

Providence St. Joseph Health

Providence St. Joseph Health Digital Commons

Articles, Abstracts, and Reports

5-1-2017

Optimal nutrition and the ever-changing dietary landscape: a conference report.

A Shao

A Drewnowski

D C Willcox

L Krämer

Christopher G Lausted

Institute for Systems Biology, Seattle, Washington, United States of America.

See next page for additional authors

Follow this and additional works at: <https://digitalcommons.psjhealth.org/publications>



Part of the [Genetics and Genomics Commons](#), and the [Medical Nutrition Commons](#)

Recommended Citation

Shao, A; Drewnowski, A; Willcox, D C; Krämer, L; Lausted, Christopher G; Eggersdorfer, M; Mathers, J; Bell, J D; Randolph, R K; Witkamp, R; and Griffiths, J C, "Optimal nutrition and the ever-changing dietary landscape: a conference report." (2017). *Articles, Abstracts, and Reports*. 1569.
<https://digitalcommons.psjhealth.org/publications/1569>

This Article is brought to you for free and open access by Providence St. Joseph Health Digital Commons. It has been accepted for inclusion in Articles, Abstracts, and Reports by an authorized administrator of Providence St. Joseph Health Digital Commons. For more information, please contact digitalcommons@providence.org.

Authors

A Shao, A Drewnowski, D C Willcox, L Krämer, Christopher G Lausted, M Eggersdorfer, J Mathers, J D Bell, R K Randolph, R Witkamp, and J C Griffiths

Optimal nutrition and the ever-changing dietary landscape: a conference report

A. Shao¹ · A. Drewnowski² · D. C. Willcox³ · L. Krämer⁴ · C. Lausted⁵ ·
M. Eggersdorfer⁶ · J. Mathers⁷ · J. D. Bell⁸ · R. K. Randolph⁹ · R. Witkamp¹⁰ ·
J. C. Griffiths¹¹

Published online: 5 May 2017

© The Author(s) 2017. This article is an open access publication

Abstract The field of nutrition has evolved rapidly over the past century. Nutrition scientists and policy makers in the developed world have shifted the focus of their efforts from dealing with diseases of overt nutrient deficiency to a new paradigm aimed at coping with conditions of excess—calories, sedentary lifestyles and stress. Advances in nutrition science, technology and manufacturing have largely eradicated nutrient deficiency diseases, while simultaneously facing the growing challenges of obesity, non-communicable diseases and aging. Nutrition research has gone through a necessary evolution, starting with a reductionist approach, driven by an ambition to understand the mechanisms responsible for the effects of individual nutrients at the

cellular and molecular levels. This approach has appropriately expanded in recent years to become more holistic with the aim of understanding the role of nutrition in the broader context of dietary patterns. Ultimately, this approach will culminate in a full understanding of the dietary landscape—a web of interactions between nutritional, dietary, social, behavioral and environmental factors—and how it impacts health maintenance and promotion.

Keywords Aging · Big data · Bioactives · Biomarkers · Dietary patterns · Dietary supplements · Longevity · Micronutrients · Obesity · Overfed · Phytonutrients · Sarcopenic obesity · Systems approaches · Undernourished · Wellness

This is the seventh CRN-International conference report. Previous conference reports were published in *Regulatory Toxicology and Pharmacology* [103], and for the last five years, in the *European Journal of Nutrition* [1, 16, 67, 70, 71].

✉ J. C. Griffiths
jgriffiths@crnusa.org

- ¹ Herbalife Nutrition, Los Angeles, CA, USA
- ² University of Washington, Seattle, WA, USA
- ³ Okinawa International University, Ginowan, Japan
- ⁴ Technische Universität Braunschweig, Brunswick, Germany
- ⁵ Institute for Systems Biology, Seattle, WA, USA
- ⁶ DSM Nutritional Products, Kaiseraugst, Switzerland
- ⁷ Newcastle University, Newcastle upon Tyne, UK
- ⁸ University of Westminster, London, UK
- ⁹ Amway Global Discovery, Buena Park, CA, USA
- ¹⁰ Wageningen University, Wageningen, The Netherlands
- ¹¹ Council for Responsible Nutrition-International, Washington, DC, USA

From reductionism to holism: the evolution of nutrition science

Compared to the classical natural sciences (chemistry, physics and mathematics) nutrition science is a relatively new discipline. In the early twentieth century, the public health challenges around nutrition mainly focused on communicable diseases and overt nutrient deficiencies, at the time major determinants of a short life expectancy. To address these concerns, the scientific focus of the time was devoted to vitamins and essential minerals to combat the effects of malnutrition.

During this time, scientists believed that single nutrients could cure or reverse particular diseases. This concept was readily accepted by the scientific and medical communities because it was simple and based on three premises [95]:

1. A simple cause–effect relationship exists between a particular nutrient and a specific effect or disease.

2. Symptoms of a specific nutrient deficiency can be physiologically explained in terms of the role played by the respective nutrient.
3. Providing the nutrient in the diet can prevent, and in many cases reverse, the deficiency disease.

This paradigm was appropriate for addressing the then challenges of overt nutrient deficiency. However, through the course of the twentieth century the situation changed rapidly, especially in developing nations. Public health and nutrition challenges, which had been rooted in deficiency, shifted to challenges resulting from excess. By the mid-to-late twentieth century, major public health challenges transitioned from communicable diseases and malnutrition as the major cause of mortality, to lifestyle-related chronic diseases [7]. These contemporary challenges are due to a combination of poor diet, reduced physical activity, rapidly aging population, rapidly expanding population, climate change and food security issues. The reductionist research model has proven to be ill-suited to address these challenges.

In response to the growing challenges of non-communicable diseases, public and private investments in science and research grew dramatically and have achieved some important beneficial milestones.

- Advances in understanding the role of diet and nutrition in the etiology of chronic diseases.
- Advances in cellular and molecular biology, and biochemistry to allow for a better understanding of macro and micronutrient metabolism and mechanisms of action.
- Advances in identification of nutrients linked to chronic disease; for example, calcium, folate, vitamin D, omega-3 fatty acids and dietary fiber.
- Advances in the discovery, study, and use of other bioactive substances found in foods, such as isoflavones, carotenoids, anthocyanins and catechins.
- Advances in the understanding of the impact of the microbiome on immunity, obesity and cognitive function.
- Advances in the various ‘omics’ technologies: genomics, transcriptomics, proteomics and metabolomics, along with epigenetics.

However, despite these advances, from a public health perspective, the emphasis on a healthy diet and lifestyle has not had the anticipated impact. Imbalanced diet and poor lifestyle have emerged as major contributors to early death [128] coinciding with a global obesity epidemic [46] and proliferation of co-morbidities such as type II diabetes [47]. The nutrition imbalance, where nutrient adequacy is worsening while energy (calorie) excess continues to rise, has

contributed to a paradox of obesity combined with under-nutrition. Populations have become overfed, but undernourished [78].

In the 1980s and 1990s, nutrition science entered an era during which the ‘magic bullet’ was sought using a reductionist approach. This has been referred to as ‘greedy reductionism’:

...in their eagerness for a bargain, in their zeal to explain too much too fast, scientists and philosophers... underestimate the complexities, trying to skip whole layers or levels of theory in their rush to fasten everything securely and neatly to the foundation [30].

Nutrients were included in drug-like randomized controlled trials, the most notable were the studies in which vitamin E and/or *beta*-carotene were given to lifelong smokers or asbestos workers, with the ill-conceived hypothesis that these nutrients could reverse decades of smoking or asbestos exposure [90, 107]. In an effort to uncover the magic bullet, scientists inappropriately studied nutrients in a drug-like context. Unlike drugs, nutrients do not function in isolation and have beneficial effects on multiple tissues and organ systems; a narrow focus on a single or ‘primary’ outcome measure is not practical and does not fit the nutritional context [50].

Perpetuated by a reductionist approach on single macro- and micronutrients, scientists have similarly spent countless resources satisfying the demand for the nutrition ‘villain’ and ‘hero.’ Heroes include substances such as anti-oxidants, fiber, protein and probiotics, while the maligned villains include saturated fat, refined carbohydrates, trans-fat, salt and sugar. In studying the effects of nutrients in isolation, premature conclusions have been reached, resulting in the inevitable ‘flip-flopping’ on whether particular nutrients are beneficial or harmful. This, in turn, has led to enormous consumer confusion and frustration. The problem with this reductionist approach is that, in emphasizing specific nutrients, it fails to take into account that food components interact in complex ways to give rise to emergent properties of diets that are not explicable at the level of individual chemical parts.

During that time span, the perspective of nutrition research has changed from a reductionist approach focusing on specific nutrients, to a complex systems-based science domain. According to the definition of nutrition science for the twenty-first century from *The Giessen Declaration*, an important international workshop sponsored by the International Union of Nutritional Sciences and the World Health Policy Forum: ‘Nutrition science is defined as the study of food systems, foods and drinks, and their nutrients and other constituents; and of their interactions within and between all relevant biological, social and environmental systems’ [108].

Nutrition research is now described as a field which integrates a variety of disciplines, including biology, physiology, sociology, economics, politics and environment [19]. Researchers are now calling for the consideration of environmental sustainability when addressing nutrition questions of public health significance [5]. To grasp these dynamic and highly intertwined connections and interactions, a systems approach is now being advocated, which incorporates transdisciplinary models for food and nutrition security. Such an approach aims to reduce complexity by revealing key interactions that influence the outcomes of interest [48]. In other words, a systems approach can help to identify common causal factors underlying the otherwise seemingly opposing challenges of malnutrition and obesity (population is undernourished, but overfed), which are occurring with more regularity around the world.

The recognition of the need for an interdisciplinary approach to nutrition research has fortunately made its way to the Federal research agenda in the United States. The *National Nutrition Research Roadmap* released by the Interagency Committee on Human Nutrition Research [56] includes highly relevant research questions targeting areas such as behavioral, lifestyle, social, cultural, economic, occupational and environmental factors that impact food choices. The report highlights the importance of environmental sustainability and social science as it pertains to maintaining healthy eating patterns.

Technology now allows use of ‘small data’ at the individual level to drive personalization of diets [129]. Handheld devices now assist consumers allowing them to understand their own nutrient and health status and to make choices. Biomarkers of nutrition status are replacing intake assessment as the basis for identifying dietary gaps. Recommendations for healthy dietary patterns are complimenting, and in some cases replacing those for specific foods and nutrients [111]. These advancements are part of the evolution from linking health benefits to specific nutrients at specific doses to understanding the broader dietary landscape that impacts health including food policy, food choices, culture, environment, dietary patterns, social/psychological factors, home/workplace/school and economy.

Despite a number of challenges and setbacks, nutrition science has developed significantly over time (and will continue). The scientific focus of nutrition had narrowed with a reductionist approach and subsequently (now) expanded to be more holistic. It is now recognized that the study of nutrition involves more than the biology of nutrients, but encompasses the integration of other scientific disciplines, including social, political and environmental sciences. Nutrition recommendations and policy need to continue to evolve in parallel with advances in science and technology to provide solutions to contemporary public health challenges.

Undernourished but overfed: the social and economic considerations

Global diets have become rich in calories but can be poor in some essential nutrients. More and more people are becoming overweight and obese while failing to meet dietary nutrient requirements. As a result, overweight and nutrient deficiencies, the latter sometimes known as ‘hidden hunger’, are both viewed as different forms of malnutrition.

Ideally, each food calorie ought to be the dietary vector for essential micronutrients, e.g., vitamins and minerals. It was long assumed that dietary nutrients and calories went together. However, that relation has now been uncoupled by the emergence in the modern food supply of low-cost empty calories from refined grains, added sugars and added fats. If low-cost fats and sweets provide dietary energy but no nutrients, then eating more food provides no further nutritional benefits. The link between energy consumption and nutrient intake is broken. The low cost of empty calories has made it possible to become undernourished but overfed.

Dietary advice has not kept pace with the basic economics of food-choice behavior. The Dietary Guidelines for Americans 2015–2020 have restated the traditional position that all nutritional needs should be met primarily from nutrient-dense foods. The recognized components of healthy eating patterns were fresh vegetables, legumes, whole fruits whole grains, fat-free or low-fat dairy, seafood, lean meats and poultry, as well as nuts and seeds. Other expert panels have gone further in recommending that all dietary nutrients should come from wholesome ‘real’ foods, preferably local, fresh, unprocessed, and organic. The 2015 Dietary Guidelines did allow that, in some cases, fortified foods and dietary supplements could be useful in providing selected nutrients that might otherwise be consumed in less than recommended amounts [111].

Underconsumed nutrients of greatest public health concern were identified as calcium, potassium, dietary fiber and vitamin D. Iron, underconsumed by adolescent girls and adult women, was another nutrient of public health concern. The suggested remedy was to shift to eating patterns with more vegetables, fruits, whole grains and dairy. Vegetables, fruit juices and dairy are important sources of potassium; vegetables and whole grains supply fiber; whereas dairy is the principal source of both calcium and vitamin D.

Although sustainable food production was an issue, the wide disparities in food prices, seemingly, were not. The nutritive value of foods is often related to their per calorie cost [33, 34]. Among the chief food sources of

potassium, the Dietary Guidelines listed low-cost baked potato but also wild Atlantic salmon, Swiss chard and pomegranate juice. Among fiber sources were a wide variety of beans and ready-to-eat cereals but also avocados, artichokes and raspberries. Calcium was provided by milk and cheese and by various fortified ready-to-eat cereals. Only pregnant women were advised to take an iron supplement when recommended by a healthcare provider. Other supplements were not mentioned.

Yet, it is food prices and diet costs that mediate the observed links between poverty, obesity and micronutrient deficiencies [26, 80]. The following arguments can be made.

First, the current hierarchy of food prices is such that energy-dense refined grains, added sugars, and vegetable oils have become the least expensive sources of concentrated dietary energy, as measured in calorie per unit cost [33, 34]. Total diets composed of energy-dense foods tend to be cheaper per calorie than low energy density diets built around the recommended healthier options [26]. In such studies, food energy density was measured in terms of kcal/100 g, whereas dietary energy density was measured in terms of cost per 1800 or 2000 kcal.

As shown in Fig. 1, energy density of foods is determined by their moisture content. This is because water provided bulk and weight but no energy and no nutrients. In general, energy-dense foods are foods that are dry. Those foods are associated with lower per calorie costs. Not by coincidence, grains, fats and sweets provided calories at low cost, as did sugar sweetened beverages.

By contrast, the more nutrient-rich meat, poultry, fish, vegetables and fruit were more expensive. Nutrient density of foods has been measured in terms of nutrients per

100 kcal, 100 g or serving size. Typically, nutrient-rich foods are those that contain more nutrients than calories. The so-called qualifying nutrients have included protein, fiber, and a variety of vitamins and minerals, including those of public-health concern. Disqualifying nutrients have typically included fat, sugar and sodium. Nutrient profiling is the technique used to separate foods that are energy-dense from those that are nutrient-rich [32, 35]. Nutrient profiling has also been used to assess nutrients per unit cost and the environmental cost of nutrient production.

The more nutrient-rich foods not only did cost more per calorie, but the price disparity between energy-dense and nutrient-rich foods has continued to grow. Tracking the cost of a market basket of 384 foods from 2004 to 2016 has shown that prices have increased the most for meat, poultry and seafood, fresh produce and whole fruit. Whereas, processed fats and sweets cost only 30% more than they did 20 years ago; the cost of fresh produce has more than doubled. As might be expected, energy-dense processed foods, including many grain-based desserts, provide palatable calories at low cost.

The shift to cheap energy-dense foods can lead to overeating due to diminished satiety signals. Whereas, vegetables and whole fruit can be bulky but low in energy, dry foods provide concentrated energy in small volume. Whereas, carrots provide 40 kcal/100, the energy density of a bar of chocolate can be several times higher, around 500 kcal/100 g. In multiple clinical studies, energy-dense foods and energy-dense diets have been linked to overeating and weight gain.

Thus, the observed links between poverty and obesity may be mediated, in part, by the low cost of energy-dense foods and reinforced by the high palatability of sugar and fat. In the US, population subgroups with less education and lower incomes had lower food expenditures and lower quality diets. Those groups were least likely to consume the recommended nutrient-rich foods and were more likely to suffer from micronutrient deficiencies, notably for the nutrients of public health concern. It has been well documented elsewhere that groups of lower socioeconomic status were also more likely to be obese [26].

As a result, populations in developed countries suffer from the dual burden of malnutrition: micronutrient deficiencies and overweight. Both conditions are linked to socioeconomic disparities and are most prevalent among population subgroups with low economic resources and high poverty rates. Improving the nutrient-to-calorie ratio, rather than focusing on calories alone, may be an effective strategy for obesity prevention and control.

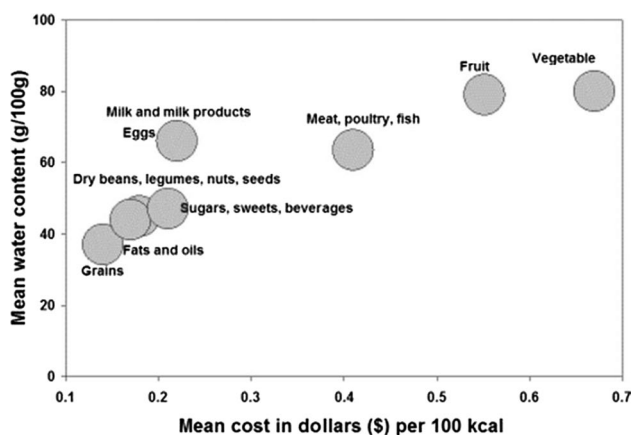


Fig. 1 Relation between mean cost per 100 kcal in US dollars and water content of foods in g/100 g. Data are for 1387 foods from the USDA Food and Nutrient Database for Dietary studies (FNDDS 2.0) aggregated by major food groups

Dietary patterns and longevity: insights from the oldest old in Okinawa

The proportion of worldwide mortality from chronic age-associated disease is projected to reach two-thirds of all deaths by 2030 [126]. This global increase in disease burden from cardiovascular disease (CVD), cancer, diabetes and other chronic age-associated diseases reflects an aging population, as well as social and economic changes, including lifestyle changes and a global nutrition transition to dietary patterns that are more energy-dense but poor in some essential nutrients (see “[Undernourished but overfed: the social and economic considerations](#)”, this article). Despite these challenges, effective public health policy and program implementation can do much to mitigate this risk and help people remain healthy as they age.

Reflecting the potential for preventive public health efforts, the World Health Organization (WHO) has estimated that 80% of coronary heart diseases and type-2 diabetes, as well as 40% of cancers could be prevented by better management of three health behaviors: dietary habits, physical activity and the use of tobacco products [126]. Of these three risk factors, dietary habits may have become the most important modifiable risk factor in many nations, as suggested by a recent study in the United States, that assessed 17 major risk factors. Composition of the diet made up the largest cluster of risk factors responsible for death (26%) and the highest percentage of disability-adjusted life years lost at 14% [83].

Because nutritional issues play such a key role in a wide range of age-associated diseases, the potential for better dietary habits to improve health outcomes in aging populations is great. The Okinawans are of special interest to this topic, as they have been celebrated as the longest lived of the Japanese since prefectural data became available, and nutritional factors may have played a key role in this healthy aging phenomenon, with low rates of cardiovascular disease (and some cancers) mirroring WHO estimates of what may be potentially achieved by preventive public health efforts [105].

Much of the longevity advantage in Okinawa is thought to be related to a healthy lifestyle and this includes the traditional diet, which is calorie-poor, yet nutritionally dense, particularly with regard to phytonutrients with potential caloric restriction (CR) mimetic properties [122]. Although many similarities exist between the traditional Okinawan and Japanese dietary patterns, including the high intake of vegetables; the abundance of soy products such as tofu, natto and miso; the low intake of fats and oils; the heavy reliance on seafood and seaweeds; and the minimal consumption of dairy products,

the traditional Okinawan diet differs in some key areas. In particular, the staple food of the Okinawan diet was neither polished white rice nor other grains but many varieties of colorful sweet potatoes.

Okinawan sweet potatoes have several interesting nutritional characteristics that may have anti-aging effects. For example, sporamin comprises most of the total protein in the edible tuberous root. The biological functions of sporamin include strong free-radical inhibitory and scavenging activity [99]. In light of the strong connection between inflammation and aging, or ‘inflammaging’, as termed by Franceschi and colleagues [40], it is of particular interest that the sweet potato has significant anti-inflammatory properties from multiple phytonutrients that include phenolic acids, flavonols, anthocyanins and carotenoids [124]. In addition to its strong anti-inflammatory effects, sweet potatoes are also good sources of B vitamins, including folate, thiamine, riboflavin, and vitamin B6. Folate and vitamin B6 help convert homocysteine into harmless cysteine. Since high homocysteine levels are associated with an increased risk of cardiovascular disease, it is also of considerable interest that low homocysteine levels accompany the low CVD mortality in Okinawa [4].

Once regarded derisively as a ‘poor farmer’s food’ by those in the upper classes, sweet potatoes now are highly recommended for their health-enhancing properties with recommendations coming from the American Heart Association, the American Cancer Society, as well as the Center for Science in the Public Interest (CSPI) [122]. In fact, the CSPI recently ranked the sweet potato as the ‘healthiest of all vegetables’, mainly for its high content of dietary fiber and protein, anti-oxidant vitamins A and C, potassium, iron, calcium and low levels of saturated fat, sodium and cholesterol [122]. Moreover, most of these nutrients are the very same that are lacking in the American (or ‘Westernized’) diet.

Traditional Okinawan cuisine revolves around steamed sweet potatoes, simmered or steamed green leafy and/or yellow root vegetables, and soy (e.g., miso soup, tofu, miso flavorings), which accompanies almost every meal. Smaller servings of fish or lean meats flavored with herbs and spices, often accompany these staples [124]. Other characteristics that define the Okinawan diet include a taste for bonito flavored broths, and the liberal use of herbs and spices in place of salt.

Ten characteristics of the traditional Okinawa diet:

1. low-caloric intake,
2. high-vegetable consumption (particularly yellow root and green leafy vegetables and seaweeds),
3. high consumption of legumes (mostly soybean),

4. moderate consumption of seafood (more in coastal areas),
5. low consumption of meat (mostly lean pork),
6. little to no dairy products,
7. low-fat intake (high mono and polyunsaturated/saturated fat ratio),
8. low-glycemic load,
9. high-fiber intake,
10. moderate alcohol consumption.

Research suggests that dietary patterns associated with a reduced risk of chronic age-associated disease are vegetable and fruit heavy (therefore phytonutrient rich) but reduced in meat, refined grains, saturated fat, sugar and salt [122]. Many characteristics of the traditional Okinawan diet are shared with other healthy dietary patterns, including traditional Asian diets (especially Japanese), native Hawaiian (taro and sweet potatoes), Mediterranean (both vegetable heavy), and the researcher designed dietary approaches to stop hypertension (DASH) and portfolio (cholesterol lowering) diets [119]. Overall, the important shared features of these healthy dietary patterns include: high intake of unrefined carbohydrates, moderate protein intake with emphasis on vegetables and legumes, fish, and lean meats as sources, and a healthy fat profile (higher in mono/polyunsaturated fats, lower in saturated fat; rich in omega-3). Additionally, the lower caloric density and higher amounts of fiber from such plant-based diets results in lower caloric intake, a lower glycemic load and a higher intake of phytonutrients; all important for minimizing risk for age-associated disease (cardiovascular disease in particular) and maximizing odds for healthy aging and longevity. The oldest old in Okinawa may also have had a longevity dividend from mild caloric restriction and an unusually high amount of foods with potential CR-mimetic effects. These mostly shared dietary characteristics reduce the risk for chronic age-associated disease and promote healthy aging and longevity through multiple mechanisms, including reduced inflammation and oxidative stress.

Recent studies of humans who practice CR (up to 15 years) have shown adaptations consistent with the model organism data, that is, lower risk for CVD and cancer, and biomarker changes suggestive of slower aging [118]. The Okinawans have long been of interest to CR researchers because they may represent the best human example of a naturally calorically restricted population (with optimal nutrition) and the study of their lifespan, healthspan, mortality and morbidity patterns could provide important information on the long-term effects of CR in humans [120]. Some researchers have argued that the Okinawans achieved a long life expectancy for genetic (or other) reasons, but the rapid disappearance of the CR phenotype, as well as the longevity disadvantage in younger Okinawans who did

not experience CR suggests otherwise [119]. Population-wide CR was over by the 1960s, and generations thereafter have had a higher BMI across all age strata, as well as more metabolic syndrome and worse cardiovascular risk factors than other Japanese. The life expectancy advantage for Okinawans, which used to be the highest in Japan for all ages, is now seen only in older ages, consistent with a residual CR-related cohort effect in older Okinawans [120].

In addition to CR, several nutritional factors are thought to be important for Okinawan longevity. For example, the health properties of particular foods from the traditional Okinawan diet may mimic the biological effects of CR, acting as caloric restriction ‘mimetics’. CR-mimetics are compounds that provide the physiological benefit of CR without the need for restriction of calories. The traditional Okinawan diet appears to be a rich source of CR-mimetics [119]. In the Okinawan language, the term *nuchi gusui*, literally means ‘food is medicine’ and reflects the cultural context wherein commonly consumed dietary items, including foods, herbs and spices are also used as folk medicines. Common items that play dual roles as both traditional medicines and foods include sweet potatoes (pulp, skin and leaves), bitter melon, multiple green leafy vegetables, ginger, turmeric, mugwort (*Artemisia vulgaris*), peppers (*Piper hancei*) and carotenoid-rich marine foods (such as seaweeds), among others [119].

We may ask why Okinawa appears to have so many food items with medicinal properties, but climate may offer a scientific basis for this phenomenon. Compounds that have potential CR-mimetic properties, such as carotenoids, flavonoids and other phytochemicals, are synthesized by plants to help scavenge free radicals formed due to stress from extremes of heat, cold, insects, UV light or other threats that are common to sub-tropical Okinawa. Murakami et al. [82] investigated typical food items from Okinawa and compared them to food items from mainland Japan and found that foods from Okinawa had, on average, stronger free radical scavenging properties. Of over one hundred food items tested for anti-oxidative and anti-nitrosative activity, many were shown to be promising anti-inflammatory agents, with implications for the prevention of inflammation-associated carcinogenesis, especially two kinds of turmeric, wild Okinawan turmeric (*Curcuma aromatica*) and Okinawan zedoary (*Curcuma zedoaria*); and ‘Botanbofu’ or ‘Sakuna’ as it is known in Okinawa (*Peucedanum japonicum*).

The most common compounds (polyphenols, flavonoids and similar compounds) can initiate a cellular stress response and induce beneficial molecular adaptations that are collectively known as ‘hormesis’ or a potentially important mechanism for the health-enhancing effects of caloric restriction [94]. Moreover, Davinelli et al. [27] proposed that phytohormetic stress resistance

also involves the activation of nuclear factor erythroid 2-related factor 2 (Nrf2) signaling, a central regulator of the adaptive response to oxidative stress, which may partly account for CR's anti-aging effects.

Hormetic phytochemicals have received much attention for their potential pro-longevity effects and ability to act as potent activators of sirtuins (upstream of FOXO3 (i.e., Forkhead Box O3, a protein coding gene, the deregulation of which is involved with tumorigenesis; and a variant is associated with longevity)), the most notable among them being resveratrol [75]. Interestingly, several of these compounds that are found in Okinawan plants, such as chebulagic acid (a resveratrol derivative) and sesquiterpenoids, seem to be even more potent than resveratrol, at least in terms of some anti-aging properties, such as the ability to inhibit reactive oxygen species (ROS) and nitric oxide (NO) [85]. Many are also potent activators of FOXO3 transcription, a key transcription factor from the insulin-IGF-1 signaling pathway [29]. FOXO3 appears essential for caloric restriction to exert its beneficial effects [121, 123], and allelic variation in the FOXO3 gene is strongly associated with human longevity [121]. In fact, this finding has been replicated in multiple independent populations and is one of only two consistently replicated genes associated with human longevity [31]. Finally, and importantly, there are numerous promising food compounds from the traditional Okinawan diet that have been shown to modulate FOXO3 expression [119] and research is currently underway with the purpose of identifying the most promising candidates with therapeutic potential.

In summary, older Okinawans consumed a diet consistent with mild caloric restriction and rich in foods with CR-mimetic effects, including the staple sweet potato, marine-based carotenoid-rich foods, such as seaweeds, and a myriad of herbs and spices. A CR-like phenotype can be witnessed in older Okinawans which includes smaller stature, less age-associated disease, longer average and maximum lifespan and longer healthspan. Moreover, the epidemiological evidence supports a link between mild caloric restriction, CR-mimetic foods commonly consumed in the traditional diet, and the healthy aging phenomenon in Okinawa [120]. Biological pathways through which caloric restriction (and CR-mimetics) operate are currently being better characterized and the insulin-IGF-1 signaling (IIS) pathway appears to be critical. FOXO3 is a major regulatory gene in the IIS pathway and acts as a transcription factor that induces changes in gene expression in many downstream target genes in response to biological stress. It is activated by CR and CR-mimetic compounds commonly found in the traditional Okinawan diet. This may have profound implications for healthy aging and longevity.

Systems approaches for defining the dietary landscape

Uptake and metabolism of food is a highly complex process and requires the coordinated interaction of cells, tissues and organs to establish homeostasis in multi-cellular organisms. Already minor disturbances in this interaction can cause disease or malfunction of the whole system. On the other hand, the metabolism has to be prepared to act on various fuel types and thus to convert various types of nutrients into energy, biomass, and at the same time to build up a storage pool for periods of fasting.

Systems approaches in the field of nutrition aim to understand the complex interplay of flux, metabolism, utilization, and regulation of nutrients from exogenous and endogenous origin in whole nutritional systems [69]. In this context, stable-isotopes are a valuable and non-invasive tool to profile the dynamics of metabolite turnover of (I) specific metabolic pathways of interest [24] and (II) on a whole organism scale [18, 52, 61]. In this contribution, we discuss the application and value of stable-isotope labeling in the field of nutrition.

To warrant the immediate and stable availability of energy even under high load conditions where complete carbohydrate oxidation is too slow or even not possible, the body must ensure that production and clearance of metabolites such as glucose, lactic acid or alanine are balanced. In humans and other mammals, the Cori cycle is a metabolic pathway that allows glucose homeostasis and involves the concerted action of central organs as it recycles alanine or lactic acid produced by glucose-consuming tissues back to glucose. Lactic acid and/or alanine produced in metabolically active tissues are transferred into the blood stream and transported to the liver. The liver takes up lactic acid and alanine, and recycles two of these molecules back to glucose in an energy-consuming fashion (gluconeogenesis). The so-assembled glucose is secreted back into the blood stream. Hepatic gluconeogenesis is, thus, essential for plasma glucose homeostasis, and when deregulated can be a cause for hyperglycemia, especially in type 2 diabetes mellitus [24].

The dynamics of gluconeogenesis *in vivo* can be revealed by oral administration of ^{13}C -stable-isotope-labeled glucose and time-resolved blood sampling. We developed a protocol to extract polar metabolites from dried blood spots (DBS) and apply gas chromatography coupled with mass spectrometry (GC-MS) to acquire mass spectrometric data for plasma metabolites. To extract pure mass-spectra from these complex data, we apply the ion-chromatographic deconvolution algorithm of our MetaboliteDetector software [53]. Finally, based on the deconvoluted mass-spectra, we correct the

compound spectra for naturally occurring stable-isotopes and determine mass isotopomer distributions (MIDs) for target metabolites. These time-resolved enrichment patterns in combination with absolute concentrations of the target metabolites both obtained from DBS sampling and GC–MS measurement are used as input for a simple mathematical ordinary differential equations (ODE) model. By solving the equation system, we are able to determine quantitative and robust values for glucose production (GP) and gluconeogenesis (GNG).

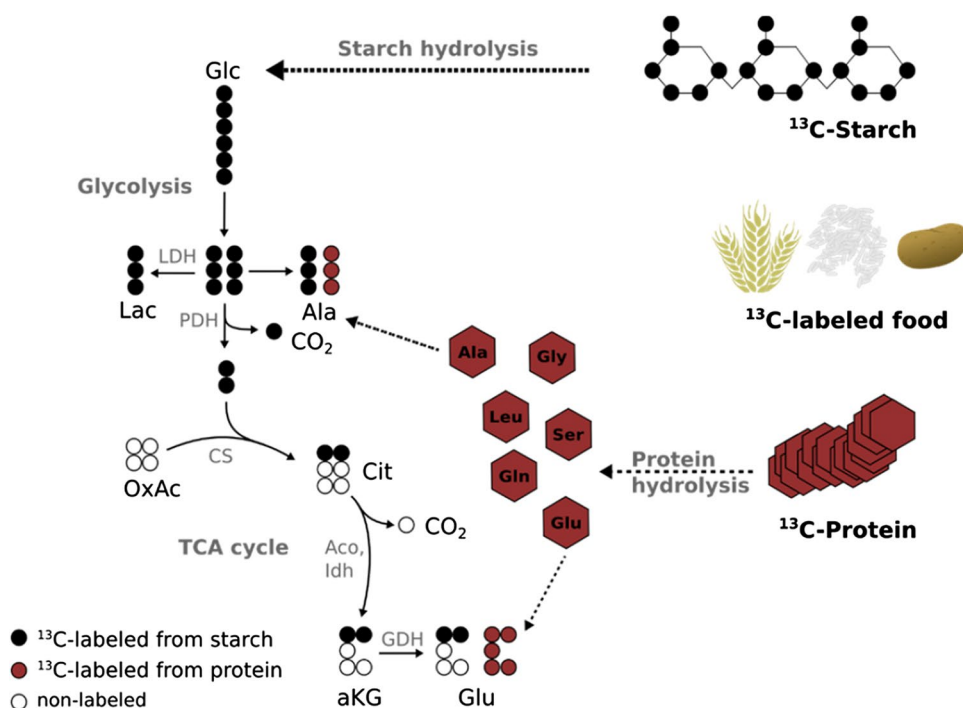
While already providing exact and quantitative metabolic turnover rates for metabolites directly involved in glucose metabolism, such a ‘small system’ does not highlight other parts of whole body homeostasis, as for example, metabolism of amino acids. To address this limitation, we are extending our focus to include quantitative flux data for more metabolites. For this purpose, we follow the fate of fully ^{13}C -labeled food products after ingestion by human individuals. The application of labeled nutrients has been reported for various applications, e.g., the investigation of postprandial response [36], and is mostly based on food products from ^{13}C -labeled plants like wheat, rice, potato or algae grown in a saturated $^{13}\text{CO}_2$ atmosphere. Due to the fact that all ^{12}C carbon atoms are substituted by ^{13}C isotopes, every molecule has an increased molecular mass. In case of wheat and rice, the major molecule types are starch and protein. Therefore, the digestion and metabolism of these bio-polymers can be traced by time-resolved plasma sampling, subsequent GC–MS measurement, and

determination of isotopic enrichment patterns of plasma metabolites of interest (Fig. 2).

As shown in Fig. 2, depending on the origin of the labeled metabolite, different mass isotopomers can be detected. For example, glutamate can be synthesized from starch hydrolysis via glucose, glycolysis, and TCA cycle metabolism and two labeled carbon atoms from glucose are conserved. In this case, the molecular mass of glutamate is increased by 2 U (M2 isotopologue). On the other hand, glutamate appears as a hydrolysis product of protein resulting in a fully labeled glutamate molecule. In this case, its molecular mass is increased by 5 U (M5 isotopologue). Combining these data with information of the composition of the applied food product, a biochemical pathway model can generate quantitative information on the dynamics of the postprandial response [36]. Such approaches to profile multifaceted systems result in the generation of high amounts of complex data. Therefore, computer-based tools and programs as well as high quality statistical analysis are increasingly important for deriving meaningful conclusions.

Nutrition is highly influencing our personal health, and it was recognized that dietary interventions in disease are important to support clinical therapy. Therefore, the generation of innovative systems approaches in the field of nutrition will be essential. As the analytical techniques and the computer tools improve, the extension of systems approaches by stable-isotope driven techniques adds invaluable information on the dynamical scale [52, 61].

Fig. 2 Fully ^{13}C -labeled food products based on, e.g., wheat, rice or potato are composed of ^{13}C -labeled starch and ^{13}C -labeled protein. The labeled hydrolysis products, labeled glucose (*black*) and free labeled amino acids (*red*) enter metabolism and enrichment patterns in the plasma metabolites, measured by GC–MS. *Lac* lactate, *Ala* alanine, *OxAc* oxaloacetate, *Cit* citrate, *aKG* alpha-ketoglutarate, *Glu* glutamate, *LDH* lactate dehydrogenase, *PDH* pyruvate dehydrogenase, *CS* citrate synthase, *IDH* isocitrate dehydrogenase, *GDH* glutamate dehydrogenase



Bringing a ‘big data’ component to the wellness discussion

As healthcare becomes more data-rich and proactive, it will focus more on maintaining and enhancing wellness rather than just reacting to disease. The Institute for Systems Biology in Seattle (ISB) recently launched the large-scale ‘100K Wellness Project’ that integrates genomics, proteomics, transcriptomics, microbiomes, and clinical chemistries along with wearable devices to monitor wellness and disease. This project has begun with a proof-of-concept pilot study in a set of 108 individuals, the ‘Pioneer 100 Project,’ completed this past year. The interpretation of these data led the participants to actionable findings in the areas of medicine, exercise, and nutrition.

Current systems for dealing with health issues remain mostly reactive, with patients reporting to their clinic once symptoms are fully developed. This is a major impediment to the early diagnosis of complex diseases. Few physiological and biochemical parameters are measured routinely, and only after long intervals, so changes might not be identified until many months after the initial stages of disease. Furthermore, parameters vary greatly depending on time of day, gender, genetics, age, exercise schedule, environment, and many other factors. These sparse measurements are then typically compared to reference ranges determined from large population averages that may not be entirely relevant. As a result, it is very difficult to accurately determine if a significant health transition has occurred in a particular individual. Today, it is not generally possible to predict when we will become ill.

What if this were not the case? Imagine if one could learn how many years he or she were away from a specific illness. No one can avoid every risk factor for every disease, but much can be done if one knows what to prioritize. Good doctors always remind their patients to eat healthy and exercise. But what are the right foods? What are the right exercises? Only by engaging healthcare consumers as active participants who provide both health-related data and insights into pathophysiology can biomedical innovation accelerate.

This new scientific wellness strategy is described as personalized, predictive, preventive, and participatory (P4) medicine. The participatory component describes patients, researchers, physicians, and the entire healthcare community joining together to make health maintenance and the practice of medicine as highly proactive activities. Implementation requires two central activities. The first is the establishment for each individual of a ‘dense, dynamic data cloud’ consisting of billions of data points. Computational analysis of these data clouds will

inform health decisions by providing actionable hypotheses chosen to optimize wellness and minimize disease. The second is patient-data integration that will uncover biological networks defining health, and that are perturbed in early disease stages. Through an understanding of these differences, fundamental insights will be gained [54].

A few highly motivated members of the nascent quantified self movement have started collecting their own data clouds. Larry Smarr (UC San Diego) is one such pioneer. After establishing solid baselines, he noticed an unexplained spike in his circulating C-reactive protein (CRP) levels—an indicator of inflammation. On sharing this with his physician, he was asked, ‘Do you have a symptom?’ and, when he answered in the negative, was sent away without treatment. Shortly after, he did indeed experience an acute, dangerous event that required urgent care [102]. Michael Snyder (Stanford University) is another. In his single test subject (not officially identified), he observed a number of distinct changes in blood biomarkers, reflecting pathways activated by viral infections and also by the onset of diabetes. Proteomic data indicated the onset of an elevated glucose response, following shortly after a respiratory syncytial virus infection [23]. Taken by itself, this could be mere coincidence, but once comparable data clouds are shared by thousands of individuals, real statistical analysis becomes possible.

The ultimate goal is to gather comprehensive genetic and physiological data, regularly, for a large number of people, over a long period of time. The objective is to observe all measurable changes that occur as individual transition between disease and healthy states and vice versa. Towards this goal, ISB recruited volunteers to the ‘Pioneer 100 Project’. This pilot project tested the feasibility of collecting and interpreting multifaceted data for multiple individuals, providing them with real-time feedback from the data, and coaching them on ways to improve their health and wellness.

ISB successfully enrolled 108 volunteers, or ‘Pioneers’, into an IRB-approved study led by Profs. Leroy Hood and Nathan Price. Pioneers ranged in age from their 20s to over 88 years and were approximately balanced between male and female. The study ran for 9 months in 2014. Each pioneer had their whole genome sequenced and provided blood, urine, saliva samples at 3-month intervals for metabolomic, proteomic, and additional chemical analysis. The gut microbiota were also surveyed three times by 16S rRNA sequencing. Participants engaged in continual self-tracking and lifestyle monitoring via digital pedometers and questionnaires. A registered dietitian served as a ‘wellness coach’ to share and interpret results with the participants. A physician reviewed all data before sharing [93].

Correlations between genetic predispositions and measurements were found for both simple traits and complex phenotypes. For example, by integrating previously identified genetic variants, each with a small effect on elevated LDL cholesterol risk, individualized risk scores were calculated. Increasing baseline LDL cholesterol levels were found to correlate with increasing genetic risk. Cohort-level statistics on all measurements also revealed that the largely ‘healthy’ volunteers had a high rate of initial lab results outside of normal range. Based on these results, all of the pioneers received actionable recommendations from their coaching phone calls. At the final time point, out-of-range measurements had dropped in diabetes, cardiovascular disease, inflammation, and nutritional categories. Two pioneers had genetic variants putting them at severe risk for hemochromatosis—a potentially life-threatening excess of blood iron. In fact, both had elevated iron and ferritin levels and neither were previously aware of this condition. By the end of the study, both had their iron levels under control and affected family members were made aware of their risk.

Even in a relatively small cohort of 108 individuals, the study uncovered proto-disease states, nutrient deficiencies, and unhealthy heavy metal levels. During the course of this study, the participants were thoroughly engaged by providing questionnaire data, biosamples for analysis, discussing the findings and the health recommendations with the wellness coach, and participating in pioneer social events. Pioneer feedback provided three consistent messages. First, participants understood that their genome does not control their health destiny. Second, the provided data enhanced their ability to take control of their health. Finally, many people are less healthy than they think—each pioneer had multiple actionable possibilities for enhancing their health.

The role of supplements to support a healthy life

According to the WHO, globally, life expectancy is increasing in the majority of countries [126]. Living longer is not the issue anymore but living longer healthy and better is the challenge. The proportion of people aged over 60 years is growing faster than any other age group. Consequently, from 2000 until 2050, the proportion of the global population over 60 years will double from approximately 11 to 22%. This can be interpreted as a success story for public health policies and for socioeconomic development, but at the same time it also challenges societies to adapt and overcome the health and economic burden of aging and non-communicable diseases. ‘Healthy’ refers to physical, mental, and social well-being as indicated in the WHO definition of health [126, 127]. Lifelong health promotion is the opportunity, with a clear emphasis on the fundamental

role of nutrition and micronutrients, to reduce risks of chronic conditions and disability, and to delay the onset of non-communicable diseases. This provides a trifold benefit: for the individual, for the society, and for healthcare system and costs [41, 100].

Most of the world’s population has inadequate intake of one or more of the essential vitamins and minerals. While vitamin and mineral deficiency, and inadequacy are often seen as problems in developing countries it is less recognized as an issue, also in developed countries [63]. Vitamins and minerals are cofactors for proteins and enzymes of metabolism or have other essential functions in our body. A varied and balanced diet should provide enough of all of the vitamins and minerals; however, an unbalanced diet provides calories but not enough of the vitamins and minerals. Insufficient vitamins and minerals result in a deficiency disease, while the consequences of inadequacy are more difficult to diagnose. The Triage theory by Dr. Bruce Ames posits that, because of recurrent shortages of vitamins and minerals during evolution, natural selection developed a strategic-rationing response to moderate shortages so that the scarce nutrient is preferentially retained by proteins that are essential for short-term survival and reproduction [6, 45]. In contrast, proteins needed for long-term health and to defend against the diseases associated with aging, are starved for vitamins, and thus are disabled. Since the damage from moderate deficiency is insidious, its importance for long-term health is not clinically apparent. Mechanistic, genetic, and epidemiological evidence suggests that the metabolic trade-off accelerates aging-associated malfunctions and diseases, such as cancer, cardiovascular disease, immune dysfunction, and cognitive decline [45].

The following example summarizes the benefit of essential nutrients for a healthy life which could result in appropriate changes, characterized by health and vitality of the individual, and a positive impact on healthcare and the attendant economic costs. Nutrition is a complex topic and so are its fields of application; nevertheless, a significant scientific and medical consensus exists as to the importance of an appropriate level of micronutrient intake throughout the life course to support growth, foster health, and prevent the onset of disease. Appropriate micronutrient intake (vitamins and minerals)—as part of a balanced diet and in combination with fortified food and supplements—encourages health and well-being [8, 43].

Micronutrient deficiencies and inadequate micronutrient intake compared to recommendations can have serious health consequences for individuals; they also have a wider impact on societies, economies, and healthcare and welfare systems. Healthcare costs are a significant part of the gross domestic product (GDP), expenditures have risen significantly; a 10% increase in non-communicable diseases results in a 0.5% reduction in annual economic growth [41,

100]. There is growing evidence from food intake surveys that a sufficient intake of micronutrients has not reached in many countries compared to the recommendations. Reasons for this are changes in lifestyle, different eating patterns with a higher share of outdoor eating and an increase of processed foods. As the insufficient intake does not result in immediate consequences for a person, the impact and long-term effects on health and well-being are often neglected [110].

A map indicating the vitamin D status in different countries and populations showed a surprisingly low vitamin D and initiated a call to act for evidence-based programs of vitamin D fortification of foods for the general population and/or supplementation in risk groups. An inadequate status in vitamin D impacts several body functions [57, 68, 113]. The classic role of vitamin D is connected to bone health: vitamin D is essential for the uptake and for the transportation of calcium, next to other factors like vitamin K. There are a number of emerging health benefits connected to an adequate vitamin D status, for example muscle strength and the risk of falling in advanced age; vitamin D strengthens the immune system and reduces the risk for multiple sclerosis. It also has an impact on the risk for cardiovascular diseases because it reduces high blood pressure, and it reduces the risk of some cancers as it inhibits or reduces cell proliferation. An analysis of published population-based studies in Europe, where the 25-hydroxy-vitamin D status was measured in blood, demonstrated for many countries a status below the desired level. There are different recommendations in place: the Institute of Medicine in the US recommends a status of 50 nmol/L or higher for the general population as optimal; the authorities in Europe, the International Osteoporosis Foundation, the Endocrine Society and other organizations recommend a status of 75 nmol/L or higher. A number of human studies indicate that a blood level of 75 nmol/L 25-hydroxy-vitamin D and higher is optimal for reducing the risk of bone fractures, improving muscle strength and reducing risk for diseases like diabetes, multiple sclerosis and some cancers. There is currently only one country in Europe which is in the desirable level of 75 nmol/L 25-hydroxy-vitamin D; this is Finland. Some countries are in the range between 50 and 75 nmol/L and the general population in countries, for example, Germany, Switzerland, and Italy are even below 50 nmol/L. For elderly, there are no data available which are representative for a complete country. An assessment for Germany has indicated that supplementation of elderly—assuming a risk reduction for fractures by 20%—would result in healthcare cost reduction of 580–780 million Euros every year [41, 68, 100].

Vitamin E is an essential nutrient; it is a powerful antioxidant and has been recognized as being vital for preserving the integrity of the cell membrane [86]. Vitamin E is

increasingly reported to be related to protecting essential fatty acids from lipid peroxidation, cognitive function, reducing the risk of Alzheimer's disease, and reducing the negative health implications of fatty liver disease. However, the intake of vitamin E is generally low across all regions worldwide; the population intakes for α -tocopherol and vitamin E are below the recommended 15 mg/day for men and women in the US as well as other Western countries, like Germany, the United Kingdom, and the Netherlands [92]. A minimum serum level of 12 μ mol/L α -tocopherol is needed to avoid deficiencies in the human body. Furthermore, results from several observational, prospective studies suggest a serum tocopherol concentration of 30 μ mol/L and above to have beneficial effects on human health in the field of cardiovascular disease, some types of cancer, and mortality. Additionally, data from the 2003–2006 National Health and Nutrition Examination Survey (NHANES) show mean α -tocopherol concentrations below the optimal concentration for the total population and non-supplement users. Apart from differences in α -tocopherol concentration between supplement and non-supplement users, a higher proportion of younger rather than older adults had suboptimal α -tocopherol concentrations. Therefore, despite low incidence of overt vitamin E deficiency many American adults have suboptimal α -tocopherol status even when supplementing their diet [86, 92].

There is a robust literature suggesting that the health benefits of the long chain omega-3 fatty acids, eicosapentaenoic acid (EPA, 20:5n3) and docosahexaenoic acid (DHA, 22:6n3) for human use. Blood levels of EPA and DHA vary across the globe, with most of the countries and regions of the world having levels that are considered as low to very low [37, 39]. The very low to low range of blood EPA and DHA may have implications for the individual as well as for the society because of higher risks for non-communicable diseases, in particular cardiovascular diseases and cognitive decline. A recent meta-analysis, to date, the most comprehensive quantitative analysis of its kind within the peer reviewed biomedical literature, showed a significant decrease of CHD risk in people with elevated TG and LDL cholesterol levels by supplementation with omega-3 fatty acids [3, 77, 89]. It is necessary to consume significantly more than 250 mg of omega-3 fatty acids per day to be classified as having a high EPA/DHA status, resulting in a blood level required to achieve cardiovascular benefits. This optimal level can be achieved by eating fatty fish rich in omega-3, like salmon or cod several times a week [89]. As many people cannot or will not have a diet heavy in marine foods, food supplements and fortified foods are the most convenient and cost effective way to ensure that the optimal level is achieved. The very low to low range of blood EPA and DHA levels may have implications on the global risk for chronic disease as well

as healthcare costs, with a recent study finding that regular consumption of omega-3 supplements in the range of 1000 mg/day could save €12.9 billion a year in healthcare costs related to CVD-related events in the European Union alone [41, 68, 100].

The consensus concerning the value of appropriate micronutrient intake is based on a robust body of evidence and several powerful nutritional and economic models. It is, however, critical to develop models that address the complexities of micronutrient interventions.

The extensive scientific knowledge currently available needs to be translated into cost-effective, practical public health solutions. These may include fortification and/or supplementation. The health and economic benefits of fortification or supplementation with certain micronutrients, e.g., vitamin A (to reduce infant mortality), iodine (to reduce goiter), vitamin D (primarily against rickets), and folic acid (primarily to reduce neural tube defects), is clear. The economic return of fortification varies from 1:24 for vitamin A fortification; to 1:46 for multivitamins (Copenhagen Consensus); up to 1:200 for iodine fortification. Many countries in the world already have mandatory fortification programs in place involving these micronutrients. The absence of such programs (for example for folic acid and vitamin D), is an unacceptable gap in our public health system and should be scrutinized in the light of the overwhelming evidence of the proven benefits of fortification/supplementation programs involving these micronutrients [8, 38, 110].

The above points should receive the urgent attention of the public health, as well as the nutrition community, policy makers, patient organizations, health insurers and other stakeholders, and next steps should be taken to give a healthy diet a higher priority. The implementation of food fortification programs, the development of nutrient-rich, energy-balanced foods and supplementation programs are all rated as important steps towards a healthy life [38]. So, in summary, healthy nutrition and reduction of healthcare costs are realistic objectives. A healthy diet including food fortification and use of supplements providing all nutrients is more influential than genetic factors to decrease morbidity, to support quality of life and healthy aging. Successful examples demonstrate that people understand the importance of eating healthy, however, that realization requires support and education. In communities where integrated programs have taken place, dramatic improvements in quality of life, healthier life and lower healthcare costs are reported. Relevant stakeholders should act together to develop and implement programs for healthy nutritional solutions to provide all essential nutrients and improve the quality of life; solutions are available.

Nutrition and healthy aging: towards the development of personalized interventions

In most countries, average human lifespan is increasing at approximately 2 years per decade. However, these additional life years are often burdened by poor health, this is not an inevitable consequence of aging. The aging process is plastic and there is good evidence that nutrition is a major determinant of how well we age [62] and of risk of age-related disease and disability [84]. The challenge is to improve eating patterns (and other lifestyle factors) to maximize healthy aging. However, conventional approaches to changing eating patterns have often produced modest improvements only [65], which are difficult to sustain. Additionally, because of optimistic bias, people are often resistant to dietary change.

The worldwide obesity epidemic attests to the difficulty in changing dietary choices in healthy ways. However, in recent years, there has been growing interest in the use of personalized nutrition (PN) approaches to help people adopt healthier eating patterns. The PN approach is predicated on the idea that individuals will be more motivated to make appropriate and sustained dietary changes if they perceive that the advice and support offered is ‘personalized’ and, therefore, more directly relevant to each individual.

The Food4Me PN intervention trial, tested the hypothesis that personalizing nutrition advice would produce bigger and more appropriate changes in eating patterns and markers of health [21]. This large-scale intervention study recruited 1609 adults (aged 18–79 years) across seven European countries and randomized them to (i) conventional dietary advice (control) or to PN advice based on: (ii) individual baseline diet; (iii) individual baseline diet plus phenotype; or (iv) individual baseline diet plus phenotype plus genotype. Participants were recruited via the internet using the Food4Me website (<http://food4me.org/>) [21, 22], the intervention was delivered via the web and by e-mail and the participants used this website to upload dietary and other data. Participants collected buccal swabs (for DNA extraction and genotyping) and blood samples (dried blood spots for metabolite assay) at home and posted them to the recruiting centers.

At 6-month follow-up, participants randomized to the PN arms of the intervention had bigger improvements in eating patterns than those in the conventional dietary advice (control) group but there was no evidence of added advantage of using phenotypic or genetic information to tailor the personalized dietary advice [22]. The Food4Me Study also showed the potential benefits of using an internet-based platform to recruit participants and to deliver PN across multiple countries. This offers opportunities for scaling-up

intervention delivery and, potentially, for improved cost-effectiveness [20].

Lifestyle-based interventions are likely to be more effective if they are offered during significant life-stage transitions when participants may be more receptive to change. In respect of healthy aging, retirement is a major intervention opportunity which has been exploited in developing living, eating, activity and planning (LEAP), an internet-based platform designed to deliver personalized advice to help individuals improve diet, physical activity and social roles/connections and so age more healthily [87]. A recently completed pilot study shows that this intervention was well-received by participants in the retirement transition with high participant retention [66]. This was further evidence that internet (digital) approaches are effective in engaging participants and in delivering PN.

Sarcopenic-obesity: a view from the inside

Modern lifestyle choices are leading to a plethora of metabolic conditions and phenotypes. Excess calorie intake, coupled with an ever increasing sedentary lifestyle has brought about an increase in overall adipose tissue mass, as well as adverse distribution. Modern imaging methods have helped to identify an ever increasing number of body composition sub-phenotypes, all of which appear to have distinct disease risk. One such phenotype is the so-called sarcopenic-obesity, with an aberrant muscle mass content and abnormal fat distribution. This phenotype, contrary to classical sarcopenia, which is especially observed in the older populations, appears to arise as a direct consequence of the current obesogenic environment that modern humans have created for themselves.

Sarcopenia describes the age-related gradual decline in muscle mass and function [96]. The consequences of sarcopenia are evident in the quality of life of many older subjects, leading to frailty, fractures, disability and mortality [76]. The etiology of sarcopenia is not fully understood, but includes physical inactivity, chronic diseases, inflammation, and poor nutrition. Currently, there is no universal method for determining sarcopenia; however, options include the use of the relative appendicular skeletal muscle mass; lowest two quintiles of relative muscle mass or ratios of appendicular fat-free mass or whole body fat-free mass to height squared or skeletal muscle/body weight $\times 100$ [25]. The lack of techniques to directly measure ‘muscle mass’ has significantly limited the objective and reproducible assessments of sarcopenia. Most studies today tend to use dual-energy X-ray absorptiometry (DXA) or bioelectrical impedance (BIA), although neither measure ‘muscle quality’. Some researchers have suggested the use of specific muscles such as the psoas or masseter

muscles as potential markers of sarcopenia [59, 116], but these have not been fully validated.

In many instances of sarcopenia, reduced muscle mass is associated with increased fat mass, a condition known as sarcopenic-obesity [10]. Sarcopenic-obesity, has been described as ‘the confluence of two epidemics’ [97], with a combination of reduced muscle mass coupled with significant metabolic dysregulation. The reported prevalence of sarcopenic-obesity varies widely, ranging from 3.6 to 94% [9]. Moreover, despite the fact that in many cases the apparent ‘volume of muscle’ may appear unchanged or even increased, there is an additional burden of obesity in the elderly as decline in physical function reduces muscle quality [114]. Recently, it has been proposed that magnetic resonance imaging (MRI) techniques could play a significant role for a more objective diagnosis of sarcopenic-obesity [81], as they allow full body composition analysis during a single examination, including ectopic fat depots.

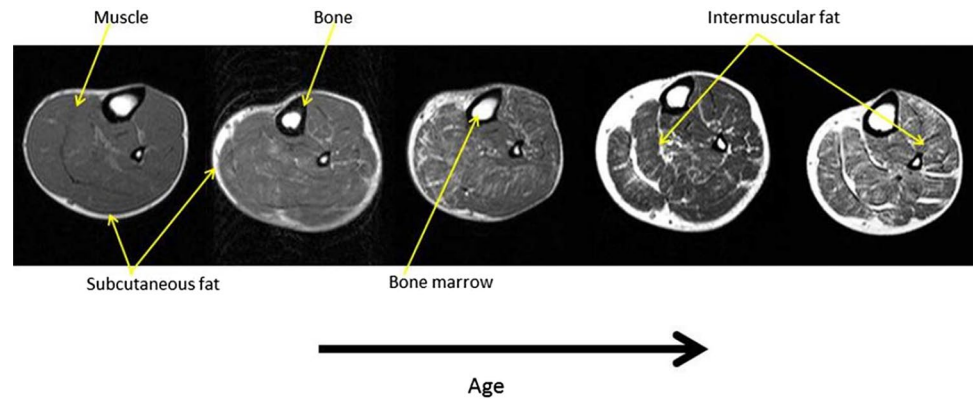
Indeed, recent studies suggest that the sarcopenic-obesity phenotype should include ectopic fat depots such as the liver, pancreas and muscle. Both intra- (within muscle cells) and extra- (between muscle cells) myocellular fat have been shown to increase as a consequence of aging, inactivity, obesity and poor dietary choices [72, 109]. This, in turn, may underpin the deterioration in ‘muscle quality’ observed with this phenotype. This is clearly illustrated in Fig. 3, where increasing fat infiltration across muscle groups can be observed with aging, in the absence of changes in ‘muscle volume’. Thus, understanding the contribution of ectopic fat accumulation may be particularly relevant to sarcopenic-obesity since there is a significant association between intermuscular fat accumulation and decline in gait speed, independent of thigh area [13]. This suggests that measurements of muscle quality/composition may be pivotal.

In conclusion, our current obesogenic environment, characterized by access to calorie rich foods coupled to uncontrolled levels of inactivity, is leading to the propagation of a sarcopenic-obesity phenotype across the population and across the generations. The advent of sophisticated high-throughput imaging systems are allowing the development of a more detailed and reproducible definition and measurement of sarcopenic-obesity, thus, determining its impact on health.

Plant-based dietary patterns, chronic disease risk, and healthy longevity: an emerging paradigm for phytonutrients and health promotion

Over the past two decades, epidemiological studies have shown a consistent inverse relationship between fruit and

Fig. 3 Axial MR images from the lower leg of 5 volunteers, showing typical changes arising as a result of the normal aging process. This includes both a reduction in the proportion of ‘muscle mass’ and the ‘quality’ of the muscle, with a visible increase in the amount of intermuscular fat deposited



vegetable intake and risk for chronic disease, and are the basis for the WHO recommendation for adults to consume at least 400 g/day fruit and vegetables. In the same timeframe, other epidemiological research has identified dietary patterns featuring an abundance and variety of plant foods as a common characteristic in populations that experience exceptionally healthy longevity. The Okinawan and Mediterranean style diets are two well-known examples (see “[Dietary patterns and longevity: insights from the oldest old in Okinawa](#)”, this article). As these two bodies of research have developed, rapid advances have been made in understanding the biology of aging. This emerging research suggests that the age-related decline in cells, tissues and organs underlies the onset of many chronic diseases. Studies in animal models suggest that interventions that slow the rate of aging hold potential to both reduce the risk for chronic disease, and also promote health. The provocative potential connections between these areas of research have been the basis for science at the bench, as well as in the clinic, to characterize the composition, assimilation and metabolism of plant food nutrient components, and their effects on biological mechanisms of aging.

Two prominent examples of science focused on the role of plant-based dietary patterns for reducing and managing chronic diseases come from Wang et al. [117] and Boeing et al. [17]. Wang et al. demonstrated an inverse relationship between the hazard ratio for all-cause mortality and fruit and vegetable consumption. Higher consumption of fruits and vegetables was significantly associated with a lower risk of all-cause mortality. There was a threshold around five servings of fruits and vegetables a day, after which the risk of all-cause mortality did not reduce further. Boeing et al. summarized the strength of evidence behind fruit and vegetable intake and specific chronic diseases. The conclusions from this work suggest that increased fruit and vegetable intake is likely protective against cardiovascular disease and cancer, and also

suggest a protective role against other chronic diseases that are less often associated with mortality.

A new paradigm for plant-based dietary patterns, phytonutrients, and health promotion comes from a selection of literature focused on four emerging areas of science: (i) plant-based dietary patterns in cohorts experiencing healthy longevity, (ii) plant-based diets and wellness outcomes, (iii) advances in understanding the biology of aging, and (iv) phytonutrient interactions with aging mechanisms.

Selected Okinawan and Mediterranean populations have been documented to experience unusually healthy longevity. Interestingly, these populations have plant-based dietary patterns that are well studied in terms of their protective effects against chronic disease. These populations not just survive to old age, they do so with high quality health until late in life. More recently, the National Geographic Society has sponsored a body of ethnographic research that features a selected few demographic groups around the world, all of which experience unusually healthy longevity similar to the Okinawan and Mediterranean populations. These have collectively been named ‘Blue Zones’. Interestingly, plant-based dietary patterns rich in fruits, vegetables, whole grains, and legumes are a feature of the Blue Zones, in addition to active lifestyle, and other health promoting behaviors.

Emerging wellness outcomes research in humans suggests that plant foods hold potential to promote health and function of different organ systems. Green leafy vegetable consumption has been associated with improvements of heart rate variability measures, which may reduce the risk of cardiovascular disease through these favorable changes in cardiac autonomic function [91]. Strong inverse associations were found between plant-based anthocyanin intake and age-related decline in lung function as determined by forced vital capacity [74]. There is evidence that increased fruit and vegetable intake has a positive impact on immune function as determined by antibody response to vaccination [45] as

well as an inverse association with 4-year weight change [14]. Vegetables having both higher fiber and lower glycemic load were more strongly inversely associated with weight change compared to lower-fiber, higher-glycemic-load vegetables. Intake of fruits, vegetables, fish, whole grains, legumes and dairy products is associated with improved bone mineral density [28].

A developing body of evidence in *in vitro* and animal model systems has identified several mechanisms that are thought to play important roles in aging. The primary mechanistic pathways result in reduced oxidative stress, suppressed low-grade chronic inflammation and induction of autophagy [101]. We have explored botanical extracts for their ability to influence one or more of these pathways. We evaluated the effects of turmeric, quercetin, and rosemary extracts on the activation of the nuclear erythroid 2-related factor 2–anti-oxidant responsive element (Nrf2–ARE) signaling pathway. Alone, each of these botanicals had modest Nrf2 activation. When combined, however, there was a surprising synergistic activation. This result is not only an example of the ability of selected phytonutrients to influence aging mechanisms favorably, but also an illustration of the potential for them to interact within a given pathway [79].

Advances in understanding the biology of aging suggest that the origins of chronic disease may lie in the age-related decline in cell, tissue and organ function. The schematic shown in Fig. 4 depicts a synthesis of current working models for the relationship between mechanisms of aging, age-related functional decline and chronic disease. Early in life, the majority of cells and tissues in the body reside at the far left hand side of this schematic. With the passage of time, however, the number of cells and tissues that exhibit compromised function rises such that there is a decline in functional capacity at the tissue and organ level. This process proceeds, if unimpeded, to pathology and the manifestation of disease.

Phytonutrients interact with aging mechanisms such that they hold potential to confer healthy longevity. Figure 4 depicts some proposed areas where phytonutrients conceivably can influence the rate of aging, but also promote functional capacity and reduce the risk for chronic disease. This

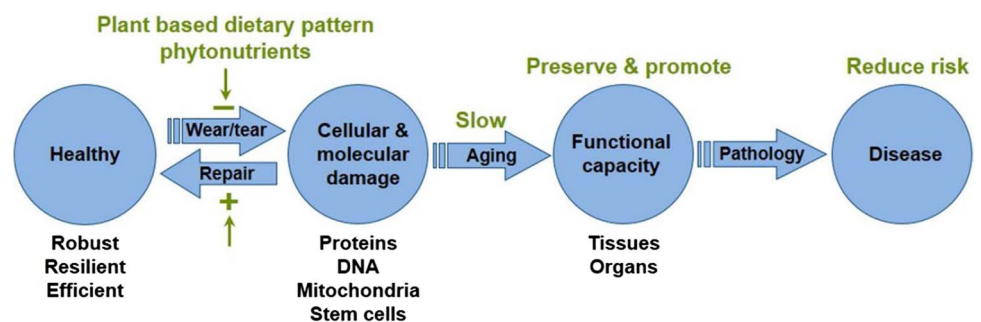
is one rendering of the emergent paradigm for the role of diet, specifically phytonutrients, to favorably influence all of these outcomes.

Evaluation of the evidence as a whole supports the role of plant-based dietary patterns in promoting healthy longevity and suggests that there is an opportunity to intercept individual health trajectories with a healthy plant-based dietary pattern, with the result being improved health and function throughout life. There are, however, important considerations in this paradigm. First, phytonutrients are not likely to act alone as individual compounds, but as a collective family of nutrient compounds. Second, dietary patterns are more powerful lenses with which one can examine healthy longevity with respect to individual foods. No single food is likely to exert healthy longevity. Third, physical activity is most likely a large influencing factor for healthy longevity. Fourth, the gastrointestinal microbiota undoubtedly play a role in the assimilation of dietary components, as well as in the manifestation of end-physiological effects.

Defining optimum nutrition: dynamic biomarkers to capture the complexity of health

Notwithstanding an increasing interest in specific diets to manage, reverse or even cure disease, the primary goal of healthy nutrition is to maintain optimal health and to prevent disorders [15, 44, 51]. Although this might only seem like a marginal difference, starting from the perspective of health, instead of disease, requires different ways of thinking and new approaches. Interestingly, many aspects of modern ‘healthcare’ are rather forms of ‘disease management’. It is perfectly understandable that this situation has evolved since health is intangible, variable, and difficult to predict. At the same time, and irrespective of its precise definition, there is general consensus that optimal health not only constitutes one of the most important values in our life, but also is a major determinant of social and economic stability. It is also clear that diet is one of the main lifestyle factors determining physical and even mental health. However, faced with difficulties to capture the effects of diet and nutritional

Fig. 4 Schematic of current working models for the relationships between mechanisms of aging, age-related functional decline and chronic disease. The proposed interactions between these mechanisms and phytonutrients are depicted by *green text*



interventions and their multifactorial character, many clinicians and policy makers struggle with the practical impact and feasibility of dietary measures. Effects of nutrition are subtle, complex and generally occurring very slowly. This complexity not only poses important scientific questions, but also fuels ongoing discussions, trends, and hypes in the field of nutrition and health. To address the knowledge gaps, several initiatives involving academic and industrial partners have been launched to develop new views and methodologies. Their focus is not only on measuring health effects, but also on trial design including the role of randomized controlled trials (RCTs), statistics, and methodology to accurately measure dietary intake. These developments are supported by the rapid progress made in -omics technologies, data management, statistics, and micro-electronics. Biomarkers for health are key in this process.

In many ways, modern medicine and nutrition science have drifted apart. Medicine has developed into a predominantly disease-oriented discipline, and in its wake the ‘one disease—one target—one drug paradigm’ has become a dominating principle. Understandably, this has consequences for thinking, training, and the way of doing research. In this respect the RCT is the golden standard to, for example, demonstrate the efficacy of therapeutic drugs. However, it is clear that nutrients behave differently from medicinal compounds. Unlike most medicines, single nutrients are involved in numerous endogenous processes and part of complex molecular networks regulating their availability, formation and degradation. As a result, their dose (or concentration) response curves, often U-shaped, are generally composed of multiple effect relationships. Acute effects or ‘quick wins’ are far less obvious than with pharmaceuticals. Another important point is that RCTs have often little usefulness or may not even be possible when studying the effect of diet and (or) specific nutrients [15, 42, 51]. To understand and quantify the way nutrition acts in improving, stabilizing or restoring health, better working definitions of ‘health’ were needed that take into account its dynamic, multi-dimensional and time-dependent character. These definitions have in common the ability to continuously adapt in varying circumstances. An example is ‘the ability to adapt and self-manage in the face of social, physical, and emotional challenges’ [55]. This paradigm dates back to the classical physiological, or even older principles of homeostasis. Nutrition plays important roles in maintaining or even strengthening a proper physiological bandwidth or flexibility (resilience). This has also been called ‘phenotypic flexibility’, being the resultant of the individual’s genotype, his/her physiological and psychological state at a particular point in time, his/her microbiota, etc. [112, 115]. Within this context, the (often gradual) onset of disease starts when and where adaptive processes fail. To measure these dynamic processes, a ‘systems

approach’ is used, meaning that multiple biomarkers, preferably of different integration levels (e.g., gene, protein, metabolite, but also a physiological response, images, etc.) are analyzed at different time points and integrated into models [11]. Such combinations of health biomarkers are often different from the classical disease biomarkers. Furthermore, ‘stress’ or ‘challenge’ tests are used to measure the flexibility and resilience of health. Such tests measure the response to a metabolic, physical, psychological or immunological stressor. Different biochemical, physiological or psychological endpoints are in use, which are indicative of specific processes [104]. For example, a metabolic challenge test measures the response to a standardized meal or beverage that provides a carbohydrate or fat ‘load’ [60, 104, 125]. Next to metabolic stress tests, a laboratory can also apply physical challenge tests [58]. These tests, applying a strenuous exercise protocol on a bicycle ergometer, generate effects on immune function and intestinal permeability, which in turn can be used to investigate potentially positive effects of dietary interventions, probiotics, etc. Other challenge tests apply vaccination, experimental infection [106] or psychological stress [73, 98].

Although these concepts are increasingly acknowledged as the way forward in nutritional science, they have not made life easier in every respect. Their development is often hampered by complexity and costs. Inter-individual differences in responses can be quite large, and standardization and data exchange between labs is not always easy. Another disadvantage is that these tests usually only provide information on the resilience of a limited domain of health (improvement) indicators such as metabolic health, intestinal health, immune health, etc. Last but not least, their extrapolation to perceivable consumer benefits often remains difficult.

An interesting development will be the possibility to move challenge test approaches out of the conventional lab environment. Thanks to the rapid developments in ‘wearable’ technologies for continuous collection of health parameters [64], it is anticipated that a new generation of challenge tests can be developed and incorporated into systems for total health monitoring with ‘real life’ challenges. Provided that technical problems with data integration will be solved, such methodology would also enable further interdisciplinary integration, for example between physiology, epidemiology, geo-informatics, microbiology, consumer behavior, etc.

The generally subtle effects of diet and nutrition on health require methods and biomarkers that capture the dynamics and complexity of physiological resilience. Technological developments are moving extremely fast, offering exciting new ways to measure systems flexibility under real life conditions. Although still many technological and IT challenges lie ahead [2, 49] these ‘living lab’ approaches

are likely to offer new ways for nutrition-related health optimization, self-empowerment, and to develop new products and services.

Conclusion

The CRN-International Scientific Symposium and this article focus on ‘Optimal Nutrition,’ a subject that is nebulous and fraught with a myriad of definitions, boundaries, caveats, and nuances. The ten recognized experts offered their personal perspectives on this most challenging theme. Definitionally, ‘optimal’ would be deemed the ‘most desirable or satisfactory, most favorable, most effective’ and would be the ‘selection of a best element (with regard to some criterion) from some set of available alternatives’, and ‘nutrition’ would be the ‘act or process of nourishing or being nourished; specifically; the sum of the processes by which an animal or plant takes in and utilizes food substances’. However, even when seeking the most desirable process of nourishment, one must then consider that what may be optimal for one person or one region, may not be translatable across the spectrum of human idiosyncrasies.

Historically, the presence or absence of a disease was the measure of how poorly or how successfully one was entering older age. However, that concept is being reevaluated and redefined and ‘functional ability’ is becoming the new rubric. A combination of the intrinsic capacity (physical and mental) that an individual can rely on, relevant environmental factors, and the interactions between the individual and these characteristics is how WHO recommends defining ‘healthy aging’ [12, 88]. A geriatric population that is characterized by high prevalence of ‘frailty’, i.e., a progressive age-related deterioration in physiological systems that result in extreme vulnerability to stressors that in turn increase the risk of multiple adverse outcomes, to the individual as well as to the personal and public healthcare systems through increased dependence, has the potential for extreme personal, familial, governmental and societal costs in time and money. In the twenty-first century, most people can expect to live well into their 60s and 70s (and even older). In 2015, the WHO estimated that global life expectancy at birth was 71.4 years (http://www.who.int/gho/mortality_burden_disease/life_tables/situation_trends/en/). However, the quality of the growing number of added years of life expectancy is highly dependent upon and affected, for better or worse, by each individual’s ‘health’.

This year’s proceedings article offers insights into systems approaches, tracking and analyzing ‘big data’, use of key nutritional and dietary supplements, personalized interventions, dealing with obesity (visible and invisible), use of dynamic biomarkers and other perspectives. It is not the ultimate answer or definitive roadmap;

however, it is highlighting opportunities and setting the stage for more robust discussion and broader and deeper recommendations.

Acknowledgements This conference report summarizes the presentations and outcomes of the meeting entitled ‘Optimal Nutrition and the Ever-Changing Dietary Landscape’ held on December 2, 2016, in Hamburg, Germany. The event was organized by the Council for Responsible Nutrition-International (www.crn-i.ch). The opinions expressed, herein, are those of the authors; this conference report is not a consensus statement; therefore, some authors may not agree with all opinions expressed. The following individuals assisted in the preparation of various portions of this conference report and are hereby acknowledged as follows: Richard Allsopp (Institute for Biogenesis Research, University of Hawaii, Honolulu, HI, USA); Gustavo Glusman (Institute for Systems Biology, Seattle, WA, USA); Leroy Hood (Institute for Systems Biology, Seattle, WA, USA); Karsten Hiller (Technische Universität Braunschweig, Abteilung für Bioinformatik und Biochemie Braunschweig Integrated Centre of Systems Biology (BRICS), Braunschweig, Germany); E Louise Thomas (Research Centre for Optimal Health, Department of Life Sciences, Faculty of Science and Technology, University of Westminster, London UK); Nathan D. Price (Institute for Systems Biology, Seattle, WA, USA); Matt Runyon (Amway Global Discovery, Buena Park, CA, USA); Makoto Suzuki (Okinawa International University, Ginowan, Japan and University of the Ryukyus, Nishihara, Japan); Bradley J Willcox (Kuakini Medical Center, Honolulu, HI, USA and Department of Geriatric Medicine, University of Hawaii, Honolulu, HI, USA).

Compliance with ethical standards

Conflict of interest A Drewnowski, D.C. Willcox, L. Krämer, C. Lausted, M. Eggersdorfer, J. Mathers, J. Bell, R.K. Randolph, R. Witkamp and J.C. Griffiths had their travel expenses reimbursed by CRN-I. J. Mathers serves on the advisory board for Estavayer Lait S.A., Switzerland. A. Shao, M. Eggersdorfer and R.K. Randolph are employees of their respective companies, Herbalife Nutrition, DSM Nutritional Products and Amway Corporation. A. Drewnowski has received grants, honoraria, and consulting fees from numerous food, beverage, and ingredient companies and from other commercial and nonprofit entities with an interest in nutrient density of individual foods, meals, and total diets.

Open Access This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.

References

1. Aggett PJ, Hathcock J, Jukes D, Richardson DP, Calder PC, Bischoff-Ferrari H, Nicklas T, Mühlebach S, Kwon O, Lewis J, Lugard MJF, Prock P (2012) Nutrition issues at Codex: health claims, nutrient reference values and WTO agreements: a conference report. *Eur J Nutr*. doi:10.1007/s00394-012-0306-8
2. Ala-Korpela M, Davey-Smith G (2016) Metabolic profiling—multitude of technologies with great research potential, but (when) will translation emerge? *Int J Epidemiol* 45:1311–1318

3. Alexander D, Miller P, Van Elswyk M, Kuratko CN, Bylsma LC (2017) A meta-analysis of randomized trials and prospective cohort studies of eicosapentaenoic and docosahexaenoic long chain omega-3 fatty acids and coronary heart disease risk. *Mayo Clin Proc* 92:15–29
4. Alftan G, Aro A, Gey KF (1997) Plasma homocysteine and cardiovascular disease mortality. *Lancet* 349(9049):397
5. Allison DB, Bassaganya-Riera J, Burlingame B, Brown AW, le Coutre J, Dickson SL, Vögele C (2015) Goals in nutrition science 2015–2020. *Front Nutr* 2:26. doi:10.3389/fnut.2015.00026
6. Ames BN (2006) Low micronutrient intake may accelerate the degenerative diseases of aging through allocation of scarce micronutrients by triage. *Proc Natl Acad Sci USA* 103:17589–17594. doi:10.1073/pnas.0608757103
7. Anjana RM, Ali MK, Pradeepa R, Deepa M, Datta M, Unnikrishnan R, Mohan V (2011) The need for obtaining accurate nationwide estimates of diabetes prevalence in India: rationale for a national study on diabetes. *Indian J Med Res* 133:369–380
8. BAPEN (2014) Malnutrition matters. British Association for Parenteral and Enteral Nutrition (BAPEN), Redditch. <http://www.bapen.org.uk/pdfs/toolkit-for-commissioners.pdf>. Accessed 23 May 2017
9. Batsis JA, Barre LK, Mackenzie TA, Pratt SI, Lopez-Jimenez F, Bartels SJ (2013) Variation in the prevalence of sarcopenia and sarcopenic obesity in older adults associated with different research definitions: dual-energy X-ray absorptiometry data from the National Health and Nutrition Examination Survey 1999–2004. *J Am Geriatr Soc* 61:974–980
10. Baumgartner RN (2000) Body composition in healthy aging. *Ann NY Acad Sci* 904:437–448
11. Bayle M-L, Wopereis S, Bouwman J, Ommen B, Scalbert A, Pujos-Guillot E (2012) Semi-targeted metabolomic approaches to validate potential markers of health for micronutrients: analytical perspectives. *Metabolomics* 8(6):1114–1129
12. Beard JR, Officer A, de Carvalho A, Sadana R, Pot AM, Michel J-P, Lloyd-Sherck P, Epping-Jordan J, Peeters GM, Mahanani WR, Thiyagarajan JA, Chatterji S (2016) The world report on ageing and health: a policy framework for healthy ageing. *Lancet* 387:2145–2154
13. Beavers KM, Beavers DP, Houston DK, Harris TB, Hue TF, Koster A, Newman AB, Simonsick EM, Studenski SA, Nicklas BJ, Kritchevsky SB (2013) Associations between body composition and gait-speed decline: results from the Health, Aging, and Body Composition study. *Am J Clin Nutr* 97:552–560
14. Bertolo ML, Mukamal KJ, Cahill LE, Hou T, Ludwig DS, Mozaffarian D, Willett WC, Hu FB, Rimm EB (2015) Changes in intake of fruits and vegetables and weight change in United States men and women followed for up to 24 years: analysis from three prospective cohort studies. *PLoS Med* 12(9):e1001878. doi:10.1371/journal.pmed.1001878
15. Biesalski HK, Aggett PJ, Anton R, Bernstein PS, Blumberg J, Heaney RP, Henry J, Nolan JM, Richardson DP, van Ommen B, Witkamp RF, Rijkers GT, Zöllner I (2011) 26th Hohenheim Consensus Conference, September 11, 2010 Scientific substantiation of health claims: evidence-based nutrition. *Nutrition* 27:S1–S20
16. Biesalski HC, Erdman JW, Hathcock J, Ellwood K, Beatty S, Johnson E, Marchioli R, Lauritzen L, Rice HB, Shao A, Griffiths J (2013) Nutrient reference values for bioactives: new approaches needed? A conference report. *Eur J Nutr*. doi:10.1007/s00394-013-0503-0
17. Boeing H, Bechthold A, Bub A, Ellinger S, Haller D, Kroke A, Leschik-Bonnet E, Müller MJ, Oberritter H, Schulze M, Stehle P, Wat B (2012) Critical review: vegetables and fruit in the prevention of chronic diseases. *Eur J Nutr* 51(6):637–663. doi:10.1007/s00394-012-0380-y
18. Brunengraber H, Kelleher JK, Des Rosiers C (1997) Applications of mass isotopomer analysis to nutrition research. *Ann Rev Nutr* 17:559–596
19. Cannon G, Leitzmann C (2005) The new nutrition science project. *Public Health Nutr* 8(6A):673–694
20. Celis-Morales C, Lara J, Mathers JC (2015) Personalising nutritional guidance for more effective behaviour change. *Proc Nutr Soc* 74(2):130–138. doi:10.1017/S0029665114001633
21. Celis-Morales C, Livingstone KM, Marsaux CF, Forster H, O'Donovan CB, Woolhead C, Macready AL, Fallaize R, Navas-Carretero S, San-Cristobal R, Kolossa S, Hartwig K, Tsigirioti L, Lambrinou CP, Moschonis G, Godlewska M, Surwillo A, Grimaldi K, Bouwman J, Daly EJ, Akujobi V, O'Riordan R, Hoonhout J, Claassen A, Hoeller U, Gundersen TE, Kaland SE, Matthews JN, Manios Y, Traczyk I, Drevon CA, Gibney ER, Brennan L, Walsh MC, Lovegrove JA, Alfredo Martinez J, Saris WH, Daniel H, Gibney M, Mathers JC (2015) Design and baseline characteristics of the Food4Me study: a web-based randomised controlled trial of personalised nutrition in seven European countries. *Genes Nutr* 10(1):450. doi:10.1007/s12263-014-0450-2
22. Celis-Morales C, Livingstone KM, Marsaux CF, Macready AL, Fallaize R, O'Donovan CB, Woolhead C, Forster H, Walsh MC, Navas-Carretero S, San-Cristobal R, Tsigirioti L, Lambrinou CP, Mavrogianni C, Moschonis G, Kolossa S, Hallmann J, Godlewska M, Surwillo A, Traczyk I, Drevon CA, Bouwman J, van Ommen B, Grimaldi K, Parnell LD, Matthews JN, Manios Y, Daniel H, Martinez JA, Lovegrove JA, Gibney ER, Brennan L, Saris WH, Gibney M, Mathers JC, Food4Me Study (2016) Effect of personalized nutrition on health-related behaviour change: evidence from the Food4me European randomized controlled trial. *Int J Epidemiol*. doi:10.1093/ije/dyw186
23. Chen R, Mias GI, Li-Pook-Than J, Jiang L, Lam HY, Chen R, Miriami E, Karczewski KJ, Hariharan M, Dewey FE, Cheng Y, Clark MJ, Im H, Habegger L, Balasubramanian S, O'Huallachain M, Dudley JT, Hillenmeyer S, Harakasingh R, Sharon D, Euskirchen G, Lacroute P, Bettinger K, Boyle AP, Kasowski M, Grubert F, Seki S, Garcia M, Whirl-Carrillo M, Gallardo M, Blasco MA, Greenberg PL, Snyder P, Klein TE, Altman RB, Butte AJ, Ashley EA, Gerstein M, Nadeau KC, Tang H, Snyder M (2012) Personal omics profiling reveals dynamic molecular and medical phenotypes. *Cell* 148(6):1293–1307
24. Consoli A, Nurjhan N, Reilly JJ Jr, Bier DM, Gerich JE (1990) Mechanisms of increased gluconeogenesis in noninsulin-dependent diabetes mellitus. *J Clin Invest* 86:2038–2045
25. Cruz-Jentoft AJ, Landi F, Schneider SM, Zúñiga C, Arai H, Boirie Y, Chen LK, Fielding RA, Martin FC, Michel JP, Sieber C, Stout JR, Studenski SA, Vellas B, Woo J, Zamboni M, Cederholm T (2014) Prevalence of and interventions for sarcopenia in ageing adults: a systematic review. Report of the International Sarcopenia Initiative (EWGSOP and IWGS). *Age Ageing* 43:748–759
26. Darmon N, Drewnowski A (2015) Contribution of food prices and diet cost to socioeconomic disparities in diet quality and health: a systematic review and analysis. *Nutr Rev* 73(10):643–660. doi:10.1093/nutrit/nuv027
27. Davinelli S, Willcox DC, Scapagnini G (2012) Extending healthy ageing: nutrient sensitive pathway and centenarian population. *Immun Ageing* 9:9
28. de Jonge EAL, Kieft-de Jong JC, de Groot LCPGM, Voortman T, Schoufour JD, Zillikens MC, Hofman A, Uitterlinden AG, Franco OH, Rivadeneira F (2015) Development of

- a food group-based diet score and its association with bone mineral density in the elderly: the Rotterdam study. *Nutrients* 7(8):6974–6990. doi:10.3390/nu7085317
29. de Magalhaes JP, Wuttke D, Wood SH, Plank M, Vora C (2012) Genome-environment interactions that modulate aging: powerful targets for drug discovery. *Pharmacol Rev* 64:88–101
 30. Dennett D (1995) Darwin's dangerous idea: evolution and the meanings of life. Simon & Schuster, New York
 31. Donlon TA, Curb JD, He Q, Grove JS, Masaki KH, Rodriguez B, Elliott A, Willcox DC, Willcox BJ (2012) FOXO3 gene variants and human aging: coding variants may not be key players. *J Gerontol A* 67:1132–1139
 32. Drewnowski A (2009) Defining nutrient density: development and validation of the Nutrient Rich Foods Index. *J Am Coll* 28(4):421S–426S
 33. Drewnowski A (2010) The cost of US foods as related to their nutritive value. *Am J Clin Nutr* 92(5):1181–1188. doi:10.3945/ajcn.2010.29300
 34. Drewnowski A (2013) New metrics of affordable nutrition: which vegetables provide most nutrients for least cost? *J Acad Nutr Diet* 113(9):1182–1187. doi:10.1016/j.jand.2013.03.015
 35. Drewnowski A, Fulgoni VL 3rd (2014) Nutrient density: principles and evaluation tools. *Am J Clin Nutr* 99(5 Suppl):1223S–1228S
 36. Eelderink C, Moerdijk-Poortvliet TC, Wang H, Schepers M, Preston T, Boer T, Vonk RJ, Schierbeek H, Priebe MG (2012) The glyceemic response does not reflect the in vivo starch digestibility of fiber-rich wheat products in healthy men. *J Nutr* 142:258–263
 37. Eggersdorfer M (2016) Mapping public health benefits of adequate omega-3 levels. http://www.nutri-facts.org/en_US/news/Mapping-Public-Health-Benefits-of-Adequate-Omega-3-Levels.html. Accessed 23 Mar 2017
 38. Eggersdorfer M, Kraemer K, Ruel M, Van Ameringen M, Biesalski HK, Bloem M, Chen J, Lateef A, Mannar V (2013) The road to good nutrition. Karger, Basel. ISBN 978-3-318-02549-1
 39. Food Supplements Europe (2016) Healthcare cost savings of omega 3 food supplements in the European Union. <http://www.foodsupplementseurope.org/sites/0023/uploads/content/hccs-omega-3/hccs-omega-3-summary.pdf>. Accessed 23 Mar 2017
 40. Franceschi C, Bonafè M, Valensin S, Olivieri F, De Luca M, Ottaviani E, De Benedictis G (2000) Inflamm-aging. An evolutionary perspective on immunosenescence. *Ann N Y Acad Sci* 908:244–254
 41. Freijer K, Tan SS, Koopmanschap MA, Meijers JM, Halfens RJ, Nuijten MJ (2013) The economic costs of disease related malnutrition. *Clin Nutr* 32:136–141
 42. Gallagher AM, Meijer GW, Richardson DP, Rondeau V, Skarp M, Stasse-Wolthuis M, Tweedie GC, Witkamp R, International Life Sciences Institute Europe Functional Foods Task Force (2011) A standardised approach towards PROving the efficacy of foods and food constituents for health CLAIMs (PROCLAIM): providing guidance. *Br J Nutr* 106(Suppl 2):S16–S28. doi:10.1017/S0007114511003618
 43. García OP, Long KZ, Rosado JL (2009) Impact of micronutrient deficiencies on obesity. *Nutr Rev* 67:559–572. doi:10.1111/j.1753-4887.2009.00228
 44. Georgiou NA, Garssen J, Witkamp RF (2011) Pharma-nutrition interface: the gap is narrowing. *Eur J Pharmacol* 651:1–8
 45. Gibson A, Edgar JD, Neville CE, Gilchrist SECM, McKinley MC, Patterson CC, Young IS, Woodside JV (2012) Effect of fruit and vegetable consumption on immune function in older people: a randomized controlled trial. *Am J Clin Nutr* 96(6):1429–1436. doi:10.3945/ajcn.112.039057
 46. Global prevalence of overweight in adults in 2014 (2015) http://gamapservr.who.int/gho/interactive_charts/ncd/risk_factors/overweight/atlas.html. Accessed 23 Mar 2017
 47. Global report on diabetes (2016) http://apps.who.int/iris/bitstream/10665/204871/1/9789241565257_eng.pdf?ua=1. Accessed 23 Mar 2017
 48. Hammond RA, Dubé L (2012) A systems science perspective and transdisciplinary models for food and nutrition security. *Proc Natl Acad Sci USA* 109(31):12356–12363. doi:10.1073/pnas.0913003109
 49. Hart T, Xie L (2016) Providing data science support for systems pharmacology and its implications to drug discovery. *Expert Opin Drug Discov* 11:241–256
 50. Heaney RP (2008) Nutrients, endpoints, and the problem of proof. *J Nutr* 138(9):1591–1595
 51. Heaney RP (2012) The nutrient problem. *Nutr Rev* 70:165–169
 52. Hellerstein MK (2004) New stable isotope-mass spectrometric techniques for measuring fluxes through intact metabolic pathways in mammalian systems: introduction of moving pictures into functional genomics and biochemical phenotyping. *Metab Eng* 6:85–100
 53. Hiller K, Hangebrauk J, Jäger C, Spura J, Schreiber K, Schomburg D (2009) MetaboliteDetector: comprehensive Analysis Tool for targeted and nontargeted GC/MS based metabolome analysis. *Anal Chem* 81:3429–3439
 54. Hood L, Price ND (2014) Demystifying disease, democratizing health care. *Sci Transl Med* 6(225):225ed5
 55. Huber M, Knottnerus JA, Green L, van der Horst H, Jadad AR, Kromhout D, Leonard B, Lorig K, Loureiro MI, van der Meer JW, Schnabel P, Smith R, van Weel C, Smid H (2011) How should we define health? *BMJ* 343:d4163
 56. Interagency Committee on Human Nutrition Research (2016) National Nutrition Research Roadmap 2016–2021: advancing nutrition research to improve and sustain health. [https://www.nal.usda.gov/sites/default/files/fnic_uploads/2016-03-30-ICHNRNRR\(2\).pdf](https://www.nal.usda.gov/sites/default/files/fnic_uploads/2016-03-30-ICHNRNRR(2).pdf). Accessed 23 Mar 2017
 57. International Osteoporosis Foundation (2017) Vitamin D. <https://www.iofbonehealth.org/osteoporosis-musculoskeletal-disorders/osteoporosis/prevention/vitamin-d>. Accessed 23 Mar 2017
 58. Janssen-Duijghuijsen LM, Mensink M, Lenaerts K, Fiedorowicz E, van Dartel DAM, Mes JJ, Luiking YC, Keijer J, Wichers HJ, Witkamp RF, van Norren K (2016) The effect of endurance exercise on intestinal integrity in well-trained healthy men. *Physiol Rep* 4(20):e12994
 59. Jones KI, Doleman B, Scott S, Lund JN, Williams JP (2015) Simple psoas cross-sectional area measurement is a quick and easy method to assess sarcopenia and predicts major surgical complications. *Colorectal Dis* 17:O20–O26
 60. Kardinaal AF, van Erk MJ, Dutman AE, Stroeve JH, van de Steeg E, Bijlsma S, Kooistra T, van Ommen B, Wopereis S (2015) Quantifying phenotypic flexibility as the response to a high-fat challenge test in different states of metabolic health. *FASEB J* 29:4600–4613
 61. Kelleher JK (2001) Flux estimation using isotopic tracers: common ground for metabolic physiology and metabolic engineering. *Metab Eng* 3:100–110
 62. Kieft-de Jong JC, Mathers JC, Franco OH (2014) Nutrition and healthy ageing: the key ingredients. *Proc Nutr Soc* 73(2):249–259. doi:10.1017/S0029665113003881
 63. Krebs-Smith SM, Guenther PM, Subar AF, Kirkpatrick SI, Dodd KW (2010) Americans do not meet federal dietary recommendations. *J Nutr* 140:1832–1838
 64. Kuehn BM (2016) Wearable biosensors studied for clinical monitoring and treatment. *JAMA* 316:255–257

65. Lara J, Hobbs N, Moynihan PJ, Meyer TD, Adamson AJ, Errington L, Rochester L, Sniehotta FF, White M, Mathers JC (2014) Effectiveness of dietary interventions among adults of retirement age: a systematic review and meta-analysis of randomized controlled trials. *BMC Med* 12:60. doi:[10.1186/1741-7015-12-60](https://doi.org/10.1186/1741-7015-12-60)
66. Lara J, O'Brien N, Godfrey A, Heaven B, Evans EH, Lloyd S, Moffatt S, Moynihan PJ, Meyer TD, Rochester L, Sniehotta FF, White M, Mathers JC (2016) Pilot randomised controlled trial of a web-based intervention to promote healthy eating, physical activity and meaningful social connections compared with usual care control in people of retirement age recruited from workplaces. *PLoS ONE* 11(7):e0159703. doi:[10.1371/journal.pone.0159703](https://doi.org/10.1371/journal.pone.0159703)
67. LeDoux MA, Appelhans KR, Braun LA, Dzedziczak D, Liu L, Osiecki H, Wyszumiala E, Griffiths JC (2015) A quality dietary supplement: before you start and after it's marketed—a conference report. *Eur J Nutr* 54:S1–S8. doi:[10.1007/s00394-014-0827-4](https://doi.org/10.1007/s00394-014-0827-4)
68. Lotters FJ, Lenoir-Wijnkoop I, Fardellone P, Rizzoli R, Rocher E, Poley MJ (2013) Dairy foods and osteoporosis: an example of assessing the health-economic impact of food products. *Osteoporos Int* 24:139–150
69. Lu DX (2014) Systems nutrition: an innovation of a scientific system in animal nutrition. *Front Biosci* 6:55–61
70. Lupton JR, Atkinson SA, Chang N, Fraga CG, Levy J, Messina M, Richardson DP, van Ommen B, Yang Y, Griffiths JC, Hathcock J (2014) Exploring the benefits and challenges of establishing a DRI-like process for bioactives. *Eur J Nutr* 53:S1–S9. doi:[10.1007/s00394-014-0666-3](https://doi.org/10.1007/s00394-014-0666-3)
71. Lupton JR, Blumberg JB, L'Abbe M, LeDoux M, Rice HB, von Schacky C, Yaktine A, Griffiths JC (2016) Nutrient reference value: non-communicable disease endpoints—a conference report. *Eur J Nutr*. doi:[10.1007/s00394-016-1195z](https://doi.org/10.1007/s00394-016-1195z)
72. Marcus RL, Addison O, Kidde JP, Dibble LE, Lastayo PC (2010) Skeletal muscle fat infiltration: impact of age, inactivity, and exercise. *J Nutr Health Aging* 14:362–366
73. McCrea CE, West SG, Kris-Etherton PM, Lambert JD, Gaugler TL, Teeter DL, Sauder KA, Gu Y, Glisan SL, Skulas-Ray AC (2015) Effects of culinary spices and psychological stress on postprandial lipemia and lipase activity: results of a randomized crossover study and in vitro experiments. *J Transl Med* 13:7
74. Mehta AJ, Cassidy A, Litonjua AA, Sparrow D, Vokonas P, Joel Schwartz J (2016) Dietary anthocyanin intake and age-related decline in lung function: longitudinal findings from the VA Normative Aging Study. *Am J Clin Nutr* 103(2):542–550. doi:[10.3945/ajcn.115.121467](https://doi.org/10.3945/ajcn.115.121467)
75. Mercken EM, Carboneau BA, Krzysik-Walker SM, de Cabo R (2012) Of mice and men: the benefits of caloric restriction, exercise, and mimetics. *Ageing Res Rev* 11:390–398
76. Metter EJ, Talbot LA, Schrager M, Conwit R (2002) Skeletal muscle strength as a predictor of all-cause mortality in healthy men. *J Gerontol A* 57:B359–B365
77. Mietus-Snyder ML, Shigenaga MK, Suh JH, Shenvi SV, Lal A, McHugh T, Olson D, Lilienstein J, Krauss RM, Gildengoren G, McCann JC, Ames BN (2012) A nutrient-dense, high-fiber, fruit-based supplement bar increases HDL cholesterol, particularly large HDL, lowers homocysteine, and raises glutathione in a 2-wk trial. *FASEB J* 26:3515–3527. doi:[10.1096/fj.11-201558](https://doi.org/10.1096/fj.11-201558)
78. Miller GD, Drewnowski A, Fulgoni V, Heaney RP, King J, Kennedy E (2009) It is time for a positive approach to dietary guidance using nutrient density as a basic principle. *J Nutr* 139(6):1198–1202. doi:[10.3945/jn.108.100842](https://doi.org/10.3945/jn.108.100842)
79. Missler SR, Rajgopal A, Roloff SJ, Scholten JD, Burns CR, Patterson JA, Pusateri DJ, Gellenbeck KW, Stern NP, Randolph RK, Lin Y, Chuang J, Kazlova V, Fast DJ (2016) Synergistic activation of the Nrf2-ARE oxidative stress response pathway by a combination of botanical extracts. *Planta Med Int Open* 3(02):e27–e30. doi:[10.1055/s-0036-158516](https://doi.org/10.1055/s-0036-158516)
80. Monsivais P, Aggarwal A, Drewnowski A (2012) Are socioeconomic disparities in diet quality explained by diet cost? *J Epidemiol Commun Health* 66(6):530–535. doi:[10.1136/jech.2010.122333](https://doi.org/10.1136/jech.2010.122333)
81. Müller MJ, Geisler C, Pourhassan M, Glüer CC, Bösby-Westphal A (2014) Assessment and definition of lean body mass deficiency in the elderly. *Eur J Clin Nutr* 68:1220–1227
82. Murakami A, Ishida H, Kobo K, Furukawa I, Ikeda Y, Yonaha M, Aniya Y, Ohigashi H (2005) Suppressing effects of Okinawan food items on free radical generation from stimulated leukocytes and identification of some active constituents: implications for the prevention of inflammation-associated carcinogenesis. *Asian Pac J Cancer Prev* 6:437–448
83. Murray CJ, US Burden of Disease Collaborators (2013) The State of US Health, 1990–2010: burden of diseases, injuries, and risk factors. *JAMA* 310(6):591–608. doi:[10.1001/jama.2013.13805](https://doi.org/10.1001/jama.2013.13805)
84. Murray CJ, Richards MA, Newton JN, Fenton KA, Anderson HR, Atkinson C, Bennett D, Bernabé E, Blencowe H, Bourne R, Braithwaite T, Brayne C, Bruce NG, Brugha TS, Burney P, Dherani M, Dolk H, Edmond K, Ezzati M, Flaxman AD, Fleming TD, Freedman G, Gunnell D, Hay RJ, Hutchings SJ, Ohno SL, Lozano R, Lyons RA, Marcenes W, Naghavi M, Newton CR, Pearce N, Pope D, Rushton L, Salomon JA, Shibuya K, Vos T, Wang H, Williams HC, Woolf AD, Lopez AD, Davis A (2013) UK health performance: findings of the Global Burden of Disease Study 2010. *Lancet* 381(9871):997–1020. doi:[10.1016/S0140-6736\(13\)60355-4](https://doi.org/10.1016/S0140-6736(13)60355-4)
85. Nguyen BC, Taira N, Tawata S (2014) Several herbal compounds in Okinawa plants directly inhibit the oncogenic/aging kinase PAK-1. *Drug Discov Ther* 8:238–244
86. NIH Office of Dietary Supplements (2017) Vitamin E. <https://ods.od.nih.gov/factsheets/VitaminE-HealthProfessional/>. Accessed 23 Mar 2017
87. O'Brien N, Heaven B, Teal G, Evans EH, Cleland C, Moffatt S, Sniehotta FF, White M, Mathers JC, Moynihan P (2016) Integrating evidence from systematic reviews, qualitative research, and expert knowledge using co-design techniques to develop a web-based intervention for people in the retirement transition. *J Med Internet Res* 18(8):e210. doi:[10.2196/jmir.5790](https://doi.org/10.2196/jmir.5790)
88. Officer A (2016) World Report on ageing and health. http://apps.who.int/iris/bitstream/10665/186463/1/9789240694811_eng.pdf. Accessed 23 Mar 2017
89. O'Keefe JH, Jacob D, Lavie CJ (2017) Omega-3 fatty acid therapy: the tide turns for a fish story. *Mayo Clin Proc* 92:1–3
90. Omenn GS, Goodman G, Thornquist M, Grizzle J, Rosenstock L, Barnhart S, Glass A (1994) The beta-carotene and retinol efficacy trial (CARET) for chemoprevention of lung cancer in high risk populations: smokers and asbestos-exposed workers. *Cancer Res* 54(7 Suppl):2038s–2043s
91. Park SK, Tucker KL, O'Neill MS, Sparrow D, Pantel S, Vokonas PS, Hu H, Schwartz J (2009) Fruit, vegetable, and fish consumption and heart rate variability: the Veterans Administration Normative Aging Study. *Am J Clin Nutr* 89(3):778–786. doi:[10.3945/ajcn.2008.26849](https://doi.org/10.3945/ajcn.2008.26849)
92. Péter S, Friedel A, Roos FF, Wyss A, Eggersdorfer M, Hoffmann K, Weber P (2016) A systematic review of global alpha-tocopherol status as assessed by nutritional intake levels and blood serum concentrations. *Int J Vitam Nutr Res* 14:1–21. doi:[10.1024/0300-9831/a000281](https://doi.org/10.1024/0300-9831/a000281)
93. Price ND, Magis AT, Earls JC, Glusman G, Levy R, Lausted CG, McDonald DT, Kusebauch U, Moss CL, Zhou Y, Qin S, Moritz RL, Brogaard K, Omenn GS, Lovejoy JC, Hood L

- (2017) Mining dense, dynamic, personal data clouds of 108 humans over 9 months. *Nat Biotech* (in press)
94. Rattan SI (2012) Rationale and methods of discovering hormetins as drugs for healthy ageing. *Expert Opin Drug Disc* 7:439–448
 95. Raubenheimer D, Simpson SJ (2016) Nutritional ecology and human health. *Annu Rev Nutr* 36:603–626. doi:10.1146/annurev-nutr-071715-051118
 96. Rosenberg IH (1997) Sarcopenia: origins and clinical relevance. *J Nutr* 127:990S–991S
 97. Roubenoff R (2004) Sarcopenic-obesity: the confluence of two epidemics. *Obes Res* 12:887–888
 98. Schrieks IC, Joosten MM, Klöpping-Ketelaars WAA, Witkamp RF, Hendriks HFJ (2016) Moderate alcohol consumption after a mental stressor attenuates the stress response. *Alcohol* 57:29–34
 99. Senthilkumar K, Manivasagan P, Venkatesan J, Kim SK (2013) Brown seaweed fucoidan: biological activity and apoptosis, growth signaling mechanism in cancer. *Int J Biol Macromol* 60:366–374
 100. Shanahan C, de Lorimier R (2013) Smart prevention—health care cost savings resulting from the targeted use of dietary supplements. <http://www.crnusa.org/sites/default/files/pdfs-hccs/SmartPrevention-fullreport0913.pdf>. Accessed 23 Mar 2017
 101. Si H, Liu D (2014) Dietary antiaging phytochemicals and mechanisms associated with prolonged survival. *J Nutr Biochem* 25(6):581–591. doi:10.1016/j.jnutbio.2014.02.001
 102. Smarr L (2012) Quantifying your body: a how-to guide from a systems biology perspective. *Biotechnol J* 7(8):980–991
 103. Somogyi A, Hathcock J, Biesalski HK, Blumberg JB, Antoine JM, Edwards G, Prock P (2011) Scientific issues related to Codex Alimentarius goals: a review of principles with examples. *Reg Toxicol Pharmacol* 60:161–164
 104. Stroeve JH, Wietmarschen H, Kremer BH, Ommen B, Wopereis S (2015) Phenotypic flexibility as a measure of health: the optimal nutritional stress response test. *Genes Nutr* 10(3):459
 105. Suzuki M, Willcox BJ, Willcox DC (2001) Implications from and for food cultures for cardiovascular disease: longevity. *Asia Pac J Clin Nutr* 10:165–171
 106. Ten Bruggencate SJ, Frederiksen PD, Pedersen SM, Floris-Vollenbroek EG, Lucas-van de Bos E, van Hoffen E, Wejse PL (2016) Dietary milk-fat-globule membrane affects resistance to diarrheagenic *Escherichia coli* in healthy adults in a randomized, placebo-controlled, double-blind study. *J Nutr* 146:249–255
 107. The Alpha-Tocopherol, Beta Carotene Cancer Prevention Study Group (1994) The effect of vitamin E and beta carotene on the incidence of lung cancer and other cancers in male smokers. *N Engl J Med* 330(15):1029–1035. doi:10.1056/NEJM199404143301501
 108. The Giessen Declaration (2005) *Public Health Nutr* 8(6A):783–786
 109. Thomas EL, Parkinson JR, Frost GS, Goldstone AP, Doré CJ, McCarthy JP, Collins AL, Fitzpatrick JA, Durighel G, Taylor-Robinson SD, Bell JD (2012) The missing risk: mRI and MRS phenotyping of abdominal adiposity and ectopic fat. *Obesity* 20:76–87
 110. Troesch B, Eggersdorfer M, Weber P (2012) 100 years of vitamins: adequate intake in the elderly is still a matter of concern. *J Nutr* 142:979–980
 111. U.S. Department of Health and Human Services and U.S. Department of Agriculture (2015) 2015–2020 dietary guidelines for Americans, 8th edn. <http://health.gov/dietaryguidelines/2015/guidelines/>. Accessed 23 Mar 2017
 112. van Ommen B, van der Greef J, Ordovas JM, Daniel H (2014) Phenotypic flexibility as key factor in the human nutrition and health relationship. *Genes Nutr* 9:423
 113. van Schoor NM, Lips P (2011) Worldwide vitamin D status. *Best Pract Res Clin Endocrinol Metab* 25(4):671–680. doi:10.1016/j.beem.2011.06.007
 114. Villareal DT, Banks M, Siener C, Sinacore DR, Klein S (2004) Physical frailty and body composition in obese elderly men and women. *Obes Res* 12:913–920
 115. Vis D, Westerhuis J, Jacobs D, van Duynhoven JM, Wopereis S, van Ommen B (2014) Analyzing metabolomics-based challenge tests. *Metabolomics* 1:50–63
 116. Wallace JD, Calvo RY, Lewis PR, Brill JB, Shackford SR, Sise MJ, Sise CB, Bansal V (2017) Sarcopenia as a predictor of mortality in elderly blunt trauma patients: comparing the masseter to the psoas using computed tomography. *J Trauma Acute Care Surg* 82:65–72
 117. Wang X, Ouyang Y, Liu J, Zhu M, Zhao G, Bao W, Hu FB (2014) Fruit and vegetable consumption and mortality from all causes, cardiovascular disease, and cancer: systematic review and dose-response meta-analysis of prospective cohort studies. *BMJ* 349:g4490. doi:10.1136/bmj.g4490
 118. Weiss EP, Fontana L (2011) Caloric restriction: powerful protection for the aging heart and vasculature. *Am J Physiol Heart Circ Physiol* 301:H1205–H1219
 119. Willcox BJ, Willcox DC (2014) Calorie restriction, calorie restriction mimetics, and healthy aging in Okinawa: controversies and clinical implications. *Curr Opin Clin Nutr Metab Care* 17:51–58
 120. Willcox BJ, Willcox DC, Todoriki H, Fujiyoshi A, Yano K, He Q, Curb JD, Suzuki M (2007) Caloric restriction, the traditional Okinawan diet, and healthy aging: the diet of the world's longest-lived people and its potential impact on morbidity and life span. *Ann N Y Acad Sci* 1114:434–455
 121. Willcox BJ, Donlon TA, He Q, Chen R, Grove JS, Yano K, Masaki KH, Willcox DC, Rodriguez B, Curb JD (2008) FOXO3A genotype is strongly associated with human longevity. *Proc Natl Acad Sci USA* 105(37):13987–13992. doi:10.1073/pnas.0801030105
 122. Willcox DC, Willcox BJ, Todoriki H, Suzuki M (2009) The Okinawan diet: health implications of a low-calorie, nutrient-dense, anti-oxidant-rich dietary pattern low in glycemic load. *J Am Coll Nutr* 28(Suppl):500S–516S
 123. Willcox DC, Willcox BJ, Yasura S, Ashitomi I, Suzuki M (2012) Gender gap in health span and life expectancy in Okinawa: health behaviours. *Asian J Gerontol Geriatr* 7:49–58
 124. Willcox DC, Scapagnini G, Willcox BJ (2014) Healthy aging diets other than the Mediterranean: a focus on the Okinawan diet. *Mech Ageing Dev* 136:148–162
 125. Wopereis S, Wolvers D, van Erk M, Gribnau M, Kremer B, van Dorsten F, Boelsma E, Garczarek U, Cnubben N, Frenken L, van der Logt P, Hendriks HF, Albers R, van Duynhoven J, van Ommen B, Jacobs DM (2013) Assessment of inflammatory resilience in healthy subjects using dietary lipid and glucose challenges. *BMC Med Genomics* 6:44
 126. World Health Organization (2005) Preventing chronic diseases: a vital investment: WHO global report. WHO Press, Geneva. http://www.who.int/chp/chronic_disease_report/contents/en/. Accessed 23 Mar 2017
 127. World Health Organization (2007) Global age-friendly cities. World Health Organisation, Geneva. http://www.who.int/ageing/publications/Global_age_friendly_cities_Guide_English.pdf. Accessed 23 Mar 2017
 128. World Health Organization (2015) Noncommunicable diseases. Fact sheet. World Health Organisation, Geneva. <http://www.who.int/mediacentre/factsheets/fs355/en/>. Accessed 23 Mar 2017
 129. Zeevi D, Korem T, Zmora N, Israeli D, Rothschild D, Weinberger A, Segal E (2015) Personalized nutrition by prediction of glycemic responses. *Cell* 163(5):1079–1094. doi:10.1016/j.cell.2015.11.001