route of infection) that lead to distinct immune and metabolic phenotypes. This and the well-characterized immune responses of LCMV infection make it a versatile tool to study immunometabolic adaptation.
Immune cells in cachexia

In response to tumorigenesis or pathogenic infections, the immune system launches cellular responses that are specifically tailored to each illness with various degrees of overlap. Although data on the link between immune cell function and cachexia are currently limited, a number of recent studies indicate the involvement of multiple immune cell types, including macrophages, neutrophils, myeloid-derived suppressor cells (MDSCs) and T cells (Fig. 3).

Innate immune cells

Studies using the PDAC mouse model of cachexia demonstrated a significant increase in myeloid immune cell infiltration into the tissue. These infiltrates targeted specific brain regions that influence feeding behaviour and/or energy metabolism, such as the hypothalamus. This region is particularly sensitive to inflammation and is highly responsive to changes in the systemic nutritional state, making it a likely target of immune-mediated metabolic regulation during cachexia. In this tumour model, an increased number of microglial cells in the hypothalamus was found to be associated with increased IL-1β and arginase 1 secretion. Reducing microglial cell numbers through treatment with a colony-stimulating factor 1 receptor (CSF1R) inhibitor exacerbated the cachectic phenotype, affecting food intake, activity and muscle atrophy, with no measured effects on tumour size. This suggests a protective role for microglial cells in mitigating cachexia during pancreatic tumour development. Interestingly, microglial cells were shown to phagocytose infiltrating CC-chemokine receptor 2 (CCR2)-positive neutrophils in the central nervous system parenchyma. In the absence of CCR2, tumour-bearing mice exhibited reduced anorexia and muscle atrophy that correlated with reduced levels of neutrophil infiltrates. Hence, it has been speculated that phagocytosis of neutrophils by microglial cells could provide a degree of protection against cachexia (Fig. 5a).

A similar protective effect of macrophages in cachexia was observed in adipose tissue in a mouse model of hepatocellular carcinoma. In that study, mice with a myeloid-specific deficiency in hypoxia-inducible factor 1α (HIF1α) — which show defective myeloid cell activation — had increased adipose tissue depletion during genetically induced hepatocellular carcinoma, and this phenotype was coupled to reduced macrophage infiltration into adipose tissue. Importantly, this defective myeloid cell activity had no influence on muscles, liver tissue or tumour size. By contrast, macrophages seemed to contribute to the tissue catabolism observed in cachexia associated with the mouse PDAC and LLC models. In both models the adipose tissue-protective effect observed following anti-IL-20 treatment (discussed earlier) was associated with reduced macrophage infiltration into adipose tissue. Moreover, the depletion of macrophages in the PDAC model using clodronate-filled liposomes resulted in reduced tumour sizes and increased body weight, muscle mass and strength coupled with a reduction in the levels of markers of proteasomal degradation. This apparent
discrepancy could potentially be explained by differences in macrophage polarization within each tissue environment (Fig. 3b,c).

Macrophages are well known for their roles in adipose tissue inflammation and metabolism during ageing and obesity1, and the nature of their interaction with the tissue is highly influenced by their polarization. In the context of obesity, there is a shift in the abundance of ‘M2-like’ macrophages to ‘M1-like’ macrophages22. In this setting, macrophages sequester noradrenaline from adipocytes by increasing noradrenaline import and degradation, which results in downregulated lipolysis and adipose tissue enlargement23,24. On the other hand, M2-like macrophages promoted lipolysis in a mouse model of cold-induced thermogenesis via the production of noradrenaline downstream of IL-4 stimulation25. Muscle biopsy samples taken from patients with pancreatic cancer showed an inverse correlation between the density of CD163+ M2-like macrophages and muscle fibre in cross-sectional areas26. In line with this, myotubes (derived from the C2C12 cell line) co-cultured with M2-polarized macrophages and exposed to tumour-conditioned medium showed a reduction in myotube thickness and protein content and an increase in the levels of proteasomal degradation markers.

In a mouse model of peritonitis-induced sepsis with concomitant loss of adipose tissue and muscle mass, single-cell RNA sequencing analysis revealed that macrophage populations increase in muscles and decrease in adipose tissue 1 day after intraperitoneal injection of faecal slurry27. This was coupled with an increase in the levels of muscle-infiltrating neutrophils, natural killer cells and T cells that remained elevated in both the muscle and the adipose tissue 1 month after injection. Although the transcriptional profiles of these cell populations showed, as expected, a role of pathways involved in infection and in the tissue damage response, further analysis focused on metabolic regulation could reveal valuable information with regard to tissue catabolism.

MDSCs constitute a heterogeneous population of myeloid cells with immunosuppressive functions28. They are found in large numbers in the tumours of patients with gastric and pancreatic cancers29,30 and in several mouse models of CAC, such as 4T1 mammary carcinoma, C26 colon adenocarcinoma and LLC31. Levels of MDSCs in the spleen and bone marrow correlate with total body weight loss, adipose tissue depletion and increased oxygen consumption rate32,33. However, the exact mechanism by which MDSC population expansion could contribute to cachexia is still unknown (Fig. 3d), and the currently available models for MDSC depletion, such as clodronate-mediated liposomal depletion, are not specific to MDSCs but also eliminate macrophages and other myeloid cells34.

Adaptive immune cells

The role of T cells has been extensively studied in the context of antitumour and infectious immunity, but whether and how they influence systemic metabolic reprogramming in the context of cachexia remains
elusive. Muscle biopsies of patients with different cancers, most of whom had gastrointestinal cancers, showed a positive correlation between the abundance of CD3−CD4+ cells (presumably CD8+ T cells) and muscle fibre cross-sectional area. In a separate cohort of patients with cancer who showed early signs of muscle impairment, circulating levels of recent thymic immigrant and effector memory CD8+ T cells were associated with increased muscle mass. Meanwhile, the abundance of regulatory T (Treg) cells and central memory T cells showed a negative correlation with muscle mass. These studies suggest that an efficient and robust antitumour CD8+ T cell response can protect against muscle catabolism during CAC, while immune suppression may have the opposite effect (Fig. 3). The reason for this protection is unclear as there is no evidence so far that uncouples the antitumour activity of the T cells from their direct catabolic effect on muscles. T cell-mediated tumour clearance may well be the predominant mechanism preventing muscle atrophy. However, another and not mutually exclusive mechanism may be that the immune cell composition within the local tissue environment alters the functionality and secretory profile of each cell type, with consequences for muscle atrophy. An indication of this could be derived from measurements of circulating immune cells, where the abundance of immune cell populations in relation to one another may also have an impact on the prognosis of patients with cancer. A high ratio of the number of circulating neutrophils to the number of circulating lymphocytes, for instance, correlates with the development of cachexia. The reason for this is unknown, and ongoing efforts aim to better understand the role of neutrophils within the circulation and the tumour microenvironment.

In the mouse model of LLC, increasing the numbers of CD4+CD44+ T cells in the spleen and lymph nodes through the adoptive transfer of these cells delayed the onset of cachexia and prevented muscle loss. The adoptive transfer of CD4+FOXP3+ Treg cells to mice infected with T. gondii reduced weight loss and prolonged survival, highlighting the regulatory function of CD4+ T cells as a potential mechanism for reducing cachexia during parasitic infection (Fig. 3). By contrast, in the chronic LCMV-CI13 infection model, loss of CD4+ T cells had no effect on the initiation of cachexia, while the activation of antigen-specific CD8+ T cells was essential to trigger IAC (Fig. 3). These differences could be related to the different types of inflammatory milieu, kinetics and immune responses that are associated with tumours, parasitic infections and viral infections. More studies are required to systematically investigate and compare how T cells affect the course of cachexia in both IAC and CAC (Box 1).

Interestingly, IAC during LCMV infection was both viral dose and strain dependent, which may be linked to differences in the associated T cell responses. Infection with LCMV-CI13 (which causes a chronic viral infection) but not with LCMV strain Armstrong (LCMV-Arm; which causes an acute infection) results in cachexia and induces T cell exhaustion, which is characterized by low levels of T cell proliferation, reduced cytokine potential and high expression of inhibitory receptors, including PD1, CTLA4 and LAG3 (Ref. 44). The exhausted phenotype is established early on during T cell priming and is seen in chronic infections and other chronic diseases such as cancer. The expression of inhibitory receptors such as PD1 and CTLA4 has been shown to alter the metabolic reliance of CD8+ T cells by promoting oxidative phosphorylation and suppressing glycolysis. This raises the question of whether the T cell exhaustion programme may be connected to the systemic immunometabolic changes preceding the development of cachexia. Likewise, cachexia-associated metabolic changes may also impact the differentiation of exhausted T cells.

In the LCMV-CI13 IAC model, the muscle microenvironment is depleted of certain cytokines, such as IL-12, IFNβ and IFNγ, which provides an environment for CD8+ T cells to thrive. These cells exhibited a lower degree of exhaustion in the muscle compared with that seen in the splenic compartment, and had a higher proliferative and cytokine-production capacity. The reduced exhaustion phenotype is facilitated by IL-15 within the muscle microenvironment. It is unclear how these immunologically distinct zones are formed and how they interplay with muscle atrophy at this stage of infection. However, one may speculate that these functional CD8+ T cells may directly take up muscle-derived amino acids to support their cytokine production and/or proliferation. In a different model of LCMV infection using LCMV strain Taub, infection of IFNγ-deficient mice, but not wild-type mice, resulted in severe weight loss that was CD8+ T cell dependent. Although it remains unclear whether the weight loss induced by LCMV-Taub infection is a manifestation of cachexia or anorexia (Box 2), it will be of interest to reconcile this observation with recent results relating to the IFNγ-shielded muscle microenvironment in the LCMV-CI13 model. In yet another variation of the LCMV model, the intracranial injection of LCMV-Arm in mice lacking MHC class I molecules results in a chronic weight loss attributed solely to anorexia.

Overall, the current evidence indicates that activation of immune cells can have both anticachectic and pro cachectic effects, and aspects other than immune cell population expansion and cytokine production can be involved. Other pertinent questions that require investigation include the pathophysiological role of cachexia itself in regulating processes relevant to infection, tumorigenesis and immune regulation, and the pathological consequences that come with chronic non-resolving cachexia (Box 3).

### Tissue catabolism downstream of immune responses

As highlighted by the consensus definition, the key pathophysiological characteristic of cachexia is the severe weight loss that occurs as a result of muscle atrophy and adipose tissue depletion. Muscle atrophy is one of the most extensively studied aspects of cachexia as it is a main contributor to functional impairment and death. It is identified by a reduction of overall muscle mass, reduced muscle fibre cross-sectional area and...
Here, we list some open questions that will be key for a better understanding of the link between immune responses and cachexia:

- Is cachexia a coordinated metabolic programme that facilitates nutrient accessibility in different disease contexts? If so, how do antitumour and/or antipathogen responses contribute to its initiation, maintenance and resolution?
- How is immune cell activation linked mechanistically to cachexia? Is tissue catabolism induced, at least in part, for the purpose of fuelling the energetically costly activation of the immune system?
- Aside from their capabilities to secrete cytokines, how do immune cells modulate tissue metabolism during cachexia?
- Do cachectic processes contribute to the ‘exhausted’ T cell phenotype that is common to cancer and chronic infections? Alternatively, is cachexia triggered downstream of T cell exhaustion?
- What are the consequences of altered inflammatory and metabolic environments during cachexia on the microbiota composition? Which feedback loops are elicited by such changes?
- What are the common and unique properties of cancer-associated cachexia and infection-associated cachexia? How can understanding this help the field to develop novel therapeutic strategies?

**Immunometabolic consequences of cachexia**

The onset of cachexia and the subsequent depletion of the energetic tissue reserves enrich the circulation with metabolic substrates that are likely to impact systemic metabolism, inflammation and immune cell activation. Using current knowledge derived from the field of immunometabolism, we discuss potential aspects of immunometabolic crosstalk during cachexia with a focus on fatty acids and amino acids as the main products of adipose tissue and muscle catabolism, respectively.

**Fatty acids and cachexia**

During cachexia, excessive adipose tissue lipolysis increases circulating levels of NEFAs and glycerol, which are then taken up and utilized by other organs. In mouse models of cachexia, metabolic cage measurements of the respiratory exchange ratio of the animal indicate a shift in the metabolic reliance from carbohydrates towards fat utilization. From the data available thus far, it is not possible to deduce whether this shift in metabolism is reactionary and occurs as result of increased exposure of cells to lipid substrates or whether it serves as a controlled step in an adaptive programme.

**NEFAs**

NEFAs can be categorized on the basis of their degree of saturation and their fatty acid chain length, both of which are properties that influence their activity and immunomodulatory function. In vitro treatment of macrophages or primary Kupffer cells with palmitate, a saturated fatty acid, shifted the cells to an M1-like phenotype engaging PPARγ–NF-kB signalling and increasing pro-inflammatory cytokine production. On the other hand, treatment with polyunsaturated fatty acids promoted an M2-like phenotype, and the cells no longer increased TNF and IL-6 production in response to saturated NEFA treatment. Along similar lines, intravenous infusion of long-chain saturated fatty acids in wild-type mice increased TNF levels in the circulation and led to the accumulation of macrophages and dendritic cells in the liver. This suggests that the composition of fatty acids released during cachexia-induced adipose tissue lipolysis could have a significant role in altering the state of innate immune cell activity, which could have profound effects on the immunopathology associated with chronic illnesses.

**NEFA-mediated modulation of cell function**

Adipose tissue undergoes depletion of its lipid droplets as a consequence of increased lipolysis coupled with a suppression of lipid uptake and lipogenesis. Adipose tissue wasting can be evaluated by measuring adipose tissue mass, adipocyte diameter to evaluate the size of the lipid droplets and circulating levels of non-esterified fatty acids (NEFAs) and glycerol as a result of triglyceride breakdown. These changes in adipose tissue metabolism occur either through transcriptional modulation or via ATGL-mediated and HSL-mediated lipolysis (Fig. 2b).

More studies are needed to understand how the local and infiltrating immune cell populations influence tissue metabolism and vice versa. Moreover, the altered circulatory environment that results from cachexia and the underlying pathology could indirectly affect the local tissue environment through central regulatory pathways, for instance by modulating sympathetic neuronal signalling. Finally, it is well known that immunological and metabolic processes converge on overlapping signalling cascades, highlighting yet another crucial area to probe the immunological mechanisms of cachexia. In the following section we consider potential immunometabolic consequences of cachexia.
Amino acids and cachexia

Alterations in amino acid profiles have been observed in the tumour microenvironment and in the circulation of cachexia-inducing tumours[18–20]. For example, the tumours of patients with colon and stomach cancers were enriched in several amino acids, including serine, tryptophan, arginine and glutamate[18]. In the case of pancreatic tumours, the amino acid profile and abundance seem to depend on the tumour stage. Tumour samples collected from patients with PDAC showed low abundance of most amino acids evaluated, with the exception of taurine, arginine and hydroxyproline[20]. Several other studies reported high abundance of amino acids in the tumour and low abundance in the serum of patients with gastric cancer[19]. Patients with colorectal cancer have high circulating concentrations of phenylalanine and low concentrations of glutamine and histidine, which correlates with systemic inflammation and poor prognosis[21]. Such varied concentrations of circulating amino acids could result from the rate of amino acid release or could be due to amino acid uptake by tumour or immune cells, such as is the case for glutamine[19,21].

Immune cells and tumour cells compete for available amino acids to support their high anabolic demands[22], which modulates the activation status of immune cells. For instance, tumour cells take up arginine and glutamate to support their growth, thereby limiting the availability of these amino acids for T cells[18,23]. Blocking glutamine uptake using an antagonist that targets the tumour microenvironment impairs tumour growth and metabolism while enabling a robust activation of antitumour T cell responses[24].

T cell priming and IL-2-mediated activation induces the expression of amino acid transporters, including SLC7A5, which mediates the import of large neutral amino acids such as phenylalanine and leucine[25]. Lack of SLC7A5 impairs CD4+ effector T cell and CD8+ effector T cell proliferation and differentiation, but does not affect the development of CD4+ Treg cells[26]. This treatment to counter T cell exhaustion.

In summary, cachexia-associated muscle depletion releases a wide range of amino acids into the circulation. Comprehensive profiling and dissection of systemic and local amino acid levels in different models of cachexia, as well as in patients with cachexia, will contribute to a better appreciation of their pathophysiological effect on inflammation, metabolism and disease outcomes.

Concluding remarks

Recent years have seen a substantial improvement in our understanding of cachexia. Yet, when viewing cachexia in the light of metabolic adaptation, the available evidence remains insufficient to definitively classify it as a maladaptive syndrome. The cachexia-associated complications that could result in fatality occur at progressive stages of disease, which renders it difficult to disentangle
what occurs as a result of cachexia as opposed to the underlying illness. Moreover, cachexia occurs in different stages, termed ‘precachexia’, ‘cachexia’ and ‘refractory cachexia’. From an evolutionary perspective, it is tempting to speculate that these stages could reflect the adaptive states of cachexia as the interplay between various trade-off mechanisms and immune responses attempting to resolve the underlying illness.

Many of the models used in the study of cachexia are also employed to examine immune cell activation and function relevant to the underlying disease. However, there is currently an information gap that disconnects what we know about the immune aspects and the metabolic aspects of these diseases. Notably, many valuable patient studies examining immune cell infiltration within the muscles and adipose tissue of patients with cachexia are limited by the variability within their selected cohorts, including different tumour types at variable stages of muscle impairment and cachexia. Such factors can influence the antitumour immune response as well as the metabolic environment in which this response occurs. Considering the difficulty of acquiring biopsy samples from patients with cachexia, such studies could benefit from new technologies that are capable of more thoroughly characterizing immune cells within a single sample. This could be achieved by single-cell RNA sequencing, which allows an unbiased characterization of the immune cell repertoire, and spatial metabolomics using imaging mass spectrometry.

In addition to the importance of cachexia in the context of disease and patient prognosis, cachexia provides a rich arena for immunological research that is full of unanswered questions. For instance, the role of tissue-resident immune cells in the development of cachexia and how these populations adapt to the altered nutrient release and inflammatory signals within their local tissue environment are largely unknown. These bidirectional immune system–tissue interactions create continuous feedback loops that allow the surveillance and modulation of supply and demand. The energy and substrate supply is derived from a finite pool that must be monitored. Supplier cells (for example, adipocytes) require the ability to modulate the level of demand by influencing the molecular and cellular pathways generating the demand. Moreover, the supply sources (lipids, carbohydrates and proteins) are not entirely interchangeable as they produce a specific repertoire of breakdown products. This results in substrate competition within the tissue microenvironment, and systemic energy redistribution. Finally, the immune responses combating the underlying illness can result in collateral damage that must be regulated to avoid excessive tissue damage.

In this case, regulation of energy and substrate availability within the circulation and in the tissue microenvironment provides compartmentalized immune niches such as that seen for T cell populations in muscle tissue.

Overall, future studies of cachexia will benefit from a more rigorous integration of immunology and the use of tools and knowledge from systems biology and evolutionary medicine. Moreover, it will be critical to pursue strategies that bridge the, at times, apparent divide between experimental animal models of cachexia and patient studies and realize the great synergistic potential of such complementary approaches. Immune cell functionality is an understudied area of cachexia research and likely a missing link that will support the development of effective therapeutic strategies against cachexia. Studying immune cell function in relation to the metabolic environment induced by cachexia will also enhance our general understanding of the metabolic regulation of immunity independently of cachexia. This and various other questions require a more holistic and interconnected understanding of the immunology of cachexia.

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by intracerebroventricular injection of recombinant human interleukin-1 in rats. Possible involvement of prostaglandin. Endocrinology 122, 1773–1779 (1988).

27. Braun, T. P. et al. Central nervous system inflammation induces muscle atrophy via activation of the hypothalamic-pituitary-adrenal axis. J. Exp. Med. 204, 1662–1672 (2007).

28. Roeland, E. J. et al. Management of cancer cachexia. ASCO guideline. J. Clin. Oncol. 34, 2438–2453 (2016).

29. Advari, S. M., Advari, P. G., Vonvile, H. M. & Jafri, S. H. Pharmacological management of cachexia in adult cancer patients: a systematic review of clinical trials. BMC Med. 15, 174 (2017).

30. Gordon, J. N. et al. Thalidomide in the treatment of cancer cachexia: a randomised placebo controlled trial. Gut 49, 560–566 (2000).

31. Venne toughness, J. E. et al. Role of thalidomide and placebo for the treatment of cancer-related anorexia-cachexia syndrome: results of a double-blind placebo-controlled randomized study. J. Palliat. Med. 15, 1059–1064 (2012).

32. Hafter, J. et al. Toxicoplasma gondii infection triggers chronic cachexia and sustained catabolic dysbiosis in mice. PLoS ONE 13, e0204695 (2018).

This study characterizes the pathophysiological changes in skeletal muscle and fat following T. gondii infection, thereby consolidating its value as a model for sustained cachexia.

33. Hunter, C. A. & Sibley, L. D. Modulation of innate immunity to Toxoplasma gondii via toll-like receptors. J. Immunol. Res. 204, 168 (2015).

34.管线, F. E. et al. Macrophage inhibitory cytokine-1 (MIC-1/GDF15): a new marker of all-cause mortality. Nat. Med. 26, 1350–1358 (2020).

35. Melchor, S. J. et al. IL-1R regulates disease tolerance and cachexia in toxoplasma gondii infection. J. Immunol. 190, 109–112 (2016).

36. Sturges, C. R. et al. TLR-independent neutrophil-mediated IFNγ is important for host resistance to intracellular pathogens. Proc. Natl. Acad. Sci. USA 110, 10711–10716 (2013).

37. Suzuki, Y., Orenlana, M. A., Schreiber, R. D. & Remington, J. S. Interferon: the major mediator of resistance against Toxoplasma gondii. Science 240, 516–518 (1988).

38. Schäffler, A. & Schölmerich, J. Innate immunity and immunometabolic disorders. Nat. Rev. Immunol. 7, 1299–1307 (2017).

39. Garfisa, G. et al. Dual role for inflammammares sensors NLPR1 and NLPR5 in murine resistance to Toxoplasma infection. J. Immunol. 190, 307–317 (2013).

40. Melcher, S. J. et al. IL-1R regulates disease tolerance and cachexia in toxoplasma gondii infection. J. Immunol. 190, 109–112 (2016).

41. Rao, S. et al. Circulating myeloid cells intrude into the central nervous system and the protective role of TGF-β1 in manipulation of neuron-glia interactions. J. Immunol. 194, 376–488 (2015).

42. O’Garra, A. Type I interferons in infectious disease. Immunity 36, 4676–4684 (2008).

43. Schmidt, D. et al. Modulation of innate immunity by Toxoplasma gondii virulence effectors. Nat. Rev. Microbiol. 10, 766–778 (2012).

44. Varnum, K. et al. Maladaptive innate immunity to Toxoplasma gondii infection. J. Immunol. 149, 109–121 (2002).

45. Toker, T. et al. Transcriptional signatures of toxoplasma gondii infection. PLoS ONE 7, e37267 (2012).

46. D’Avossa, M. A. et al. Macrophages contribute to obesity by importing catecholamine catabolism in macrophages blunts lipolysis during ageing. Nature 550, 119–123 (2017).

47. Nguyen, K. D. et al. Alternatively activated macrophages produce catecholamines to sustain adaptive thermogenesis. Nature 480, 104–108 (2011).

48. Cho, D. S., Schmitt, R. E., Dasgupta, A., Durkacz, A. M. & Sfikakis, P. P. Neutrophil-mediated resolution of post-sepsis skeletal muscle and adipose tissue microenvironments. J. Cachexia Sarcopenia Muscle 11, 1351–1365 (2020).

49. Gabrilovich, D. I. & Nagaraj, S. Myeloid-derived suppressor cells as regulators of the immune system. Nat. Rev. Immunol. 9, 162–174 (2009).

50. Khaleel, Y. S., Ammann, B. J. & Elkord, E. Increased levels of granulocytic myeloid-derived suppressor cells in peripheral blood and tissue of pancreatic cancer patients. J. Immunol Res. 2016, 878987 (2014).

51. Onuki, S. et al. Circulating myeloid-derived suppressor cells are increased and correlate to immune suppression, inflammation and hypoproteinemia in patients with cancer. Oncol. Rep. 28, 455–468 (2012).

52. Cuenca, A. G. A. L. et al. Novel role for tumor-induced expansion of myeloid-derived cells in cancer cachexia. J. Immunol. 192, 6111–6119 (2014).

53. Anzuorri-Barrera, A. et al. Immunohistochemical phenotyping of T cells, granulocytes, and phagocytes in the muscle of cancer patients: association with radiochemically defined mortality and morbidity. J. Clin. Oncol. 37, 1–13 (2019).

54. Narsale, A. et al. Cancer-driven changes link T cell frequency to muscle strength in people with cancer: a pilot study. J. Cachexia Sarcopenia Muscle 10, 827–843 (2019).

55. Barker, T. et al. Muscle injury activates resident macrophages to promote antibody-mediated inhibition of primary and metastatic tumor growth. J. Immunol. 195, 1351–1365 (2015).

56. Suriben, R. et al. Antibody-mediated inhibition of tumor cells to enable cell cycle progression. Nature 525, 550–559 (2015).

57. Wang, Z., Zhao, C., Moya, R. & Davies, J. D. A novel role for CD4 T cells in the control of cancer. J. Immunol. 187, 4676–4686 (2008).

58. Olgren, J. F. et al. Adaptive transfer of CD4+ Foxp3+ regulatory T cells to C57BL/6J mice during acute infection with Toxoplasma gondii downmodulates the exacerbated Th1 immune response. Microbes Infect. 17, 586–595 (2015).

59. Wherry, E. J. & Kurachi, M. Molecular and cellular insights into T cell exhaustion. Nat. Rev. Immunol. 15, 486–499 (2015).

60. Sullivan, B. M., Tejero, J. R., De La Torre, J. C. & Oldstone, M. B. A. Early virus-host interactions dictate the course of a persistent infection. PLoS Pathog. 11, 1004588 (2015).

61. McLane, L. M., Abdel-Hakeem, M. S. & Wherry, E. J. T cell exhaustion in people with viral infection and cancer. Annu. Rev. Immunol. 37, 457–495 (2019).

62. Patsouklis, N. et al. PD-1 alters T cell metabolic reprogramming by inhibiting the pentose phosphate pathway and promoting lipolysis and fatty acid oxidation. Nat. Commun. 6, 6692 (2015).

63. Wu, J. et al. Skeletal muscle antioxidants antagonize CD4 T cell exhaustion. Sci. Adv. 6, eaba5345 (2020).

64. Nansen, A. et al. Compromised virus control and augmented perforin-mediated immunopathology in IFN-gamma-deficient mice infected with lymphocytic choriomeningitis virus. J. Immunol. 165, 6114–6122 (2000).

65. Hildeman, D. A. & Muller, D. Innate immune response loss in intracranial LCMV infection initiated by the absence of effector IL-1β. Viral Immunol. 13, 275–285 (2000).

66. Cohen, S., Nathan, J. A. & Goldberg, A. L. Muscle wasting in disease: molecular mechanisms and promising therapies. Nat. Rev. Drug Discov. 14, 58–74 (2014).

67. Sandri, M. Protein breakdown in cancer cachexia. Semin. Cell Dev. Biol. 21, 276–285 (2010).

68. Aversa, Z. et al. Autophagy is induced in the skeletal muscle of cachectic cancer patients. Sci. Rep. 6, 1–10 (2016).

69. Penna, F. et al. Autophagy exacerbates muscle wasting in cancer cachexia and impairs mitochondrial function. Nat. Commun. 10, 3417 (2019).

70. Bossola, M., Marzetti, E., Rosa, F. & Pozzilli, F. Skeletal muscle regeneration in cancer cachexia. Clin. Exp. Pharmacol. Physiol. 43, 522–527 (2016).
Harizi, H., Corcuff, J. B. & Gualde, N. Arachidonic-acid-derived eicosanoids: roles in biology and immunopathology. Trends Mol. Med. 14, 461–469 (2008).

Hirayama, A. et al. Quantitative metabolome profiling of colon and stomach cancer microenvironment by capillary electrophoresis time-of-flight mass spectrometry. Cancer Res. 69, 4918–4925 (2009).

Hughes, S. et al. Review of metabolomic profiling of gastric cancer and esophageal cancer. Cancer Biol. Med. 17, 181–198 (2020).

Hirakaa, N. et al. Tissue amino acid profiles are characteristic of tumor type, malignant phenotype, and tumor progression in pancreatic tumors. Sci. Rep. 9, 9816 (2019).

Sirmir, P. et al. Alterations in serum amino-acid profile in the progression of colorectal cancer: associations with systemic inflammation, tumour stage and patient survival. Br. J. Cancer 120, 238–246 (2019).

Lemos, H., Huang, L., Prendergast, G. C. & Mellor, A. L. Immune control by amino acid catabolism during tumorigenesis and therapy. Nat. Rev. Cancer 19, 162–175 (2019).

Czyszekova-Kuzmicz, M. et al. Small extracellular vesicles containing arginase-1 suppress T cell responses and promote tumor growth in ovarian carcinoma. Nat. Commun. 10, 5000 (2019).

Leone, R. D. et al. Glutamine blockade induces divergent metabolic programs to overcome tumor immune evasion. Science 366, 1013–1021 (2019).

Sinclair, L. V. et al. Control of amino-acid transport by antigen receptors coordinates the metabolic reprogramming essential for T cell differentiation. Nat. Immunol. 14, 500–508 (2013).

Ma, E. H. et al. Serine is an essential metabolite for effector T cell expansion. Cell Metab. 25, 345–357 (2017).

Klysz, D. et al. Glutamine-dependent α-ketoglutarate production regulates the balance between T helper 1 cell and regulatory T cell generation. Sci. Signal. 8, rno7 (2015).

Wang, K. et al. Glutamine supplementation suppresses herpes simplex virus reactivation. J. Clin. Investig. 127, 2626–2630 (2017).

Rodriguez, P. C. et al. Regulation of T cell receptor CD3γ chain expression by l-arginine. J. Biol. Chem. 277, 21125–2129 (2002).

Rodriguez, P. C., Quinones, D. G. & Ochoa, A. C. l-arginine availability regulates Tlymphocyte cell-cycle progression. Blood 109, 1568–1575 (2007).

Fletcher, M. et al. l-Arginine depletion blunts antitumor T cell responses and promotes tumor growth in ovarian carcinoma. Cancer Res. 75, 275–285 (2015).

Shen, B. et al. Proteomic and metabolic characterization of COVID-19 patient sera. Cell 182, 59–72 e15 (2020).

Lercher, A. et al. Type I interferon signaling disrupts the hepatic urea cycle and alters systemic metabolism to suppress T cell function. Immunity 51, 1074–1087 e9 (2019).

Tito Faja, A. W. Lo Price, value, and the cost of cancer drugs. Lancet Oncol. 17, 927–960 (2016).

Wang, A., Luan, H. H. & Medzhitov, R. An evolutionary perspective on immunometabolism. Science 363, eaar5392 (2019).

Alexandrov, T. Spatial metabolomics and imaging mass spectrometry in the age of artificial intelligence. Annu. Rev. Biochem. Data Sci. 3, 61–87 (2020).

Kahn, C. R., Wang, G. & Lee, K. Y. Altered adipose tissue metabolism and its role in cancer cachexia in mice. EMBO Rep. 20, 2944–2951 (2019).

Teijaro, J. R. et al. Persistent LCMV infection is controlled by blockade of type I interferon signaling. Science 340, 207–211 (2013).

Althaus, C. L., Ganusov, V. V. & De Boer, R. J. Dynamics of CD8 T cell responses during acute and chronic lymphocytic choriomeningitis virus infection. J. Immunol. 179, 2946–2957 (2007).

Rouse, B.T. & Seshawat, S. Immunity and immunopathology to viruses: what decides the outcome? Nat. Rev. Immunol. 10, 514–526 (2010).

Kelley, K. W. et al. Cytokine-induced sickness behavior. Brain Behav. Immun. 17, 112–118 (2003).

Dantzer, R. et al. Unexpected role of interferon-γ in regulating neuronal connectivity and social behaviour. Nature 535, 429–429 (2016).

Dantzer, R. et al. Unleashing the rage: the role of the gut microbiome, immunity, and neuroinflammation in the pathophysiology of eating disorders. Nutrients 13, 1–19 (2021).

Bolmont, L. et al. Role for intestinal TLR5-driven inflammatory response during activity-based anorexia. Sci. Rep. 6, 35813 (2016).

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