

## БЕЛЕЖКИ КЪМ

### ЖЕНИТЕ, ХРАНАТА И ХОРМОНИТЕ

на д-р Сара Готфрид

#### **ВЪВЕДЕНИЕ: ЕЗИКЪТ НА ХОРМОНИТЕ**

1. Diabetes Prevention Program Research Group, "10-Year Follow-Up of Diabetes Incidence and Weight Loss in the Diabetes Prevention Program Outcomes Study," *The Lancet* 374, no. 9702 (2009): 1677–86; R. B. Goldberg et al., "Targeting the Consequences of the Metabolic Syndrome in the Diabetes Prevention Program," *Arteriosclerosis, Thrombosis, and Vascular Biology* 32, no. 9 (2012): 2077–90; Diabetes Prevention Program Research Group, "Long-Term Effects of Lifestyle Intervention or Metformin on Diabetes Development and Microvascular Complications over 15-year Follow-Up: The Diabetes Prevention Program Outcomes Study," *The Lancet Diabetes & Endocrinology* 3, no. 11 (2015): 866–75.
2. P. Garrido et al., "Proposal for the Creation of a National Strategy for Precision Medicine in Cancer: A Position Statement of SEOM, SEAP, and SEFH," *Clinical and Translational Oncology* 20, no. 4 (2018): 443–47; G. Gonzalez-Hernandez et al., "Advances in Text Mining and Visualization for Precision Medicine," *Biocomputing* 23 (2018): 559–65; C. A. L. Wicklund et al., "Clinical Genetic Counselors: An Asset in the Era of Precision Medicine," *American Journal of Medical Genetics, Part C: Seminars in Medical Genetics* 178, no. 1 (2018): 63–67; "Precision Medicine," *National Institutes of Health*, <https://olao.od.nih.gov/content/precision-medicine>. Достъпно на 18 септември 2020 година.
3. "The Truth Is Out There, Somewhere," *Lancet* 396, no. 10247 (2020): 291.
4. T. N. Seyfried et al., "Role of Glucose and Ketone Bodies in the Metabolic Control of Experimental Brain Cancer," *British Journal of Cancer* 89, no. 7 (2003): 1375–82; L. M. Rodrigues et al., "The Action of  $\beta$ -hydroxybutyrate on the Metabolism, and Global Histone H3 Acetylation of Spontaneous Mouse Mammary Tumours: Evidence of a  $\beta$ -hydroxybutyrate Paradox," *Cancer & Metabolism* 5, no. 1 (2017): 4–17; C. Bartmann et al., "Beta-hydroxybutyrate (3-OHB) Can Influence the Energetic Phenotype of Breast Cancer Cells but Does Not Impact Their Proliferation and the Response to Chemotherapy or Radiation," *Cancer & Metabolism* 6, no. 1 (2018): 8; M. Chen et al., "An Aberrant SREBP-Dependent Lipogenic Program Promotes Metastatic Prostate Cancer," *Nature Genetics* 50, no. 2 (2018): 206–18; G. Kolata, "High-Fat Diet May Fuel Spread of Prostate Cancer," *The New York Times*, 16.01.2018 година <https://>

- [www.nytimes.com/2018/01/16/health/fat-diet-prostate-cancer.html](http://www.nytimes.com/2018/01/16/health/fat-diet-prostate-cancer.html). Достъпно на 15 август 2018 година.; J. Sremanakova et al., "A Systematic Review of the Use of Ketogenic Diets in Adult Patients with Cancer," *Journal of Human Nutrition and Dietetics* 3, no. 6 (2018): 793–802.
5. G. Bonuccelli et al., "Ketones and Lactate 'Fuel' Tumor Growth and Metastasis: Evidence That Epithelial Cancer Cells Use Oxidative Mitochondrial Metabolism," *Cell Cycle* 9, no. 17 (2010): 3506–14; U. E. Martinez-Outschoorn et al., "Ketones and Lactate Increase Cancer Cell 'Stemness,' Driving Recurrence, Metastasis, and Poor Clinical Outcome in Breast Cancer: Achieving Personalized Medicine Via Metabolo-Genomics," *Cell Cycle* 10, no. 8 (2011): 1271–86.
  6. S. E. Swithers, "Artificial Sweeteners Produce the Counterintuitive Effect of Inducing Metabolic Derangements," *Trends in Endocrinology & Metabolism* 24, no. 9 (2013): 431–41.
  7. J. S. Volek et al., "Cardiovascular and Hormonal Aspects of Very-Low-Carbohydrate Ketogenic Diets," *Obesity Research* 12, no. S11 (2004): 115S-123S; J. S. Volek et al., "Comparison of Energy-Restricted Very-Low-Carbohydrate and Low-Fat Diets on Weight Loss and Body Composition in Overweight Men and Women," *Nutrition & Metabolism* 1, no. 13 (2004): 1–13; H. M. Dashti et al., "Long-Term Effects of Ketogenic Diet in Obese Subjects," *Molecular and Cellular Biochemistry* 286, no. 1–2 (2006): 1–9; **G. Ruaño, "Physiogenomic Analysis of Weight Loss Induced by Dietary Carbohydrate Restriction,"** *Nutrition & Metabolism* 3, no. 1 (2006): 3–20; K. Durkalec-Michalski et al., "Effect of a Four-Week Ketogenic Diet on Exercise Metabolism in CrossFit-Trained Athletes," *Journal of the International Society of Sports Nutrition* 16, no. 1 (2019): 16.
  8. По-високият прием на мазнини при жените е в условията на относително висок прием на въглехидрати (51% от общите дневни калории), а не при кетогенна диета. S. L. Mumford et al., "Dietary Fat Intake and Reproductive Hormone Concentrations and Ovulation in Regularly Menstruating Women," *The American Journal of Clinical Nutrition* 103, no. 3 (2016): 868–77.
  9. J. M. Wilson et al., "The Effects of Ketogenic Dieting on Body Composition, Strength, Power, and Hormonal Profiles in Resistance Training Males," *The Journal of Strength and Conditioning Research* (2017); A. R. Kuchkuntla et al., "Ketogenic Diet: An Endocrinologist Perspective," *Current Nutrition Reports* 8, no. 4 (2019): 402–10.
  10. Когато става въпрос за здравословна чревна микробиота, това, което има значение, са видът, качеството, количеството и произходът на храната. Чревните бактерии използват хранителните вещества в храната, която консумирате, за основни биологични функции (като регулиране на имунната система) и след това създават изходни метаболитни продукти, които влияят на вашата физиология – включително енергиен

баланс, сигнализиране на глюкозата, възпаление и загуба на мазнини. Едно от най-важните хранителни вещества за здрав микробиом са въгледехидратите, достъпни за микробиотата, като пребиотичните фибри. Ключът към успеха на Протокола на Готфрид е да се набавят достатъчни количества от тези специфични въгледехидрати, за да се подхранват добрите бактерии.

F. Bäckhed et al., "The Gut Microbiota as an Environmental Factor That Regulates Fat Storage," *Proceedings of the National Academy of Sciences of the United States of America* 101, no. 44 (2004): 15718–723; M. Rescigno, "Intestinal Microbiota and Its Effects on the Immune System," *Cellular Microbiology* 16, no. 7 (2014): 1004–13; L. Geurts et al., "Gut Microbiota Controls Adipose Tissue Expansion, Gut Barrier, and Glucose Metabolism: Novel Insights into Molecular Targets and Interventions Using Prebiotics," *Beneficial Microbes* 5, no. 1 (2014): 3–17; B. O. Schroeder et al., "Signals from the Gut Microbiota to Distant Organs in Physiology and Disease," *Nature Medicine* 22, no. 10 (2016): 1079–89; K. Makki, "The Impact of Dietary Fiber on Gut Microbiota in Host Health and Disease," *Cell Host & Microbe* 23, no. 6 (2018): 705–15.

11. R. de Cabo et al., "Effects of Intermittent Fasting on Health, Aging, and Disease," *New England Journal of Medicine* 381, no. 26 (2019): 2541–51.
12. Имайте предвид, че периодичното гладуване увеличава неврогенезата – продължаващия растеж и развитие на нови нервни клетки (тоест неврони, които допринасят за функции като учене, емоционална регулация и памет). В. Malinowski, "Intermittent Fasting in Cardiovascular Disorders — An Overview," *Nutrients* 11, no. 3 (2019): 673; A. L. Mindikoglu, Intermittent Fasting from Dawn to Sunset for 30 Consecutive Days Is Associated with Anticancer Proteomic Signature and Upregulates Key Regulatory Proteins of Glucose and Lipid Metabolism, Circadian Clock, DNA Repair, Cytoskeleton Remodeling, Immune System, and Cognitive Function in Healthy Subjects," *Journal of Proteomics* 217 (2020): 103645; S. H. Baik et al., "Intermittent Fasting Increases Adult Hippocampal Neurogenesis," *Brain and Behavior* 10, no. 1 (2020): e01444.
13. Периодичното гладуване понижава общия холестерол, LDL холестерола (познат като „лош“ холестерол) и серумните триглицериди. Освен това то води до намаляване на натрупването на мазнини в черния дроб и мастните тъкани. G. M. Tinsley et al., "Effects of Intermittent Fasting on Body Composition and Clinical Health Markers in Humans," *Nutrition Reviews* 73, no. 10 (2015): 661–74; A. Bener et al., "Effect of Ramadan Fasting on Glycemic Control and Other Essential Variables in Diabetic Patients," *Annals of African Medicine* 17, no. 4 (2018): 196; A. R. Rahbar et al., "Effects of Intermittent Fasting During Ramadan on Insulin-like Growth Factor-1, Interleukin 2, and Lipid Profile in Healthy Muslims," *International Journal of Preventive Medicine* 10, no. 7 (2019): 1–6; S. Ebrahimi et al., "Ramadan

- Fasting Improves Liver Function and Total Cholesterol in Patients with Nonalcoholic Fatty Liver Disease,” *International Journal for Vitamin and Nutrition Research* (2019).
14. B. H. Goodpaster et al., “Metabolic Flexibility in Health and Disease,” *Cell Metabolism* 25, no. 5 (2017): 1027–36
  15. <https://www.cdc.gov/media/releases/2020/s0917-adult-obesity-increasing.html>. Достъпно на 29 септември 2020 година.
  16. N. Stefan et al., “Causes, Characteristics, and Consequences of Metabolically Unhealthy Normal Weight in Humans,” *Cellular Metabolism* 26, no. 2 (2017): 292–300; N. Stefan et al., “Obesity and Impaired Metabolic Health in Patients with COVID-19,” *Nature Reviews Endocrinology* (2020): 1–2.
  17. S. Y. Tartof, et al. “Obesity and Mortality Among Patients Diagnosed With COVID- 19: Results from an Integrated Health Care Organization,” *Annals of Internal Medicine* (2020): M20-3742; W. Dietz et al., “Obesity and Its Implications for COVID- 19 Mortality,” *Obesity* (Silver Spring) 28, no. 6 (2020): 1005; A. Simonnet et al., “High Prevalence of Obesity in Severe Acute Respiratory Syndrome Coronavirus- 2 (SARS-CoV-2) Requiring Invasive Mechanical Ventilation,” *Obesity* (Silver Spring) 28, no. 7 (2020): 1195–99; Erratum in *Obesity* (Silver Spring) 28, no. 10 (2020): 1994; B. M. Popkin et al., “Individuals with Obesity and COVID-19: A Global Perspective on the Epidemiology and Biological Relationships,” *Obesity Reviews* (2020) Aug 26:10.1111/obr.13128.
  18. L. Gupta et al., “Ketogenic Diet in Endocrine Disorders: Current Perspectives,” *Journal of Postgraduate Medicine* 63, no. 4 (2017): 242–51.

## **1. ИСТИНАТА ЗА ХОРМОНИТЕ, ЖЕНИТЕ И ТЕГЛОТО**

1. Емили Хукър, моят изключителен акупунктурист, е стигнала до следните прозрения за традиционната китайска медицина (ТКМ) и хормоните при жените: „Макар текстовете на традиционната китайска медицина да не признават изрично перименопаузата, смята се, че женският жизнен цикъл протича на седемгодишни фази, а 42 години е възрастта, в която shao yang (жлъчния мехур, свързан с черния дроб) започва да запада. При все това в ТКМ също така се твърди, че косата на жената побелява в тази фаза, което изглежда отживелица и потенциално основано на начина на живот. Точно както пишеш, че кортизоловата дисрегулация е в основата на много модели, същото може да се каже и за стагнацията на чернодробната енергия чи и да, въздишането определено е индикация, че има компонент на такава стагнация. С течение на времето стагнацията на чернодробната енергия чи има потенциал да причини множество други дисбаланси, често започващи

с дефицит на чи в далака. Това може да е в основата на различни метаболитни проблеми, макар и рядко като единствена диагноза. Така или иначе това няма отношение към точността на оценката ти за Мелиса, при която със сигурност има стагнация на чернодробната енергия чи. Между другото в момента имам 44-годишна пациентка с почти идентична картина. Нивата на кортизола ѝ са високи, прогестеронът и тестостеронът са ниски, щитовидната жлеза е недостатъчно активна и тя също е качила около 9 – 10 килограма през последната година. Нейната диагноза според ТКМ е стагнация на чернодробната енергия чи с дефицит на чи и ян в далака, което води до повишаване на влагата в организма.“ За да научите повече от Емили, отидете на следния интернет адрес: [emilyhookeracupuncture.com](http://emilyhookeracupuncture.com).

2. P. K. Whelton et al., “ACC/AHA/AAPA/ABC/ACPM/AGS/APhA/ASH/ASPC/NMA/PCNA Guideline for the Prevention, Detection, Evaluation, and Management of High Blood Pressure in Adults: Executive Summary: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines,” *Hypertension* 71, no. 6 (2018): 1269–1324.
3. B. V. Howard et al., “Insulin Resistance and Weight Gain in Postmenopausal Women of Diverse Ethnic Groups,” *International Journal Obesity and Related Metabolism Disorder* 28, no. 8 (2004): 1039–47; O. T. Hardy et al., “What Causes the Insulin Resistance Underlying Obesity?” *Current Opinion Endocrinology, Diabetes, and Obesity* 19, no. 2 (2012): 81–87; H. U. Moon et al., “The Association of Adiponectin and Visceral Fat with Insulin Resistance and  $\beta$ -Cell Dysfunction,” *Journal of Korean Medical Science* 34, no. 1 (2018): e7; J. Fatima et al., “Association of Sonographically Assessed Visceral and Subcutaneous Abdominal Fat with Insulin Resistance in Prediabetes,” *Journal of the Association Physicians of India* 67, no. 4 (2019): 68–70.
4. J. Rezzonico et al., “Introducing the Thyroid Gland as Another Victim of the Insulin Resistance Syndrome,” *Thyroid* 18, no. 4 (2008): 461–64; C. Anil et al., “Metformin Decreases Thyroid Volume and Nodule Size in Subjects with Insulin Resistance: A Preliminary Study,” *Medical Principles and Practice* 25, no. 3 (2016): 233–36; C. Sallorenzo et al., “Prevalence of Pancreatic Autoantibodies in Non-Diabetic Patients with Autoimmune Thyroid Disease and Its Relation to Insulin Secretion and Glucose Tolerance,” *Archives of Endocrinology and Metabolism* 61, no. 4 (2017): 361–66; P. Zhu et al., “Thyroid-Stimulating Hormone Levels Are Positively Associated with Insulin Resistance,” *Medical Science Monitor* 24, no. 1 (2018): 342–47; U. Mousa et al., “Fat Distribution and Metabolic Profile in Subjects with Hashimoto’s Thyroiditis,” *Acta Endocrinologica* 14, no. 1 (2018): 105–12; X. Zhang et al., “Effect of Insulin on Thyroid Cell Proliferation, Tumor Cell Migration, and Potentially Related Mechanisms,” *Endocrine Research* 44, nos. 1–2 (2019): 55–70; X. He et al., “Role of Metformin in the Treatment of Patients with

- Thyroid Nodules and Insulin Resistance: A Systematic Review and Meta-Analysis," *Thyroid* 29, no. 3 (2019): 359–67.
5. A. Verma et al., "Hypothyroidism and Obesity? Cause or Effect," *Saudi Medical Journal* 29, no. 8 (2008): 1135–38; R. Song et al., "The Impact of Obesity on Thyroid Autoimmunity and Dysfunction: A Systematic Review and Meta-Analysis," *Frontiers in Immunology* 10, no. 1 (2019): 2349.
  6. R. C. Kessler et al., "Lifetime and 12-Month Prevalence of DSM-III-R Psychiatric Disorders in the United States," *Archives of General Psychiatry* 51, no. 1 (1994): 8–19; R. C. Kessler et al., "Posttraumatic Stress Disorder in the National Comorbidity Survey," *Archives of General Psychiatry* 52, no. 12 (1995): 1048–60; M. Altemus et al., "Sex Differences in Anxiety and Depression Clinical Perspectives," *Frontiers in Neuroendocrinology* 35, no. 3 (2014): 320–30.
  7. L. Fan et al., "Non-linear Relationship Between Sleep Duration and Metabolic Syndrome: A Population-Based Study," *Medicine (Baltimore)* 99, no. 2 (2020): e18753.
  8. B.L.Fredrickson et al., "Objectification Theory: Toward Understanding Women's Lived Experiences and Mental Health Risks," *Psychology of Women Quarterly* 21, no. 2 (1997): 173–206; C. Rollero et al., "Self-Objectification and Personal Values: An Exploratory Study," *Frontiers in Psychology* 8, no. 1 (2017): 1055; R. Kahalon et al., "Experimental Studies on State Self-Objectification: A Review and an Integrative Process Model," *Frontiers in Psychology* 9, no. 1 (2018): 1268.
  9. L. M. Schaefer et al., "Self-Objectification and Disordered Eating: A Meta-Analysis," *The International Journal of Eating Disorders* 51, no. 6 (2018): 483–502.
  10. L. Cheng, "The Commercialization of Female Bodies in Consumer Society," *Journal of Humanity* 9, no. 1 (2015): 123–25.
  11. M. P. J. Vanderpump, "The Epidemiology of Thyroid Disease," *British Medical Bulletin* 99, no. 1 (2011): 39–51; R. Hoermann et al., "Recent Advances in Thyroid Hormone Regulation: Toward a New Paradigm for Optimal Diagnosis and Treatment," *Frontiers in Endocrinology* 8, no. 1 (2017): 364; A. G. Jubly et al., "Clinical Challenges in Thyroid Disease: Time for a New Approach?" *Maturitas* 87, no. 1 (2016): 72–78.
  12. D. M. Roesch, "Effects of Selective Estrogen Receptor Agonists on Food Intake and Body Weight Gain in Rats," *Physiology & Behavior* 87, no. 1 (2006): 39–44; A. L. Hirschberg, "Sex Hormones, Appetite, and Eating Behaviour in Women," *Maturitas* 71, no. 3 (2012): 248–56; L. Asarian et al. "Sex Differences in the Physiology of Eating," *American Journal of Physiology-Regulatory, Integrative, and Comparative Physiology* 305, no. 11 (2013): R1215–67.
  13. G. D. Miller et al., "Basal Growth Hormone Concentration Increased Following a Weight Loss Focused Dietary Intervention in Older Overweight and Obese Women," *The Journal of Nutrition, Health, & Aging* 16, no. 2 (2012): 169–74.

14. M. Devaki et al., "Chronic Stress-Induced Oxidative Damage and Hyperlipidemia Are Accompanied by Atherosclerotic Development in Rats," *Stress* 16, no. 2 (2013): 233–43; S. N. Kales et al., "Firefighters and On-Duty Deaths from Coronary Heart Disease: A Case Control Study," *Environmental Health* 2, no. 1 (2003): 14; M. Kumari et al., "Chronic Stress Accelerates Atherosclerosis in the Apolipoprotein E Deficient Mouse," *Stress* 6, no. 4 (2003): 297–99; H. E. Webb et al., "Stress Reactivity to Repeated Low-Level Challenges: A Pilot Study," *Applied Psychophysiology Biofeedback* 36, no. 4 (2011): 243–50.
15. P. M. Peeke et al., "Hypercortisolism and Obesity," *Annals of New York Academy of Science* 771, no. 1 (1995): 665–76; S. Paredes et al., "Cortisol: The Villain in Metabolic Syndrome?" *Revista da Associacao Medica Brasileira* (1992) 60, no. 1 (2014): 84–92; J. Q. Purnell et al., "Enhanced Cortisol Production Rates, Free Cortisol, and 11beta-HSD-1 Expression Correlate with Visceral Fat and Insulin Resistance in Men: Effect of Weight Loss," *American Journal of Physiology Endocrinology and Metabolism* 296, no. 2 (2000): E351–57; A. Tchernof et al., "Pathophysiology of Human Visceral Obesity: An Update," *Physiological Reviews* 93, no. 1 (2013): 359–404.
16. M. J. McAllister et al., "Exogenous Carbohydrate Reduces Cortisol Response from Combined Mental and Physical Stress," *International Journal of Sports Medicine* 37, no. 14 (2016): 1159–65.
17. C. J. Ley et al., "Sex- and Menopause-Associated Changes in Body-Fat Distribution," *The American Journal of Clinical Nutrition* 55, no. 5 (1992): 950–54.
18. L. M. Brown et al., "Central Effects of Estradiol in the Regulation of Food Intake, Body Weight, and Adiposity," *The Journal of Steroid Biochemistry and Molecular Biology* 122, nos. 1–3 (2010): 65–73.
19. Ley et al., "Sex- and Menopause-Associated Changes."
20. Q. Cao et al., "Waist-Hip Ratio as a Predictor of Myocardial Infarction Risk: A Systematic Review and Meta-Analysis," *Medicine* 97, no. 30 (2018); V. A. Benites-Zapata et al., "High Waist-to-Hip Ratio Levels Are Associated with Insulin Resistance Markers in Normal-Weight Women," *Diabetes Metabolic Syndrome* 13, no. 1 (2019): 636–42.
21. B. Tramunt et al., "Sex Differences in Metabolic Regulation and Diabetes Susceptibility," *Diabetologia* 63, no. 3 (2020): 453–61.
22. V. Regitz-Zagrosek et al., "Gender Aspects of the Role of the Metabolic Syndrome as a Risk Factor for Cardiovascular Disease," *Gender Medicine* 4 (2007): S162–77; E. Gerds et al., "Sex Differences in Cardiometabolic Disorders," *Nature Medicine* 25, no. 11 (2019): 1657–66.
23. S. V. Ahn et al., "Sex Difference in the Effect of the Fasting Serum Glucose Level on the Risk of Coronary Heart Disease," *Journal of Cardiology* 71, no. 2 (2018): 149–54.

24. G. A. Greendale et al., "Changes in Body Composition and Weight During the Menopause Transition," *JCI Insight* 4, no. 5 (2019).
25. A. M. Goss et al., "Longitudinal Associations of the Endocrine Environment on Fat Partitioning in Postmenopausal Women," *Obesity (Silver Spring)* 20, no. 5 (2012): 939–44; S. Ballestri et al., "NAFLD as a Sexual Dimorphic Disease: Role of Gender and Reproductive Status in the Development and Progression of Nonalcoholic Fatty Liver Disease and Inherent Cardiovascular Risk," *Advances in Therapy* 34, no. 6 (2017): 1291–326.
26. J. R. Guthrie et al., "Weight Gain and the Menopause: A 5-Year Prospective Study," *Climacteric: The Journal of the International Menopause Society* 2, no. 3 (1999): 205–11.
27. S. L. Crawford et al., "A Longitudinal Study of Weight and the Menopause Transition: Results from the Massachusetts Women's Health Study," *Menopause (New York, N.Y.)* 7, no. 2 (2000): 96–104.
28. S. C. Ho et al., "Menopausal Transition and Changes of Body Composition: A Prospective Study in Chinese Perimenopausal Women," *International Journal of Obesity* (2005) 34, no. 8 (2010): 1265–74.
29. F. Fery et al., "Hormonal and Metabolic Changes Induced by an Isocaloric Isoproteic Ketogenic Diet in Healthy Subjects," *Diabète & Métabolisme* 8, no. 4 (1982): 299–305; E. Kose et al., "Changes of Thyroid Hormonal Status in Patients Receiving Ketogenic Diet Due to Intractable Epilepsy," *Journal of Pediatric Endocrinology & Metabolism* 30, no. 4 (2017): 411–16; Y. J. Lee et al., "Longitudinal Change in Thyroid Hormone Levels in Children with Epilepsy on a Ketogenic Diet: Prevalence and Risk Factors," *Journal of Epilepsy Research* 7, no. 2 (2017): 99–105.
30. За пълен списък на хипотиреоидните симптоми вижте в моята книга „Хормонален баланс“.
31. J. Sirven et al., "The Ketogenic Diet for Intractable Epilepsy in Adults: Preliminary Results," *Epilepsia* 40, no. 12 (1999): 1721–26.
32. R. M. Kwan et al., "Effects of a Low Carbohydrate Isoenergetic Diet on Sleep Behavior and Pulmonary Functions in Healthy Female Adult Humans," *Journal of Nutrition* 116, no. 12 (1986): 2393–402.
33. A. A. Prather et al., "Poor Sleep Quality Potentiates Stress-Induced Cytokine Reactivity in Postmenopausal Women with High Visceral Abdominal Adiposity," *Brain, Behavior, Immunity* 35, no. 1 (2014): 155–62; S. K. Sweatt et al., "Sleep Quality Is Differentially Related to Adiposity in Adults," *Psychoneuroendocrinology* 98, no. 1 (2018): 46–51.
34. В това експериментално изследване на кетогенната диета при плъхове изследователите са използвали микрокомпютърна томография и хистоморфометрични анализи на дисталната бедрена кост. Те открили, че обемът на трабекуларните кости, серумният IGF-I и маркерът за образуване на кости P1NP са по-ниски при мъжки плъхове на диета с



- ниско съдържание на въглехидрати и високо съдържание на мазнини. A. Zengin et al., "Low- Carbohydrate, High-Fat Diets Have Sex-Specific Effects on Bone Health in Rats," *European Journal of Nutrition* 55, no. 7 (2016): 2307–20.
35. G. K. Schwalfenberg, "The Alkaline Diet: Is There Evidence That an Alkaline pH Diet Benefits Health?" *Journal of Environmental and Public Health* 2012, no. 1 (2012): 727630.
  36. B. E. Millen et al., "The 2015 Dietary Guidelines Advisory Committee Scientific Report: Development and Major Conclusions," *Advances in Nutrition* 7, no. 3 (2016): 438–44; Q. Qian, "Dietary Influence on Body Fluid Acid-Base and Volume Balance: The Deleterious 'Norm' Furthers and Cloaks Subclinical Pathophysiology," *Nutrients* 10, no. 6 (2018): 778.
  37. Киселинен и алкален се отнасят до рН: киселините имат ниско рН (по-малко от 7), а високото рН (над 7) е признак за алкалност. рН на кръвта е 7,4, но храните могат да оставят кисела или алкална пепел. L. Frassetto et al., "Diet, Evolution and Aging — The Pathophysiologic Effects of the Post-Agricultural Inversion of the Potassium-to-Sodium and Base-to-Chloride Ratios in the Human Diet," *European Journal of Nutrition* 40, no. 5 (2001): 200–213; M. Konner et al., "Paleolithic Nutrition: Twenty-Five Years Later," *Nutrition in Clinical Practice* 25, no. 6 (2010): 594–602. J. R. Buendia et al., "Longitudinal Effects of Dietary Sodium and Potassium on Blood Pressure in Adolescent Girls," *JAMA Pediatrics* 169, no. 6 (2015): 560–68; A. Sebastian et al., "Postulating the Major Environmental Condition Resulting in the Expression of Essential Hypertension and Its Associated Cardiovascular Diseases: Dietary Imprudence in Daily Selection of Foods in Respect of Their Potassium and Sodium Content Resulting in Oxidative Stress-Induced Dysfunction of the Vascular Endothelium, Vascular Smooth Muscle, and Perivascular Tissues," *Medical Hypotheses* 119, no. 1 (2018): 110–19.
  38. S. T. Reddy et al., "Effect of Low-Carbohydrate High-Protein Diets on Acid-Base Balance, Stone-Forming Propensity, and Calcium Metabolism," *American Journal of Kidney Diseases* 40, no. 2 (2002): 265–74; E. H. Kossoff et al., "Dietary Therapies for Epilepsy," *Biomed Journal* 36, no. 1 (2013): 2–8.
  39. L. Frassetto et al., "Potassium Bicarbonate Reduces Urinary Nitrogen Excretion in Postmenopausal Women," *The Journal of Clinical Endocrinology & Metabolism* 82, no. 1 (1997): 254–59; L. Frassetto et al., "Long-term Persistence of the Urine Calcium-Lowering Effect of Potassium Bicarbonate in Postmenopausal Women," *The Journal of Clinical Endocrinology & Metabolism* 90, no. 2 (2005): 831–34; J. A. Wass et al., "Growth Hormone and Memory," *The Journal of Endocrinology* 207, no. 2 (2010): 125–26; G. K. Schwalfenberg, "The Alkaline Diet: Is There Evidence That an Alkaline pH Diet Benefits Health?" *Journal of Environmental and Public Health* 2012, no. 1 (2012): 727630.

40. R. Solianik et al., "Two-Day Fasting Evokes Stress, but Does Not Affect Mood, Brain Activity, Cognitive, Psychomotor, and Motor Performance in Overweight Women," *Behavioural Brain Research* 338, no. 1 (2018): 166–72.
41. R. Solianik et al., "Effect of 48H Fasting on Autonomic Function, Brain Activity, Cognition, and Mood in Amateur Weight Lifters," *Biomed Research International* 2016, no. 1 (2016): 1503956.

## **2. КАК ХОРМОНЪТ НА РАСТЕЖА ВИ ПОДДЪРЖА СЛАБИ**

1. J. D. Veldhuis et al., "Somatotrophic and Gonadotrophic Axes Linkages in Infancy, Childhood, and the Puberty-Adult Transition," *Endocrine Reviews* 27, no. 2 (2006): 101–40; J. D. Veldhuis, "Aging and Hormones of the Hypothalamo-Pituitary Axis: Gonadotrophic Axis in Men and Somatotrophic Axes in Men and Women," *Ageing Research Reviews* 7, no. 3 (2008): 189–208.
2. E. Corpas et al., "Human Growth Hormone and Human Aging," *Endocrine Reviews* 14, no. 1 (1993): 20–39; A. Bartke, "Growth Hormone and Aging: Updated Review," *The World Journal of Men's Health* 37, no.1 (2019): 19–30.
3. S. Perrini et al., "Metabolic Implications of Growth Hormone Therapy," *Journal of Endocrinological Investigation — Supplements* 31, no. 9 (2008): 79–84; S. Perrini et al., "Abnormalities of Insulin-like Growth Factor-I Signaling and Impaired Cell Proliferation in Osteoblasts from Subjects with Osteoporosis," *Endocrinology* 149, no. 3 (2007): 1302–13; K. R. Short et al., "Enhancement of Muscle Mitochondrial Function by Growth Hormone," *The Journal of Clinical Endocrinology & Metabolism* 93, no. 2 (2008): 597–604; N. Møller et al., "Effects of Growth Hormone on Glucose, Lipid, and Protein Metabolism in Human Subjects," *Endocrine Reviews* 30, no. 2 (2009): 152–77.
4. L. I. Arwert et al., "The Relation Between Insulin-Like Growth Factor I Levels and Cognition in Healthy Elderly: A Meta-Analysis." *Growth hormone & IGF Research* 15, no. 6 (2005): 416–422.
5. U. J. Lewis, "Growth Hormone: What Is It and What Does It Do?" *Trends in Endocrinology & Metabolism* 3, no. 4 (1992): 117–21; M. B. Ranke et al., "Growth Hormone — Past, Present, and Future." *Nature Reviews Endocrinology* 14, no. 5 (2018): 285–300.
6. F. Mourkioti et al., "IGF-1, Inflammation, and Stem Cells: Interactions During Muscle REGEneration," *Trends in Immunology* 26, no. 10 (2005): 535–42; C. P. Velloso, "Regulation of Muscle Mass by Growth Hormone and IGF-I," *British Journal of Pharmacology* 154, no. 1 (2008): 557–68, M. E. Molitch et al., "Evaluation and Treatment of Adult Growth Hormone Deficiency: An Endocrine Society Clinical Practice Guideline," *The Journal of Clinical Endocrinology & Metabolism* 96, no. 6 (2011): 1587–609.

7. A. L. Cardoso et al., "Towards Frailty Biomarkers: Candidates from Genes and Pathways Regulated in Aging and Age-Related Diseases," *Ageing Research Reviews* 47, no. 1 (2018): 214–77.
8. G. Vab den Berg et al., "An Amplitude-Specific Divergence in the Pulsatile Mode of Growth Hormone (GH) Secretion Underlies the Gender Difference in Mean Growth Hormone Concentrations in Men and Premenopausal Women," *Journal of Clinical Endocrinology and Metabolism* 81, no. 7 (1996): 2460–67; J. O. Jørgensen et al., "Sex Steroids and the Growth Hormone/Insulin-like Growth Factor-I Axis in Adults," *Hormone Research in Paediatrics* 64, Suppl. 2 (2005): 37–40.
9. G. Norstedt et al., "Secretory Rhythm of Growth Hormone Regulates Sexual Differentiation of Mouse Liver," *Cell* 36, no. 4 (1984): 805–12.
10. F. Roelfsema et al., "Growth-Hormone Dynamics in Healthy Adults Are Related to Age and Sex, and Strongly Dependent on Body Mass Index," *Neuroendocrinology* 103, nos. 3–4 (2016): 335–44; J. P. Span et al., "Gender Difference in Insulin-Like Growth Factor I Response to Growth Hormone (GH) Treatment in Growth Hormone-Deficient Adults: Role of Sex Hormone Replacement," *Journal of Clinical Endocrinology Metabolism* 85, no. 3 (2000): 1121–25.
11. A. Eliakim et al., "Effect of Gender on the Growth Hormone-IGF-I Response to Anaerobic Exercise in Young Adults," *Journal of Strength and Conditioning Research* 28, no. 12, (2014): 3411–15.
12. M. Russell-Aulet et al., "Aging-Related Growth Hormone Decrease Is a Selective Hypothalamic Growth Hormone-Releasing Hormone Pulse Amplitude Mediated Phenomenon," *The Journals of Gerontology, Series A: Biological Sciences and Medical Sciences* 56, no. 2 (2001): M124–29.
13. N. Vahl et al., "Abdominal Adiposity and Physical Fitness Are Major Determinants of the Age-Associated Decline in Stimulated Growth Hormone Secretion in Healthy Adults," *The Journal of Clinical Endocrinology & Metabolism* 81, no. 6 (1996): 2209–215.
14. M. Misra et al., "Lower Growth Hormone and Higher Cortisol Are Associated with Greater Visceral Adiposity, Intramyocellular Lipids, and Insulin Resistance in Overweight Girls," *American Journal of Physiology-Endocrinology and Metabolism* 295, no. 2 (2008): E385–92.
15. I. Fukuda et al., "Serum Adiponectin Levels in Adult Growth Hormone Deficiency and Acromegaly," *Growth Hormone & IGF Research* 14, no. 6 (2004): 449–54; R. Stawerska et al., "Relationship Between IGF-I Concentration and Metabolic Profile in Children with Growth Hormone Deficiency: The Influence of Children's Nutritional State as Well as the Ghrelin, Leptin, Adiponectin, and Resistin Serum Concentrations," *International Journal of Endocrinology* 2017 (2017); E. Witkowska-Sędek et al., "The Associations Between the Growth Hormone/ Insulin-like Growth Factor-1 Axis, Adiponectin, Resistin,

- and Metabolic Profile in Children with Growth Hormone Deficiency Before and During Growth Hormone Treatment,” *Acta Biochimica Polonica* 65, no. 2 (2018): 333–40.
16. Z. P. Li et al., “Study of the Correlation Between Growth Hormone Deficiency and Serum Leptin, Adiponectin, and Visfatin Levels in Adults,” *Genetics and Molecular Research: GMR* 13, no. 2 (2014): 4050–56.
  17. J. D. Veldhuis et al., “Distinctive Inhibitory Mechanisms of Age and Relative Visceral Adiposity on Growth Hormone Secretion in Pre- and Postmenopausal Women Studied Under a Hypogonadal Clamp,” *The Journal of Clinical Endocrinology & Metabolism* 90, no. 11 (2005): 6006–13.
  18. Li et al., “Study of the Correlation Between Growth.”
  19. Пълният списък на хормоните, които участват в регулирането на растежния хормон и/или на IGF-1, включва естроген, кортизол (тоест адреналокортикотропен хормон), хормони на щитовидната жлеза (по-конкретно контролният хормон за секрецията на щитовидната жлеза, тиреотропин-освобождаващ хормон), лутеинизиращ хормон, фоликулостимулиращ хормон, човешки хорион гонадотропин (хормона на бременността), инсулин, други растежни фактори (например тромбоцитен растежен фактор [PDGF], епидермален растежен фактор [EGF] и факторите на растеж на фибробласти [FGFs]), комбинирани с възраст, пол, хранене и други фактори от начина на живот. A. Kasprzak et al., “Insulin-like Growth Factor (IGF) Axis in Cancerogenesis.” *Mutation Research/Reviews in Mutation Research* 772, no. 1 (2017): 78–104.
  20. Препоръчвам на моите пациенти пълен хормонален панел, включително тиреостимулиращ хормон, или TSH, свободен T3, свободен T4, реверсивен T3, антитела срещу тиреоидната пероксидаза и тиреоглобулинови антитела. На база симптомите може да се наложат и допълнителни изследвания на щитовидната жлеза. Моля, обсъдете резултатите с лекар.
  21. E. Giovannucci et al., “Nutritional Predictors of Insulin-like Growth Factor I and Their Relationships to Cancer in Men,” *Cancer Epidemiology, Biomarkers, & Prevention* 12, no. 2 (2003): 84–89.
  22. S. C. Larsson et al., “Association of Diet with Serum Insulin-like Growth Factor I in Middle-Aged and Elderly Men,” *The American Journal of Clinical Nutrition* 81, no. 5 (2005): 1163–67.
  23. M. Holmes et al., “Dietary Correlates of Plasma Insulin-like Growth Factor I and Insulin-like Growth Factor Binding Protein 3 Concentrations,” *Cancer Epidemiology, Biomarkers, & Prevention* 11, no. 9 (2002): 852–61.
  24. S. M. Phillips et al., “Dietary Protein for Athletes: From Requirements to Optimum Adaptation,” *Journal of Sports Sciences* 29, Suppl. 1 (2011): S29–38; M. Huecker et al., “Protein Supplementation in Sport: Source, Timing, and Intended Benefits,” *Current Nutrition Reports* 8, no. 4 (2019): 382–396.

25. K. Zhu et al., "The Effects of a Two-Year Randomized, Controlled Trial of Whey Protein Supplementation on Bone Structure, IGF-1, and Urinary Calcium Excretion in Older Postmenopausal Women," *Journal of Bone and Mineral Research* 26, no. 9 (2011): 2298–306.
26. J. M. Bauer et al., "Effects of a Vitamin D and Leucine-Enriched Whey Protein Nutritional Supplement on Measures of Sarcopenia in Older Adults, The PROVIDE Study: A Randomized, Double-Blind, Placebo-Controlled Trial," *Journal of the American Medical Directors Association* 16, no. 9 (2015): 740–47; M. Rondanelli et al., "Whey Protein, Amino Acids, and Vitamin D Supplementation with Physical Activity Increases Fat-Free Mass and Strength, Functionality, and Quality of Life and Decreases Inflammation in Sarcopenic Elderly," *The American Journal of Clinical Nutrition* 103, no. 3 (2016): 830–40; S. Verlaan et al., "Sufficient Levels of 25-Hydroxyvitamin D and Protein Intake Required to Increase Muscle Mass in Sarcopenic Older Adults — The PROVIDE Study," *Clinical Nutrition* 37, no. 2 (2018): 551–57.
27. E. Castellero et al., "Comparison of the Effects of the n-3 Polyunsaturated Fatty Acid Eicosapentaenoic and Fenofibrate on the Inhibitory Effect of Arthritis on IGF1," *Journal of Endocrinology* 210, no. 3 (2011): 361–68.
28. По пътя на омега-3 няколко проблема могат да нарушат регулацията на ензим, наречен делта-5-десатураза, което води до производство на повече арахидонова киселина, а тя е възпалителна. Тези проблеми включват есенциална хипертония, сърдечносъдови заболявания, инсулинова резистентност, затлъстяване и метаболитен синдром.  
C. Russo et al., "Increased Membrane Ratios of Metabolite to Precursor Fatty Acid in Essential Hypertension," *Hypertension* 29, no. 4 (1997): 1058–63; B. Vessby, "Dietary Fat, Fatty Acid Composition in Plasma and the Metabolic Syndrome," *Current Opinion in Lipidology* 14, no. 1 (2003): 15–19; T. Domei et al., "Ratio of Serum n-3 to n-6 Polyunsaturated Fatty Acids and the Incidence of Major Adverse Cardiac Events in Patients Undergoing Percutaneous Coronary Intervention," *Circulation Journal* 76, (2012): 423–29; K. Inoue et al., "Low Serum Eicosapentaenoic Acid/Arachidonic Acid Ratio in Male Subjects with Visceral Obesity," *Nutrition & Metabolism* 10, no. 1 (2013): 25; E. Warensjö et al., "Fatty Acid Composition and Estimated Desaturase Activities Are Associated with Obesity and Lifestyle Variables in Men and Women," *Nutrition, Metabolism, and Cardiovascular Diseases* 16, no. 2 (2006): 128–36.
29. B. Yang et al., "Ratio of n-3/n-6 PUFAs and Risk of Breast Cancer: A Meta-Analysis of 274135 Adult Females from 11 Independent Prospective Studies," *BMC Cancer* 14, no. 1 (2014): 105.
30. T. J. Merimee et al., "Diet-Induced Alterations of Growth Hormone Secretion in Man," *Journal of Clinical Endocrinology Metabolism* 42, no. 5 (1976): 931–37; K. Y. Ho et al., "Fasting Enhances Growth Hormone Secretion and

- Amplifies the Complex Rhythms of Growth Hormone Secretion in Man," *Journal of Clinical Investigation* 81, no. 4 (1988): 968–75; H. Nørrelund et al., "Modulation of Basal Glucose Metabolism and Insulin Sensitivity by Growth Hormone and Free Fatty Acids During Short-Term Fasting," *European Journal Endocrinology* 150, no. 6 (2004): 779–87; H. Nørrelund, "The Metabolic Role of Growth Hormone in Humans with Particular Reference to Fasting," *Growth Hormone & IGF Research* 15, no. 2 (2005): 95–122.
31. H. E. Bergan et al., "Nutritional State Modulates Growth Hormone–Stimulated Lipolysis," *General and Comparative Endocrinology* 217–218 (2015): 1–9.
  32. B. D. Horne et al., "Relation of Routine, Periodic Fasting to Risk of Diabetes Mellitus, and Coronary Artery Disease in Patients Undergoing Coronary Angiography," *American Journal of Cardiology* 109, no. 11 (2012): 1558–62.
  33. R. Gatti et al., "IGF-I/IGFBP System: Metabolism Outline and Physical Exercise." *Journal of Endocrinological Investigation* 35, no. 7 (2012): 699–707.
  34. J. Leppäluoto et al., "Heat Exposure Elevates Plasma Immunoreactive Growth Hormone–Releasing Hormone Levels in Man," *The Journal of Clinical Endocrinology & Metabolism* 65, no. 5 (1987): 1035–38; J. Sirviö et al., "Adenohypophyseal Hormone Levels During Hyperthermia," *Endocrinologie* 25, no. 1 (1987): 21–23; K. Kukkonen-Harjula et al., "How the Sauna Affects the Endocrine System," *Annals of Clinical Research* 20, no. 4 (1988): 262–66; K. Kukkonen-Harjula et al., "Haemodynamic and Hormonal Responses to Heat Exposure in a Finnish Sauna Bath," *European Journal of Applied Physiology and Occupational Physiology* 58, no. 5 (1989): 543–50; D. Jezová et al., "Sex Differences in Endocrine Response to Hyperthermia in Sauna," *Acta Physiologica Scandinavica* 150, no. 3 (1994): 293–98.
  35. R. Lammintausta et al., "Change in Hormones Reflecting Sympathetic Activity in the Finnish Sauna," *Annals of Clinical Research* 8, no. 4 (1976): 266–71.
  36. M. Välimäki et al., "Effect of Ethanol on Serum Concentrations of Somatomedin C and the Growth hormone (GH) Secretion Stimulated by the Releasing Hormone (GHRH)," *Alcohol and Alcoholism* 1 (1987): 557–59; L. Dees et al., "Effects of Ethanol During the Onset of Female Puberty," *Neuroendocrinology* 51, no. 1 (1990): 64–69; M. Välimäki et al., "The Pulsatile Secretion of Gonadotropins and Growth Hormone, and the Biological Activity of Luteinizing Hormone in Men Acutely Intoxicated with Ethanol," *Alcoholism: Clinical and Experimental Research* 14, no. 6 (1990): 928–31; N. Rachdaoui et al., "Pathophysiology of the Effects of Alcohol Abuse on the Endocrine System," *Alcohol Research: Current Reviews* 38, no. 2 (2017): 255–76.
  37. A. De Spiegeleer et al., "Pharmacological Interventions to Improve Muscle Mass, Muscle Strength, and Physical Performance in Older People: An Umbrella Review of Systematic Reviews and Meta-Analyses," *Drugs & Aging* 35, no. 8 (2018): 719–34.

38. A. R. Martineau et al., "Vitamin D Supplementation to Prevent Acute Respiratory Tract Infections: Systematic Review and Meta-Analysis of Individual Participant Data," *British Medical Journal* (2017): 356: i6583.
39. M. R. Blackman et al., "Growth Hormone and Sex Steroid Administration in Healthy Aged Women and Men: A Randomized Controlled Trial," *Journal of the American Medical Association* 288, no. 18 (2002): 2282–92; H. Liu et al., "Systematic Review: The Safety and Efficacy of Growth Hormone in the Healthy Elderly," *Annals of Internal Medicine* 146, no. 2 (2007): 104–15.
40. S. M. Orme et al., "Mortality and Cancer Incidence in Acromegaly: A Retrospective Cohort Study," *The Journal of Clinical Endocrinology & Metabolism* 83, no. 8 (1998): 2730–34; W. E. Sonntag et al., "Adult-Onset Growth Hormone and Insulin-Like Growth Factor I Deficiency Reduces Neoplastic Disease, Modifies Age-Related Pathology, and Increases Life Span," *Endocrinology* 146, no. 7 (2005): 2920–32; A. J. Swerdlow et al., "Cancer Risks in Patients Treated with Growth Hormone in Childhood: The SAGhE European Cohort Study," *The Journal of Clinical Endocrinology & Metabolism* 102, no. 5 (2017): 1661–72.
41. J. Berlanga-Acosta et al., "Synthetic Growth Hormone–Releasing Peptides (GHRPs): A Historical Appraisal of the Evidences Supporting Their Cytoprotective Effects," *Clinical Medicine Insights: Cardiology* 11, no. 1 (2017); J. T. Sigalos et al., "Growth Hormone Secretagogue Treatment in Hypogonadal Men Raises Serum Insulin-Like Growth Factor-1 Levels," *American Journal Men's Health* 11, no. 6 (2017): 1752–57; J. T. Sigalos et al., "The Safety and Efficacy of Growth Hormone Secretagogues," *Sexual Medicine Reviews* 6, no. 1 (2018).
42. B. C. Nindl et al., "Insulin-like Growth Factor I as a Biomarker of Health, Fitness, and Training Status," *Medicine and Science in Sports and Exercise* 42, no. 1 (2010): 39–49.

### **3. ТЕСТОСТЕРОНЪТ НЕ Е САМО ЗА МЪЖЕ**

1. C. Longcope, "Adrenal and Gonadal Androgen Secretion in Normal Females," *Clinics in Endocrinology and Metabolism* 15, no. 2 (1986): 213–28.
2. S. L. Davison et al., "Androgen Levels in Adult Females: Changes with Age, Menopause, and Oophorectomy," *The Journal of Clinical Endocrinology & Metabolism* 90, no. 7 (2005): 3847–53.
3. За да изследвате вашия тестостерон, препоръчвам да попитате лекар за кръвно изследване, което включва общ и свободен тестостерон (тоест количеството, което е биологично налично, за да окаже въздействие върху вашите клетки). Можете да изследвате в урина DHEA, тестостерон и свързаните хормони надолу по веригата.
4. Davison et al., "Androgen Levels in Adult Females."

5. N. Orentreich et al., "Age Changes and Sex Differences in Serum Dehydroepiandrosterone Sulfate Concentrations Throughout Adulthood," *Journal of Clinical Endocrinology Metabolism* 59, no. 3 (1984): 551–55.
6. Davison et al., "Androgen Levels in Adult Females"; R. Haring et al., "Age-Specific Reference Ranges for Serum Testosterone and Androstenedione Concentrations in Women Measured by Liquid Chromatography-Tandem Mass Spectrometry," *Journal of Clinical Endocrinology Metabolism* 97, no. 2 (2012): 408–15.
7. C. M. Coenen et al., "Changes in Androgens During Treatment with Four Low- Dose Contraceptives," *Contraception* 53, no. 3 (1996): 171–76; Y. Zimmerman et al., "The Effect of Combined Oral Contraception on Testosterone Levels in Healthy Women: A Systematic Review and Meta-Analysis," *Human Reproductive Update* 20, no. 1 (2014): 76–105; N. Zethraeus et al., "Combined Oral Contraceptives and Sexual Function in Women — A Double-Blind, Randomized, Placebo- Controlled Trial," *Journal of Clinical Endocrinology Metabolism* 101, no. 11 (2016): 4046–53; S. Both et al., "Hormonal Contraception and Female Sexuality: Position Statements from the European Society of Sexual Medicine (ESSM)," *Journal of Sexual Medicine* 16, no. 11 (2019): 1681–95.
8. Статините изчерпват CoQ10, селен, селенопротеини, омега-3 мастни киселини, токофероли и токотриеноли, K<sub>2</sub>, други мастрозтворими витамини, Неме А карнитин, слободен Т3, креатин, мед и цинк. P. H. Langsjoen et al., "The Clinical Use of HMG CoA-Reductase Inhibitors and the Associated Depletion of Coenzyme Q10: A Review of Animal and Human Publications," *Biofactors* 18, nos. 1–4 (2003): 101–11; C. R. Harper et al., "Evidence- Based Management of Statin Myopathy," *Current Atherosclerosis Reports* 12, no. 5 (2010): 322–30; H. Qu et al., "Effects of Coenzyme Q10 on Statin-Induced Myopathy: An Updated Meta-Analysis of Randomized Controlled Trials," *Journal of the American Heart Association* 7, no. 19 (2018): e009835.
9. J. Y. Shin et al., "Are Cholesterol and Depression Inversely Related? A Meta-Analysis of the Association Between Two Cardiac Risk Factors," *Annals of Behavioral Medicine* 36, no. 1 (2008): 33–43; G. Corona et al., "The Effect of Statin Therapy on Testosterone Levels in Subjects Consulting for Erectile Dysfunction," *Journal of Sexual Medicine* 7, no. 4, part 1 (2010): 1547–56; E. J. Giltay et al., "Salivary Testosterone: Associations with Depression, Anxiety Disorders, and Antidepressant Use in a Large Cohort Study," *Journal of Psychosomatic Research* 72, no. 3 (2012): 205–13; G. Roberto et al., "Statin-Associated Gynecomastia: Evidence Coming from the Italian Spontaneous ADR Reporting Database and Literature," *European Journal of Clinical Pharmacology* 68, no. 6 (2012): 1007–11; C. M. Schooling et al., "The Effect of Statins on Testosterone in Men and Women: A Systematic Review And Meta-Analysis of Randomized Controlled Trials," *BMC Medicine* 11, no. 1 (2013): 57.



10. L. Mernone et al., "Psychobiological Factors of Sexual Functioning in Aging Women — Findings from the Women 40+ Healthy Aging Study," *Frontiers in Psychology* 10, no. 1 (2019): 546.
11. S. R. Davis et al., "Circulating Androgen Levels and Self-Reported Sexual Function in Women," *Journal of the American Medical Association* 294, no. 1 (2005): 91–96.
12. R. Basson et al., "Role of Androgens in Women's Sexual Dysfunction," *Menopause* 17, no. 5 (2010): 962–71; S. Wåhlin-Jacobsen et al., "Is There a Correlation Between Androgens and Sexual Desire in Women?" *Journal of Sexual Medicine* 12, no. 2 (2015): 358–73.
13. Mernone et al., "Psychobiological Factors of Sexual Functioning."
14. S. R. Davis et al., "Global Consensus Position Statement on the Use of Testosterone Therapy for Women," *The Journal of Clinical Endocrinology and Metabolism* 104, no. 10 (2019): 4660–66.
15. C. Bentley et al., "Dehydroepiandrosterone: A Potential Therapeutic Agent in the Treatment and Rehabilitation of the Traumatically Injured Patient," *Burns & Trauma* 7, no. 26 (2019).
16. H. E. Nagels et al., "Androgens (dehydroepiandrosterone or testosterone) for Women Undergoing Assisted Reproduction," *Cochrane Database of Systemic Reviews* 11, no. 1 (2015).
17. G. P. Williams, "The Role of Oestrogen in the Pathogenesis of Obesity, Type 2 Diabetes, Breast Cancer, and Prostate Disease," *European Journal of Cancer Prevention* 19, no. 4 (2010): 256–71; J. McHenry et al., "Sex Differences in Anxiety and Depression: Role of Testosterone," *Front Neuroendocrinology* 35, no. 1 (2014): 42– 57; F. Saad, "The Emancipation of Testosterone from Niche Hormone to Multi- System Player," *Asian Journal of Andrology* 17, no. 1 (2015): 58–60; L. Y. Hui et al., "Association Between MKP-1, BDNF, and Gonadal Hormones with Depression on Perimenopausal Women," *Journal of Women's Health* 25, no. 1 (2016): 71–77; S. Rovira-Llopis et al., "Low Testosterone Levels Are Related to Oxidative Stress, Mitochondrial Dysfunction, and Altered Subclinical Atherosclerotic Markers in Type 2 Diabetic Male Patients," *Free Radical Biology & Medicine* 108, no. 1 (2017): 155–62; H. O. Santos, "Ketogenic Diet and Testosterone Increase: Is the Increased Cholesterol Intake Responsible? To What Extent and Under What Circumstances Can There Be Benefits?" *Hormones (Athens)* 16, no. 3 (2017): 150–60.
18. X. Zhang et al., "Postmenopausal Plasma Sex Hormone Levels and Breast Cancer Risk over 20 Years of Follow-Up," *Breast Cancer Research and Treatment* 137, no. 3 (2013): 883–92; R. T. Fortner et al., "Premenopausal Endogenous Steroid Hormones and Breast Cancer Risk: Results from the Nurses' Health Study II," *Breast Cancer Research* 15, no. 2 (2013): R19; Endogenous Hormones and Breast Cancer Collaborative Group et al., "Sex Hormones and Risk of Breast Cancer in Premenopausal Women: A Collaborative Reanalysis

- of Individual Participant Data from Seven Prospective Studies,” *The Lancet Oncology* 14, no. 10 (2013): 1009–19; R. Kaaks et al., “Premenopausal Serum Sex Hormone Levels in Relation to Breast Cancer Risk, Overall and by Hormone Receptor Status — Results from the EPIC Cohort,” *International Journal of Cancer* 134, no. 8 (2014): 1947–57; R. Glaser et al., “Testosterone and Breast Cancer Prevention,” *Maturitas* 82, no. 3 (2015): 291–95; K. A. Bertrand et al., “Circulating Hormones and Mammographic Density in Premenopausal Women,” *Hormones & Cancer* 9, no. 2 (2018): 117–27.
19. J. L. Shifren et al., “Transdermal Testosterone Treatment in Women with Impaired Sexual Function After Oophorectomy,” *New England Journal of Medicine* 343, no. 10 (2000): 682–88; R. Goldstat et al., “Transdermal Testosterone Therapy Improves Well-Being, Mood, and Sexual Function in Premenopausal Women,” *Menopause* 10, no. 5 (2003): 390–98; E. J. Hermans et al., “Exogenous Testosterone Attenuates the Integrated Central Stress Response in Healthy Young Women,” *Psychoneuroendocrinology* 32, nos. 8–10 (2007): 1052–61; K. K. Miller et al., “Low-Dose Transdermal Testosterone Augmentation Therapy Improves Depression Severity in Women,” *CNS Spectrums* 14, no. 12 (2009): 688–94.
  20. B. C. Trainor et al. “Testosterone Promotes Paternal Behaviour in a Monogamous Mammal via Conversion to Oestrogen,” *Proceedings of the Royal Society of London, Series B: Biological Sciences* 269, no. 1493 (2002): 823–29.
  21. Метаболитните проблеми, свързани със СПКЯ, включват непоносимост към глюкоза, метаболитен синдром и диабет от втори тип. L. J. Moran et al., “Impaired Glucose Tolerance, Type 2 Diabetes, and Metabolic Syndrome in Polycystic Ovary Syndrome: A Systematic Review and Meta-Analysis,” *Human Reproduction Update* 16, no. 4 (2010): 347–63; N. S. Kakoly et al., “Ethnicity, Obesity, and the Prevalence of Impaired Glucose Tolerance and Type 2 Diabetes in PCOS: A Systematic Review and Meta-Regression,” *Human Reproduction Update* 24, no. 4 (2018): 455–67.
- Проблемите с психичното здраве, свързани със СПКЯ, включват тревожност, депресия, неудовлетвореност от тялото и по-ниско качество на живот. S. Elsenbruch et al., “Quality of Life, Psychosocial Well-Being, and Sexual Satisfaction in Women with Polycystic Ovary Syndrome,” *The Journal of Clinical Endocrinology & Metabolism* 88, no. 12 (2003): 5801–7; M. J. Himelein et al., “Depression and Body Image Among Women with Polycystic Ovary Syndrome,” *Journal of Health Psychology* 11, no. 4 (2006): 613–25; L. M. Pastore et al., “Depression Symptoms and Body Dissatisfaction Association Among Polycystic Ovary Syndrome Women,” *Journal of Psychosomatic Research* 71, no. 4 (2011): 270–76; A. F. Nasiri et al., “The Experience of Women Affected by Polycystic Ovary Syndrome: A Qualitative Study from Iran,” *International Journal of Endocrinology and Metabolism* 12, no. 2 (2014); C. Kaczmarek et al., “Health-related Quality of Life in Adolescents and Young

- Adults with Polycystic Ovary Syndrome: A Systematic Review,” *Journal of Pediatric and Adolescent Gynecology* 29, no. 6 (2016): 551–57.
22. L. J. Moran et al., “Sex Hormone Binding Globulin, but Not Testosterone, Is Associated with the Metabolic Syndrome in Overweight and Obese Women with Polycystic Ovary Syndrome,” *Journal of Endocrinological Investigation* 36, no. 11 (2013): 1004–10.
  23. X. Zhang et al., “The Effect of Low Carbohydrate Diet on Polycystic Ovary Syndrome: A Meta-Analysis of Randomized Controlled Trials,” *International Journal of Endocrinology* 2019, no. 4386401 (2019): 1–14.
  24. J. C. Mavropoulos et al., “The Effects of a Low-Carbohydrate, Ketogenic Diet on the Polycystic Ovary Syndrome: A Pilot Study,” *Nutrition & Metabolism (Lond)* 2, no. 35 (2005); G. Muscogiuri et al., “Current Insights into Inositol Isoforms, Mediterranean, and Ketogenic Diets for Polycystic Ovary Syndrome: From Bench to Bedside,” *Current Pharmaceutical Design* 22, no. 36 (2016): 5554–57; R. K. Stocker et al., “Ketogenic Diet and Its Evidence-Based Therapeutic Implementation in Endocrine Diseases,” *Praxis (Bern 1994)* 108, no. 8 (2019): 541–53 (Статията е на немски. Издателят предоставя абстракт на немски.)
  25. L. Chen et al., “Sugar-Sweetened Beverage Intake and Serum Testosterone Levels in Adult Males 20–39 Years Old in the United States,” *Reproductive Biology Endocrinology* 16, no. 1 (2018).
  26. Обърнете внимание, че напитката е съдържала 30 грама глюкоза и 30 грама протеин. A. Schwartz et al., “Acute Decrease in Serum Testosterone After a Mixed Glucose and Protein Beverage in Obese Peripubertal Boys,” *Clinical Endocrinology* 83, no. 3 (2015): 332–38.
  27. Обърнете внимание, че напитката е съдържала 75 грама глюкоза. L. M. Caronia et al., “Abrupt Decrease in Serum Testosterone Levels After an Oral Glucose Load in Men: Implications for Screening for Hypogonadism,” *Clinical Endocrinology* 78, no. 2 (2013): 291–96.
  28. N. M. Wedick et al., “The Effects of Caffeinated and Decaffeinated Coffee on Sex Hormone–Binding Globulin and Endogenous Sex Hormone Levels: A Randomized Controlled Trial,” *Nutrition Journal* 11, no. 1 (2012): 86.
  29. K. C. Schliep et al., “Serum Caffeine and Paraxanthine Concentrations and Menstrual Cycle Function: Correlations with Beverage Intakes and Associations with Race, Reproductive Hormones, and Anovulation in the BioCycle Study,” *The American Journal of Clinical Nutrition* 104, no. 1 (2016): 155–63.
  30. R. L. Ferrini et al., “Caffeine Intake and Endogenous Sex Steroid Levels in Postmenopausal Women: The Rancho Bernardo Study,” *American Journal of Epidemiology* 144, no. 7 (1996): 642–44.
  31. D. Hang et al., “Coffee Consumption and Plasma Biomarkers of Metabolic and Inflammatory Pathways in US Health Professionals,” *The American Journal of Clinical Nutrition* 109, no. 3 (2019): 635–47.

32. T. Hu et al., "Testosterone-Associated Dietary Pattern Predicts Low Testosterone Levels and Hypogonadism," *Nutrients* 10, no. 11 (2018): 1786.
33. Wilson et al., "The Effects of Ketogenic Dieting."
34. K. Z. de Souza et al., "Efficacy of *Tribulus terrestris* for the Treatment of Hypoactive Sexual Desire Disorder in Postmenopausal Women: A Randomized, Double-Blinded, Placebo-Controlled Trial," *Menopause* 23, no. 11 (2016): 1252–56; F. B. C. Vale et al., "Efficacy of *Tribulus terrestris* for the Treatment of Premenopausal Women with Hypoactive Sexual Desire Disorder: A Randomized Double-Blinded, Placebo-Controlled Trial," *Gynecological Endocrinology* 34, no. 5 (2018): 442–45; S. Palacios et al., "Effect of a Multi-Ingredient-Based Food Supplement on Sexual Function in Women with Low Sexual Desire," *BMC Women's Health: London* 19, no. 1 (2019): 58.
35. E. Steels et al., "Efficacy of a Proprietary Trigonella Foenum-Graecum L. Of-Husked Seed Extract in Reducing Menopausal Symptoms in Otherwise Healthy Women: A Double-Blind, Randomized, Placebo-Controlled Study," *Phytotherapy Research* 31, no. 9 (2017): 1316–22; S. Begum et al., "A Novel Extract of Fenugreek Husk (Fenusmart™) Alleviates Postmenopausal Symptoms and Helps to Establish the Hormonal Balance: A Randomized, Double-Blind, Placebo-Controlled Study," *Phytotherapy Research* 30, no. 11 (2016): 1775–84; A. Rao et al., "Influence of a Specialized Trigonella Foenum-Graecum Seed Extract (Libifem), on Testosterone, Estradiol, and Sexual Function in Healthy Menstruating Women: A Randomised Placebo-Controlled Study," *Phytotherapy Research* 29, no. 8 (2015): 1123–30.
36. de Souza et al., "Efficacy of *Tribulus terrestris*"; Vale et al., "Efficacy of *Tribulus terrestris*."
37. T. Takeuchi et al., "Serum Bisphenol A Concentrations Showed Gender Differences, Possibly Linked to Androgen Levels," *Biochemical and Biophysical Research Communications* 291, no. 1 (2002): 76–78; A. Tomza-Marciniak et al., "Effect of Bisphenol A on Reproductive Processes: A Review of In Vitro, In Vivo, and Epidemiological Studies," *Journal of Applied Toxicology* 38, no. 1 (2018): 51–80; Y. Hu et al., "The Association Between the Environmental Endocrine Disruptor Bisphenol A and Polycystic Ovary Syndrome: A Systematic Review and Meta-Analysis," *Gynecological Endocrinology* 34, no. 5 (2018): 370–77; A. Konieczna et al., "Serum Bisphenol A Concentrations Correlate with Serum Testosterone Levels in Women with Polycystic Ovary Syndrome," *Reproductive Toxicology* 82, no. 1 (2018): 32–37.
38. L. Le Corre et al., "BPA, an Energy Balance Disruptor," *Critical Reviews Food Science and Nutrition* 55, no. 6 (2015): 769–77; S. Legeay et al., "Is Bisphenol A an Environmental Obesogen?" *Fundamental & Clinical Pharmacology* 31, no. 6 (2017): 594–609; J. J. Heindel et al., "Environmental Obesogens: Mechanisms and Controversies," *Annual Review of Pharmacology Toxicology* 59, no. 1 (2019): 89–106; B. S. Rubin et al., "The Case for BPA

as an Obesogen: Contributors to the Controversy,” *Front Endocrinology (Lausanne)* 10, no. 30 (2019).

39. R. H. W. van Lunsen et al., “Maintaining Physiologic Testosterone Levels During Combined Oral Contraceptives by Adding Dehydroepiandrosterone: II. Effects on Sexual Function. A Phase II Randomized, Double-Blind, Placebo-Controlled Study,” *Contraception* 98, no. 1 (2018): 56–62.
40. C. S. Scheffers et al., “Dehydroepiandrosterone for Women in the Peri- or Postmenopausal Phase,” *Cochrane Database Systemic Reviews* 1 (2015).

#### **4. ПАРАДОКСЪТ НА КЕТОДИЕТАТА**

1. E. Vining et al., “A Multicenter Study of the Efficacy of the Ketogenic Diet,” *Archives of Neurology* 55, no. 11 (1998): 1433–37; D. R. Nordli et al., “Experience with the Ketogenic Diet in Infants,” *Pediatrics* 108, no. 1 (2001): 129–33; K. Tran et al., “Can You Predict an Immediate, Complete, and Sustained Response to the Ketogenic Diet?” *Epilepsia* 46, no. 4 (2005): 580–82; E. Neal et al., “The Ketogenic Diet for the Treatment of Childhood Epilepsy: A Randomised Controlled Trial,” *The Lancet Neurology* 7, no. 6 (2008): 500–506; L. Shah et al., “How Often Is Antiseizure Drug-Free Ketogenic Diet Therapy Achieved?” *Epilepsy & Behavior* 93, no. 1 (2019): 29–31; B. Gilbert, “Benefits and Complications of the Ketogenic Diet for Epilepsy,” *Neurology Advisor*, <https://www.neurologyadvisor.com/topics/epilepsy/benefits-andcomplications-of-theketogenic-diet-for-epilepsy/>. Достъпно на 27.10.2019 година.
2. T. J. W. McDonald et al., “Ketogenic Diets for Adult Neurological Disorders,” *Neurotherapeutics* 15, no. 4 (2018): 1018–31; M. Rusek et al., “Ketogenic Diet in Alzheimer’s Disease,” *International Journal of Molecular Sciences* 20, no. 16 (2019): 3892; G. M. Broom et al., “The Ketogenic Diet as a Potential Treatment and Prevention Strategy for Alzheimer’s Disease,” *Nutrition* 60 (2019): 118–21; R. Nagpal et al., “Modified Mediterranean-Ketogenic Diet Modulates Gut Microbiome and Short-Chain Fatty Acids in Association with Alzheimer’s Disease Markers in Subjects with Mild Cognitive Impairment,” *EBioMedicine* 47 (2019): 529–42.
3. H. Y. Chung et al., “Rationale, Feasibility, and Acceptability of Ketogenic Diet for Cancer Treatment,” *Journal of Cancer Prevention* 22, no. 3 (2017): 127–134; D. D. Weber et al., “Ketogenic Diet in the Treatment of Cancer — Where Do We Stand?” *Molecular Metabolism* 33 (2020): 102–21.
4. B. D. Hopkins et al., “Suppression of Insulin Feedback Enhances the Efficacy of PI3K Inhibitors,” *Nature* 560, no. 7719 (2018): 499–503.
5. A. F. Luat et al., “The Ketogenic Diet: A Practical Guide for Pediatricians,” *Pediatric Annals* 45, no. 12 (2016): e446–50; G. Muscogiuri et al., “The

- Management of Very Low-Calorie Ketogenic Diet in Obesity Outpatient Clinic: A Practical Guide,” *Journal of Translational Medicine* 17, no. 1 (2019): 356.
6. A. Johannessen et al., “Prolactin, Growth Hormone, Thyrotropin, 3, 5, 3'-Triiodothyronine, and Thyroxine Responses to Exercise After Fat- and Carbohydrate-Enriched Diet,” *The Journal of Clinical Endocrinology & Metabolism* 52, no. 1 (1981): 56–61; L. J. McCargar et al., “Dietary Carbohydrate-to-Fat Ratio: Influence on Whole-Body Nitrogen Retention, Substrate Utilization, and Hormone Response in Healthy Male Subjects,” *The American Journal of Clinical Nutrition* 49, no. 6 (1989): 1169–78; J. Langfort et al., “Effect of Low-Carbohydrate-Ketogenic Diet on Metabolic and Hormonal Responses to Graded Exercise in Men,” *Journal of Physiology and Pharmacology: An Official Journal of the Polish Physiological Society* 47, no. 2 (1996): 361–71; F. Q. Nuttall et al., “The Metabolic Response to a High-Protein, Low-Carbohydrate Diet in Men with Type 2 Diabetes Mellitus,” *Metabolism* 55, no. 2 (2006): 243–51; A. E. Lima-Silva et al., “Low Carbohydrate Diet Affects the Oxygen Uptake on-Kinetics and Rating of Perceived Exertion in High-Intensity Exercise,” *Psychophysiology* 48, no. 2 (2011): 277–84; A. Zajac et al., “The Effects of a Ketogenic Diet on Exercise Metabolism and Physical Performance in Off-Road Cyclists,” *Nutrients* 6, no. 7 (2014): 2493–508; K. D. Hall et al., “Energy Expenditure and Body Composition Changes After an Isocaloric Ketogenic Diet in Overweight and Obese Men,” *The American Journal of Clinical Nutrition* 104, no. 2 (2016): 324–33; S. Vargas et al., “Efficacy of Ketogenic Diet on Body Composition During Resistance Training in Trained Men: A Randomized Controlled Trial,” *Journal of the International Society of Sports Nutrition* 15, no. 1 (2018): 31.
  7. A. M. Johnstone et al., “Effects of a High-Protein Ketogenic Diet on Hunger, Appetite, and Weight Loss in Obese Men Feeding Ad Libitum,” *The American Journal of Clinical Nutrition* 87, no. 1 (2008): 44–55.
  8. S. R. Nymo et al., “Timeline of Changes in Appetite During Weight Loss with a Ketogenic Diet,” *International Journal of Obesity* 41, no. 8 (2017): 1224–31.
  9. Hall et al., “Energy Expenditure and Body Composition.”
  10. Vargas et al., “Efficacy of Ketogenic Diet.”
  11. M. J. Sharman et al., “A Ketogenic Diet Favorably Affects Serum Biomarkers for Cardiovascular Disease in Normal-Weight Men,” *The Journal of Nutrition* 132, no. 7 (2002): 1879–85.
  12. K. K. Ryan et al., “Dietary Manipulations That Induce Ketosis Activate the HPA Axis in Male Rats and Mice: A Potential Role for Fibroblast Growth Factor-21,” *Endocrinology* 159, no. 1 (2017): 400–413.
  13. C. M. Young et al., “Effect on Body Composition and Other Parameters in Obese Young Men of Carbohydrate Level of Reduction Diet,” *The American Journal of Clinical Nutrition* 24, no. 3 (1971): 290–96; S. B. Hulley et al., “Lipid and Lipoprotein Responses of Hypertriglyceridaemic Outpatients to a

- Low-Carbohydrate Modification of the AHA Fat-Controlled Diet,” *The Lancet* 300, no. 7777 (1972): 551–55; B. Fagerberg et al., “Weight-Reducing Diets: Role of Carbohydrates on Sympathetic Nervous Activity and Hypotensive Response,” *International Journal of Obesity* 8, no. 3 (1984): 237–43; J. W. Helge et al., “Prolonged Adaptation to Fat-Rich Diet and Training: Effects on Body Fat Stores and Insulin Resistance in Man,” *International Journal of Obesity* 26, no. 8 (2002): 1118–24; J. S. Volek et al., “Body Composition and Hormonal Responses to a Carbohydrate-Restricted Diet,” *Metabolism-Clinical and Experimental* 51, no. 7 (2002): 864–70; R. H. Stimson et al., “Dietary Macronutrient Content Alters Cortisol Metabolism Independently of Body Weight Changes in Obese Men,” *The Journal of Clinical Endocrinology & Metabolism* 92, no. 11 (2007): 4480–84; A. R. Lane et al., “Influence of Dietary Carbohydrate Intake on the Free Testosterone: Cortisol Ratio Responses to Short-Term Intensive Exercise Training,” *European Journal of Applied Physiology* 108, no. 6 (2010): 1125–31; K. Pilis et al., “Three-Year Chronic Consumption of Low-Carbohydrate Diet Impairs Exercise Performance and Has a Small Unfavorable Effect on Lipid Profile in Middle-Aged Men,” *Nutrients* 10, no. 12 (2018): 1914; H. S. Waldman et al., “Effects of a 15-day Low Carbohydrate, High-Fat Diet in Resistance-Trained Men,” *The Journal of Strength & Conditioning Research* 32, no. 11 (2018): 3103–11; M. M. Michalczyk et al., “Anaerobic Performance After a Low-Carbohydrate Diet (LCD) Followed by 7 Days of Carbohydrate Loading in Male Basketball Players,” *Nutrients* 11, no. 4 (2019): 778.
14. L. Stern et al., “The Effects of Low-Carbohydrate Versus Conventional Weight Loss Diets in Severely Obese Adults: One-Year Follow-Up of a Randomized Trial,” *Annals of Internal Medicine* 140, no. 10 (2004): 778–85; I. Shai et al., “Weight Loss with a Low-Carbohydrate, Mediterranean, or Low-Fat Diet,” *New England Journal of Medicine* 359, no. 3 (2008): 229–41; N. Iqbal et al., “Effects of a Low-Intensity Intervention That Prescribed a Low-Carbohydrate vs. a Low-Fat Diet in Obese, Diabetic Participants,” *Obesity* 18, no. 9 (2010): 1733–38.
  15. Volek et al., “Cardiovascular and Hormonal Aspects”; Volek et al., “Comparison of Energy-Restricted”; Dashti et al., “Long Term Effects of Ketogenic Diet”; Ruaño, “Physiogenomic Analysis of Weight Loss”; Durkalec-Michalski et al., “Effect of a Four-week Ketogenic Diet.”
  16. R. L. Williams et al., “Effectiveness of Weight Loss Interventions — Is there a Difference Between Men and Women? A Systematic Review,” *Obesity Review* 16, no. 2 (2015): 171–86.
  17. D. J. Millward et al., “Sex Differences in the Composition of Weight Gain and Loss in Overweight and Obese Adults,” *British Journal of Nutrition* 111, no. 5 (2014): 933–43.
  18. A. Wirth et al., “Gender Differences in Changes in Subcutaneous and Intra-Abdominal Fat During Weight Reduction: An Ultrasound Study,” *Obesity Research* 6, no. 6 (1998): 393–99.

19. За мъжете е с 64% по-малко вероятно да осъзнават теглото си („възприемане на теглото“), 61% по-малко вероятно да изпитват неудовлетвореност от теглото си и 45% по-малко вероятно да се опитат да отслабнат. Тенденцията при мъжете, които са се опитвали да отслабнат, е с 40% по-голяма вероятност от жените да отслабнат с 4 или повече килограма за една година, да поддържат постигнатото тегло и да увеличат упражненията. S. A. Tsai et al., “Gender Differences in Weight- Related Attitudes and Behaviors Among Overweight and Obese Adults in the United States,” *American Journal of Men’s Health* 10, no. 5 (2016): 389–98.
20. A. Furnham et al., “Body Image Dissatisfaction: Gender Differences in Eating Attitudes, Self-Esteem, and Reasons for Exercise,” *Journal of Psychology* 136, no. 6 (2002): 581–96.
21. E. Johansson et al., “Obesity and Labour Market Success in Finland: The Difference Between Having a High BMI and Being Fat,” *Economics and Human Biology* 7, no. 1 (2009): 36–45.
22. Диагностични диапазони за кортизол и глюкоза в кръвта. Измервам кортизола в кръвта и в урината. Измервам глюкозата в кръвта и интерстициалното пространство с устройство за непрекъснат мониторинг на глюкозата. Ето как Американската диабетна асоциация определя преддиабета: гранична гликемия, измерена с някоя от трите мерки – плазмена глюкоза на гладно 100 – 125 mg/dL (5,6 – 6,9 mmol/L), плазмена глюкоза на втория час 140 – 199 mg/dL (7,8 – 11,0 mmol/L) или хемоглобин A1C 5,7 – 6,4% (39 – 46 mmol/mol).  
J. S. Yudkin, “‘Prediabetes’: Are There Problems with This Label? Yes, the Label Creates Further Problems!” *Diabetes Care* 39, no. 8 (2016): 1468–71; American Diabetes Association, “2. Classification and Diagnosis of Diabetes: Standards of Medical Care in Diabetes — 2018,” *Diabetes Care* 41, no. 1 (2018): S13–27.
23. D. E. Laaksonen et al., “Serum Fatty Acid Composition Predicts Development of Impaired Fasting Glycaemia and Diabetes in Middle-aged Men,” *Diabetic Medicine* 19, no. 6 (2002): 456–64; R. M. Van Dam et al., “Dietary Fat and Meat Intake in Relation to Risk of Type 2 Diabetes in Men,” *Diabetes Care* 25, no. 3 (2002): 417–24; G. Riccardi et al., “Dietary Fat, Insulin Sensitivity, and the Metabolic Syndrome,” *Clinical Nutrition* 23, no. 4 (2004): 447–56; A. Shaheen et al., “A Hypothetical Model to Solve the Controversy over the Involvement of UCP2 in Palmitate-Induced  $\beta$ -cell Dysfunction,” *Endocrine* 54, no. 2 (2016): 276–83; M. Mazidi et al., “Dietary Food Patterns and Glucose/Insulin Homeostasis: A Cross-sectional Study Involving 24,182 Adult Americans,” *Lipids in Health and Disease* 16, no. 1 (2017): 192; M. Rapoport et al., “Triglycerides, Free Fatty Acids, and Glycemic Control: An Unresolved Puzzle,” *The Israel Medical Association Journal* 20, no. 6 (2018): 385–87; A. Julibert et al., “Total and Subtypes of Dietary Fat Intake and Its Association with Components of the Metabolic Syndrome in a Mediterranean Population at High Cardiovascular Risk,” *Nutrients* 11, no. 7 (2019): 1493.



24. J. Y. Lee et al., "Saturated Fatty Acids, but Not Unsaturated Fatty Acids, Induce the Expression of Cyclooxygenase-2 Mediated Through Toll-Like Receptor 4," *Journal of Biological Chemistry* 276, no. 20 (2001): 16683–89; J. M. Fernández-Real et al., "Insulin Resistance, Inflammation, and Serum Fatty Acid Composition." *Diabetes Care* 26, no. 5 (2003): 1362–68; K. M. Ajuwon et al., "Palmitate Activates the NF- $\kappa$ B Transcription Factor and Induces IL-6 and TNF $\alpha$  Expression in 3T3-L1 Adipocytes," *The Journal of Nutrition* 135, no. 8 (2005): 1841–46; C. Klein-Platat et al., "Plasma Fatty Acid Composition Is Associated with the Metabolic Syndrome and Low-Grade Inflammation in Overweight Adolescents," *The American Journal of Clinical Nutrition* 82, no. 6 (2005): 1178–84; A. R. Weatherill et al., "Saturated and Polyunsaturated Fatty Acids Reciprocally Modulate Dendritic Cell Functions Mediated Through TLR4," *The Journal of Immunology* 174, no. 9 (2005): 5390–97; S. Santos et al., "Systematic Review of Saturated Fatty Acids on Inflammation and Circulating Levels of Adipokines," *Nutrition Research* 33, no. 9 (2013): 687–95; J. E. Kaikkonen et al., "High Serum n6 Fatty Acid Proportion Is Associated with Lowered LDL Oxidation and Inflammation: The Cardiovascular Risk in Young Finns Study," *Free Radical Research* 48, no. 4 (2014): 420–26; C. Harris et al., "Associations Between Fatty Acids and Low-Grade Inflammation in Children from the MELISSAplus Birth Cohort Study," *European Journal of Clinical Nutrition* 71, no. 11 (2017): 1303–11; D. M. Rocha et al., "The Role of Dietary Fatty Acid Intake in Inflammatory Gene Expression: A Critical Review," *São Paulo Medical Journal* 135, no. 2 (2017): 157–68.
25. L. Arab et al., "Biomarkers and the Measurement of Fatty Acids," *Public Health Nutrition* 5, no. 6a (2002): 865–71; C. Kasapis et al., "The Effects of Physical Activity on Serum C-Reactive Protein and Inflammatory Markers: A Systematic Review," *Journal of the American College of Cardiology* 45, no. 10 (2005): 1563–69; M. Gleeson et al., "The Anti-Inflammatory Effects of Exercise: Mechanisms and Implications for the Prevention and Treatment of Disease," *Nature Reviews Immunology* 11, no. 9 (2011): 607–15; B. Ruiz-Núñez et al., "Lifestyle and Nutritional Imbalances Associated with Western Diseases: Causes and Consequences of Chronic Systemic Low-Grade Inflammation in an Evolutionary Context," *The Journal of Nutritional Biochemistry* 24, no. 7 (2013): 1183–1201; S. Santos et al., "Fatty Acids Derived from a Food Frequency Questionnaire and Measured in the Erythrocyte Membrane in Relation to Adiponectin and Leptin Concentrations," *European Journal of Clinical Nutrition* 68, no. 5 (2014): 555–60; M. Mazidi et al., "Impact of the Dietary Fatty Acid Intake on C-Reactive Protein Levels in US Adults," *Medicine* 96, no. 7 (2017): e5736.
26. Rodrigues et al., "The Action of  $\beta$ -hydroxybutyrate."
27. G. Beccuti et al., "Sleep and Obesity," *Current Opinion in Clinical Nutrition and Metabolic Care* 14, no. 4 (2011): 402–12.

## 5. КАК ДА ЗАПОЧНЕТЕ И КАКВО ДА ЯДЕТЕ

1. T. L. Stanley et al., "Effects of Growth Hormone-Releasing Hormone on Visceral Fat, Metabolic, and Cardiovascular Indices in Human Studies," *Growth Hormone & IGF Research* 25, no. 2 (2015): 59–65.
2. L. R. Squire et al., "Conscious and Unconscious Memory Systems," *Cold Spring Harbor Perspectives in Biology* 7, no. 3 (2015): a021667; J. Goodman et al., "Memory Systems and the Addicted Brain," *Frontiers in Psychiatry* 7 (2016): 24; M. M. Torregrossa et al., "Neuroscience of Learning and Memory for Addiction Medicine: From Habit Formation to Memory Reconsolidation," *Progress in Brain Research* 223 (2016): 91–113; J. Goodman et al., "The Dorsolateral Striatum Selectively Mediates Extinction of Habit Memory," *Neurobiology of Learning and Memory* 136 (2016): 54–62; L. Mang et al., "The Influence of Mood and Attitudes Towards Eating on Cognitive and Autobiographical Memory Flexibility in Female University Students," *Psychiatry Research* 269 (2018): 444–49.
3. E. Patrono et al., "Transitionality in Addiction: A 'Temporal Continuum' Hypotheses Involving the Aberrant Motivation, the Hedonic Dysregulation, and the Aberrant Learning," *Medical Hypotheses* 93 (2016): 62–70.
4. V. Voon, "Cognitive Biases in Binge Eating Disorder: The Hijacking of Decision Making," *CNS Spectrums* 20, no. 6 (2015): 566–73.
5. J. Goodman et al., "Enhancing and Impairing Extinction of Habit Memory Through Modulation of NMDA Receptors in the Dorsolateral Striatum," *Neuroscience* 352 (2017): 216–25.
6. Mang et al., "The Influence of Mood and Attitudes."
7. За цели храни вземете общите въглехидрати и извадете фибрите (в грамове), за да определите нетните си въглехидрати. За преработени храни извадете фибрите (в грамове) и захарния алкохол (в грамове от общите въглехидрати в грамове), за да определите нетните си въглехидрати.
8. Лабораторните изследвания за инсулинова резистентност отчитат както нарушена глюкоза на гладно, така и нарушен глюкозен толеранс. Това са диагностичните критерии за оптимално състояние, гранично състояние, преддиабет и диабет, които използвам в медицинската си практика въз основа на най-добрите доказателства. Трябва да знаете обаче, че много организации, от Американската диабетна асоциация до Световната здравна организация, използват критерии, за които липсва консенсус.
  - Глюкоза на гладно: оптимална 70 – 85, гранична 86 – 99, преддиабет 100 – 125, диабет над 126 mg/dL.
  - Непрекъснат глюкозен монитор с оптимална средна глюкоза под 100 и стандартно отклонение, по-малко от 15 mg/dL (според мен,

въз основа на клиничния ми опит и предпочитания от мен начин за диагностициране на проблеми с инсулина).

- Постпрандиална глюкоза 2 часа след хранене 140 – 199 mg/dL, диабет над 199 mg/dL.
- Оптимален гликиран хемоглобин A1C по-малко от 5%, граничен 5,0 – 5,6%, преддиабет 5,7 – 6,4%, диабет над 6,4%.

World Health Organization, “Definition and Diagnosis of Diabetes Mellitus and Intermediate Hyperglycaemia: Report of a WHO/IDF Consultation” (2006):1–50; N. Bansal, “Prediabetes Diagnosis and Treatment: A Review,” *World Journal of Diabetes* 6, no. 2 (2015): 296; W. C. Y. Yip et al., “Prevalence of Pre-Diabetes Across Ethnicities: A Review of Impaired Fasting Glucose (IFG) and Impaired Glucose Tolerance (IGT) for Classification of Dysglycaemia,” *Nutrients* 9, no. 11 (2017): 1273; Z. Punthakee et al., “Classification and Diagnosis of Diabetes, Prediabetes, and Metabolic Syndrome,” *Canadian Journal of Diabetes* 42 (2018): S10– 15; American Diabetes Association, “2. Classification and Diagnosis of Diabetes: Standards of Medical Care in Diabetes — 2018,” *Diabetes Care* 41, no. 1 (2018): S13–27.

9. A. Hozawa et al., “Association Between Body Mass Index and All-Cause Death in Japanese Population: Pooled Individual Participant Data Analysis of 13 Cohort Studies,” *Journal of Epidemiology* 29, no. 12 (2019): 457–63; M. D. Rahman et al., “Trend, Projection, and Appropriate Body Mass Index Cut-Off Point for Diabetes and Hypertension in Bangladesh,” *Diabetes Research and Clinical Practice* 126 (2017): 43–53.
10. Този онлайн калкулатор се предлага от Център за контрол на заболяванията: [https://www.cdc.gov/healthyweight/assessing/bmi/adult\\_bmi/english\\_bmi\\_calculator/bmi\\_calculator.html](https://www.cdc.gov/healthyweight/assessing/bmi/adult_bmi/english_bmi_calculator/bmi_calculator.html). Достъпно на 20 май 2020 година.
11. Когато обучавам пациент относно Протокола на Готфрид, питам дали иска основния, или усъвършенствания подход. За хора, които имат достъп до съответните устройства, може да бъде полезно измерването на по-разширени показатели, включително следните:
  - Ежедневно измерване на кръвна захар и кетони (чрез изследване с убождане на пръста), за да се изчисли глюкозо-кетонния индекс.
  - Непрекъснат мониторинг на глюкозата.
  - Bluetooth кантар за състава на тялото.
  - Метаболитна скорост в покой.
  - Изпълнение на упражнения за оценка на аеробната издръжливост като VO2 max – показва максималната скорост на консумация на кислород, който тялото ни е способно да абсорбира, транспортира и консумира.
12. Едно от най-добрите описания на кетогенното съотношение е налично в тази книга: Jacob Wilson and Ryan Lowery, *The Ketogenic Bible* (Las Vegas: Victory Belt, 2017), 39–40.

13. Протеините са смесени в техния кетогенен ефект. Защо? Някои градивни елементи на протеините, наречени аминокиселини, са кетогенни, докато други са антикетогенни. Примери за кетогенни аминокиселини са левцин и лизин. Пример за антикетогенна аминокиселина е аланинът.
14. I. A. Cohen, "A Model for Determining Total Ketogenic Ratio (TKR) for Evaluating the Ketogenic Property of a Weight-Reduction Diet," *Medical Hypotheses* 73, no. 3 (2009): 377–81.
15. Young et al., "Effect on Body Composition."
16. S. H. Duncan et al., "Reduced Dietary Intake of Carbohydrates by Obese Subjects Results in Decreased Concentrations of Butyrate and Butyrate-Producing Bacteria in Feces," *Applied and Environmental Microbiology* 73, no. 4 (2007): 1073–78.
17. За да изчислите нетните въглехидрати в преработените храни, извадете фибрите и количеството на захарните алкохоли. Например за една трета от средно авокадо това значи 4 грама въглехидрати, по-малко от 3 грама фибри (4 грама – 3 грама = 1 грам), тоест 1 грам нетни въглехидрати. За любимия ми сладкиш това са общо 13 грама въглехидрати на всяка порция брауни, по-малко от 5 грама фибри, по-малко от 7 грама захарни алкохоли (в този случай еритритол и алулоза), тоест 1 грам нетни въглехидрати.
18. J. Rehm et al., "Alcohol Use and Cancer in the European Union," *European Addiction Research* (2020): 1–8; S. Parida et al., "Microbial Alterations and Risk Factors of Breast Cancer: Connections and Mechanistic Insights," *Cells* 9, no. 5 (2020): 1091.
19. X. Yao et al., "Change in Moderate Alcohol Consumption and Quality of Life: Evidence from 2 Population-Based Cohorts," *CMAJ* 191, no. 27 (2019): E753–60.
20. M. Venkatesh et al., "Dietary Oil Composition Differentially Modulates Intestinal Endotoxin Transport and Postprandial Endotoxemia," *Nutrition & Metabolism* 10, no. 1 (2013): 6.
21. A. Dagfinn et al., "Nut Consumption and Risk of Cardiovascular Disease, Total Cancer, All-Cause and Cause-Specific Mortality: A Systematic Review and Dose-Response Meta-Analysis of Prospective Studies," *BMC Medicine* 14, no. 1 (2016): 207; C. Guo-Chong et al., "Nut Consumption in Relation to All-Cause and Cause-Specific Mortality: A Meta-Analysis of 18 Prospective Studies," *Food & Function* 8, no. 11 (2017): 3893–905.
22. M. P. St-Onge et al., "Consumption of a Functional Oil Rich in Phytosterols and Medium-Chain Triglyceride Oil Improves Plasma Lipid Profiles in Men," *The Journal of Nutrition* 133, no. 6, (2003): 1815–20; J. R. Han et al., "Effects of Dietary Medium-Chain Triglyceride on Weight Loss and Insulin Sensitivity in a Group of Moderately Overweight Free-Living Type 2 Diabetic Chinese Subjects," *Metabolism* 56, no. 7 (2007): 985–91; M. P. St-Onge et al., "Medium-

- Chain Triglyceride Oil Consumption as Part of a Weight Loss Diet Does Not Lead to an Adverse Metabolic Profile When Compared to Olive Oil," *Journal of the American College of Nutrition* 27, no. 5 (2008): 547–52.
23. K. Mumme et al., "Effects of Medium-Chain Triglycerides on Weight Loss and Body Composition: A Meta-Analysis of Randomized Controlled Trials," *Journal of the Academy of Nutrition and Dietetics* 115, no. 2 (2015): 249–63.
  24. T. Maher et al., "A Comparison of the Satiating Properties of Medium-Chain Triglycerides and Conjugated Linoleic Acid in Participants with Healthy Weight and Overweight or Obesity," *European Journal of Nutrition* (2020): 1–13.
  25. Ето допълнителни подробности за МСТ:  
 МСТ маслото увеличава кетоните с 19% и може да помогне за превключване на метаболитния превключвател от изгаряне на глюкоза към изгаряне на мазнини.(C. Vandenberghe et al., "Medium-Chain Triglycerides Modulate the Ketogenic Effect of a Metabolic Switch," *Frontiers in Nutrition* 7 (2020): 3–6.  
 Освен това в ограничени проучвания на болестта на Алцхаймер някои показатели за когнитивните способности се подобряват с добавяне на МСТ масло към диетата (K. I. Avgerinos et al., "Medium-Chain Triglycerides Induce Mild Ketosis and May Improve Cognition in Alzheimer's Disease: A Systematic Review and Meta-Analysis of Human Studies," *Ageing Research Reviews* [2019]: 101,001), а използването на кетоните от мозъка може да се удвои. (E. Croteau et al., "Ketogenic Medium-Chain Triglycerides Increase Brain Energy Metabolism in Alzheimer's Disease," *Journal of Alzheimer's Disease* 64, no. 2 [2018]: 551–61).  
 МСТ маслото също „говори“ с хормоните и може да ви помогне да станете по-чувствителни към инсулина и да намалите адипонектина си в рамките на шест седмици, поне според едно малко неконтролирано проучване (D. D. Thomas et al., "Effects of Medium-Chain Triglycerides Supplementation on Insulin Sensitivity and Beta Cell Function: A Feasibility Study," *PLoS One* 14, no. 12 (2019)).
  26. Смята се, че механизмът на увреждане е чрез повишаване на бактериалните токсини от червата, известно като ендотоксемия, пет до осем часа, след като го изядете. P. Dandona et al., "Macronutrient Intake Induces Oxidative and Inflammatory Stress: Potential Relevance to Atherosclerosis and Insulin Resistance," *Experimental & Molecular Medicine* 42, no. 4 (2010): 245–53; F. Biobaku et al., "Macronutrient-Mediated Inflammation and Oxidative Stress: Relevance to Insulin Resistance, Obesity, and Atherogenesis," *The Journal of Clinical Endocrinology & Metabolism* 104, no. 12 (2019): 6118–28.
  27. Когато консумирате наситени или трансмазнини заедно с рафинирани въглехидрати, механизмът на увреждане е комбинация от оксидативен стрес, неразрешено възпаление, ендотоксемия, повишена експресия на SOCS-3 и TLR4, което блокира IRS-1 и PI3K пътищата, предизвиквайки

- инсулинова резистентност (според лична комуникация с д-р Марк Хюстеън и Dandona et al., "Macronutrient Intake").
28. Y. Bao et al., "Association of Nut Consumption with Total and Cause-Specific Mortality," *The New England Journal of Medicine* 369, no. 21 (2013): 2001–11.
  29. C. Smith-Spangler et al., "Are Organic Foods Safer or Healthier Than Conventional Alternatives? A Systematic Review," *Annals of Internal Medicine* 157, no. 5 (2012): 343–66; S. Watson, "Organic Food No More Nutritious Than Conventionally Grown Food," *Harvard Women's Health Watch*, September 5, 2012, <https://www.health.harvard.edu/blog/organic-food-no-more-nutritious-than-conventionally-grown-food-201209055264>. Достъпно на 6.05.2020 година.
  30. A. S. Abargouei et al., "Effect of Dairy Consumption on Weight and Body Composition in Adults: A Systematic Review and Meta-Analysis of Randomized Controlled Clinical Trials," *International Journal of Obesity* 36, no. 12 (2012): 1485–93.
  31. California Avocados, "Avocado Nutritional Information," <https://www.californiaavocado.com/nutrition/nutrients>. Достъпно на 6.05.2020 година.
  32. S. Kim et al., "Effects of Growth Hormone on Glucose Metabolism and Insulin Resistance in Humans," *Annals of Pediatric Endocrinology & Metabolism* 22, no. 3 (2017): 145.
  33. L. A. Frohman, "Growth Hormone," *Encyclopedia of Neuroscience*, vol. 1 (London: Academic Press, 2009).
  34. Diabetes Teaching Center at the University of California, San Francisco, "Blood Sugar & Other Hormones," <https://dtc.ucsf.edu/types-of-diabetes/type1/understanding-type-1-diabetes/how-the-body-processes-sugar/blood-sugar-other-hormones/>
  35. R. Lanzi et al., "Elevated Insulin Levels Contribute to the Reduced Growth Hormone (GH) Response to GH-Releasing Hormone in Obese Subjects," *Metabolism* 48, no. 9 (1999): 1152–56; J. Xu et al., "Crosstalk Between Growth Hormone and Insulin Signaling," *Vitamins & Hormones* 80 (2009): 125–53; H. Qiu et al., "Influence of Insulin on Growth Hormone Secretion, Level, and Growth Hormone Signaling," *Sheng Li Xue Bao* 69, no. 5 (2017): 541–56.
  36. M. La Merrill et al., "Toxicological Function of Adipose Tissue: Focus on Persistent Organic Pollutants," *Environmental Health Perspectives* 121, no. 2 (2013): 162–69.
  37. Вашият черен дроб обработва токсините на две фази. В първата фаза той превръща мастноразтворимите токсини във водоразтворими вещества. В края на втората фаза черният дроб отделя тези водоразтворими токсини чрез урината, изпражненията, потта и други телесни течности. В протокола за детоксикация този двуетапен процес трябва да се управлява в обратен ред: оптимизиране на фаза II, преди да се

насочат усилията към фаза I. Това е една от причините програмите за детоксикация и „прочистване“ да са противоречиви и да е възможно да ви накарат да се чувствате зле. Ако токсините се отстраняват от тъканите с по-висока скорост, отколкото се изхвърлят от тялото, това кара човек да се чувства ужасно и дори може да има тежки последици.

38. J. Obert et al., “Popular Weight Loss Strategies: A Review of Four Weight Loss Techniques,” *Current Gastroenterology Reports* 19, no. 12 (2017): 61.
39. M. S. Duchowny, “Food for Thought: The Ketogenic Diet and Adverse Effects in Children,” *Epilepsy Currents* 5, no. 4 (2005): 152–54.
40. G. Zong et al., “Consumption of Meals Prepared at Home and Risk of Type 2 Diabetes: An Analysis of Two Prospective Cohort Studies,” *PLoS Medicine* 13, no. 7 (2016): e1002052.

## **6. ДЕТОКСИКАЦИЯ, СЪОБРАЗЕН С ЦИРКАДНИТЕ РИТМИ ФАСТИНГ И СПРАВЯНЕ С ЕВЕНТУАЛНИ ПРОБЛЕМИ**

1. M. M. Hetherington et al., “Understanding the Science of Portion Control and the Art of Downsizing,” *Proceedings of the Nutrition Society* 77, no. 3 (2018): 347–55.
2. J. J. Meidenbauer et al., “The Glucose Ketone Index Calculator: A Simple Tool to Monitor Therapeutic Efficacy for Metabolic Management of Brain Cancer,” *Nutrition & Metabolism* 12, no. 1 (2015): 1–7.
3. Y. Li, “Exogenous Stimuli Maintain Intraepithelial Lymphocytes Via Aryl Hydrocarbon Receptor Activation,” *Cell* 147, no. 3 (2011): 629–40.
4. A. Paoli et al., “Effect of Ketogenic Mediterranean Diet with Phytoextracts and Low Carbohydrates/High-Protein Meals on Weight, Cardiovascular Risk Factors, Body Composition, and Diet Compliance in Italian Council Employees,” *Nutrition Journal* 10, no. 1 (2011): 112; A. Paoli et al., “Long Term Successful Weight Loss with a Combination Biphasic Ketogenic Mediterranean Diet and Mediterranean Diet Maintenance Protocol,” *Nutrients* 5, no. 12 (2013): 5205–17; A. Paoli et al., “Ketogenic Diet and Phytoextracts,” *Scientific Advisory Board* 21, no. 4 (2010): 24–29; A. Paoli et al., “Ketogenic Diet Does Not Affect Strength Performance in Elite Artistic Gymnasts,” *Journal of the International Society of Sports Nutrition* 9, no. 1 (2012): 34; A. Paoli et al., “Effects of n-3 Polyunsaturated Fatty Acids ( $\omega$ -3) Supplementation on Some Cardiovascular Risk Factors with a Ketogenic Mediterranean Diet,” *Marine Drugs* 13, no. 2 (2015): 996–1009; G. Bosco et al., “Effects of the Ketogenic Diet in Overweight Divers Breathing Enriched Air Nitrox,” *Scientific Reports* 8, no. 1 (2018): 1–8; A. Paoli et al., “Effects of a Ketogenic Diet in Overweight Women with Polycystic Ovary Syndrome,” *Journal of Translational Medicine* 18, no. 1 (2020): 1–11.

5. Y. Aitbali et al., "Glyphosate Based-Herbicide Exposure Affects Gut Microbiota, Anxiety, and Depression-Like Behaviors in Mice," *Neurotoxicology and Teratology* (2018); I. Argou-Cardozo et al., "Clostridium Bacteria and Autism Spectrum Conditions: A Systematic Review and Hypothetical Contribution of Environmental Glyphosate Levels," *Medical Sciences* 6, no. 2 (2018): 29; C. E. Gallegos et al., "Perinatal Glyphosate-Based Herbicide Exposure in Rats Alters Brain Antioxidant Status, Glutamate and Acetylcholine Metabolism, and Affects Recognition Memory," *Neurotoxicity Research* (2018): 1–12; P. Good. "Evidence the US Autism Epidemic Initiated by Acetaminophen (Tylenol) Is Aggravated by Oral Antibiotic Amoxicillin/Clavulanate (Augmentin) and Now Exponentially by Herbicide Glyphosate (Roundup)," *Clinical Nutrition ESPEN* 23 (2018): 171–83; L. N. Nielsen et al., "Glyphosate Has Limited Short-Term Effects on Commensal Bacterial Community Composition in the Gut Environment Due to Sufficient Aromatic Amino Acid Levels," *Environmental Pollution* 233 (2018): 364–76.
6. J. J. Gildea et al., "Protection Against Gluten-Mediated Tight Junction Injury with a Novel Lignite Extract Supplement," *Journal of Nutrition & Food Sciences* 6, no. 547 (2016): 2; J. J. Gildea et al., "Protective Effects of Lignite Extract Supplement on Intestinal Barrier Function in Glyphosate-Mediated Tight Junction Injury," *Journal of Clinical Nutrition & Dietetics* 3, no. 1 (2017).
7. A. Di Ciaula et al., "Diet and Contaminants: Driving the Rise to Obesity Epidemics?" *Current Medicinal Chemistry* 26, no. 19 (2019): 3471–82; L. A. Hoepner, "Bisphenol A: A Narrative Review of Prenatal Exposure Effects on Adipogenesis and Childhood Obesity Via Peroxisome Proliferator-Activated Receptor Gamma," *Environmental Research* 173 (2019): 54–68; Rubin et al., "The Case for BPA as an Obesogen"; R. Chamorro-Garcia et al., "Current Research Approaches and Challenges in the Obesogen Field," *Frontiers in Endocrinology* 10 (2019): 167; J. J. Heindel, "History of the Obesogen Field: Looking Back to Look Forward," *Frontiers in Endocrinology* 10 (2019): 14.
8. K. Katoh et al., "Suppressing Effects of Bisphenol A on the Secretory Function of Ovine Anterior Pituitary Cells," *Cell Biology International* 28, no. 6 (2004): 463–69.
9. A. B. Javurek et al., "Effects of Exposure to Bisphenol A and Ethinyl Estradiol on the Gut Microbiota of Parents and Their Offspring in a Rodent Model," *Gut Microbes* 7, no. 6 (2016): 471–85; J. Xu et al., "Developmental Bisphenol A Exposure Modulates Immune-Related Diseases," *Toxics* 4, no. 4 (2016): 23; K. P. Lai et al., "Bisphenol A Alters Gut Microbiome: Comparative Metagenomics Analysis," *Environmental Pollution* 218 (2016): 923–30; L. Reddivari et al., "Perinatal Bisphenol A Exposure Induces Chronic Inflammation in Rabbit Offspring via Modulation of Gut Bacteria and Their Metabolites," *MSystems* 2, no. 5 (2017); Y. Malaisé et al., "Gut Dysbiosis and Impairment of Immune System Homeostasis in Perinatally Exposed Mice to Bisphenol A Precede Obese Phenotype Development," *Scientific Reports* 7, no. 1 (2017): 1–12;



- J. A. DeLuca et al., "Bisphenol-A Alters Microbiota Metabolites Derived from Aromatic Amino Acids and Worsens Disease Activity During Colitis," *Experimental Biology and Medicine* 243, no. 10 (2018): 864–75; T. R. Catron et al., "Host Developmental Toxicity of BPA and BPA Alternatives Is Inversely Related to Microbiota Disruption in Zebrafish," *Toxicological Sciences* 167, no. 2 (2019): 468–83.
10. K. Oishi, "Effect of Probiotics, Bifidobacterium Breve, and Lactobacillus Casei on Bisphenol A Exposure in Rats," *Bioscience, Biotechnology, and Biochemistry* 72, no. 6 (2008): 1409–15; S. Song et al., "The Anti-Allergic Activity of Lactobacillus Plantarum L67 and Its Application to Yogurt," *Journal of Dairy Science* 99, no. 12 (2016): 9372–82.
  11. A. A. Ismail et al., "Chronic Magnesium Deficiency and Human Disease: Time for Reappraisal?" *QJM: An International Journal of Medicine* 111, no. 11 (2018): 759–63; M. S. Razzaque, "Magnesium: Are We Consuming Enough?" *Nutrients* 10, no. 12 (2018): 1863; J. L. Workinger et al., "Challenges in the Diagnosis of Magnesium Status," *Nutrients* 10, no. 9 (2018): 1202.
  12. Специалисти по функционална медицина може да намерите на следния адрес: <https://www.ifm.org/find-a-practitioner/>. Достъпно на 16 декември 2020 година.
  13. J. Hussain et al., "Clinical Effects of Regular Dry Sauna Bathing: A Systematic Review," *Evidence-Based Complementary and Alternative Medicine* (2018).
  14. C. P. Oliveira et al., "N-Acetylcysteine and/or Ursodeoxycholic Acid Associated with Metformin in Non-Alcoholic Steatohepatitis: An Open-Label Multicenter Randomized Controlled Trial," *Arquivos de Gastroenterologia* 56, no. 2 (2019): 184–90; D. Thakker et al., "N-Acetylcysteine for Polycystic Ovary Syndrome: A Systematic Review and Meta-Analysis of Randomized Controlled Clinical Trials," *Obstetrics and Gynecology International* (2015).
  15. A. M. Fulghesu et al., "N-Acetyl-Cysteine Treatment Improves Insulin Sensitivity in Women with Polycystic Ovary Syndrome," *Fertility and Sterility* 77, no. 6 (2002): 1128–35; G. Oner et al., "Clinical, Endocrine, and Metabolic Effects of Metformin vs. N-acetyl-cysteine in Women with Polycystic Ovary Syndrome," *European Journal of Obstetrics & Gynecology and Reproductive Biology* 159, no. 1 (2011): 127–31.
  16. A. Elnashar et al., "N-Acetyl Cysteine vs. Metformin in Treatment of Clomiphene Citrate-Resistant Polycystic Ovary Syndrome: A Prospective Randomized Controlled Study," *Fertility and Sterility* 88, no. 2 (2007): 406–9.
  17. S. Ebrahimpour-Koujan et al., "Lower Glycemic Indices and Lipid Profile Among Type 2 Diabetes Mellitus Patients Who Received Novel Dose of Silybum Marianum (L.) Gaertn.(silymarin) Extract Supplement: A Triple-Blinded Randomized Controlled Clinical Trial," *Phytomedicine* 44 (2018): 39–44.

18. S. Rahmani et al., "Treatment of Non-Alcoholic Fatty Liver Disease with Curcumin: A Randomized Placebo-Controlled Trial," *Phytotherapy Research* 30, no. 9 (2016): 1540–48; Y. Panahi et al., "Efficacy and Safety of Phytosomal Curcumin in Non-Alcoholic Fatty Liver Disease: A Randomized Controlled Trial," *Drug Research* 67, no. 04 (2017): 244–51; R. Goodarzi et al., "Does Turmeric/Curcumin Supplementation Improve Serum Alanine Aminotransferase and Aspartate Aminotransferase Levels in Patients with Nonalcoholic Fatty Liver Disease? A Systematic Review and Meta-Analysis of Randomized Controlled Trials," *Phytotherapy Research* 33, no. 3 (2019): 561–70; F. Mansour-Ghanaei et al., "Efficacy of Curcumin/Turmeric on Liver Enzymes in Patients with Non-Alcoholic Fatty Liver Disease: A Systematic Review of Randomized Controlled Trials," *Integrative Medicine Research* 8, no. 1 (2019): 57–61; A. Ghaffari et al., "Turmeric and Chicory Seed Have Beneficial Effects on Obesity Markers and Lipid Profile in Non-Alcoholic Fatty Liver Disease (NAFLD)," *International Journal for Vitamin and Nutrition Research* (2019).
19. K. Gabel et al., "Effects of 8-hour Time Restricted Feeding on Body Weight and Metabolic Disease Risk Factors in Obese Adults: A Pilot Study," *Nutrition and Healthy Aging* 4, no. 4 (2018): 345–53, <https://content.iospress.com/articles/nutrition-and-healthy-aging/nha170036>.
20. M. N. Harvie, "The Effects of Intermittent or Continuous Energy Restriction on Weight Loss and Metabolic Disease Risk Markers: A Randomized Trial in Young Overweight Women," *International Journal of Obesity (London)* 35 (2011): 714–27; S. Gil et al., "A Smartphone App Reveals Diurnal Eating Patterns in Humans That Can Be Modulated for Health Benefits," *Cell Metabolism* 22, no. 5 (2015): 789–98; G. M. Tinsley et al., "Effects of Intermittent Fasting on Body Composition."
21. A. Chaix et al., "The Effects of Time-Restricted Feeding on Lipid Metabolism and Adiposity," *Adipocyte* 4, no. 4 (2015): 319–24; H. Chung et al., "Time-Restricted Feeding Improves Insulin Resistance and Hepatic Steatosis in a Mouse Model of Postmenopausal Obesity," *Metabolism-Clinical and Experimental* 65, no. 12 (2016): 1743–54.
22. A. Chaix et al., "Time-Restricted Feeding Is a Preventative and Therapeutic Intervention Against Diverse Nutritional Challenges," *Cell Metabolism* 20, no. 6 (2014): 991–1005; R. Antoni et al., "Effects of Intermittent Fasting on Glucose and Lipid Metabolism," *Proceedings of the Nutrition Society* 76, no. 3 (2017): 361–68.
23. G. C. Melkani et al., "Time Restricted Feeding for Prevention and Treatment of Cardiometabolic Disorders," *The Journal of Physiology* 595, no. 12 (2017): 3691–700.
24. R. E. Patterson et al., "Intermittent Fasting and Human Metabolic Health," *Journal of the Academy of Nutrition and Dietetics* 115, no. 8 (2015): 1203–12; C.

- R. Marinac et al., "Prolonged Nightly Fasting and Breast Cancer Prognosis," *JAMA Oncology* 2, no. 8 (2016): 1049–55; L. A. Smith et al., "Translating Mechanism-Based Strategies to Break the Obesity–Cancer Link: A Narrative Review," *Journal of the Academy of Nutrition and Dietetics* 118, no. 4 (2018): 652–67.
25. E. N. Manoogian et al., "Circadian Rhythms, Time-Restricted Feeding, and Healthy Aging," *Ageing Research Reviews* 39 (2017): 59–67.
  26. J. T. Haas et al., "Fasting the Microbiota to Improve Metabolism?" *Cell Metabolism* 26, no. 4 (2017): 584–85; R. Kivelä et al., "White Adipose Tissue Coloring by Intermittent Fasting," *Cell Research* 27, no. 11 (2017): 1300–1301; G. Li et al., "Intermittent Fasting Promotes White Adipose Browning and Decreases Obesity by Shaping the Gut Microbiota," *Cell Metabolism* 26, no. 4 (2017): 672–85.
  27. S. Eslami et al., "Annual Fasting; The Early Calories Restriction for Cancer Prevention," *BioImpacts: BI* 2, no. 4 (2012): 213–15; A. Zarrinpar et al., "Diet and Feeding Pattern Affect the Diurnal Dynamics of the Gut Microbiome," *Cell Metabolism* 20, no. 6 (2014): 1006–17; A. Chaix et al. "The Effects of Time-Restricted Feeding"; J. L. Kaczmarek et al., "Complex Interactions of Circadian Rhythms, Eating Behaviors, and the Gastrointestinal Microbiota and Their Potential Impact on Health," *Nutrition Reviews* 75, no. 9 (2017): 673–82; Li et al., "Intermittent Fasting Promotes White"; R. E. Patterson et al., "Metabolic Effects of Intermittent Fasting," *Annual Review of Nutrition* 37 (2017): 371–93; E. Beli et al., "Restructuring of the Gut Microbiome by Intermittent Fasting Prevents Retinopathy and Prolongs Survival in db/db Mice," *Diabetes* (2018): db180158.
  28. S. Panda, "Circadian Physiology of Metabolism." *Science* 354, no. 6315 (2016): 1008–15.
  29. M. P. Mattson et al., "Impact of Intermittent Fasting on Health and Disease Processes," *Ageing Research Reviews* 39 (2017): 46–58; B. K. Shin et al., "Intermittent Fasting Protects Against the Deterioration of Cognitive Function, Energy Metabolism, and Dyslipidemia in Alzheimer's Disease-Induced Estrogen Deficient Rats," *Experimental Biology and Medicine* 234, no. 4 (2018): 334–43.
  30. M. Hatori et al., "Time-Restricted Feeding Without Reducing Caloric Intake Prevents Metabolic Diseases in Mice Fed a High-Fat Diet," *Cell Metabolism* 15, no. 6 (2012): 848–60.
  31. K. M. Pursey et al., "Neural Responses to Visual Food Cues According to Weight Status: A Systematic Review of Functional Magnetic Resonance Imaging Studies," *Frontiers in Nutrition* 1 (2017): 7–18.
  32. Научих този трик от д-р Джефри Бекер на лекция, която той изнесе в Института по интегративна психиатрия, където съм преподавател. Той препоръчва по 1 чаена лъжичка МСТ масло на работа или половин високопротеиново барче следобед, за да предотвратите пристъпите на непреодолимо желание за алкохол.

33. M. C. Houston, "Treatment of Hypertension with Nutraceuticals, Vitamins, Antioxidants, and Minerals," *Expert Review of Cardiovascular Therapy* 5, no. 4 (2007): 681–91; S. T. Sinatra et al., *Nutritional and Integrative Strategies in Cardiovascular Medicine* (Boca Raton, FL: CRC Press, 2015); L. Rochette et al., "Alpha-Lipoic Acid: Molecular Mechanisms and Therapeutic Potential in Diabetes," *Canadian Journal of Physiology and Pharmacology* 93, no. 12 (2015): 1021–27; S. Kucukgoncu et al., "Alpha-Lipoic Acid (ALA) as a Supplementation for Weight Loss: Results from a Meta-Analysis of Randomized Controlled Trials," *Obesity Reviews* 18, no. 5 (2017): 594–601.
34. Rochette et al., "Alpha-Lipoic Acid."
35. K. H. Weylandt et al., "Omega-3 Fatty Acids and Their Lipid Mediators: Towards an Understanding of Resolvin and Protectin Formation," *Prostaglandins & Other Lipid Mediators* 97, nos. 3–4 (2012): 73–82; R. Ramaswami et al., "Fish Oil Supplementation in Pregnancy," *New England Journal of Medicine* 375, no. 26 (2016): 2599–601.
36. Yang et al., "Ratio of N-3/N-6 PUFAs"; C. J. Fabian et al., "Omega-3 Fatty Acids for Breast Cancer Prevention and Survivorship," *Breast Cancer Research* 17, no. 1 (2015): 1–11.
37. M. Houston, *Personalized and Precision Integrative Cardiovascular Medicine* (Philadelphia: Lippincott Williams & Wilkins, 2019); Z. Ilyas et al., "The Effect of Berberine on Weight Loss in Order to Prevent Obesity: A Systematic Review," *Biomedicine & Pharmacotherapy* 127 (2020): 110137; M. Rondanelli et al., "Polycystic Ovary Syndrome Management: A Review of the Possible Amazing Role of Berberine," *Archives of Gynecology and Obstetrics* (2020): 1–8.
38. C. N. Serhan, "Pro-Resolving Lipid Mediators Are Leads for Resolution Physiology," *Nature* 510, no. 7503 (2014): 92–101; C. N. Serhan et al., "Resolvins in Inflammation: Emergence of the Pro-Resolving Superfamily of Mediators," *The Journal of Clinical Investigation* 128, no. 7 (2018): 2657–69; P. C. Norris et al., "Identification of Specialized Pro-Resolving Mediator Clusters from Healthy Adults After Intravenous Low-Dose Endotoxin and Omega-3 Supplementation: A Methodological Validation," *Scientific Reports* 8, no. 1 (2018): 1–13.
39. C. C. Douglas et al., "Role of Diet in the Treatment of Polycystic Ovary Syndrome," *Fertility and Sterility* 85, no. 3 (2006): 679–88; Gottfried, *The Hormone Cure*, M. McGrice et al., "The Effect of Low Carbohydrate Diets on Fertility Hormones and Outcomes in Overweight and Obese Women: A Systematic Review," *Nutrients* 9, no. 3 (2017): 204; L. Barrea et al., "Source and Amount of Carbohydrate in the Diet and Inflammation in Women with Polycystic Ovary Syndrome," *Nutrition Research Reviews* 31, no. 2 (2018): 291–301; L. Barrea et al., "Adherence to the Mediterranean Diet, Dietary Patterns, and Body Composition in Women with Polycystic Ovary Syndrome (PCOS)," *Nutrients* 11, no. 10 (2019): 2278.

40. Mavropoulos et al., "The Effects of a Low-Carbohydrate"; Gottfried, *The Hormone Cure*; D. Kulak et al., "Should the Ketogenic Diet Be Considered for Enhancing Fertility?" *Maturitas* 74, no. 1 (2013): 10–13; Muscogiuri et al., "Current Insights into Inositol Isoforms"; M. Melanie et al., "The Effect of Low Carbohydrate Diets on Fertility Hormones and Outcomes in Overweight and Obese Women: A Systematic Review," *Nutrients* 9, no. 3 (2017): 204; M. Caprio et al., "Very-Low-Calorie Ketogenic Diet (VLCKD) in the Management of Metabolic Diseases: Systematic Review and Consensus Statement from the Italian Society of Endocrinology (SIE)," *Journal of Endocrinological Investigation* 42, no. 11 (2019): 1365–86; Paoli et al., "Effects of a Ketogenic Diet in Overweight Women."
41. Gottfried, *The Hormone Cure*; M. J. Carvalho et al., "Controversial Association Between Polycystic Ovary Syndrome and Breast Cancer," *European Journal of Obstetrics & Gynecology and Reproductive Biology* 243 (2019): 125–32.
42. A. Balen, "Polycystic Ovary Syndrome and Cancer," *Human Reproduction Update* 7, no. 6 (2001): 522–25; Gottfried, *The Hormone Cure*; C. C. Shen et al., "A Nationwide Population-Based Retrospective Cohort Study of the Risk of Uterine, Ovarian, and Breast Cancer in Women with Polycystic Ovary Syndrome," *The Oncologist* 20, no. 1 (2015): 45; F. Shobeiri et al., "The Association Between Polycystic Ovary Syndrome and Breast Cancer: A Meta-Analysis," *Obstetrics & Gynecology Science* 59, no. 5 (2016): 367–72.
43. Gottfried, *The Hormone Cure*; J. Barry et al., "Risk of Endometrial, Ovarian, and Breast Cancer in Women with Polycystic Ovary Syndrome: A Systematic Review and Meta-Analysis," *Human Reproduction Update* 20, no. 5 (2014): 748–58; M. Gottschau et al., "Risk of Cancer Among Women with Polycystic Ovary Syndrome: A Danish Cohort Study." *Gynecologic Oncology* 136, no. 1 (2015): 99–103; H. R. Harris et al., "Polycystic Ovary Syndrome and Risk of Endometrial, Ovarian, and Breast Cancer: A Systematic Review," *Fertility Research and Practice* 2, no. 1 (2016): 14; D. C. Ding et al., "Association Between Polycystic Ovarian Syndrome and Endometrial, Ovarian, and Breast Cancer: A Population-Based Cohort Study in Taiwan," *Medicine* 97, no. 39 (2018).
44. Gottfried, *The Hormone Cure*; Barry et al., "Risk of Endometrial, Ovarian, and Breast Cancer"; Shen et al., "A Nationwide Population-Based Retrospective"; Gottschau et al., "Risk of Cancer Among Women"; Harris et al., "Polycystic Ovary Syndrome"; Ding et al., "Association Between Polycystic Ovarian Syndrome."

## 7. ПИРЕХОД

1. R. R. Wing et al., "Weight Gain at the Time of Menopause," *Archives of Internal Medicine* 151, no. 1 (1991): 97–102; G. M. Van Dijk et al., "The Association Between Vasomotor Symptoms and Metabolic Health in Peri- and Postmenopausal Women: A Systematic Review," *Maturitas* 80, no. 2

- (2015): 140–47; P. Tuomikoski et al., “Vasomotor Symptoms and Metabolic Syndrome,” *Maturitas* 97 (2017): 61–65; S. Sayan et al., “Relationship Between Vasomotor Symptoms and Metabolic Syndrome in Postmenopausal Women,” *Journal of International Medical Research* 46, no. 10 (2018): 4157–66.
2. V. D. Longo et al., “Fasting, Circadian Rhythms, and Time-Restricted Feeding in Healthy Lifespan,” *Cell Metabolism* 23, no. 6 (2016): 1048–59.
  3. F. Coucke, “Food Intolerance in Patients with Manifest Autoimmunity: Observational Study,” *Autoimmunity Reviews* 17, no. 11 (2018): 1078–80.
  4. T. C. Wallace et al., “Dairy Intake Is Not Associated with Improvements in Bone Mineral Density or Risk of Fractures Across the Menopause Transition: Data from the Study of Women’s Health Across the Nation,” *Menopause* 27, no. 8 (2020): 879–86.
  5. A. Trichopoulou et al., “Healthy Traditional Mediterranean Diet: An Expression of Culture, History, and Lifestyle.” *Nutrition Reviews* 55, no. 11 (1997): 383–89; S. Dernini, “The Erosion and the Renaissance of the Mediterranean Diet: A sustainable Cultural Resource,” *Quaderns de la Mediterrania* 16 (2011): 75–82; T. I. González, “The Mediterranean Diet: Consumption, Cuisine, and Food Habits,” in *MediTERRA 2012: The Mediterranean Diet for Sustainable Regional Development*, ed. F. Mombiola (Paris: CIHEAM Sciences/Presses de Sciences Po, 2012), 115–32; N. R. Sahyoun et al., *Historical Origins of the Mediterranean Diet, Regional Dietary Profiles, and the Development of the Dietary Guidelines* (Totowa, NJ: Humana Press, 2016), 43–56; C. M. Lăcătușu et al., “The Mediterranean Diet: From an Environment-Driven Food Culture to an Emerging Medical Prescription,” *International Journal of Environmental Research and Public Health* 16, no. 6 (2019): 942.
  6. M. De Lorgeril et al., “Mediterranean Alpha-Linolenic Acid-Rich Diet in Secondary Prevention of Coronary Heart Disease,” *The Lancet* 343, no. 8911 (1994): 1454–59; M. De Lorgeril et al., “Mediterranean Diet, Traditional Risk Factors, and the Rate of Cardiovascular Complications After Myocardial Infarction: Final Report of the Lyon Diet Heart Study,” *Circulation* 99, no. 6 (1999): 779–85; R. Estruch et al., “Retraction and Republication: Primary Prevention of Cardiovascular Disease with a Mediterranean Diet,” *New England Journal of Medicine* 368 (2013): 1279–90; M. Sotos-Prieto et al., “Assessing Validity of Self-Reported Dietary Intake Within a Mediterranean Diet Cluster Randomized Controlled Trial among US Firefighters,” *Nutrients* 11, no. 9 (2019): 2250.
  7. J. Salas-Salvado et al., “Effect of a Mediterranean Diet Supplemented with Nuts on Metabolic Syndrome Status: One-Year Results of the PREDIMED Randomized Trial,” *Archives of Internal Medicine* 168, no. 22 (2008): 2449–58; M. T. Mitjavila et al., “The Mediterranean Diet Improves the Systemic Lipid and DNA Oxidative Damage in Metabolic Syndrome Individuals: A Randomized, Controlled Trial,” *Clinical Nutrition* 32, no. 2 (2013): 172–78;

- N. Di Daniele et al., "Impact of Mediterranean Diet on Metabolic Syndrome, Cancer, and Longevity," *Oncotarget* 8, no. 5 (2017): 8947–79; M. Finicelli et al., "Metabolic Syndrome, Mediterranean Diet, and Polyphenols: Evidence and Perspectives," *Journal of Cell Physiology* 234, no. 5 (2019): 5807–26.
8. O. Ajala et al., "Systematic Review and Meta-Analysis of Different Dietary Approaches to the Management of Type 2 Diabetes," *American Journal of Clinical Nutrition* 97, no. 3 (2013): 505–16; J. Salas-Salvadó et al., "Prevention of Diabetes with Mediterranean Diets: A Subgroup Analysis of a Randomized Trial," *Annals of Internal Medicine* 160, no. 1 (2014): 1–10; Публикуван коригиран вариант е наличен в 169, no. 4 (2018): 271–272.
  9. Shai et al., "Weight Loss with a Low-Carbohydrate"; F. M. Sacks et al., "Comparison of Weight-Loss Diets with Different Compositions of Fat, Protein, and Carbohydrates." *New England Journal of Medicine* 360, no. 9 (2009): 859–73; C. Haro et al., "Two Healthy Diets Modulate Gut Microbial Community Improving Insulin Sensitivity in a Human Obese Population," *Journal of Clinical Endocrinology Metabolism* 101, no. 1 (2016): 233–42.
  10. N. Di Daniele et al., "Impact of Mediterranean Diet on Metabolic Syndrome, Cancer, and Longevity," *Oncotarget* 8, no. 5 (2017): 8947–79.
  11. E. Toledo et al., "Mediterranean Diet and Invasive Breast Cancer Risk Among Women at High Cardiovascular Risk in the PREDIMED Trial: A Randomized Clinical Trial," *JAMA Internal Medicine* 175, no. 11 (2015): 1752–60.
  12. E. H. Martínez-Lapiscina et al., "Mediterranean Diet Improves Cognition: The PREDIMED-NAVARRA Randomised Trial," *Journal of Neurology and Neurosurgery Psychiatry* 84, no. 12 (2013): 1318–25; A. Knight et al., "A Randomised Controlled Intervention Trial Evaluating the Efficacy of a Mediterranean Dietary Pattern on Cognitive Function and Psychological Wellbeing in Healthy Older Adults: The MedLey Study," *BMC Geriatrics* (2015): 15:55; C. Valls-Pedret et al., "Mediterranean Diet and Age-Related Cognitive Decline: A Randomized Clinical Trial," *JAMA Internal Medicine* 175, no. 7 (2015): 1094–103.
  13. J. E. de la Rubia Ortí et al., "Improvement of Main Cognitive Functions in Patients with Alzheimer's Disease After Treatment with Coconut Oil Enriched Mediterranean Diet: A Pilot Study," *Journal of Alzheimer's Disease* 65, no. 2 (2018): 577–87.
  14. S. I. Katz et al., "Randomized-Controlled Trial of a Modified Mediterranean Dietary Program for Multiple Sclerosis: A Pilot Study," *Multiple Sclerosis Related Disorders* 36 (2019): 101403.
  15. O. Ajala et al., "Systematic Review and Meta-Analysis."
  16. J. L. Steiner et al., "Impact of Alcohol on Glycemic Control and Insulin Action," *Biomolecules* 5, no. 4 (2015): 2223–46; M. B. Esser et al., "Peer Reviewed: Prevalence of Alcohol Dependence Among US Adult Drinkers, 2009–2011," *Preventing Chronic Disease* 11 (2014); R. W. Wilsnack et al., "Gender

Differences in Binge Drinking: Prevalence, Predictors, and Consequences,” *Alcohol Research: Current Reviews* 39, no. 1 (2018): 57–76.

17. K. Bogusz et al., “Prevalence of Alcohol Use Disorder Among Individuals Who Binge Eat: A Systematic Review and Meta-Analysis,” *Addiction* (2020). doi: 10.1111/add.15155.