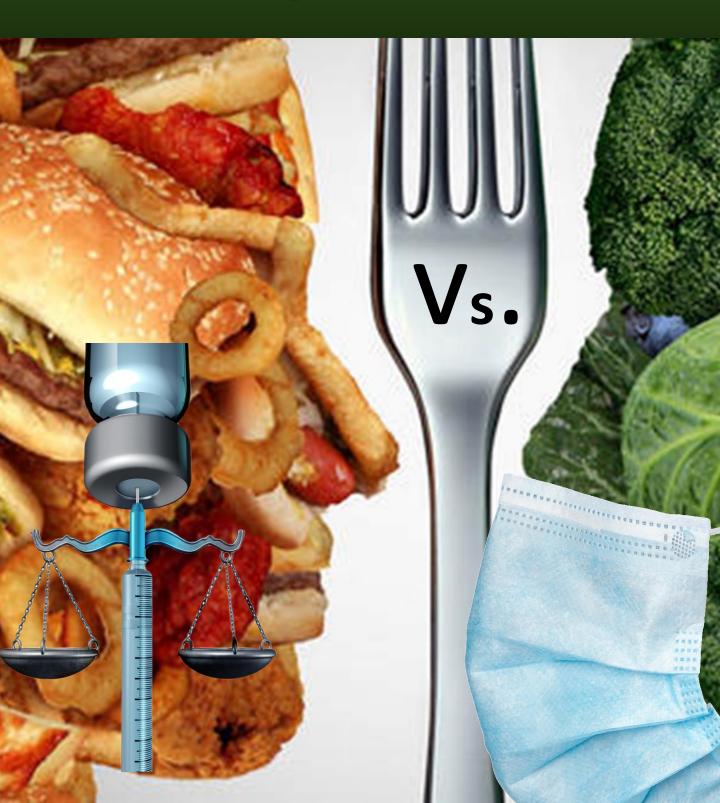
The Plant-Based Mandate Option - Bill #34-21





SUPPORT WITH PROPOSED FRIENDLY AMENDMENTS <u>PLANT-BASED MANDATE OPTION</u> Expedited Bill 34-21 COVID-19 Vaccination Required October 18, 2021

Council President: Tom Hucker Council Vice President: Gabe Albornoz Montgomery County Council Stella B. Werner Council Office Building 100 Maryland Avenue Rockville, MD 20850

President, Vice President and entire Montgomery County Council (Council), thank you for the opportunity to share our support of Expedited Bill 34-21 with "friendly amendments" (Amendments) that our medical advisory council are confident will help to achieve the Council's expressed interest in reducing severe Covid, and even its spread, in employees.

The HBCU College of Plant-Base Lifestyle Medicine (HBCUCPLM) is a new Maryland 501c3 born out of the pandemic in August 2020, whose mission is to educate and train medical professionals of color and HBCU health science students and grads in the practice of prescribing whole plant-based "Food as Medicine" and lifestyle interventions knowns as THE TEN LAWS of Plant-based Lifestyle Medicine (THE TEN LAWS). THE TEN LAWS is the evidence based theraputic use of whole food plant-based nutrition and lifestyle interventions for the prevention, treatment, and reversal of chronic degenerative diseases and to reduce the severity of communicable diseases in communities of color. There are over 100 peer reviewed studies on the benefits of a plant-based diet and lifestyle interventions in treating and even reversing chronic diseases and optimizing immunity in persons of all ages, ethnicities and genders.¹

While Bill 34-21 focuses on mandating employee vaccination as the single medical intervention capable of reducing severe Covid and its spread, the mandate fails to address what the Centers for Disease Control (CDC) has determined are key "risk factors" (pre-existing chronic disease), which have been identified as the underlying causes for severe Covid outcomes and death.²

PRO MANDATE ALTERNATIVE - ALL SHOULD BE ACCOUNTABLE FOR REDUCING SEVERE COVID

Consequently, the proposed friendly amendments are aimed at reducing the primary risk factors, particularly for those who choose to remain unvaccinated based on "sincerely held religious beliefs". At the same time, the Amendments seek to provide an "alternative mandate" that offers a <u>form of</u> <u>exemption</u> to religious objectors to the vaccine (which Judge David Hurd of the U.S. District Court for the Northern District of New York citizens have constitutional right to seek³) that also requires the unvaccinated to <u>take personal action</u> for reducing severe Covid and its spread by participating in THE TEN LAWS of Plant-Based Lifestyle Medicine. Several scientific studies have demonstrated that a plant-based diet can reduce severe Covid by 73%, which will be discuss later in the memorandum of support.

¹ See Plant-based Research Database - <u>https://plantbasedresearch.org/</u>

² See CDC Report - Underlying Medical Conditions Associated with Higher Risk for Severe COVID-19: Information for Healthcare Providers, Updated Oct. 14, 2021 - <u>https://www.cdc.gov/coronavirus/2019-ncov/hcp/clinical-</u>care/underlyingconditions.html

³ See Memorandum & Order for Injunction from Judge Hurd in Dr.'s A-Z v. NY Gov. Hochul - <u>https://eadn-wc01-1479010.nxedge.io/cdn/wp-content/uploads/2021/09/Dr.-A-v.-Hochul-Order-GRANTING-PI-10-12-21.pdf</u>



<u>RELIGIOUS EXEMPTION NOT NEEDED – ALTERNATIVE MANDATE REDUCES MARGINALIZING</u> <u>SINCERE RELIGIOUS OBJECTORS</u>

While the HBCUCPLM supports everyone's right to seek religious exemptions, we are concerned that those who obtain religious exemptions who are not already on a plant-based diet will do nothing (in addition to masking or testing) to reduce the risks of severe Covid in the workplace by remaining unvaccinated without addressing their own pre-existing risk factors for contracting severe Covid and dying. On the other hand, we are also concerned that many in the very small religious communities who do practice a plant-based lifestyle, Hindus, Buddhist, Seventh-Day Adventist and some Muslims may get terminated despite the fact that their employment does not pose a safety risk to the County because the scientific evidence establishes that those who practice a plant-based diet have a 73% reduce chance of getting severe covid or even contracting the disease. (The studies will be discussed later in the memorandum.)

Because religious exemptions pursuant to Title VII does allow employers to reject religious exemption and accommodations based on a claim that any accommodation could place a "de minimis amount" of undue burden on the workplace, it is our position that religious exemptions, including medical exemptions can be a breeding ground for workplace discrimination that will not meet the objectives of the Council, which is to reduce severe Covid in the workplace.

To avoid the unnecessary process of the County having to try to evaluate whether an unvaccinated employee should or should not receive a religious exemption or medical exemption, the HBCUCPLM proposes the following amendments as the best solution to meet the Council's true objectives without the risk of the Country treating religious groups or medical conditions differently and harming employees and the County.

PROPOSED FRIENDLY AMENDMENT

The HBCUCPLM proposes the following friendly amendments to Bill 34-21, wherein the vaccine mandate should eliminate the medical exemption and add the mandate that unvaccinated employees who elect not to take the vaccine regardless of the reason – whether religious or medical, are required to complete a plant-based lifestyle medicine training course so that they will change their diet, which is a primary determinate of Covid severity and death. The specific amendment language is as follows: (See Draft Amendment redlines attached as **Exhibit A**):

- 1. Add to section 33-22(b)(3) the following:
 - a. "Or alternatively complete THE TEN LAWS of Plant-Based Lifestyle Medicine 28-day online training course;
- 2. Add to Section 33022(c)(1)(A) the following:
 - a. attend the first online THE TEN LAWS of Plant-Based Lifestyle Medicine course and report weekly attendance;
- 3. Add to Section 33022(c)(1)(b) B. All employees enrolled in the online THE TEN LAWS of Plantbased Lifestyle Medicine are required to:
 - a. test once a week for Covid-19 utilizing free testing services by the County.
 - b. provide electronic proof of course attendance
 - c. within thirty (30) days from completing the course until the need for the Covid vaccine mandate is declared ended by the County Executive, submit monthly grocery receipts to the Human Resource Department to show compliance with the plant-based dietary plans learned in the course; and
 - d. at the option of the employee, disclose any positive changes in medical condition due to participating in the training by submitting a video, photo or written narrative.
- 4. Add a new Section that states: The Director of Department of Health and Human Services shall approve all instructors that will provide THE TEN LAWS of Plant-Based Lifestyle Medicine training.



- (C) "All instructors, at minimum, shall have practiced a whole food plant-based lifestyle for at least three (3) years and are either a licensed physician in any state, a PhD in Nutrition, Dietetics, Naturopath, Preventative Health, or Naturopath, a registered nurse or certified plant-based lifestyle medicine coach."
- 5. Eliminate the medical exemption. If the Plant-Based Mandate option is added to the bill, there is not need for a medical exemption. Having the medical exemption without a religious exemption leaves the bill subject to constitutional challenge as in the New York case decided last week.

COMPELING INTEREST FOR THE AMENDMENT

According to the CDC, Covid-19 vaccines are effective at helping to protect against severe disease and death caused by the Covid-19 virus and variants. Nowhere on the CDC website regarding the benefits of the vaccine does it say that those who are vaccinated cannot "cause" the spread of the virus that causes Covid-19.⁴ Most important it states plainly on the OSHA website⁵, (which is the organization directed by President Biden to create workplace safety standards for private and public employers), the following:

"However, <u>preliminary evidence</u> suggests that fully vaccinated people who do become infected with the Delta variant can be infectious and can spread the virus to others."

According to Dr. Christina Parks, PhD, a molecular biology scientist, (experienced in MRNA vaccine research), experts have always been aware that vaccines, in general, do not STOP the spread of viruses. At hearing before the Michigan legislators, Dr. Parks testified on August 19, 2021, stating that neither the Covid vaccines, nor any other vaccine, are designed to "prevent" transmission of a virus. Dr. Parks explained that vaccines are merely created to reduce symptoms.⁶ This testimony is consistent with the statements on the websites for the CDC and OSHA regarding the new Covid vaccine's ability to reduce severity.

Based on the forgoing evidence, there is technically no difference between vaccinated and unvaccinated employees in the workplace as it relates to stopping the spread of Covid-19, which is why the CDC updated its website to require vaccinated to people to continue to wear masks, particularly indoors.

Moreover, the death of <u>General Colin L. Powell</u>, former U.S. Secretary of State and Chairman of the Joint Chiefs of Staff, also establishes that the Covid vaccines do not stop the spread and possibly may not reduce the severity of Covid. According to the CNN, report, Secretary Power died due to complications from Covid 19," according to the Powell family post on Facebook, <u>noting he was fully vaccinated</u>.

PLANT-BASED LIFESTYLE MEDICINE MUST BE ADDED TO THE ARSENAL AGAINST COVID

Based on these undisputed facts, the County has a compelling interest in mandating THE TEN LAWS of Plant-Based Lifestyle Medicine as an additional intervention to mitigate against severe Covid and death in the County's population.

In a very recent review in the *British Journal of Nutrition* it was reported that people with optimal levels of micronutrients may be more resilient to COVID-19.⁷ Micronutrients are <u>vitamins</u> and minerals that

- ⁶ See <u>Testimony of Dr. Christina Parks, PhD</u> Michigan State Legislature Hearing -Aug. 19, 2021.
- ⁷ See Nutritional status of micronutrients as a possible and modifiable risk factor for COVID-19: a UK perspective -British Journal of Nutrition (2021), 125, 678–684 - <u>https://www.cambridge.org/core/services/aop-cambridge-</u> <u>core/content/view/35B4C4BC5B0FBD132370128EC03FE309/S000711452000330Xa.pdf/div-class-title-nutritional-</u> <u>status-of-micronutrients-as-a-possible-and-modifiable-risk-factor-for-covid-19-a-uk-perspective-div.pdf</u>

⁴ See CDC Website – Key Things To Know about the Covid Vaccines - <u>https://www.cdc.gov/coronavirus/2019-ncov/vaccines/keythingstoknow.html</u>

⁵ See OSHA website - Protecting Workers: Guidance on Mitigating and Preventing the Spread of COVID-19 in the Workplace <u>https://www.osha.gov/coronavirus/safework</u>



people obtain from their diet. Human bodies also produce <u>vitamin D</u> in response to exposure to sunlight. Plant foods contain many vitamins and minerals essential for a healthy immune system, such as <u>zinc</u>, <u>selenium</u>, and vitamins A, C, and E. Selenium is a trace mineral that benefits immune system health and cognitive function. According to the CDC however,⁸ only one in 10 adults in the United States eat enough fruits or vegetables.

Furthermore, according to a recent study of a few thousand healthcare workers published June 7, 2021, in BMJ Nutrition Prevention & Health,⁹ it was established that those medical workers who ate a 100% plant-based diet had a <u>73% reduction in Covid severity</u>. The study also showed that those who eat a "pescatarian diet" which is a predominantly plant-based diet, with some consumption of only fish, has an approximate 51% reduction in Covid-19 severity. That same study also showed that those who were on a predominantly animal flesh diet had an approx. 45% increase in Covid-19 severity.

In a second study performed by Massachusetts General Hospital just published September 8, 2021, it was stated that a healthy plant-based diet was also linked to a lower risk of "getting" Covid-19 and a lower risk of severe symptoms. Lastly, a third study of approximately 600,000 individuals was published in June 24, 2021, also concluded that a plant-based diet was associated with lower risk and severity of Covid-19.¹⁰

While more and more studies are being done and have been done on the macronutrients in plants and there effectiveness in preventing severe Covid, legislators have a compelling interest for insuring these no risk Interventions are implemented in conjunction with the other interventions including masks and testing.

TESTING ALONG WITH ALTERNATIVE PREVENTATIVE STRATEGIES ARE EFFECTIVE AT STOPING THE SPREAD OF COVID-19 AND REDUCES COVID SEVERITY

While the proposed alternative mandate relieves the unvaccinated from taking the covid vaccine if employees choose the option, the option does not or should not relieve ALL employees from periodic testing and mask wearing. According to the Safer Federal Workforce Task Force report of April 7, 2021, it states that screen testing is useful to detect and stop transmission of Covid, which the report states as follows:

Screening Testing of asymptomatic persons without known or suspected exposure to SARS-CoV-2 Viral testing of asymptomatic workers without known or suspected exposure to SARS-CoV-2 (screening testing) may be useful to detect SARS-CoV-2 early and stop transmission quickly, particularly in areas with community COVID-19 indicators in the moderate to high categorizations (Table 2, Table 3). Persons with asymptomatic or presymptomatic SARS-CoV-2 infection are significant contributors to SARS-CoV-2 transmission. Screening testing should be used as an addition to, not as a replacement for, other prevention strategies.

While the report indicates that the screening should be used as an addition to other prevention strategies, the other preventative strategy is the plant-based mandate option, wherein the goal is to convert the unvaccinated to a 100% whole food plant-based diet to eliminate the risks of Covid.

In summary, weekly testing is an effective method for early detection of Covid that reduces the spread of the virus along with the preventative strategy of a plant-based diet and THE TEN LAWS which the medical evidenced contained in the above medical literature establishes are effective "preventative strategies" that should be added to the bill.

⁸ See CDC - <u>https://www.cdc.gov/nccdphp/dnpao/division-information/media-tools/adults-fruits-vegetables.html</u>

⁹ See Plant-based diets, pescatarian diets and COVID-19 severity: a population-based case–control study in six countries – BMJ Journal Jun 2021 - <u>https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8219480/</u>

¹⁰ See Diet may affect risk and severity of COVID-19 - September 8, 2021, <u>https://www.sciencedaily.com/releases/2021/09/210908180530.htm</u>



WHAT ARE THE TENS LAW OF PLANT-BASED LIFESTYLE MEDICINE?

THE TEN LAWS is an acronym our organization has developed to describe the wholistic approach that we follow to promote health by promoting the lifestyle laws of nature, which when followed collectively on a daily basis, the scientific literature reveals are effective at preventing, reducing and reversing common degenerative disease. The law are as follows: T = law of Trust in natural laws as revealed either by an intelligent creator or revealed according to one's faith or belief system - daily everyone exercises trusts in something in order to function in this world and that trust or belief impacts health positively – violation of the law of trust results in poor emotional states depression and fear that impact health negatively according to the scientific literature, H = The law of Hygiene, which requires each person to wash themselves and keep a clean environment to maintain health – violation of this law result in sickness and disease, E= The law of Exercise – which requires all person to engage in daily physical movement to maintain proper blood flow – violation of this law results in obesity and other diseases that result in premature death, T = law of **Temperance**, this natural law informs each person that he or she should refrain from ingesting harmful toxic substances, like smoking, alcohol, illicit drugs other illicit behaviors, which such substances are ingested one natural system is harmed, E = law of Excellent Air – no one can live without air, and poor air quality can be harmful to one's health and contribute to sickness and disease, N = law of natural whole food plan-based food – no person can go more than 50 days without food and when someone is starving they need food that is nutrient dense to survive and only living plant-based food provide all the micro and macronutrients that body needs, L = Law of rest, scientific studies shows that lost rest reduces life expectancy and contributes to mental health problems, A = law of an attitude of gratitude, studies have shown that a grateful heart works like a medicine to the mind and contributes to preventing mental health diseases, W= law of Water – a person can only go two weeks without water before they die – water is essential to all health, and S = Law of Sunshine – nothing can live or survive on the earth without sunshine, including humans – daily we must obtain sunshine to maintain adequate amounts of sunshine for the body to make Vit D an essential vitamin needed for proper respiratory function and more.

The course will provide attendees with live physician discussions regard the scientific research, medical journal articles and clinical studies that support the effectiveness of following THE TEN LAWS along with practical implementation tools, schedules, meal plans and groups discussions about real cases involving success stories.

In addition, attendees are educated about the importance of following each one of the lifestyle laws on a daily basis based on the anatomy and physiology of their body. Attendees do not just focus on "food", attendees are taught to manage all aspect of wellness from a wholistic approach focusing on mind, body and spirit as they define their spiritual life.

The proposed instructors who will teach this course may call their course by a different name, but the subject matter will be the same. Below are proposed medical professionals that can provide the online course along with links to their websites that provide details regarding how the courses are structured to optimize the learning experience to result in lasting behavior modifications that can reduce severe Covid and other chronic diseases.

CAN THE TEN LAWS OF PLANT-BASED LIFESTYLE MEDICINE AFFECT THE HEALTH STATUS OF THE UNVACCINATED IN THE SHORT-TERM? YES!!!!

While many believe it takes years before bad food can affect your health, compromise your immune system, and result in deadly chronic disease and death. A recent study demonstrated in the block buster documentary "The Game Changers" shows that every meal a person eats has a positive or negative impact on their blood flow and overall health within hours. This evidence further supports the need for the unvaccinated to participate in THE TEN LAWS of plant-based lifestyle medicine to receive immediate benefits from a lifestyle change.

<u>CLICK HERE TO WATCH THE</u> <u>3 MIN VIDEO TO SEE THE IMPACT OF A</u> <u>PLANT-BASED MEAL v. AN ANIMAL BASED MEAL</u> <u>https://www.hbcuplantbasedlifestyle.com/Evidence%20of%20Impact%20of%20Animal%20Food%20On%20Blood-</u> <u>%20Game%20Changers%20Clip%20-%20BILl%2034-21.mp4</u>

> 14216 DUNWOOD VALLEY DR., BOWIE MD 20721 www.HBCUPlantBasedLifestyle.com - Phone: 602-326-8663



IMPLEMENTATION & COST SAVINGS

The HBCUCPLM is happy to announce that our advisory board members have the tools to provide THE TEN LAWS of plant-based lifestyle medicine ready for the County to implement. Below are two of our advisory board members and their programs which are currently available for the County's employees to commence once the bill is passed with the friendly amendments.

DR. BATER MONTGOMERY

Dr. Baxter Montgomery, one of our HBCUCPLM board members, is a board-certified Cardiologist and Electro physicist and has for over 15 years practiced plant-based lifestyle medicine at the Montgomery Heart & Wellness Clinic where he serves as the Medical Director. Dr. Montgomery particularly serves communities of color and has completed medical research studies published in acclaimed journals on the benefits of a plant-based diet. (See Dr. Montgomery's three (3) clinical studies attached as **Exhibit A** along with his CV and he will be starting another study this November) Dr. Montgomery provides online plant-based lifestyle medicine group trainings, as well as one on one care, that reverse heart disease (still the number # killer over Covid), diabetes, lowers blood pressure and helps patients get off of medications. To learn more about Dr. Montgomery – click here to watch a quick overview of how Dr. Montgomery has used plant-based interventions on patients in critical care in the ICU.¹¹ Dr. Montgomery offers his 30 day version of THE TEN LAWS online plant-based lifestyle courses offered weekly in the evenings at the total cost of \$599, which includes a year access and support which County employees can enrolled at any time.

Finally, Dr. Montgomery has treated more than several dozen patients who have tested positive for Covid during this past year, and he has not lost one patient to sever Covid or death. In the event any of the unvaccinated employees do test positive for Covid, Dr. Montgomery has a very aggressive protocol that he uses to help patients that test positive. However, by having the County's employees enroll in his online course as soon as possible, they can start the detox to improve their immune system so that they have better protections as they continue to work. <u>Click here to view</u> the course that best fits the Counties unvaccinated employee needs.¹²

DR. RUBY LATHON

Another HBCUPLM board member that is available to implement the mandate is Ruby Lathon, PhD. She is a board-certified holistic nutritionist with an inspiring powerful story of how she recovered from thyroid cancer through natural treatments focused on a whole food, plant-based diet. Dr. Lathon was featured in the hit documentary, <u>What the Health</u>. Dr. Lathon lives in Montgomery County, has an office in D.C. and the only plant-based African American nutritionist in the entire state that exclusively practice plant-based lifestyle medicine nutrition education. To help individuals with the learning curve of changing to a plant-based diet, Dr. Lathan provides weekly in-person and virtual group cooking classes. However, during the beginning of the Covid pandemic, Dr. Lathon also launched an affordable whole food Plant-Based meal delivery service called <u>Ruby Reds Vegan</u>. See Dr. Lathon's CV attached as Exhibit B.

With these two awesome plant-based lifestyle medicine practitioners ready to work with the County to implement the proposed plant-base mandated option, we hope that the Council will strongly consider all of the upsides of this very friendly amendment.

It is important to note that, there are less than 25 board certified plant-based lifestyle medicine physicians of color in the entire United States out of over 55,000 physicians of color. These two medical professionals represent the very tiny community of practitioners of color that live and practice this form of medicine.

¹¹ See Dr. Baxter Montgomery – The Plant-Based Doctor https://www.youtube.com/watch?v=KMidCUcDdJc&t=158s

¹² See Dr. Montgomery Weekly Plant-Based Food Rx Healthy Lifestyle Series - <u>https://www.online.montgomeryheart.com/lifestyle</u>



It is the mission of the HBCCPLUM to create a pipeline of future plant-based physicians of color in the state of Maryland and particularly in Montgomery County by focusing on training efforts on the states four HBCUs.

Therefore, one of our additional goals is to develop a Center of Plant-Based Lifestyle Medicine and Business Innovation in Montgomery County patterned after the clinical facility operated by Dr. Baxter Montgomery. However, we seek to add with the clinic an incubator/accelerator space for continued research and plantbased business development so that we can create a farm, to medical professional to table supply chain solution that also increases the availability of fresh whole plant-based foods in Maryland's "food deserts".

POTENTIAL IMPLEMENATION COST TO COUNTY

Please note that if County were to pay for the course for all its unvaccinated employees as an employee benefit, Dr. Montgomery would provide a special rate for the County so that the impact is not drastic unexpected budgetary expense to the County. However, it is our understanding that Montgomery County received Federal Cares/Covid County Relief Funding which allows for expenditures for Covid mitigation education. If those funds are available, the proposed plant-based mandate option would have a ZEOR fiscal impact on the Counties budget.

POTENTIAL COST TO THE EMPLOYEE WITH UPSIDE BENEFITS

Alternatively, if the County is not in the position to fund employee's participation in the program, the County could still mandate the plant-based option and have the unvaccinated employees pay for the course on their own, thereby saving THE County from experiencing an unexpected budget item. This implementation option is most closely akin to what happened in the Supreme Court Jacobson v. Massachusetts 1905 vaccine mandate case. In that case, the Supreme Court upheld a fine levied against Mr. Jacobson failing to get vaccinated under the state's mandatory vaccination program. In this case, however, the cost of the program could be charged to the employee without infringing on any constitutional rights of the employees. Because the plant-based mandate option is truly narrowly tailored to the Government interest of reducing severe Covid based on substantive medical research that shows that such a course is truly beneficial to the reduction of Covid severity, the County should prevail in any court challenge. The course is not a "fine" per se, but rather a "safety" program that benefits the entire health of the employee directly. As mentioned before, it may be a small price that employees will pay rather than have the threat of loosing a job or going against their sincerely held religious beliefs.

LONG TERM COST SAVINGS TO THE COUNTY

Lastly, if the County does decide to pass the plant-based mandate option and pay the cost of the course, the County will realize a huge long term cost savings on the healthcare cost for the unvaccinated. According to the Maryland Diabetes Association, the state of Maryland spends approximately <u>\$7 Billion on diabetes</u> care for its residents. If any of the unvaccinated employees required to take the training do suffer from diabetes, the County could save thousands if not millions of dollars if any of unvaccinated employees comply with the program and their diabetes is reversed and they are able to get off of their insulin or whatever medication they may be on. According to the Maryland Diabetes Association, insulin for 1 year for 1 employee is between \$8,000 and \$12,000 per year. If just 45 unvaccinated employees are on insulin are required to take the plant-based lifestyle medicine course and they are able to get off their medication, the cost of the entire online training for all 800 employees would be paid for and year after year the County will save millions.

CONCLUSION

The HBCUCPLM respectfully requests that the Council accept the proposed friendly amendments and pass Bill 34-21 with the Plant-Based Mandate Option. This proposal is a WIN/WIN for the County on multiple levels and for the unvaccinated employees who want to keep their jobs and would value the opportunity to take such a valuable course, this bill amendment could save their lives and improve their overall health.



To pass the bill without a religious exemption and only with a medical exemption which does nothing to address the underlying causes of severe Covid and Covid related deaths, will only open the door for litigation and tons of money spent on lawyers rather than on employees that deserve the opportunity to "choose" an option that can truly have a positive impact on the reduction of severe Covid in their lives.

If the Council would like to receive more information about the programs, please contact Jo Saint-George, Chair of the HBCUCPLM at cell #602-326-8663 or by email at exec@HBCUPlantbasedlifestyle.com

Thank you for your attention to this proposal and your vote.

Sincerely,

Jo Saint-George (I am a resident of Gaithersburg, Maryland) Chair

w/Attachments

Supported by:

Dr. Baxter Montgomery Dr. Ruby Lathon Dr. Theodor Watkins Dr. John St. Rose

FRIENDLY AMENDMENT

Expedited Bill No. 34-21 Concerning: Personnel

| | Concernin | g: Personnel | and | Human |
|-----------|------------|----------------|---------|-----------|
| | Resou | arces - COVID |)-19 Va | ccination |
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Proposed by the HBCU Colleg

Plant-Based Lifstyle Medicin

COUNTY COUNCIL FOR MONTGOMERY COUNTY,

Lead Sponsors: Councilmembers Riemer and

AN EXPEDITED ACT to:

(1) require the vaccination of County employees against COVID-19; or alternatively (2) require THE TEN LAWS of Plant-Based Lifestyle Medicine 28-day mandatory online training option (32) permit medical accommodations to the COVID-19 vaccination requirements;

 $(\underline{43})$ exempt the COVID-19 vaccination requirements from collective bargaining; and

By amending

Montgomery County Code Chapter 33, Personnel and Human Resources

| Boldface | Heading or defined term. |
|------------------------------|---|
| Underlining | Added to existing law by original bill. |
| [Single boldface brackets] | Deleted from existing law by original |
| bill. Double underlining | Added by amendment. |
| [[Double boldface brackets]] | Deleted from existing law or the bill by amendment. |
| * * * | Existing law unaffected by bill. |
| | - |

The County Council for Montgomery County, Maryland approves the following

| 1 | Sec 1 | . Section 33-22 is amended as follows: |
|------------------------|--|--|
| 2 | 33-22. [Res | served.] COVID-19 Vaccination |
| 3 | <u>(a)</u> | Definitions. For purposes of this section, the following words have the |
| 4 | | meanings indicated. |
| 5 | | <u>COVID-19</u> Vaccine means a vaccine authorized or approved by |
| 6 | | the federal Food and Drug Administration to prevent or reduce |
| 7 | | the transmission of SARS-CoV-2. |
| 8 | | Employee means an individual employed by the County, regardless of |
| 9 | | the individual's merit system status or representation by an |
| 10 | | employee organization. |
| 11 | | Fully vaccinated means having received all doses of a COVID-19 |
| 12 | | vaccine. |
| 13 | <u>(b)</u> | Vaccination Required. As a condition of employment by the County, |
| 14 | | an |
| 15 16 | | (1) <u>be fully vaccinated and provide to the County proof of</u> <u>vaccination</u> |
| 17 | (c) | (2) <u>under subsection (c); or complete the 28-day THE TEN LAWS</u> <u>of Plant-based Lifestyle Medicine online course</u> <u>Procedures; remedies for</u> |
| 19 | <u>(C)</u> | (1) Within 7 days after notification by the County to an employee of |
| 20 | | the requirements of this section, the employee must: |
| 20 | | (A) provide to the County proof that the employee is fully |
| 21 22 23 24 25 | See rest of additional language on last page. | (A) provide to the county proof that the employee is fully vaccinated; or attend the first online THE TEN LAWS of Plant-based Lifestyle Medicine Course and complete the course (B) apply for an accommodation under subsection (d) must be |

- 2 -

F:\LAW\BILLS\21xx Employee Vaccination Requirement\Bill

| 2 | 5 | | <u>(3)</u> | Within 7 days after being placed on unpaid leave under |
|----|---|------------|-------------|---|
| 2 | 7 | | | paragraph |
| 2 | 3 | | | (A) provide to the County proof that the employee has |
| 29 |) | | | received |
| 30 |) | | | (B) <u>at least one dose of a COVID-19 vaccine; or</u> |
| 3 | l | | <u>(4)</u> | An employee under subparagraph (3)(A) must provide to |
| 32 | 2 | | | the |
| 3. | 3 | | | County, within 40 days of being placed on unpaid leave, proof |
| 34 | 1 | | (5) | that the employee is fully vaccinated. |
| 3 | 5 | | | An employee who fails to comply with paragraphs (3) or (4) of |
| 30 | 5 | | | this subsection, or with paragraph (3) of subsection (d), |
| 3' | 7 | <u>(d)</u> | <u>Heal</u> | th-based |
| 3 | 3 | | <u>(1)</u> | An employee may apply for an accommodation to the |
| 3 |) | | | requirements of this section based on the health of the |
| 40 |) | | <u>(2)</u> | employee. The Director of Human Resources, or the |
| 4 | l | | | Director's designee, must approve an application for an |
| 42 | 2 | | | accommodation if the accommodation is required for the |
| 43 | 3 | | | health of the employee, as documented by a licensed |
| 44 | 1 | | <u>(3)</u> | physician. |
| 4 | 5 | | | Within 7 days after the denial of an application for an |
| 4 | 5 | | | accommodation under paragraph (1), the employee must provide |
| 4′ | 7 | | | to the County proof that the employee has received at least one |
| 4 | 3 | | | dose of a COVID-19 vaccine. Within 40 days after the denial of |
| 49 |) | | | the application, the employee must provide to the County proof |
| 50 |) | <u>(e)</u> | Exer | mption from Collective Bargaining. The requirements and Formatted: Indent: Left: 0" |
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F:\LAW\BILLS\21xx Employee Vaccination Requirement\Bill

EXPEDITED BILL NO. 34-

| 52 53 54 55 56 | legislat takes e | receipts to the Human Resources Department to show compliance with the plant-based dietary plans learned in the course; and iii) at the option of the employee, disclose any positive medical condition changes as a result of participating in the training by submitting a video, photos or written narrative, C. The Director of Department of Health and Human Services shall approve all instructors that will provide THE TEN LAWS of Plant-Based Lifestyle Medicine training, which minimum requirements shall be that the instructor lives the whole food plant-based lifestyle and is either a licensed physician in any state, a PhD in Nutrition, Dietitics, Naturalpath, Preventative health, or | Formatted: Indent: Left: 1", First line: 0.5" |
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BAXTER DELWORTH MONTGOMERY, MD The Plant-Based Physician <u>Montgomery Heart & Wellness</u> <u>Video Bio</u>

| EXPERIENCE: | Clinical Assistant Professor The University of Texas Health Science Center Department of Medicine Division of Cardiology/Clinical Cardiac Electrophysiology |
|-----------------------------|--|
| | President and CEO Houston Associates of Cardiovascular Medicine, PA. (1997-Present) |
| | Executive Director The Johnsie and Aubary Montgomery Institute of Medical Education and Research (a 501(c) 3 nonprofit organization) |
| BIRTHPLACE: | Houston, Texas United States of America |
| OFFICE ADDRESS: | 10480 South Main Street Houston, Texas 77025 (713) 599-1144 phone (713) 599-1199 fax bmontgomery@drbaxtermontgomery. com |
| UNDERGRADUATE EDUCATION: | William Marsh Rice University Houston, Texas Bachelor's Degree in Biochemistry (1986) |
| GRADUATE EDUCATION: | The University of Texas Medical Branch at Galveston Galveston, Texas Doctor or Medicine |
| RESIDENCY: | Baylor College of Medicine Houston, Texas Internal Medicine |
| FELLOWSHIP: | The University of Texas Health Science Center at Houston Houston, Texas Cardiovascular Diseases Clinical Cardiac Electrophysiology |

| CERTIFICATION: | Diplomate of the American Board of Internal Medicine, Cardiovascular Diseases |
|------------------------|--|
| | Diplomate of the American Board of Internal Medicine, Clinical Cardiac Electrophysiology |
| LICENSURE: | Texas State Board of Medical Examiners (Since 1999) Permit Number H9549 |
| HOSPITAL APPOINTMENTS: | |

Attending Physician Memorial Hermann Hospital - The Texas Medical Center Houston, Texas

Attending Physician The Heart and vascular Institute Memorial Hermann Hospital - The Texas Medical Center Houston, Texas

Consulting Physician Select Specialty Hospital - Heights Houston, Texas

TEACHING RESPONSIBILITES:

Teaching Faculty for Cardiology Fellows and Clinical Advanced Nurse Practitioners The Heart and Vascular Institute Memorial Hermann Hospital - The Texas Medical Center 1997 - Present

Cardiovascular Disease Lecturer GlaxoSmithKline, Inc. 2000 - Present

Cardiovascular Disease Lecturer Novartis, Inc. 2006 - Present

Cardiovascular Disease Lecturer Boston Scientific, Inc. 2006 - Present

Co-Director and Lecturing Faculty Cardiology Concepts for Non-Cardiologists (An Annual Houston Area Educational Symposium) JAM Institute, Inc. 2006 - 2008

Steering Committee Member and Lecturing Faculty *Close the Gap* Boston Scientific, Inc. 2006 - Present

RESEARCH:

CLINICAL STUDIES:

ALLHAT: Antihypertensive and Lipid-Lowering Treatment to Prevent Heart Attack Trial. ALLHAT ALLHAT was a blinded, randomized trial that investigated the relative efficacy of different classes of antihypertensive agents in reducing stroke, illness and death from cardiovascular diseases. A subgroup of patients with hyperlipidemia was randomized comparing Pravastatin compared to usual care. A Houston Site - Principal Investigator (1998)

INVEST: The International Verapamil SR/Trandolapril Study. INVEST was a randomized controlled clinical trial comparing a calcium antagonist treatment strategy (Isoptin® SR) with a non calcium antagonist treatment strategy for the control of hypertension in a primary care coronary artery disease patient population.

A Houston Site - Principal Investigator (2000)

INVEST SUB-STUDY: This study was a sub-study of the INVEST patient population designed to evaluate the impact of genetic differences on pharmacokinetics.

A Houston Site - Principal Investigator (2000)

The Safety and Efficacy of PNU-182716 Versus Rosiglitazone: This was a one-year, randomized, double blind, parallel group, and active comparator study.

A Houston Site - Principal Investigator (2000)

FACTOR: Fenofibrate and Cerivastatin Trial Optimizing Response. FACTOR was a multicenter, randomized, double blind, placebo controlled, parallel group, study of the safety and efficacy of Cerivastatin in combination with Fenofibrate compared to Cerivastatin alone, Fenofibrate alone and placebo in a population of Type 2 Diabetic Men and Women.

Grant Sponsor - Bayer 2001 A Houston Site - Principal Investigator **ADHERE:** ADHERE was a national registry of patients admitted to hospitals with acute decompensated congestive heart failure. **A Houston Site - Principal Investigator (2001)**

STELID TM AND STELIX TM LEADS STUDY: This study was a

safety and efficacy study of steroid-eluting cardiac pacing leads. Grant Sponsor - Ella Medical 2002

ARRHYTHMIA PATHWAY STUDY: This was a patient registry study designed to assess the efficacy of a clinical algorithm for identifying and assessing patients at risk of sudden cardiac arrest.

Grant Sponsor - Medtronic, Inc. 2002

A Houston Site - Principal Investigator

RAPIDO CATHETER STUDY: This study was to evaluate the efficacy of a left ventricular defibrillator-pacemaker lead delivery system. **Grant Sponsor - Guidant, Inc. 2003**

A Houston Site - Principal Investigator

PROTOS HEART RATE DISTRIBUTION STUDY: This was a clinical study designed to compare the heart rate distribution in patients undergoing pacemaker implants requiring heart rate response therapy. This study compared the heart rate distribution of accelerometer rate response therapy to the BIOTRONIK Closed Loop System therapy.

Grant Sponsor - Biotronik, Inc. 2003

A Houston Site - Principal Investigator

CSPP100A2404 - A 54 week, randomized, double-blind, parallel-group, multicenter study evaluating the long-term gastrointestinal (GI) safety and tolerability of Aliskiren (300 mg) compared to Ramipril (10 mg) in patients with essential hypertension.

Sponsored by Novartis, since April 4, 2008.

A Houston Site - Principal Investigator

CSPP100AUS03 - An 8 week Prospective, Multicenter, Randomized, Double-Blind, Active Control, Parallel Group Study to Evaluate the Efficacy and Safety of Aliskiren HCTZ versus Amlodipine in African American Patients with Stage 2 Hypertension. Sponsored by Novartis, since August 2008. **A Houston Site - Principal Investigator**

CSPP100A2409- An 8 week randomized, double-blind, parallel-group, multicenter, active-controlled dose escalation study to evaluate the

efficacy and safety of Aliskiren HCTZ (300/25 MG) compared to Amlodipine (10 mg) in patients with satage 2 systolic hypertension and diabetes mellitus. Sponsored by Novartis, since December 2008. **A Houston Site - Principal Investigator**

SPAIOOAUSOI - An 8 week randomized, double-blinded, parallel-group, multicenter, active-controlled dose escalation study to evaluate the efficacy and safety of Aliskiren Administered in Combination with Amlodipine (150/5 mg, 300/10 mg) versus Amlodipine alone (5 mg, 10 mg) in African American patient with Stage 2 Hypertension. Sponsored by Novartis, since February 2009.

CLAF237B22OI- A multicenter, randomized, double-blind study to evaluate the efficacy and long-term safety of vildagliptin modifies release (MR) as monotherapy in patients with type 2 diabetes. Sponsored by Novartis, since February 2009.

A Houston Site - Principal Investigator

CLAF237B2224 - A multi-center, randomized, double-blind study to evaluate the efficacy and long-term safety of vildagliptin modified release (MR) as add-on therapy to metformin in patients with type 2 diabetes. Sponsored by Novartis, since February 2009. **A Houston Site - Principal Investigator**

Galaxy study: An aftermarket registry of one of the Biotronik implantable cardioverter defibrillators ICD leads (2009 to present) **A Houston Site - Principal Investigator**

Paradigm study: A multicenter, randomized, double-blind, parallel group, active-controlled study to evaluate the efficacy and safety of LCZ696 compared to enalapril on morbidity and mortality in patients with chronic heart failure and reduced ejection fraction. 2009 -2014 **A Houston Site - Principal Investigator**

BASIC RESEARCH:

In Rapid Separation of Mitochondria from Extra- mitochondrial Space Applied to Rat Heart Mitochondria. An abstract presented at an NIH sponsored student research poster session, Univ. of Texas Medical Branch, Galveston, TX, June 17, 1987.

Regulation of the Adenine Nucleotide Pool-Size of Heart Mitochondria by the ADP/ATP Translocase. Abstract and poster presented at the Galveston-Houston Conference for Cardiovascular Research, Univ. of Texas, Medical Branch, Galveston, TX, February 26, 1988.

The Adenine Nucleotide Pool-Size of Heart Mitochondria is Regulated by the ADP/ATP Translocase. Abstract presented at the 29th Annual National Student Research Forum, University of Texas Medical Branch, Galveston Texas, April 6-8, 1988.

Increased Frequency of the Deletion Allele of the ACE Gene in African-Americans Compared to Caucasians. This study evaluated the prevalence of the deletion allele of the ACE gene in a population of African Americans compared to Caucasians. The findings were presented at the annual meeting of the American College of Cardiology in March of 1996.

Determination of the effect of Calcium infusion on CGRP mRNA Production. A pilot study investigating a possible mechanism by which calcium supplementation may increase CGRP (Calcitonin gene-related peptide, a potent peripheral vasodilator) content in afferent neurons of Sprague Dawley rats, 1990.

PUBLICATIONS:

Montgomery, B, D, MD. A Review of Microanatomy for Medical Students, 1987, chapter 1-8.

Baxter D. Montgomery, MD, Elizabeth A. Putnam, Ph.D., John Reveille, MD, Dianna M. Milewicz. MD, Ph.D.: Increased Frequency of the Deletion Allele of the ACE Gene in African-Americans Compared to Caucasians. (Abstract) J. American College of Cardiology March, 1996

Doyle, N.M., <u>Monga, M.</u>, **Montgomery, B.**, Dougherty, A.H.: Arrhythmogenic right ventricular cardiomyopathy with implantable cardioverter defibrillator placement in pregnancy. J Mat Fetal Neo Med 18:141-4, 2005

Baxter D. Montgomery, MD_Co-Author of Dreams of the nation Book: "Improving Health" with focus on strengthening the food and health connection and replacing unnatural foods from our diet and replacing them with natural foods as a way of reversing illness. 2009

Montgomery, Baxter D: The Food Prescription for Better Health, Houston: Delworth Publishing, 2011

Montgomery,B.D, MD, Effects of the Montgomery Food Prescription on Clinical Biomarkers of Cardiovascular Disease. Plant-based diet can improve clinical biomarkers associated with cardiovascular disease. This study was submitted to the 10th annual Texas A&M University System Pathways Student Research Symposium 2012.

Baxter D. Montgomery, MD Co-Author of the book Rethink Food: About the need for revolutionary change in how to address chronic illness with optimal nutrition.2014

CLINICAL PRESENTATIONS:

Clinical Concepts for Non Cardiologist, Director and Faculty. An educational symposium held for primary care and other non-cardiology specialists in the Houston area. October 2006

Patients at Risk for Sudden Cardiac Arrest Dinner Symposium at the Houston Forum June, 2007

Clinical Concepts for Non Cardiologist, Director and Faculty. An educational symposium held for primary care and other non-cardiology specialists in the Houston area. October 2007

Clinical Concepts for Non Cardiologist, Director and Faculty. An educational symposium held for primary care and other non-cardiology specialists in the Houston area. October 2008

Houston Town Hall Meeting, Director and Faculty. Health summit on the benefits of a healthy nutritional lifestyle for the management of chronic illnesses held for both health care professional and the general public in the Houston area. 2009

Houston Town Hall Meeting, Director and Faculty. Health summit on the benefits of a healthy nutritional lifestyle for the management of chronic illnesses held for both health care professional and the general public in the Houston area. 2010

Houston Health Summit (Town Hall Meeting), Director and Faculty. Health summit on the benefits of a healthy nutritional lifestyle for the management of chronic illnesses held for both health care professional and the general public in the Houston area. 2011 Houston Health Summit (Town Hall Meeting), Director and Faculty. Health summit on the benefits of a healthy nutritional lifestyle for the management of chronic illnesses held for both health care professional and the general public in the Houston area. 2012

Houston Health Summit (Town Hall Meeting), Director and Faculty. Health summit on the benefits of a healthy nutritional lifestyle for the management of chronic illnesses held for both health care professional and the general public in the Houston area. 2013

PROFESSIONAL APPOINTMENTS:

Clinical Assistant Professor of Medicine, University of Texas Health Science Center - Houston 1996 - Present

Steering Committee Member, Boston Scientific Close the Gap Initiative 2005 - Present Scientific/Medical Board of Advisors, Nutritional Excellence, Inc. 2007 -Present

Medical Board of Directors, Twelve Oaks Medical Center Independent Physician's Association 2005 - Present

Medical Executive Committee (Twelve Oaks Hospital), Member at Large 2002 - 2006

Patient Safety Committee (Twelve Oaks Hospital), Chairman 2002 - 2004

Physician Peer Review Committee (Twelve Oaks Hospital) 2002 - 2005

Medical Director, SCCI (Specialized Complex Care) Hospital, 2003 - 2005

Physician Relation Council Advisory Board, Unicare, 2002 - 2004

Aldine Education Foundation: The mission of the Aldine Education Foundation is to provide community-based support to the Aldine Independent School District in pursuit of excellence in teaching, innovation in the classroom and superior learning opportunities for all students.

CLINICAL INTERESTS:

Nutritional Lifestyle Interventions for the Management of Chronic Illnesses Cardiac Pacing and Electrophysiology Diastolic and Systolic Heart Failure Hypertensive Heart Disease Cardiovascular Exercise Physiology Basic Echocardiography Nuclear Cardiology Diagnostic Cardiac Catheterization Cardiovascular Wellness and Nutrition

PROFESSIONAL ASSOCIATIONS:

American College of Cardiology (Elected as Fellow of the College in January, 1999) American Heart Association Heart Rhythm Society (North American Society of Pacing and Electrophysiology, NASPE) American College of Physicians Harris County Medical Society Houston Medical Forum

HONORS AND AWARDS:

Benjamin Spock Award for Compassion in Medicine - 2010

America's Top Physicians - 2007

Cumulative evaluation of "Superior" performance by senior house staff and faculty during first year of residency (Baylor College of Medicine), 1990

Outstanding Young Men of America, 1988

Kempner Award (University of TX Medical Branch) 1986-87 and 1987-88

Academic Scholarship (University of TX Medical Branch) 1986-87

Who's Who Among American Colleges and Universities (Rice University) 1986

Franz Brotzen Outstanding Senior Award (Rice University) 1986

Jones College Service Award (Rice University) 1986 and 1985

100 Black Men of Metropolitan Houston (Awarded in 2012) for the dedication to the improvement of the community.

Physicians Committee for Responsible Medicine- Member of Advisory Board- Current.

ACTIVITIES:

Gardening Scouting Physical Conditioning

DR. RUBY D. LATHON

Email: Ruby@RubyLathon.com 99 Blair Alley SW Suite W962; Washington, DC 20024 Mobile (202) 709-7829

SUMMARY: Dr. Ruby Lathon is a certified holistic nutritionist and inspires with a powerful story of recovering from thyroid cancer through alternative treatment focused on a whole food, plant-based diet. Dr. Lathon worked as a researcher and an award-winning engineer, and now teaches others how to re-engineer their health and live disease free. In the technical arena Dr. Lathon is a highly skilled professional with a wide range of experience in strategic planning and management for resolving long-standing problems and creating solutions that improve operational efficiency. Dr. Lathon has a rich mix of technical analysis, program management, business development, and business management.

| EDUCATION | Holistic Nutrition Certification, 8/2013 | Holistic Nutritionist, Holistic Nutrition Practitioner |
|-----------|--|--|
| | Washington Institute of Natural Medicine, Wash DC | American Naturopathic & Holistic Association |
| | Ph.D., Industrial & Systems Engineering, 8/2000 | Major: Operations Research |
| | University of Alabama in Huntsville | Minor: Engineering Management, Statistics |
| | Dissertation: The Use of Clustering Analysis & Feature E | Extraction for the Reduction of Very Large Data Sets |
| | M.S., Industrial & Systems Engineering, 5/95 | Major: Operations Research |

University of Alabama in Huntsville Minor: Artificial Intelligence, Statistics Thesis: An Intelligent Strategy Discriminator for an Automated Guided Vehicle System

B.S., Computer Science, Cum Laude, 5/92 Oakwood University, Huntsville, AL

EXPERIENCE

5/2010 - Present **DIRECTOR/HOLISTIC NUTRITIONIST, Roadmap to Holistic Health, LLC, Washington DC** Roadmap to Holistic Health, LLC was founded in 2010, and is headquartered in Washington, D.C. Through the leadership of Dr. Ruby Lathon, Ph.D., Roadmap to Holistic Health specializes in health and wellness consultations, whole foods nutrition education, cooking instruction, seminars, and workshops. It works with individuals and groups to deliver scalable and customized nutrition programs. Dr. Lathon is a sought-after speaker in the health and wellness industry. Dr. Lathon's personal experiences uniquely qualify Roadmap to Holistic Health as an expert in a lifestyle-centered, holistic approach to overall health and wellness. www.RubyLathon.com

10/10-6/2016 STRATEGIC PLANNING OFFICER, Government of the District of Columbia, Department of Human Services, Washington, DC.

Implement strategies to redesign the DC public assistance program, including the development of community, and inter and intra-agency relationships. Oversee, plan, and coordinate various aspects of research, design, and program implementation. Provide strategic analysis including analyzing and evaluating the public assistance employment program. Survey and analyze program performance, such as organizational structures; process flow and work systems. Perform independent research and analysis of other state and local public benefits assistance programs to ensure use of best practice standards and innovations in the agency's public assistance program. Collaborate with local and national officials to develop strategies to improve the quality and effectiveness of services. Review and assist in the preparation of legislation that affects agency programs.

12/08 – 10/10 NUTRITION POLICY MANAGER, Physicians Committee for Responsible Medicine, Wash, DC. Developed and managed child nutrition public policy initiatives with a focus on implementing campaigns to promote healthful diets and effect change at state and federal levels. Designed and implemented campaigns to advocate healthful diets to the federal government, non-government organizations, nonprofit health organizations, educational institutions, and the general public through writing, speaking, advertising campaigns, and media outreach. Analyzed legislation and regulations and developed strategic partnerships to further organizational goals. Developed a successful national grassroots campaign. Prepared public education materials and lobbied Members of Congress and state legislators to formulate legislation on critical child nutrition issues. Promoted organization and its mission in professional settings.

- 5/03 12/09 VICE PRESIDENT, EMT Inc. (Engineering, Management & Technology), Albuquerque, NM. Lead efforts in business and technology development and established and operated division offices in Albuquerque, NM. Assisted in development of policies, procedures, and standards and in the development of the long and short-term organizational goals and strategic plans. Responsible for marketing, planning, and management of technical research initiatives. Lead successful development of flagship logistics support software product, OptSim®, including conceptual design, development, testing and marketing. Major Achievements: Highly instrumental in significantly increasing company revenue (>30%) through the cultivation of strategic partnerships, developed all marketing/sales materials, provided increased organizational structure, built and managed the Systems Engineering Division from the ground up.
- 11/98 5/03 SENIOR MEMBER TECHNICAL STAFF, Sandia National Laboratories, Albuquerque NM. Emerging Threats Division, Systems Reliability Department. Project Manager for the Support Enterprise Model Program, a program designed to facilitate instantaneous "situation awareness" of the entire operating, support, and logistics environment for the Joint Strike Fighter and other large, complex systems. Served as Laboratory Project Manager of the Demand Activated Manufacturing Architecture Project that was part of the American Textile Partnership (AMTEX). This project represented a nationwide effort focused on increasing the competitiveness of the fiber, textile, apparel and retail industries. Responsibilities included project management and planning, development of collaboration guidelines, coordinating tasks among four national labs, two commercial software vendors, and four major textile sectors, from the project planning phase through pilot implementation. Other responsibilities include overseeing the development and implementation of a large-scale textile supply chain simulation tool. Other Projects: Project/Technical Lead for the Lockheed Martin P-3C Orion Performance Based Logistics Program; Task Lead for research/development of Information Assurance Toolkit for the DARPA Information Assurance Science & Engineering Tools Program; Spare parts optimization, modeling and simulation, and algorithm development for multi-echelon supply and support systems.

8/94 – 10/98 **RESEARCH FELLOW, NASA Marshall Space Flight Center**, Huntsville AL.

Astrionics Laboratory, Software Simulation Division. Major Project: Worked in cooperation with Lockheed Martin to develop a prototype Automated Diagnostic System for the main propulsion system of the Reusable Launch Vehicle modeled in Gensym's G2 development system. Responsible for development and implementation of the neural network model for the solenoid valve signature traces as well as development of the test conductor control modules for automated test and checkout.

8/92-8/94 **GRADUATE RESEARCH FELLOW, Intelligent Systems Laboratory,** Univ. of AL, Huntsville. Assisted in various research projects in the areas of simulation modeling and expert system design and development. Modeling platforms included AutoMod by AutoSimulations, LISPS and CLIPS. Responsibilities included knowledge acquisition, literary research, and software development. Major Project: Chrysler Huntsville Electronics Division, Automated Guided Vehicle simulation development and analysis.

TEACHING
EXPERIENCE5/2006SEMINAR INSTRUCTOR, Huntsville AL Elementary School Leadership Teams, "Developing Culture
Through Vision, Mission & Values", May 2006.5/98-7/98INSTRUCTOR, Oakwood University, Huntsville, AL; LEAP Adult Degree Continuing Education Program.
Course: Statistics & Business Research Methods.8/94-8/96SUBSTITUTE TEACHER, Huntsville City Schools, Huntsville AL. Took on responsibilities of absent teachers
ranging from kindergarten through 10th grade, throughout the Huntsville City School System.3/91-5/92

| 1/91-3/91 | COMPUTER LAB ASSISTANT , Business & Information Systems, Oakwood University, Huntsville AL. Tutored and assisted students with computer programs, monitored students in progress, responsible for daily lab operations. Utilized knowledge of programming languages and analytical and problem solving skills. TEACHING ASSISTANT , Business & Information Systems, Oakwood University, Huntsville AL. Numerous responsibilities including grading test papers and class work, electronically documenting grades and general clerical duties. Utilized knowledge of programming languages and general computer skills. |
|--|---|
| HONORS & AWARDS | 2007 Girl Scouts of Chaparral Council, Inc. Hall of Fame: Women in Science, Technology, & Engineering Technologist of the Year Award, National Society of Engineers, March 2002 Technology All-Star in Government and Defense Award, Career Communications Group (CCG), July 2001 NASA Graduate Student Researchers Fellowship, 1994-1998; National Science Foundation Assistantship, 1992-1994; Who's Who Among American Colleges & Universities, 1997; General Conference of SDA Regional Scholarship, 1996,1997; ASRS/AGVS User's Association Student Scholarship, 1995,1996; Scholastic Achievement Award (UAH), 1995; ITT Hartford Insurance Student Scholarship, 1991; Scholastic Achievement Scholarship Oakwood University, 1988-1990; National Deans List & Honor Roll. |
| ASSOCIATIONS & COMMUNITY INVOLVEMENT | Board Chair, Vice President - Endorphin Power Company (Substance Abuse Recovery Center) (2005-2007) Del Norte Rotary Club, Paul Harris Fellow, International Committee Matching Grants Chair (2005 – 2009) President, Strategic Action Forum, Albuquerque Think Tank on Public Policy (2006 – 2009) Vice President, National Society of Engineers Alumni Extension-ABQ (99-2001); Coordinated quarterly Professional Development Workshops, tutored students for regional/national competitions and SAT/ACT exams, coordinated a variety of community outreach programs and fundraising events. Graduate Studies Representative - University Judicial Board, University of Alabama, Huntsville Youth Motivation Task Force Consultant, Oakwood University Licensed Foster Parent (2000) Vice President, UAH Graduate Student Association, 96-98 Youth Mentor, CO-ARMM (Coalition for At-Risk Males) Youth Mentor Program, 91 – 97 |
| PROFESSIONAL DEVELOPMENT | Selected for participation in the 2007 Leadership Albuquerque Class. Leadership Albuquerque is the Greater Albuquerque Chamber of Commerce Leadership program developed to enhance the skills of veteran leaders and prepare aspiring leaders to "make a difference" in the economic health and quality of life of the community by teaching how they can take a leadership role in the voluntary (nonprofit) and public (government) sectors. Selected by Sandia National Lab's (SNL) Vice President for the Employee Development, Growth, & Education (<i>EDGE</i>) Program, October 2000. Program's goal is to create next generation leaders by targeting and focusing attention on high-performing managers and staff. Selected for SNL's first Business Development Scholarship Program, February 2001. On-going Training: <i>Social Styles Training Course</i>: Developing communication skills, managing diversity, developing and motivating others, May 2001. Developed, coordinated, and obtained funding for the First Annual Spring Symposium focusing on academic excellence and advancement at the University of Alabama in Huntsville, April 1998. Presented workshop on "Making Professional & Dynamic Presentations". |
| SELECTED CONFERENCE PAPERS | "Optimization Strategies for Complex Supply Chains", R. Lathon, Proceedings of the IIE Annual Conference, Orlando FL, May 2006; Proceedings of Simulation Solutions Annual Conference, Atlanta, GA, March 2005 – Invited (Committee Chair). |
| | "Achieving the Objectives of Performance Based Logistics Modeling", R. Lathon, Proceedings of the 2004 Simulation Solutions Annual Conference, Orlando FL, March 2004 (Committee Chair). |
| | "Supply Chain Optimization via Modeling and Simulation", R. Lathon, Proceedings of the 2002 Defense Manufacturing Conference, Dallas, TX, December 2002. |
| | "The Use of Clustering Analysis and Feature Extraction for the Reduction of Very Large Data Sets, Analyzed Via a RBF Neural Network", R. Lathon, American Institute of Aeronautics and Astronautics (AIAA) Journal, (98-5182), November 1998. |

"An Application of G2 & Neuron-Line to An Automated Propulsion Diagnostic System", R. Lathon, J. Patterson, Proceedings of the Gensym Users Society Worldwide Conference, Paris, France, April 1997. Also in the Proceedings of the NASA University Research Center Technical Conference, Huntsville, AL, Feb 1998.

"Theories-in-use Versus Espoused Theories", C. Bell-Roundtree, R. Lathon, J. Westbrook, D. Utley, Proceedings of the 17th Annual American Society of Engineering Management (ASEM) Conference, University of Texas at Dallas, October 1996.

"Negotiation Among Scheduling Agents to Achieve Global Productions Goals", R. Lathon, A. Claassen, D. Rochowiak, L. Interrante, in Proceedings of the 1994 IEEE International Conference on Systems, Man and Cybernetics, San Antonio, TX, October 1994.

"Intelligent, Dynamic Scheduling of AGV's: Simulation, OR, and AI," L. Interrante, L. Shields, R. Lathon, N. Romero, Proceedings of the CICHME Colloquium, April 1994.

"Design of an Intelligent Manufacturing Scheduling System", L. Interrante, A. Claassen, R. Lathon, Proceedings of the 3rd IERC, January 1994.

SECURITY CLEARANCE: Held DOE Top Secret Clearance (Q) – 2/1999 through 9/2007; Reinvestigation -2005.



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A defined, plant-based diet as a potential therapeutic approach in the treatment of heart failure: A clinical case series



Rami S. Najjar^a, Baxter D. Montgomery^{b,c,*}

^a Department of Nutrition, Georgia State University, Atlanta, Georgia

^b Houston Cardiac Association, Houston, Texas, United States

^c University of Texas Health Science Center, Houston, Texas, United States

| ARTICLE INFO | A B S T R A C T |
|--|---|
| <i>Keywords:</i> Diet Vegan Cardiovascular diseases Nutrition therapy Complementary therapies | <i>Background:</i> Individuals diagnosed with congestive heart failure (CHF) have a 50% five-year mortality rate and approximately 650,000 new cases of CHF are diagnosed annually. Plant-based diets are known to improve plasma lipid concentrations, reduce blood pressure, and as part of a lifestyle intervention, lead to the regression of atherosclerotic lesions. However, a paucity of data exists with regards to plant-based diets in the treatment of CHF. <i>Methods:</i> Three patients diagnosed with CHF opted to undergo a dietary intervention consisting of a defined plant-based diet as an adjunct to standard medical treatment for CHF. Cardiac magnetic resonance imaging was performed. Patients' consumed the defined plant-based diet for an average of "79 days. <i>Results:</i> Follow-up cardiac magnetic resonance images revealed a 92% increase in ejection fraction [mean \pm standard deviation for all data] (22.0 \pm 6.9% vs 42.2 \pm 18.4%), 21% reduction in left ventricular mass (214 \pm 90 g vs 170 \pm 102 g), 62% increase in stroke volume (55.8 \pm 24.3 cc vs 90.3 \pm 30.6 cc) and a 17% increase in cardiac output (3.6 \pm 1.2 L/min vs 4.2 \pm 1.6 L/min). In patient 1, 90–95% ostial stenosis of the left anterior descending artery nearly completely regressed following the dietary intervention. All patients subjectively reported significant clinical improvements, including less angina, shortness of breath and fatigue. <i>Conclusion:</i> As an adjunct treatment, a defined plant-based diet may contribute to the reversal of cardiac morphological and functional abnormalities in the setting of CHF. |

1. Introduction

Congestive heart failure (CHF) independently increases the risk of mortality by 50% within the first five years of diagnosis.¹ Cardiac remodeling due to increased left ventricular pressure, increased reactive oxygen species (ROS), decreased antioxidant enzymatic activity, and decreased nitric oxide (NO), may also contribute to structural remodeling of the myocardium, promoting the development of CHF.²

Plant-based diets have emerged as effective therapeutic interventions to treat and even reverse coronary atherosclerosis.^{3,4} Both interventional and observational evidence suggests that plant-based diets may decrease the incidence and severity of CHF.⁵ These positive effects may be due to decreased saturated fat and dietary cholesterol intakes, which may reduce serum cholesterol,^{6,7} as well as increased phytonutrient consumption, such as antioxidants, which can reduce oxidative stress and inflammation. Indeed, previous investigations utilizing plantbased diets have demonstrated reduced inflammation, body weight and blood pressure.^{8,9}

Current pharmacological therapies to treat CHF rely on modifying hemodynamics to reduce cardiac work as well as modifying cardiac signaling via neurohormonal means.¹⁰ While these drugs prolong survival and decrease hospitalizations, these therapies have not definitely been shown to improve cardiac function and morphology. Despite compelling evidence suggesting that plant-based diets may be beneficial in the treatment of CHF, it has yet to be demonstrated in the clinical setting.⁵ Presented are a case series of 3 patients with CHF and reduced ejection fraction (EF) who underwent a defined, plant-based dietary intervention to treat CHF without surgical interventions.

2. Methods

2.1. Patient presentations

A 46-year-old female (Patient 1) presented with complaints of mild

* Corresponding author at: Houston Cardiac Association, Houston, Texas, United States.

E-mail addresses: rnajjar1@student.gsu.edu (R.S. Najjar), bjsam05@hotmail.com (B.D. Montgomery).

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Table 1 Baseline characteristics.

| Patient | Patient 1 | Patient 2 | Patient 3 |
|--------------------------|------------------|--|---|
| Gender | Female | Male | Male |
| Ethnicity | African American | African American | African American |
| Age (y) | 46 | 58 | 70 |
| Smoking status | No | Quit 1–5 y | Quit > 20 y |
| Alcohol consumption | Occasional | No | Occasional |
| Diet | Regular | Regular | Regular |
| Exercise | No | No | No |
| BMI (kg/m ²) | 37.9 | 30.1 | 33.7 |
| SBP (mmHg) | 149 | 174 | 128 |
| DBP (mmHg) | 85 | 94 | 84 |
| HR (beats/min) | 74 | 69 | 57 |
| Medical history | None | Hypertension | Hypertension |
| | | Type II diabetes | Hypercholestrolemia |
| | | Kidney disease | Cardiac arrythmia |
| Medications | None | furosemide 40 mg, 1 tablet 2x/day | amiodarone 200 mg, 1 tablet 1x/day |
| | | Tribenzor [olmesartan medoxomil, amlodipine & hydrochlorothiazide] 5 mg-25 mg- 40 mg, 1 tablet 1x/day | furosemide 20 mg, 4 tablets 2x/day |
| | | One-A-Day Men 50 Plus, 1 tablet 1x/day | metoprolol tartrate 50 mg, 0.5 tablets 2x/ day |
| | | | potassium chloride 8 mEq, 4 tablets 1x/da simvastatin 80 mg, 0.5 tables 1x/day |
| | | | |

simvastatin 80 mg, 0.5 tables 1x/day finasteride 5 mg, 1 tablet 1x/day isosorbide dinitrate 40 mg, 1 tablet 3x/day lisinopril 20 mg, 1 tablet 2x/day ferrous sulfate 325 mg, 1 tablet 2x/day

Abbreviations: BMIbody mass index; SBPsystolic blood pressure; DBPdiastolic blood pressure; HRheart rate.

Table 2

Clinical and pharmacological changes.

| | Baseline | Final |
|------------------------------|----------|-------|
| BMI (kg/m ²) | | |
| Patient 1 | 37.9 | 34.2 |
| Patient 2 | 30.1 | 26.1 |
| Patient 3 | 33.7 | 32 |
| SBP (mmHg) | | |
| Patient 1 | 149 | 123 |
| Patient 2 | 174 | 158 |
| Patient 3 | 128 | 124 |
| DBP (mmHg) | | |
| Patient 1 | 85 | 82 |
| Patient 2 | 94 | 92 |
| Patient 3 | 84 | 73 |
| HR (beats/min) | | |
| Patient 1 | 74 | 61 |
| Patient 2 | 69 | 50 |
| Patient 3 | 57 | 64 |
| Ejection Fraction (%) | | |
| Patient 1 | 24.9 | 50 |
| Patient 2 | 27.1 | 55.6 |
| Patient 3 | 14.2 | 21.2 |
| LV Mass (g) | | |
| Patient 1 | 117.4 | 94 |
| Patient 2 | 295.18 | 286 |
| Patient 3 | 231 | 130 |
| Stroke Volume (cc) | | |
| Patient 1 | 46.6 | 100 |
| Patient 2 | 83.5 | 115 |
| Patient 3 | 37.5 | 56.1 |
| Cardiac output (L/min) | | |
| Patient 1 | 4.7 | 4.6 |
| Patient 2 | 4 | 5.6 |
| Patient 3 | 2.2 | 2.4 |
| Perscription medications (n) | | |
| Patient 1 | 0 | 6 |
| Patient 2 | 2 | 2 |
| Patient 3 | 8 | 9 |

Abbreviations: BMI, body mass index; SBP, systolic blood pressure.

chest pain, fatigue, night palpitations and shortness of breath induced by physical activity. She was not taking any medications at the time of her office visit and had no previous medical diagnoses as indicated in Table 1 detailing baseline patient characteristics. A baseline physical examination revealed a normal heart rate, normal first and second heart sounds and normal cardiac amplitude. The heart rhythm was regular and no murmurs, gallops or rubs were identified. An electrocardiogram (EKG) revealed nonspecific ST and T wave abnormalities. Based on the presented symptoms and abnormal EKG findings, cardiac magnetic resonance imaging (MRI) and a coronary angiogram were ordered. Findings from the cardiac MRI revealed a left ventricular (LV) mass of 117 g and an EF of 22.1% (Table 2). The coronary angiogram revealed a 90%-95% ostial left anterior descending coronary artery (LAD) stenosis with diffuse left main disease. The left main coronary artery was notably small compared to the LAD and circumflex arteries. This finding was consistent with the likelihood of diffuse atherosclerosis in the left main coronary artery.

A 58-year-old male (Patient 2) complained of chest pain, shortness of breath, low energy levels and edema of the lower extremities. Patient 2 reported taking furosemide (40 mg), tribenzor [olmesartan medoxomil, amlodipine & hydrochlorothiazide] (5 mg–25 mg-40 mg) and a multivitamin (Table 1). He had been previously diagnosed with hypertension, diabetes, and kidney disease. At the time of his office visit, his heart rate was normal with no abnormal sounds. An echocardiogram was performed which indicated an estimated EF of 20–25%, mild to moderate LV hypertrophy, severe LV dilation, left and right atrial enlargement and mild pulmonary and tricuspid valve regurgitation. A cardiac MRI was ordered which revealed an EF of 27.1%, and an LV mass of 295 g.

Lastly, a 70-year-old male (Patient 3) presented with complaints of shortness of breath at rest, dyspnea with minimal physical exertion, orthopnea, profuse diaphoresis, fatigue and chest pain at rest and with exertion. Patient 3 had been previously diagnosed with hypertension, hypercholesterolemia and cardiac arrhythmia. He had been prescribed 8 different medications (Table 1) to manage these conditions. His EKG showed sinus bradycardia with occasional premature ventricular complexes. The QRS duration on EKG was mildly increased. His echocardiogram estimated his EF to be 20–25% in addition to LV

enlargement, mild-moderate LV hypertrophy, restrictive diastolic dysfunction, moderate left atrial enlargement, thickened aortic and mitral valves as well as mild to moderate mitral and tricuspid valve regurgitation. A cardiac MRI also revealed an LV mass of 231 g and an EF of 14.2%.

2.2. Intervention

Each patient was prescribed a defined, plant-based dietary intervention (DPBD) within levels 0-4b in The Food Classification System described elsewhere.⁶ The composition of the DPBD consisted of raw fruits, vegetables, avocado, seeds, with small amounts of raw oats and buckwheat. Patients were advised to eliminate the consumption of all animal products, cooked foods, free oils, soda, alcohol, and coffee.

Patient 1 was prescribed nebivolol 5 mg (1 tablet once per day), valsartan 160 mg (1 tablet once per day). ranolazine 500 mg (1 tablet twice per day), rosuvastatin 10 mg (1 tablet once per day), clopidogrel bisulfate 75 mg (1 tablet once per day) and diazepam 5 mg (1 tablet twice per day). Patient 2 was prescribed nebivolol 5 mg (1 tablet once per day) in place of furosemide. Patient 3 was prescribed spironolactone 25 mg (tablet once per day) and remained on his current medications.

3. Results

The DPBD was followed for 53, 88 and 95 days by Patient 1, Patient 2 and Patient 3, respectively. Each patient was mostly compliant with the nutritional intervention without adverse reactions. Overall, morphological and functional parameters of the heart improved for all 3 patients (Fig. 1). Patient 1 reported having more energy and less chest discomfort within 2 weeks of the intervention, and greater exercise tolerance within 4 weeks with full compliance. Her body mass index, blood pressure and heart rate (Table 2) dramatically improved. Her EF improved by 100%, cardiac stroke volume improved by 115%, LV mass

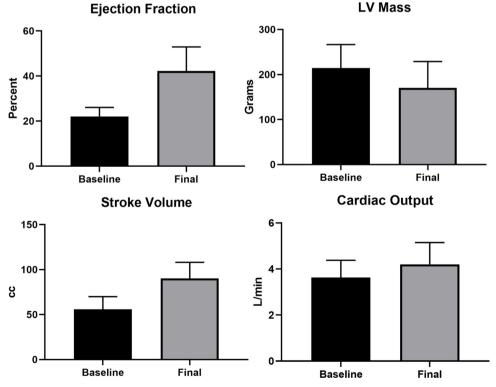
decreased by 20% and her cardiac output was relatively unchanged. Cardiac MRI footage for Patient 1 demonstrated a clear visual improvement in LV function (Supplementary video file 1).

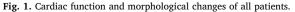
Additionally, stenosis of the ostial LAD coronary artery nearly completely regressed after initiating the DPBD (Fig. 2). Patient 2 reported feeling better within 4 weeks of the DPBD. He was mostly compliant and did not require any significant medication changes during his course of treatment. He experienced complete resolution of his symptoms within 5 weeks, including resolution of angina and shortness of breath. His EF improved by 105%, LV mass regressed by 3%, stroke volume improved by 38%, and cardiac output improved by 40%.

Patient 3 experienced a more complex clinical course. He was admitted to the hospital with decompensated heart failure 4 days after his initial evaluation in our clinic prior to starting the dietary intervention. He started the nutritional intervention during this hospitalization. He had decreased shortness of breath and chest discomfort and continued to have subjective improvements until 6 weeks after initiation of his dietary treatment; he suffered a clinical stroke with resolution of his symptoms in 48 h and a subsequent transient ischemic attack 2 days later. His follow-up cardiac MRI was performed during this hospitalization. In addition to having sustained improvement in his heart failure symptoms, he was found to have a 50% improvement in EF, a 44% regression in LV mass, a 19% improvement in cardiac stroke volume, and a 9% improvement in his cardiac output.

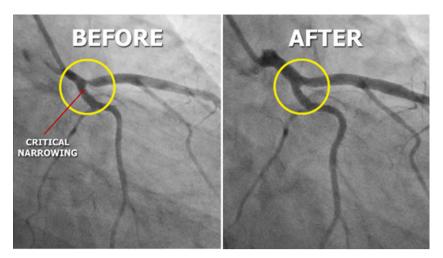
4. Discussion

This dietary intervention has previously been shown to significantly reduce blood pressure, heart rate and systemic inflammation.^{8,9} These hemodynamic and biochemical changes suggest a possible mechanisms by which the DPBD improves cardiac function. There was a significant reduction in LV mass observed in each subject, including a 101 g regression seen in Patient 3. This large reduction in LV mass could be due





Legend: Mean cardiac function and morphology of all 3 patients at baseline and final as determined by cardiac magnetic resonance imaging. Error bars are standard error of the mean.



to a reduction in intramyocardial edema, possibly due to reduced leukocyte infiltration causing a decrease in reactive oxygen and nitrogen species which may have ameliorated the degradation of the extracellular matrix and decreased collagen deposition.¹¹

In general, the consumption of animal based foods are associated with increased oxidative stress and inflammation in humans, while plant-based foods have an inverse association.¹² These positive redox effects associated with consuming plants could result in the higher bioavailability of nitric oxide, resulting in vasodilation and a reduction in blood pressure likely due to reduced systemic vascular resistance (SVR).¹³ With a reduction in SVR, stroke volume would increase, improving cardiac output and possibly reducing heart rate. None of these clinical improvements would be expected to occur with the standard medical treatments for CHF. Multicenter drug trials have not definitely shown improvements in EF, nor have these investigations demonstrated changes in physiological function of the heart to the extent that was examined here.^{10,14} Hence, the DPBD resulted in both stabilization and partial reversal of advanced cardiovascular disease across a broad age spectrum of patients with differing clinical courses. Indeed, previous investigations have demonstrated that a plant-based diet can reverse coronary atherosclerosis, however, a paucity of data exists with regards to plant-based diets in the treatment of CHF.3,

5. Conclusion

In the standard treatment of CHF, such dramatic and rapid improvements in heart morphology and function would be deemed highly improbable. However, the findings in this case series demonstrate that a plant-based diet as an adjunct to standard medical therapies may reverse certain pathophysiologic processes in heart failure. This intervention provides an outline for a potential novel therapy for heart failure with reduced ejection fraction. A larger case series or a prospective clinical trial utilizing this plant-based dietary intervention is needed to confirm these findings.

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Fig. 2. Coronary angiogram changes for Patient 1. Legend: Baseline coronary angiogram (left) showing diffusely small left main coronary artery with a 90–95% ostial stenosis of the left anterior descending artery. Follow up angiogram (right) showing increased luminal size of left main coronary artery with a near-total regression of ostial left anterior descending artery lesion.

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Appendix A. Supplementary data

Supplementary material related to this article can be found, in the online version, at doi:https://doi.org/10.1016/j.ctim.2019.06.010.

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CLINICAL INVESTIGATIONS



PREVENTION WILEY CLINICAL

A defined, plant-based diet utilized in an outpatient cardiovascular clinic effectively treats hypercholesterolemia and hypertension and reduces medications

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Background: Cardiovascular disease (CVD) is a major economic burden in the United States. CVD risk factors, particularly hypertension and hypercholesterolemia, are typically treated with drug therapy. Five-year efficacy of such drugs to prevent CVD is estimated to be 5%. Plantbased diets have emerged as effective mitigators of these risk factors.

Hypothesis: The implementation of a defined, plant-based diet for 4 weeks in an outpatient clinical setting may mitigate CVD risk factors and reduce patient drug burden.

Methods: Participants consumed a plant-based diet consisting of foods prepared in a defined method in accordance with a food-classification system. Participants consumed raw fruits, vegetables, seeds, and avocado. All animal products were excluded from the diet. Participant anthropometric and hemodynamic data were obtained weekly for 4 weeks. Laboratory biomarkers were collected at baseline and at 4 weeks. Medication needs were assessed weekly. Data were analyzed using paired-samples t tests and 1-way repeated-measures ANOVA.

Results: Significant reductions were observed for systolic (-16.6 mmHg) and diastolic (-9.1 mmHg) blood pressure (P < 0.0005), serum lipids (P \leq 0.008), and total medication usage (P < 0.0005). Other CVD risk factors, including weight (P < 0.0005), waist circumference (P < 0.0005), heart rate (P = 0.018), insulin (P < 0.0005), glycated hemoglobin (P = 0.002), and high-sensitivity C-reactive protein (P = 0.001) were also reduced.

Conclusion: A defined, plant-based diet can be used as an effective therapeutic strategy in the clinical setting to mitigate cardiovascular risk factors and reduce patient drug burden.

KEYWORDS

Biomarkers, General Clinical Cardiology/Adult, Hypertension, Preventive Cardiology, Vegetarian Diet

1 | INTRODUCTION

Cardiovascular disease (CVD) is a major economic burden to the United States. Currently, 17% of all healthcare expenditures go toward CVD care.¹ Projections are expected to rise, as 40.5% of the US population may have some form of CVD by 2030, leading to a near tripling in medical care costs, from \$273 billion to \$818 billion. CVD has been the leading cause of death in the United States since 1950.² The standard of clinical care in the primary prevention of CVD is to reduce CVD risk factors, particularly through lipid-lowering and antihypertensive drug therapy.³ It has been estimated that nearly

40% of the population has high serum low-density lipoprotein cholesterol (LDL-C).⁴ In addition, approximately one-third of individuals age 40 to 59 years are estimated to be hypertensive.⁵ Of those with hypertension (HTN), 76% are on medications to reduce blood pressure, but only 52% achieve blood-pressure control. The highest drug prices in the world are found within the United States. On average, per capita spending on prescription drugs in the United States is \$858, compared with an average of \$400 in 19 other industrialized countries.6

Patients' opinion of the efficacy of drug therapy in CVD prevention is often inflated multifold.^{7,8} It has been estimated that high-risk patients have a < 5% chance of benefiting from cardioprotective drugs within the next 5 years. Moreover, most patients wish to take drugs at a benefit threshold of \geq 20% over 5 years.⁹ Thus, if patients were aware of the actual benefit of cardioprotective drugs, many patients may not be willing to take such medications.

Based on growing evidence,¹⁰⁻¹⁵ it has been recommended that physicians encourage patients to consume plant-based diets.¹⁶ The aim of this investigation was to evaluate the effectiveness of a defined, plant-based diet as an adjunct to or replacement of prescription drugs in the treatment of hypercholesterolemia and HTN in an outpatient clinical setting.

2 | METHODS

2.1 | Study population

All subjects were registered new patients of a cardiovascular center. The study intervention was carried out in an outpatient clinical setting. All participants provided written informed consent after the study protocol and procedure had been fully explained. The study was approved by the Texas Woman's University Institutional Review Board.

Baseline characteristics of the patients are shown in Table 1. All participants were age 32 to 69 years with HTN, elevated LDL-C, and excess body weight. HTN was defined as systolic blood pressure (SBP) ≥140 mmHg or diastolic blood pressure (DBP) ≥90 mmHg. Elevated LDL-C was considered to be a serum LDL-C concentration ≥ 100 mg/dL, and excess body weight was defined as a body mass index ≥25 kg/m².

Exclusion criteria included current tobacco use, current drug abuse, excessive alcohol use (defined as >2 glasses of wine or alcohol equivalent per day for men or >1 glass of wine or alcohol equivalent for women), a current cancer diagnosis, an ongoing clinically defined infection, a mental disability that would prevent the participant from following the study protocol, an estimated glomerular filtration rate < 60 mg/dL, current pregnancy or lactation, a hospitalization within the past 6 months, and previous exposure to the nutrition program.

2.2 | Screening

Eligibility was determined through initial screening of participants who expressed interest in participating in the intervention. Demographics, lifestyle habits, anthropometrics, and hemodynamics were used to determine the eligibility of participation for each subject. A trained medical assistant measured blood pressure, heart rate, and body weight. Medical history and lifestyle habits were obtained by the medical assistant and/or nurse practitioner. Fasting blood was collected by a licensed phlebotomist. The clinical care of all patients was overseen by a board-certified cardiologist.

2.3 | Weekly visits

After subjects were screened for study inclusion, follow-up appointments were arranged for study enrollment. Participants were instructed to attend 4 follow-up weekly office visits in addition to a baseline assessment. Baseline weight, blood pressure, heart rate,
 TABLE 1
 Baseline patient demographics and clinical diagnoses

| | Participants, n = 31 |
|---|----------------------|
| Maan aga y | - |
| Mean age, y | 53.4 (32–69) |
| Sex | 10 (22) |
| M | 10 (33) |
| F | 21 (67) |
| Race/ethnicity | 05 (00) |
| African American | 25 (80) |
| Hispanic | 3 (10) |
| White | 3 (10) |
| BMI, kg/m ² | 37.5 ± 8.3 |
| 25-29.9 (overweight) | 6 (19) |
| 30-34.9 (obese class 1) | 6 (19) |
| 35-39.9 (obese class 2) | 10 (33) |
| ≥40 | 9 (29) |
| Current diagnoses | |
| CAD | 10 (33) |
| T2DM | 8 (27) |
| Arthritis | 7 (23) |
| Prediabetes | 5 (17) |
| Medications, n | |
| BP medications, total | 49 |
| ACEI | 5 |
| ARB | 11 |
| Central antiadrenergic | 1 |
| Cardioselective (\(\beta1\)-blocker | 6 |
| Noncardioselective (β 1)-blocker | 2 |
| ССВ | 9 |
| Potassium-sparing diuretic | 1 |
| Thiazide diuretic | 14 |
| Other prescription drugs, total | 33 |
| Biguanide | 2 |
| Sulfonylurea | 3 |
| Dipeptidylpeptidase-4 inhibitor | 1 |
| Insulin | 2 |
| NSAID | 1 |
| Biologic immune suppressant | 1 |
| Statin | 2 |
| Bronchodilator/steroid inhaler | 5 |
| Thyroid drugs | 3 |
| Xanthine oxidase inhibitor | 2 |
| PPI | 1 |
| Antiplatelet | 1 |
| Antianginal | 2 |
| Digitalis glycoside | 1 |
| Vasodilator | 1 |
| Other | 5 |
| Total medications | 82 |
| | |

Abbreviations: ACEI, angiotensin-converting enzyme inhibitor; ARB, angiotensin II receptor blocker; BMI, body mass index; BP, blood pressure; CAD, coronary artery disease; CCB, calcium channel blocker; F, female; M, male; NSAID, nonsteroidal anti-inflammatory drug; PPI, proton pump inhibitor; SD, standard deviation; T2DM, type 2 diabetes mellitus. Unless otherwise noted, data are presented as n (%) or mean \pm SD (range).

waist circumference, medications, and biochemical indicators were documented. A baseline 24-hour dietary recall was conducted by a trained nutritionist with the utilization of food models to verify portion sizes of foods and beverages consumed. Nutrient intake was analyzed by the Nutrition Data System for Research software, version 2016 (University of Minnesota, Minneapolis).

Follow-up visits (weeks 1–4) consisted of obtaining weight, blood pressure, heart rate, and waist circumference. Medications were assessed and adjusted as needed by the medical doctor or nurse practitioner during the follow-up visits. The final visit (week 4) consisted of a second 24-hour dietary recall and a second collection of fasting blood to assess biochemical measures.

2.4 | Medications

Medications were documented following the conclusion of each office visit. All medications listed at baseline were chronic stable medications (>3 months), except for medications changed during the baseline office visit as outlined in the protocol below. All other medication changes were documented in the medication tracking of this study. No lipid-lowering medications were added at the onset or during the study. The medication needs-assessment protocol is as follows:

 Baseline: All nonessential medications and supplements were discontinued. Additionally, diuretics were discontinued in patients who were clinically euvolemic. Insulin, sulfonylureas, and other potential glucose-lowering medications were either removed or the dosage was decreased in patients whose glucose levels were routinely below 250 mg/dL. All baseline medications are indicated in Table 1.

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- Week 1 follow-up: If a patient's blood pressure was low and the patient had symptoms of dizziness or fatigue associated with low blood pressure, then blood pressure medications were decreased by 25% to 50%. Other medications were reviewed with consideration of removal based on patient needs (eg, hypoglycemics).
- Week 2 follow-up: The patients' clinical response to the diet was reevaluated and medication adjustments were made according to their clinical response. Medications primarily prescribed for symptom management were assessed (eg, sleep, allergies, mood disorders, pain) and discontinued if necessary.
- Weeks 3 and 4 follow-up: Based on the patients' clinical response to the dietary intervention, changes were made to the medications as needed for the remainder of the intervention.

2.5 | Dietary protocol

Participants were instructed to follow a defined plant-based dietary intervention for 4 weeks. A food classification system using a scale of 0 to 10 was devised to create a simple, reproducible way of prescribing a nutritional regimen to patients in the clinical setting (Table 2). Participants were instructed to consume foods within this food classification system. Food levels 0 through 4B were permitted, whereas all other food levels were excluded. Briefly, food levels 0 through 4B exclude all animal products, with the exception of honey. Cooked foods, free oils, soda, alcohol, and coffee were also excluded. Emphasized were raw fruits and vegetables, with avocado and raw seeds

| Food Level | Description |
|------------|--|
| 0 | Liquids including water, tea, unpasteurized fruit and vegetable juices, and blended fruit and vegetable smoothies. These foods would be consumed raw, except for tea, which can be steeped in hot water. |
| 1 | Raw fruits and vegetables with a low glycemic index (<56) |
| 2 | Raw fruits and vegetables with a medium to low GI (56–70) |
| 3 | Raw fruits and vegetables with a high GI (>70) |
| 4A | Plant foods that are raw with a high fat content (≥20% of caloric content from fat), such as raw seeds and avocados |
| 4B | Plant foods that are dehydrated to temperatures ≤160°F |
| 4C | Plant foods that are dried, dehydrated, or warmed (dry-heat cooking) at 160°F–175°F, or steamed or boiled for a short duration (steaming, <4 min; boiling, <10 min). Includes lightly steamed, soaked, sprouted, dehydrated, or warmed fruits, vegetables, legumes or beans, and grains. Heated foods with >20% of calories from fat are excluded. |
| 5 | Foods that are warmed, dried, or dehydrated at 175°F to 200°F, and steamed or boiled for a medium duration (steaming, 4–10 min; boiling, 10–45 min). Typical foods include greens, beans and legumes, and starches, including grains, bean or mixed- vegetable soups, and other fruit and vegetables boiled for up to 45 min or oven-warmed (at 155°F–200°F). Heated foods with >20% of calories from fat are excluded. |
| 6 | Foods that are baked, warmed, dried, or dehydrated at >200°F, or steamed or boiled for a long duration (steaming, >10 min; boiling, >45 min). Heated foods with >20% of calories from fat are excluded. |
| 7 | Fish with low mercury content lightly steamed or poached for ≤8 min. Processed plant foods with preservatives or additives, free oils, and heated foods with >20% calories from fat are included. |
| 8 | Same as level 7, except also includes wild-game meats, low-mercury fish lightly steamed or poached for >8 min, and plant-based foods that are grilled or heavily processed. May also include carbohydrates with white flour or white rice, or natural foods that have been stripped of their natural components. |
| 9 | Animal-based foods that include domestically raised animals (excluding beef and pork) and plant-based foods that are sautéed, stir-fried, medium-fried or deep-fried in oil. Other animal-based foods include all other types of fish. May also include foods containing dairy products. |
| 10 | All other types of animal-based foods, and plant-based foods prepared in any way. May include processed foods of any kind. |

 TABLE 2
 The food classification system

Abbreviations: F, Fahrenheit; GI, glycemic index. Food classification levels 0 through 4B were permitted for consumption during the dietary intervention; levels >4B were excluded from the intervention. Sodium consumption was low, although the food provided to patients contained small amounts of sea salt.

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provided as condiments. All meals and snacks were provided at no cost to the participants for the full duration of the 4-week intervention. Vitamin, herbal, and mineral supplements were to be discontinued unless otherwise clinically indicated. Participants were not advised to alter their exercise habits, nor were exercise habits monitored

Participants were free to consume foods outside of what was provided, as long as the foods fell within food levels 0 through 4B. No caloric targets were prescribed, nor were any macronutrient restrictions advocated; participants were free to consume food ad libitum. Participants were also instructed to track dietary adherence with a daily adherence-assessment tool. Participants indicated in writing each day whether they were "100% on the diet" or "ate anything off of the diet." The number "1" was assigned to an adherent day, and "0" was assigned to a nonadherent day. Scores after 4 weeks could therefore range from 0 to 28 points for each participant. Evaluation of the adherence-assessment tool was conducted during each weekly follow-up visit by a trained nutritionist.

2.6 **Biochemical measures**

After a 12-hour fast during the baseline and final office visits, the following serum biomarkers were obtained: total cholesterol, LDL-C, high-density lipoprotein cholesterol, triglycerides, insulin, glucose, glycated hemoglobin (HbA_{1c}), and high-sensitivity C-reactive protein (hs-CRP). These specific biomarkers of interest were analyzed by either True Health Diagnostics (Frisco, TX) or Singulex (Alameda, CA), depending on the subject's insurance. The same company that analyzed the baseline laboratory tests for a participant was used for the follow-up testing to ensure assay consistency.

Serum lipids were measured by enzymatic colorimetric assay, and insulin was measured by a no-competitive sandwich-type enzymelinked immunosorbent assay with electrochemical detection for both True Health Diagnostics and Singulex. Glucose was measured by an enzymatic reference method with hexokinase using colorimetric detection, and hs-CRP was measured by a particle-enhanced immunoturbidometric assay for both Singulex and True Health Diagnostics. HbA_{1c} was measured by a turbidometric inhibition immunoassay for Singulex. Boronate affinity chromatography was used by True Health Diagnostics for HbA_{1c}.

2.7 Statistical analysis

Paired-samples t tests were used for the analysis of biochemical and nutrient intake means. A one-way repeated-measures ANOVA was used to analyze the means for anthropometric, hemodynamic, and medication data. Significance was set at a P value of < 0.05. SPSS version 24 (IBM Corp., Armonk, NY) was used for data analysis.

3 RESULTS

3.1 | Demographics

During screening, a total of 65 patients showed interest in participating in the study; however, 30 patients did not meet inclusion

criteria or were excluded. Two individuals were unable to participate due to scheduling conflicts. Although 33 participants initially enrolled into the study, 2 participants were either lost to contact (n = 1) or no longer wished to follow the dietary protocol (n = 1). One participant refused to complete the final 24-hour dietary recall during week 4 due to time availability. Thus, a total of 31 participants provided clinical data and 30 participants provided nutrient intake data

Based on clinical diagnoses and medical history, 33% of participants had coronary artery disease and 44% were either prediabetic (HbA_{1c} 5.7%-6.4%) or had diabetes mellitus (HbA_{1c} \ge 6.5%; (Table 1). The average body mass index was 37.5 kg/m² \pm 8.3 kg/m², and approximately 81% of the participants were obese.

3.2 | Nutrient intake

Nutrient intake of participants on the defined, plant-based diet significantly changed after 4 weeks (Table 3). Significant reductions in energy intake, saturated fat as a percent of energy, dietary cholesterol, protein as a percent of energy, total fat, monounsaturated and polyunsaturated fat as a percent of energy, trans fat, vitamin D, vitamin B12, calcium, zinc, and sodium were observed after 4 weeks. Carbohydrate intake as a percent of energy, vitamin A, vitamin C, folate, dietary fiber, magnesium, and potassium intake increased significantly after 4 weeks. Patients anecdotally reported overall satisfaction with the food provided during the clinical follow-ups, and no significant symptoms of increased hunger were reported.

3.3 | Clinical variables

Anthropometric and hemodynamic characteristics, as well as medications, changed significantly ($P \le 0.018$) from baseline to 4 weeks (Table 4). Adherence was well maintained over the 4-week period. Overall, participants were noncompliant for 3.6 out of 28 days. There were no significant differences between subjects with 100% adherence and lower-adherent subjects. Participants lost on average a total of 6.7 kg (14.7 lbs.) after 4 weeks on the defined plant-based diet (Table 4). SBP and DBP decreased by 16.6 mmHg and 9.1 mmHg, respectively. The reduction in blood pressure was accompanied with a decreased use of blood pressure medications (decreased 33% by week 4). Additionally, those taking hypoglycemic drugs, including insulin, reduced medication usage by 87%. Overall, total medication usage decreased 40% by week 4.

3.4 | **Biomarkers**

All biochemical changes were significant ($P \le 0.037$) at 4 weeks compared with baseline, with the exception of the total cholesterol to high-density lipoprotein cholesterol ratio (P = 0.068) and glucose (P = 0.25; Table 5). Although fasting glucose was not significantly reduced, HbA_{1c} was significantly reduced (P = 0.002).

The distribution of high-interest clinical variable changes during the intervention are displayed in Supporting Information, Figure, in the online version of this article.

TABLE 3 Nutrient intake^b



| | Baseline | Final | Change, % ^a | P Value ^c |
|----------------------------------|-------------------------------------|-------------------------------------|---------------------------|----------------------|
| Energy, Kcal | $\textbf{2053} \pm \textbf{873}$ | 1369 ± 488 | -33 (-683 ± 808) | <0.0005 |
| Fat, % of energy | $\textbf{36.4} \pm \textbf{10.4}$ | $\textbf{19.0} \pm \textbf{8.9}$ | -48 (-17.3 \pm 12.8) | <0.0005 |
| Saturated fat, % of energy | $\textbf{11.6} \pm \textbf{4.5}$ | $\textbf{3.8} \pm \textbf{2.7}$ | -67 (-7.7 ± 5.5) | <0.0005 |
| Monounsaturated fat, % of energy | 13.2 ± 4.8 | $\textbf{7.0} \pm \textbf{3.9}$ | -47 (-6.2 ± 5.4) | <0.0005 |
| Polyunsaturated fat, % of energy | $\textbf{8.4} \pm \textbf{5.6}$ | $\textbf{5.4} \pm \textbf{2.7}$ | -36 (-3.0 \pm 3.5) | <0.0005 |
| Omega-6, g | $\textbf{18.5} \pm \textbf{11.1}$ | $\textbf{6.0} \pm \textbf{4.7}$ | -67 (-12.4 \pm 10.6) | <0.0005 |
| Omega-3, g | $\textbf{2.11} \pm \textbf{1.60}$ | $\textbf{2.14} \pm \textbf{1.95}$ | 1 (0.03 \pm 2.16) | 0.92 |
| Omega-6/omega-3 ^d | $\textbf{9.8}\pm\textbf{3.7}$ | $\textbf{4.3}\pm\textbf{3.0}$ | -56 (-5.5 ± 3.8) | <0.0005 |
| Trans fat, g | $\textbf{2.25} \pm \textbf{1.97}$ | $\textbf{0.04} \pm \textbf{0.09}$ | -99 (-2.21 \pm 2.00) | <0.0005 |
| Cholesterol, mg | $\textbf{295.4} \pm \textbf{211.7}$ | $\textbf{12.2} \pm \textbf{56.2}$ | -96 (-283.2 \pm 214.8) | <0.0005 |
| Carbohydrate, % of energy | $\textbf{46.3} \pm \textbf{14.0}$ | $\textbf{72.6} \pm \textbf{11.3}$ | 57 (26.3 ± 17.0) | <0.0005 |
| Protein, % of energy | $\textbf{16.5} \pm \textbf{6.4}$ | $\textbf{7.5} \pm \textbf{2.1}$ | -54% (-9.0 \pm 6.1) | <0.0005 |
| Total fiber, g | $\textbf{20.4} \pm \textbf{11.9}$ | $\textbf{51.0} \pm \textbf{17.7}$ | 150 (30.6 \pm 17.8) | <0.0005 |
| Total vitamin A activity, IU | 8265 ± 9258 | $\textbf{33387} \pm \textbf{19052}$ | 303 (25 121 \pm 21 876) | <0.0005 |
| Vitamin D, IU | $\textbf{159.1} \pm \textbf{154.3}$ | 12.3 ± 30.4 | -92 (-146.8 \pm 161.8) | <0.0005 |
| Vitamin E, mg | $\textbf{9.9}\pm\textbf{6.3}$ | 10.5 ± 5.6 | 6 (0.6 ± 6.4) | 0.60 |
| Vitamin C, mg | $\textbf{87.7} \pm \textbf{108.8}$ | $\textbf{412.7} \pm \textbf{164.7}$ | 370 (325.0 \pm 197.3) | <0.0005 |
| Vitamin B12, µg | $\textbf{4.0} \pm \textbf{1.9}$ | $\textbf{0.3}\pm\textbf{0.8}$ | -92 (-3.6 ± 2.3) | <0.0005 |
| Folate, µg | 298 ± 229 | $\textbf{741} \pm \textbf{298}$ | 115 (343 \pm 329) | <0.0005 |
| Iron, mg | $\textbf{15.4} \pm \textbf{7.2}$ | $\textbf{15.3} \pm \textbf{6.9}$ | -1 (-0.1 ± 9.9) | 0.97 |
| Calcium, mg | $\textbf{796} \pm \textbf{438}$ | 566 ± 279 | –29 (–229 \pm 527) | 0.024 |
| Sodium, mg | $\textbf{3730} \pm \textbf{1783}$ | 839 ± 778 | -76 (-2891 \pm 1776) | <0.0005 |
| Magnesium, mg | $\textbf{288.1} \pm \textbf{119.9}$ | $\textbf{488.1} \pm \textbf{186.0}$ | 69 (200.0 \pm 178.0) | <0.0005 |
| Zinc, mg | $\textbf{12.2} \pm \textbf{5.9}$ | $\textbf{7.8} \pm \textbf{3.4}$ | -76 (-4.4 ± 7.0) | 0.002 |
| Potassium, mg | $\textbf{2668} \pm \textbf{1190}$ | 5078 ± 1758 | 90 (2410 \pm 1764) | <0.0005 |

Data are presented as mean \pm standard deviation unless otherwise indicated.

 $^{\rm a}$ Data are presented as percent change (mean \pm standard deviation).

^b Data are for subjects who completed 24-hour recalls at both baseline and 4 weeks and do not include dietary supplements (n = 30).

^c Paired samples *t* tests for within-group comparisons of changes from baseline to final values.

^d Values indicate a ratio.

TABLE 4 Change of anthropometrics, hemodynamics, medications, and adherence over 4 weeks

| | Baseline | Week 1 | Week 2 | Week 3 | Week 4 | P Value ^a |
|------------------------------|-----------------------------------|---------------------------------------|-----------------------------------|-----------------------------------|-----------------------------------|----------------------|
| Weight, kg, mean \pm SE | 108.1 ± 5.1 | 105.4 ± 4.8^{b} | 103.9 ± 4.8^{b} | 102.6 ± 4.7^{b} | 101.4 ± 4.7^{b} | <0.0005 |
| BMI, kg/m ² | $\textbf{37.5} \pm \textbf{1.4}$ | $\textbf{36.5} \pm \textbf{1.4}^{b}$ | $36.0\pm1.4^{\text{b}}$ | $35.6 \pm \mathbf{1.4^{b}}$ | $35.2 \pm \mathbf{1.4^{b}}$ | <0.0005 |
| WC, cm | $\textbf{111.9} \pm \textbf{2.5}$ | 109.2 ± 2.5^{b} | 107.6 ± 2.5^{b} | 106.3 ± 2.5^{c} | 105.3 ± 2.5^{b} | <0.0005 |
| SBP, mm Hg | 146.6 ± 2.8 | $\textbf{131.9} \pm \textbf{2.8}^{b}$ | 127.0 ± 2.4 | 129.5 ± 1.9 | 130.0 ± 2.3 | <0.0005 |
| DBP, mm Hg | $\textbf{91.2} \pm \textbf{1.3}$ | 81.5 ± 1.4^{b} | $\textbf{79.0} \pm \textbf{1.3}$ | $\textbf{82.1} \pm \textbf{1.2}$ | $\textbf{82.1} \pm \textbf{1.2}$ | <0.0005 |
| BP medications | $\textbf{1.6} \pm \textbf{1.1}$ | $\textbf{1.6} \pm \textbf{1.0}$ | 1.4 ± 1.0^{d} | 1.1 ± 1.0^{d} | $\textbf{1.0} \pm \textbf{0.1}$ | <0.0005 |
| Heart rate, bpm | $\textbf{69.8} \pm \textbf{1.8}$ | $\textbf{71.8} \pm \textbf{1.9}$ | 68.4 ± 1.7 | 68.1 ± 1.7 | $\textbf{66.2} \pm \textbf{1.2}$ | 0.018 |
| Other prescription drugs | $\textbf{1.0} \pm \textbf{1.4}$ | $\textbf{1.0} \pm \textbf{1.4}$ | $\textbf{0.9} \pm \textbf{1.5}$ | $\textbf{0.6}\pm\textbf{0.9}$ | $\textbf{0.5}\pm\textbf{0.9}$ | 0.008 |
| Total medications | $\textbf{2.6} \pm \textbf{2.0}$ | $\textbf{2.7} \pm \textbf{2.0}$ | 2.3 ± 2.0^{d} | $\textbf{1.8} \pm \textbf{1.6}$ | $\textbf{1.6} \pm \textbf{1.3}$ | <0.0005 |
| Adherence, d/wk ^e | - | $\textbf{6.32} \pm \textbf{0.19}$ | $\textbf{6.03} \pm \textbf{0.25}$ | $\textbf{6.06} \pm \textbf{0.27}$ | $\textbf{5.96} \pm \textbf{0.27}$ | 0.531 |

Abbreviations: ANOVA, analysis of variance; BMI, body mass index; BP, blood pressure; DBP, diastolic blood pressure; SBP, systolic blood pressure; SE, standard error; WC, waist circumference.

^a Repeated-measures 1-way ANOVA with a Greenhouse–Geisser correction due to violation of Mauchly's test of sphericity (P > 0.05).

^b $P \leq 0.001$ compared with previous week.

^c $P \leq 0.01$ compared with previous week.

^d P ≤ 0.05 compared with previous week (all pairwise comparisons were determined by post hoc analysis with a Bonferroni adjustment).

^e Measured by weekly adherence-assessment tool. Values represent the number of days on average that adherence was 100% out of 1 week (7 days).

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4 | DISCUSSION

Four weeks of a defined, plant-based dietary intervention resulted in clinically significant reductions in SBP, DBP, blood pressure medication usage, total medication usage, and serum lipids. Statistically significant reductions were also observed for other CVD risk factors, including body weight, heart rate, waist circumference, insulin, HbA_{1c}, and hs-CRP. This intervention demonstrated that a plant-based diet can be used effectively in the clinical setting with profound results. Additionally, subjects were able to transition from a standard American diet to the plant-based diet outlined in this intervention with good adherence. Physician advice can significantly impact the dietary choices of patients,¹⁷ as demonstrated in this trial.

Although weight was reduced, this likely did not contribute fully to the reduction in blood pressure. A recent Cochrane review of randomized trials lasting ≥24 weeks examined the effects of weight loss on blood pressure and concluded that a 4-kg reduction in weight resulted in a 4.5-mmHg and 3-mmHg reduction in SBP and DBP, respectively.¹⁸ Results from this review would underestimate expected outcomes of this trial. In comparison, participants in the present study lost 6.7 kg and reduced SBP and DBP by 16.6 mmHg and 9.1 mmHg, respectively. These findings are striking considering that blood pressure medications were reduced by 33% by week 4 and blood pressure nearly normalized. Participants' blood pressure was better even when discontinuing medications, which may indicate superiority of the dietary intervention over drug therapy. The reduction in blood pressure by this nutritional intervention was due to a variety of contributing factors, which may include a reduction in hs-CRP $(-2.4 \pm 3.7 \text{ mg/L})^{19}$ and increased consumption of nitrates,²⁰ potassium,²¹ and magnesium.²² Increased dietary fiber,²³ phytosterols.²⁴ and polyphenols²⁵ also likely contributed to reduced serum lipids in addition to the exclusion of animal-based foods.²⁶

It is interesting to note that fasting blood glucose was not significantly reduced (P = 0.25), yet HbA_{1c} was significantly reduced (P = 0.002). It is likely that reduced postprandial glucose fluctuations accounted for this decrease in HbA_{1c}, although this was not directly

TABLE 5 Change in biochemical variables after 4 weeks

| | Baseline | Final | Change | P Value ^a |
|-----------------------|------------------------------------|------------------------------------|------------------------------------|----------------------|
| TC, mg/dL | $\textbf{216.6} \pm \textbf{34.2}$ | $\textbf{182.7} \pm \textbf{29.9}$ | -33.8 ± 25.9 | <0.0005 |
| LDL-C, mg/dL | 143.0 ± 28.9 | 118.4 ± 26.4 | $\textbf{-24.6} \pm \textbf{21.3}$ | <0.0005 |
| HDL-C, mg/dL | $\textbf{54.8} \pm \textbf{9.4}$ | $\textbf{49.5} \pm \textbf{10.6}$ | $\textbf{-5.2}\pm\textbf{6.2}$ | < 0.0005 |
| TC/HDL ^b | $\textbf{4.04} \pm \textbf{0.88}$ | $\textbf{3.81} \pm \textbf{0.88}$ | $\textbf{-0.22}\pm0.64$ | 0.068 |
| TG, mg/dL | $\textbf{124.1} \pm \textbf{58.1}$ | 104.5 ± 53.6 | $\textbf{-19.6} \pm \textbf{38.4}$ | 0.008 |
| Insulin, uIU/mL | 14.6 ± 7.6 | 10.3 ± 7.6 | -4.2 ± 5.1 | <0.0005 |
| Glucose, mg/dL | $\textbf{90.1} \pm \textbf{12.0}$ | $\textbf{87.1} \pm \textbf{4.7}$ | $\textbf{-2.9} \pm \textbf{14.0}$ | 0.25 |
| HbA _{1c} , % | $\textbf{5.9} \pm \textbf{0.5}$ | $\textbf{5.7} \pm \textbf{0.3}$ | -0.2 \pm 0.3 | 0.002 |
| hs-CRP, mg/L | $\textbf{7.8} \pm \textbf{6.4}$ | $\textbf{5.3} \pm \textbf{4.7}$ | -2.4 ± 3.7 | 0.001 |

Abbreviations: HbA_{1c}, glycated hemoglobin; HDL-C, high-density lipoprotein cholesterol; hs-CRP, high-sensitivity C-reactive protein; LDL-C, low-density lipoprotein cholesterol; SD, standard deviation; TC, total cholesterol; TG, triglycerides. Data are presented as mean \pm SD; n = 31.

- ^a Paired-samples *t* tests for within-group comparisons of changes from baseline to final values.
- ^b Values indicate a ratio.

tested. It has been previously demonstrated that HbA_{1c} < 7% is mostly influenced by postprandial glucose.²⁷ The average HbA_{1c} of this sample was 5.9%; therefore, postprandial blood glucose would likely play a more significant role.

Other similar plant-based dietary trials have also demonstrated reduced CVD risk factors. In a 4-week randomized trial comparing a low-fat, plant-based diet to an American Heart Association diet, Macknin et al²⁸ reported significant reductions in weight $(3.64 \pm 3.41 \text{ kg})$, SBP $(7.96 \pm 12.28 \text{ mmHg})$, and LDL-C $(27.00 \pm 26.72 \text{ mg/dL})$ compared with baseline in adults on the plant-based diet. Bloomer et al²⁹ conducted a trial in which subjects consumed a plant-based diet for 3 weeks. Despite normal baseline clinical indicators, large reductions were observed in LDL-C (22.3 mg/dL), SBP (8.8 mmHg), and DBP (5.2 mmHg).

Jenkins et al³⁰ fed 3 weight-maintaining diets for 2 weeks that were low in saturated fat to participants with elevated LDL-C (~115 mg/dL at baseline). The dietary groups included a conventional low-fat diet, a vegetarian diet high in complex carbohydrates, and a raw vegan diet similar to that of the present study. Significant differences in changes of serum LDL-C were observed between these dietary groups. The conventional low-fat diet reduced LDL-C by 8 mg/ dL, the starch-based vegetarian diet reduced LDL-C by 27 mg/dL, and the raw vegan diet reduced LDL-C by 38 mg/dL (P < 0.001). Thus, a raw plant-based diet may result in greater reductions in serum lipids than one that includes cooked complex carbohydrates.

4.1 | Study strengths and limitations

Several strengths of the present study should be noted. First, the utilization of the food classification system allows for reproducibility in other clinical practices and trials, as the food selection type, preparation, and degree of processing is detailed. Second, the utilization of a prescribed nutrition program in an outpatient cardiovascular clinic allows for the close assessment of the patient's clinical response to the diet. This was facilitated by weekly office visits that allowed for medication weaning as needed. In addition, the provision of food to participants helped facilitate adherence to the dietary protocol. Although there were no statistical differences between high- and low-adherent subjects, a lack of statistical power may be present due to a reduced sample size when groups were divided based on adherence. Additionally, strict adherence standards may also have required a larger sample size for statistical significance to be apparent between groups. A single bite or drink of any food outside of the prescribed diet counted against adherence for the day, even if the remainder of the day represented complete dietary compliance. Lastly, the range of reported dependent variables represents meaningful clinical indicators often evaluated in cardiology practices across the United States. These clinical indicators are most commonly used in the assessment of CVD risk. Thus, this study has real-world applicability in the clinical setting.

Limitations of the current study include the small sample size, lack of a control group, and short duration of follow-up. Although the sample size was small, the large effect sizes indicate that the sample size was more than sufficient for adequate power of the primary endpoints. Further research is needed to determine whether medications, serum lipids, and blood pressure would continue to decrease if the diet were consumed for an extended period of time. In addition, extended trials are needed to assess long-term adherence to the diet. Lastly, inclusion of periodic postprandial glucose testing during the intervention may help establish a potential relationship between postprandial glucose fluctuations and reduced HbA_{1c}.

5 | CONCLUSION

A defined plant-based diet can be used as an effective therapeutic approach in the clinical setting in the treatment of HTN, hypercholesterolemia, and other cardiovascular risk factors while simultaneously reducing overall medication usage. Patients may find this therapeutic approach preferable to conventional and costly drug therapy. Further replication trials are needed with larger sample sizes, control groups, and other dietary comparison groups.

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Conflicts of interest

The authors declare no potential conflicts of interest.

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SUPPORTING INFORMATION

Additional Supporting Information may be found online in the supporting information tab for this article.

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CLINICAL INVESTIGATIONS



Consumption of a defined, plant-based diet reduces lipoprotein(a), inflammation, and other atherogenic lipoproteins and particles within 4 weeks

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Background: Lipoprotein(a) [Lp(a)] is a highly atherogenic lipoprotein and is minimally effected by lifestyle changes. While some drugs can reduce Lp(a), diet has not consistently shown definitive reduction of this biomarker. The effect of consuming a plant-based diet on serum Lp(a) concentrations have not been previously evaluated.

Hypothesis: Consumption of a defined, plant-based for 4 weeks reduces Lp(a).

Methods: Secondary analysis of a previous trial was conducted, in which overweight and obese individuals (n = 31) with low-density lipoprotein cholesterol concentrations >100 mg/dL consumed a defined, plant-based diet for 4 weeks. Baseline and 4-week labs were collected. Data were analyzed using a paired samples t-test.

Results: Significant reductions were observed for serum Lp(a) ($-32.0 \pm 52.3 \text{ nmol/L}$, P = 0.003), apolipoprotein B ($-13.2 \pm 18.3 \text{ mg/dL}$, P < 0.0005), low-density lipoprotein (LDL) particles ($-304.8 \pm 363.0 \text{ nmol/L}$, P < 0.0005) and small-dense LDL cholesterol ($-10.0 \pm 9.2 \text{ mg/dL}$, P < 0.0005). Additionally, serum interleukin-6 (IL-6), total white blood cells, lipoprotein-associated phospholipase A2 (Lp-PLA2), high-sensitivity c-reactive protein (*hs*-CRP), and fibrinogen were significantly reduced ($P \le 0.004$).

Conclusions: A defined, plant-based diet has a favorable impact on Lp(a), inflammatory indicators, and other atherogenic lipoproteins and particles. Lp(a) concentration was previously thought to be only minimally altered by dietary interventions. In this protocol however, a defined plant-based diet was shown to substantially reduce this biomarker. Further investigation is required to elucidate the specific mechanisms that contribute to the reductions in Lp(a) concentrations, which may include alterations in gene expression.

KEYWORDS

general clinical cardiology/adult, lipoproteins, preventive cardiology, vegetarian diet

1 | INTRODUCTION

Lipoprotein(a) [Lp(a)] is an atherogenic lipoprotein structurally similar to low-density lipoprotein cholesterol (LDL-C), although synthesis occurs through independent pathways. Key differences include the linkage of apolipoprotein B100 (Apo-B) to apolipoprotein(a) on the LDL surface.^{1,2} It has been estimated that expression of the genomic region encoding apolipoprotein(a) (LPA gene) accounts for approximately 90% of plasma Lp(a) concentrations.³ Elevated Lp(a) is independently associated with cardiovascular disease,⁴ and the LPA gene

was observed to have the strongest genetic link to cardiovascular disease.⁵ Individuals with Lp(a) plasma concentrations >20 mg/dL have twice the risk of developing cardiovascular disease and approximately 25% of the population may have this plasma concentration.⁶ The mode of action by which Lp(a) exerts its atherogenic effect is likely similar to that of LDL-C, by deposition in the sub-endothelial space and uptake by macrophages mediated via the VLDL receptor.⁷ Lp(a) is particularly atherogenic due to its unique property of being a carrier of oxidized phospholipids, in addition to its higher binding affinity to negatively charged endothelial proteoglycans.⁸ Lp(a) can facilitate

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endothelial dysfunction when concentrations are elevated likely due to this effect.⁹

While PCSK9 inhibitors, high dose atorvastatin, ezetimibe and niacin have resulted in significant reductions in Lp(a).¹⁰⁻¹² lifestyle interventions have not reliably demonstrated reduced Lp(a) to a clinically significant degree. Interestingly, even high saturated fat and high cholesterol diets known to induce hypercholesterolemia have had little influence on plasma Lp(a) concentrations.¹³ Despite the lack of evidence in the literature indicating a relationship between diet and Lp(a) concentrations, a defined, plant-based has not been previously evaluated with respect to its potential effect to reduce Lp(a). Previous investigations have found that a very-high fiber diet comprised of vegetables, fruits and nuts can reduce LDL-C by 33% and Apo-B by 26%,¹⁴ although Lp(a) was not measured. Since such a diet can result in dramatic reductions in LDL-C and Apo-B, secondary analysis of a previously published investigation¹⁵ employing a similar plant-based diet were analyzed to evaluate if Lp(a) could be significantly reduced after 4 weeks among other inflammatory indicators and atherogenic lipoproteins and particles.

2 | METHODS

2.1 | Study population

Participants were subjects of a previous study in which written informed consent was obtained to draw blood for analysis.¹⁵ Laboratory reports for each subject included biomarkers used for clinical purposes, and selected biomarkers are included in the present investigation. The study protocol was approved by the Texas Woman's University Institutional Review Board, Houston.

The study protocol has been previously described.¹⁵ Briefly, all participants were registered new patients of a cardiovascular center and were hypertensive (systolic blood pressure \geq 140 mmHg or diastolic blood pressure \geq 90 mmHg), had elevated LDL-C (\geq 100 mg/dL) and excess body weight (body mass index \geq 25 kg/m²) at baseline. Exclusionary criteria included current tobacco use, current drug abuse, excessive alcohol use (>2 glasses of wine or equivalent for men or > 1 glass of wine or equivalent for woman), a current cancer diagnosis, an ongoing clinically defined infection, a mental disability that would prevent a participant from following the study protocol, an estimated glomerular filtration rate < 60 mg/dL, current pregnancy or lactation, a hospitalization within the past 6 months, and previous exposure to the nutrition program.

2.2 | Intervention

Participants were instructed to consume a defined, plant-based diet for 4 weeks ad-libitum which included the consumption of foods within a food classification system.¹⁵ These foods fell within food levels 0 to 4b of the food classification system (Table S1, Supporting information). Briefly, excluded were animal products, cooked foods, free oils, soda, alcohol, and coffee. Allowed for consumption were raw fruits, vegetables, seeds, and avocado. Small amounts of raw buckwheat and oats were also permitted. Vitamin, herbal, and mineral supplements were to be discontinued unless otherwise clinically indicated. All meals and snacks were provided to subjects, although they were free to consume food on their own within food levels 0 to 4b. In addition, subjects were not advised to alter their exercise habits. Adherence was measured daily as previously described¹⁵ with an adherence assessment tool. Participants indicated in writing each day whether they were adherent. Dietary recalls (24-hour) were conducted by a trained nutritionist at baseline and at 4 weeks. Nutrient intake was analyzed by the Nutrition Data System for Research software (University of Minnesota, version 2016). No lipid lowering medications were altered throughout the intervention.

2.3 | Measures

After a 12-hour fast, the following plasma biomarkers were obtained at baseline and after 4-weeks: total cholesterol (Total-C), LDL-C, highdensity lipoprotein cholesterol (HDL-C), triglycerides, LDL particles (LDL-P), small-dense low-density lipoprotein cholesterol (sdLDL-C), Apo-B, high-density lipoprotein 2 cholesterol (HDL2-C), apolipoprotein A-1 (Apo A-1), and Lp(a). Additionally, high-sensitivity c-reactive protein (hs-CRP), endothelin, interleukin-6 (IL-6), tumor necrosis factor alpha (TNF-a), lipoprotein-associated phospholipase A2 (Lp-PLA2), myeloperoxidase, fibrinogen, troponin-I, N-terminal pro b-type natriuretic peptide (NT-proBNP), total white blood cell count (WBC), neutrophil count, lymphocyte count, monocyte count, eosinophil count, and basophil count were documented. These specific biomarkers of interest were analyzed by either True Health Diagnostics (Frisco, Texas) or Singulex (Alameda, California) depending on the subject's health insurance. The same company that analyzed the baseline labs for a participant was used for the follow-up labs to ensure consistency.

2.4 | Data analysis

Paired samples t-tests were used for the analysis of biochemical measures at baseline and 4-weeks, and significance was confirmed with non-parametric tests. Significance was determined to be a *P* value less than 0.05. spss (version 24) was used for data analysis.

3 | RESULTS

Baseline demographics are indicated in Table 1. Subjects represent a sample that was 81% obese with multiple clinical diagnoses. Two-thirds of subjects were women and 80% were African American.

Adherence to the dietary intervention was approximately 87% over the course of the 4 weeks as measured by the daily adherence assessment tool. Food group consumption is indicated in Table 2 at baseline and 4-weeks. Notably, total fruit consumption increased from 1.3 \pm 2.0 servings to 11.8 \pm 10.4 servings (808% increase, *P* < 0.0005) and total vegetable consumption increased 2.7 \pm 2.0 servings to 16.0 \pm 9.2 servings (493% increase, *P* < 0.0005). Additionally, total animal product consumption decreased from 7.9 \pm 4.7 servings to 0.4 \pm 1.4 servings (95% decrease, *P* = 0.001). The consumption of avocados, dark-green vegetables, deep-yellow vegetables, tomatoes,

| TABLE 1 | Baseline | characteristics and | clinical | diagnoses |
|---------|----------|---------------------|----------|-----------|
|---------|----------|---------------------|----------|-----------|

| | Participants ^a |
|--|---------------------------|
| n | 31 |
| Age (years) | 53.4 (32-69) |
| Sex | |
| Male | 10 (33%) |
| Female | 21 (67%) |
| Race, ethnicity | |
| African American | 25 (80%) |
| Hispanic | 3 (10%) |
| White | 3 (10%) |
| Mean BMI (kg/m²) | $37.5\pm\ 8.3$ |
| Overweight (25-29.9 kg/m ²) | 6 (19%) |
| Obesity class 1 (30-34.9 kg/m ²) | 6 (19%) |
| Obesity class 2 (35-39.9 kg/m ²) | 10 (33%) |
| Obesity class 3 (≥40 kg/m²) | 9 (29%) |
| Current diagnoses | |
| Coronary artery disease | 10 (33%) |
| Type II diabetes mellitus | 8 (27%) |
| Arthritic condition | 7 (23%) |
| Pre-diabetes | 5 (17%) |

Abbreviation: BMI, body mass index.

^a Data are mean (range) unless otherwise indicated.

and other vegetables also significantly increased ($P \le 0.006$). A decreased consumption of white potatoes, fried potatoes, total grains, refined grains, whole grains, added oils, added animal fat, red meat, white meat, eggs, and dairy were also observed ($P \le 0.027$). The consumption of sweets (5% decrease, P = 0.90) and the consumption of nuts/seeds (17% increase, P = 0.736) did not significantly change between baseline and 4-weeks.

Body weight, BMI, total cholesterol, LDL-C, HDL-C, and triglycerides (Table 3) were significantly reduced after 4-weeks of the dietary intervention ($P \le 0.008$). Lp(a) was also significantly reduced (-32.0 \pm 52.3 nmol/L, P = 0.003). In addition, LDL-P, sdLDL-C, Apo-B, HDL2-C, and Apo A-1 were significantly reduced ($P \le 0.03$). Of the atherogenic lipoproteins, sdLDL-C had the greatest relative reduction of approximately 30% (Figure 1). Lp(a) reduced 16% which was proportional to the decrease in Total-C, triglycerides and LDL-P.

Of the inflammatory indicators, *hs*-CRP, IL-6, Lp-PLA2, and fibrinogen significantly decreased ($P \le 0.004$) (Table 4). The WBC, neutrophil, lymphocyte, monocyte, eosinophil and basophil count also significantly decreased ($P \le 0.033$). Interestingly, no statistically significant changes were observed for endothelin-1, TNF-a, myeloperoxidase, troponin-I, or NT-proBNP ($P \ge 0.056$) between baseline and 4-weeks.

TABLE 2 Number of food group servings at baseline and 4-weeks^a

| Food group | Serving size | Baseline ^b | Final ^b | Change ^c | P ^d |
|-------------------------------------|--|---------------------------------|---------------------------------|------------------------|----------------|
| Fruits, total | 1/2 cup chopped, $1/4$ cup dried or 1 medium piece | $\textbf{1.3} \pm \textbf{2.0}$ | 11.8 ± 10.4 | 808% (10.5 \pm 10.8) | <0.0005 |
| Avocado | 1/2 cup chopped | $\textbf{0.1}\pm\textbf{0.2}$ | $\textbf{0.9} \pm \textbf{0.9}$ | 800% (0.8 \pm 0.9) | <0.0005 |
| Vegetables, Total | 1/2 cup chopped or 1 cup raw leafy | $\textbf{2.7} \pm \textbf{2.0}$ | 16.0 ± 9.2 | 493% (13.3 \pm 9.2) | <0.0005 |
| Dark-green vegetables | 1/2 cup chopped or 1 cup raw leafy | $\textbf{0.7} \pm \textbf{1}$ | 5.2 ± 3.8 | 643% (4.5 \pm 4.0) | <0.0005 |
| Deep-yellow vegetables | 1/2 cup chopped | $\textbf{0.2}\pm\textbf{0.4}$ | $\textbf{1.2} \pm \textbf{1.1}$ | 500% (1.0 \pm 1.3) | <0.0005 |
| Tomatoes | 1/2 cup chopped | 0.4 ± 0.5 | 1.7 ± 2.4 | 325% (1.3 \pm 2.4) | 0.006 |
| Other vegetables | 1/2 cup chopped | 1.4 ± 1.2 | $\textbf{7.9} \pm \textbf{6.6}$ | 464% (6.5 \pm 6.3) | <0.0005 |
| White Potatoes ^e | 1/2 cup chopped or 1 medium baked potato | $\textbf{0.3}\pm\textbf{0.7}$ | $\textbf{0.0}\pm\textbf{0.0}$ | –100% (–0.3 \pm 0.7) | 0.03 |
| Fried potatoes | 1/2 cup chopped or 70 g french fries | $\textbf{0.5}\pm\textbf{0.9}$ | $\textbf{0.1}\pm\textbf{0.3}$ | –80% (–0.4 \pm 0.9) | 0.027 |
| Grains, Total | 1 slice of bread or halfcup cooked cereal | 5.7 ± 3.5 | $\textbf{0.7}\pm\textbf{0.9}$ | –88% (–5.0 \pm 3.6) | <0.0005 |
| Refined grains | 1 slice of bread or half cup cooked cereal | $\textbf{3.8} \pm \textbf{2.7}$ | $\textbf{0.2}\pm\textbf{0.7}$ | –95% (–3.6 \pm 3.0) | <0.0005 |
| Whole grains | 1 slice of bread or half cup cooked cereal | $\textbf{1.9} \pm \textbf{2.6}$ | 0.5 ± 0.7 | –74% (–1.4 \pm 2.7) | 0.007 |
| Sweets ^f | 4 g of sugar, 1 tbsp honey or 2 tbsp syrup | $\textbf{1.8} \pm \textbf{2.3}$ | 1.7 ± 1.5 | –5% (–0.1 \pm 2.7) | 0.90 |
| Nuts/seeds | 1/2 oz | 1.2 ± 3.0 | 1.4 ± 1.6 | 17% (0.2 \pm 3.4) | 0.736 |
| Added oils | 1 tsp | $\textbf{3.2}\pm\textbf{3.5}$ | $\textbf{0.1}\pm\textbf{0.2}$ | –97% (–3.1 \pm 3.5) | <0.0005 |
| Added animal fat | 1 tsp | 1.3 ± 2.3 | $\textbf{0.0}\pm\textbf{0.1}$ | –100% (–1.3 \pm 2.3) | 0.005 |
| Animal products, Total ^g | 1 oz | $\textbf{7.9} \pm \textbf{4.7}$ | $\textbf{0.4} \pm \textbf{1.4}$ | –95% (–7.5 \pm 5.3) | 0.001 |
| Red meat | 1 oz | $\textbf{2.1} \pm \textbf{2.9}$ | $\textbf{0.1}\pm\textbf{0.2}$ | –95% (–2.0 \pm 3.0) | <0.0005 |
| White meat | 1 oz | $\textbf{3.9} \pm \textbf{3.7}$ | $\textbf{0.2} \pm \textbf{1.1}$ | –95% (–3.7 \pm 4.1) | <0.0005 |
| Eggs | 1 large egg | 0.5 ± 0.7 | $\textbf{0.0}\pm\textbf{0.1}$ | –100% (–0.5 \pm 0.7) | 0.002 |
| Dairy | 1 cup of milk/yogurt or 1.5 oz of cheese | 1.5 ± 1.6 | $\textbf{0.1}\pm\textbf{0.3}$ | –93% (–1.4 \pm 1.7) | <0.0005 |

^a Data are for subjects who completed 24-h recalls at both baseline and 4-weeks (n = 30).

 $^{\rm b}$ Data are listed in serving size and are presented as mean \pm SD.

 $^{
m c}$ Data indicated as % change (mean \pm SD).

^d Paired samples *t*-tests for within-group comparisons of changes from baseline to final values.

^e Excludes fried potatoes.

^f Includes honey, candy, or other added sugars.

^g Excludes added animal fat.

WILEY CARD

TABLE 3 Atherogenic lipoproteins and particles at baseline and 4-weeks

| | Baseline ^a | Final ^a | Change ^b | P ^c |
|-----------------------------|-------------------------------------|-------------------------------------|-------------------------|----------------|
| Weight (kg) | 108.1 ± 28.6 | 101.4 ± 26.3 | -6% (-6.6 \pm 3.6) | <0.0005 |
| BMI (kg/m ²) | $\textbf{37.5} \pm \textbf{8.3}$ | 35.2 ± 7.8 | -6% (-2.2 \pm 1.1) | <0.0005 |
| Total-C (mg/dL) | $\textbf{216.6} \pm \textbf{34.2}$ | $\textbf{182.7} \pm \textbf{29.9}$ | –16% (–33.8 \pm 25.9) | <0.0005 |
| LDL-C (mg/dL) | 143.0 ± 28.9 | 118.4 ± 26.4 | -17% (-24.6 \pm 21.3) | <0.0005 |
| HDL-C (mg/dL) | 54.8 ± 9.4 | 49.5 ± 10.6 | –9% (–5.2 \pm 6.2) | <0.0005 |
| Triglycerides (mg/dL) | $124.1~\pm~58.1$ | 104.5 ± 53.6 | –16% (–19.6 \pm 38.4) | 0.008 |
| Lp(a) (nmol/L) ^d | 200.7 ± 150.0 | $\textbf{168.8} \pm \textbf{126.7}$ | –16% (–32.0 \pm 52.3) | 0.003 |
| Apo-B (mg/dL) | 115.2 ± 24.5 | 101.9 ± 17.7 | –11% (–13.3 \pm 18.3) | <0.0005 |
| LDL-P (nmol/L) ^e | $\textbf{1891} \pm \textbf{586}$ | 1586 ± 508 | –16% (–305 \pm 363) | <0.0005 |
| sdLDL-C (mg/dL) | $\textbf{33.7} \pm \textbf{11.5}$ | 23.7 ± 8.7 | –30% (–10.0 \pm 9.2) | <0.0005 |
| HDL2-C (mg/dL) | $\textbf{17.4} \pm \textbf{9.8}$ | $\textbf{15.6} \pm \textbf{9.9}$ | –10% (–1.8 \pm 4.5) | 0.030 |
| Apo A-1 (mg/dL) | $\textbf{189.7} \pm \textbf{150.7}$ | 160.2 ± 126.5 | -14% (-27.0 \pm 19.6) | <0.0005 |

Abbreviations: Apo A-1, apolipoprotein A-1; Apo-B, apolipoprotein B100; BMI, body mass index; HDL-C, high-density lipoprotein cholesterol; HDL2-C, high-density lipoprotein-2 cholesterol; LDL-C, low-density lipoprotein cholesterol; LDL-P, low-density lipoprotein particles; Lp(a), lipoprotein(a); sdLDL-C, small-dense low-density lipoprotein cholesterol; total-C, total cholesterol.

^a Mean \pm SD (*n* = 31 unless otherwise indicated).

 $^{\rm b}$ Data indicated as % change (mean \pm SD).

^c Paired samples *t*-tests for within-group comparisons of changes from baseline to final values.

^d n = 28 due to premature coagulation of sample (n = 1) and incompatible units (mg/dL) when merging laboratory results (n = 2).

^e n = 29 due to premature coagulation of samples.

4 | DISCUSSION

The consumption of a defined, plant-based diet resulted in a significant reduction in Lp(a) after 4 weeks; thus, the study hypothesis was accepted. The reduction in Lp(a) was profound and is one of the largest reductions due to lifestyle reported in the literature. The magnitude of change was comparable to other leading medical therapies, such as niacin (~20% reduction) and PCSK9 inhibitors (~25% reduction).¹² It is important to note that this dietary intervention rapidly reduced Lp(a) by 16% in only 4 weeks, whereas shorter duration

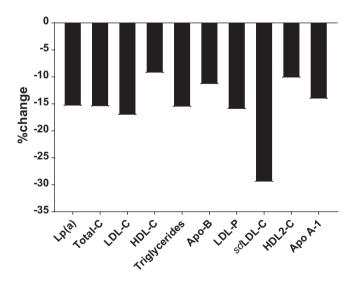


FIGURE 1 Percent change of atherogenic lipoproteins and particles from baseline to 4-weeks. All variable changes indicated are significant (P < 0.05). Lp(a), lipoprotein(a); Total-C, total cholesterol; LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol; Apo-B, apolipoprotein B100; LDL-P, lowdensity lipoprotein particles; sdLDL-C, small-dense low-density lipoprotein cholesterol; HDL2-C, high-density lipoprotein-2 cholesterol; Apo A-1, apolipoprotein A-1

niacin and PCSK9 inhibitor drug trials typically lasted 8 to 12 weeks. It should also be noted that niacin may reduce inflammation, such as *hs*-CRP, by 15% after 3 months, although PCSK9 inhibitors do not.^{16,17} After 4 weeks, the dietary intervention reduced *hs*-CRP by 30.7%. In addition, IL-6, Lp-PLA2, fibrinogen, and white blood cells were significantly reduced, as were sdLDL-C, LDL-P, and Apo-B, all of which represent a systemic, cardio-protective effect.^{18–24} Thus, the use of this single dietary approach in the clinical setting, vs multiple drug therapy, may be an appropriate tool in treating complex patients with a myriad of elevated CVD-related biomarkers.

Elevated Apo A1, HDL-C, and HDL2-C are associated with reduced cardiovascular disease risk.^{24,25} While these HDL fractions were significantly reduced in this trial, this is a common phenomenon observed when consuming plant-based diets. A systematic review and meta-analysis of plant-based observational and clinical trials found that while HDL-C was significantly reduced compared to those consuming non-vegetarian diets, LDL-C and total-C were also reduced.²⁶ Despite reductions in HDL-C, those who consumed plant-based diets had a 25% reduced incidence of ischemic CVD compared with non-vegetarian counterparts.²⁷

Lp(a) concentrations in the present study represent a high-risk population.²⁸ This may be explained by the higher proportion of African Americans in this sample, as African Americans may have higher Lp(a) concentrations compared with Caucasians.²⁹ An evaluation of 532 359 patients found that an Lp(a) concentration > 50 mg/dL was common among patients.³⁰ This range roughly corresponds to the mean nmol/L Lp(a) concentration observed in the present study.

4.1 | Effect of weight loss on plasma Lp(a) concentrations

An energy restricted diet was found to independently reduce serum Lp(a) in those with baseline concentrations >20 mg/dL, but not <20 mg/dL.³¹ Further studies have found that weight loss may not

TABLE 4 Inflammatory and other cardiovascular indicators at baseline and 4-weeks



| | Baseline ^a | Final ^a | Change ^b | Pc |
|--|-------------------------------------|-------------------------------------|-----------------------------|---------|
| hs-CRP (mg/dL) | 7.8 ± 6.4 | 5.4 ± 4.7 | –30.7% (–2.4 \pm 3.7) | 0.001 |
| Endothelin (pg/mL) ^d | $\textbf{2.2}\pm\textbf{0.7}$ | $\textbf{2.2}\pm\textbf{0.8}$ | 0% (0.0 \pm 0.7) | 0.916 |
| IL-6 (pg/mL) ^d | $\textbf{2.6} \pm \textbf{1.4}$ | 2.0 ± 1.0 | –23.1% (–0.6 \pm 1.0) | 0.001 |
| TNF- α (pg/mL) ^d | $\textbf{2.0}\pm\textbf{0.9}$ | $\textbf{2.2}\pm\textbf{0.9}$ | 10.0% (0.2 \pm 0.6) | 0.096 |
| Lp-PLA ₂ (ng/mL) ^d | $\textbf{252.3} \pm \textbf{136.3}$ | $\textbf{210.7} \pm \textbf{119.1}$ | -16.4% (-41.6 \pm 64.6) | 0.001 |
| Myeloperoxidase (pmol/L) ^e | 124.1 ± 58.1 | 104.5 ± 53.6 | -23.0% (-28.5 \pm 66.1) | 0.056 |
| Fibrinogen (mg/dL) ^f | $\textbf{561.4} \pm \textbf{112.2}$ | 530.1 ± 102.9 | -5.6% (-31.3 \pm 50.7) | 0.004 |
| NT-proBNP (pg/mL) ^d | 65.2 ± 71.2 | 69.4 ± 75.9 | 6.2% (4.1 \pm 23.2) | 0.337 |
| Total WBC (K/μL) ^d | $\textbf{6.3} \pm \textbf{2.0}$ | 4.8 ± 1.3 | –22.2% (–1.4 \pm 1.1) | <0.0005 |
| Neutrophils (K/µL) ^d | 3.5 ± 1.4 | $\textbf{2.5}\pm\textbf{0.9}$ | -28.6% (-1.0 \pm 0.8) | <0.0005 |
| Lymphocytes (K/µL) ^d | $\textbf{1.9} \pm \textbf{0.7}$ | $\textbf{1.6} \pm \textbf{0.6}$ | -15.8% (-0.3 \pm 0.4) | <0.0005 |
| Monocytes (K/µL) ^d | $\textbf{0.46}\pm\textbf{0.12}$ | 0.38 ± 0.09 | –15.2% (–0.07 \pm 0.1) | <0.0005 |
| Eosinophils (K/µL) ^d | $\textbf{0.18} \pm \textbf{0.11}$ | $\textbf{0.15}\pm\textbf{0.11}$ | -16.6% (-0.03 \pm 0.07) | 0.033 |
| Basophils (K/µL) ^d | $\textbf{0.029} \pm \textbf{0.016}$ | $\textbf{0.024} \pm \textbf{0.015}$ | –17.2% (–0.005 \pm 0.010) | 0.016 |

Abbreviations: hs-CRP, high-sensitivity c-reactive protein; IL-6, interleukin-6; Lp-PLA₂, lipoprotein-associated phospholipase A2; NT-proBNP, N-terminal pro b-type natriuretic peptide; TNF- α , tumor necrosis factor-alpha; WBC, white blood cells.

^a Mean \pm SD (*n* = 31 unless otherwise indicated).

 $^{\rm b}$ Data indicated as % change (mean \pm SD).

^c Paired samples *t*-tests for within-group comparisons of changes from baseline to final values.

^d n = 30 due to premature coagulation of samples.

e n = 25 due to premature coagulation of samples.

f n = 27 due to premature coagulation of samples.

independently reduce Lp(a) concentrations. A pooled analysis of cohorts found that as weight loss ensued, Lp(a) concentrations surprisingly increased.³² Baseline Lp(a) concentrations on average between the four cohorts analyzed were approximately 40 mg/dL, well above the >20 mg/dL threshold reported in the initial study.³¹ Other investigations examining the effect of weight loss on Lp(a) concentration have not demonstrated a relationship between these two variables.^{33,34} Interestingly, the emphasis on consuming plant-based foods, even with a calorie restricted diet, did not result in Lp(a) reductions compared with a calorie restricted red meat centered diet.³⁵ The plant-centered diet in this trial³⁵ still contained a significant number of calories derived from animal-based sources in addition to processed plant foods. Also, both diets contained similar quantities of dietary fiber, a measure of plant-food intake. Based on these weight loss trials, Lp(a) concentration is likely not influenced by weight reduction.

4.2 | Effect of diet on plasma Lp(a) concentrations

Other trials using diets emphasizing plant-based foods have not demonstrated similar results. A low-fat and low-saturated fat diet with an increased intake of fruits and vegetables interestingly increased Lp(a) concentrations.³⁶ Subjects consumed four to five servings of fruits or berries and five to six servings of vegetables daily for 5 weeks and all food was provided. It is important to note that subjects still consumed animal products throughout the intervention³⁶ which included dairy products and lean meats. The fiber content (40 g vs 51 g in the present study) was not as high as would be expected when consuming a higher quantity of plantfoods, and the number of fruits and vegetables did not meet the levels observed in the present study (11.8 servings of fruits and

16 servings of vegetables). Based on this data, it is probable that exclusively increasing fruit and vegetable intake is not sufficient to elicit reduced Lp(a) concentrations.

It has also been reported that a low-carbohydrate, high-fat diet (45% carbohydrate, 40% fat) may have a favorable impact on Lp(a) concentrations compared with a high-carbohydrate, low-fat diet (65% carbohydrate, 20% fat), although it is unclear as to what precisely was consumed on either of these diets.³⁷ In addition, the differences were small, as only a 2.17 mg/dL difference was observed between both groups, and baseline Lp(a) concentrations were <20 mg/dL. The Omni Heart Trial also found that replacing calories from carbohydrates and protein with unsaturated fats produced a smaller increase in Lp(a) comparatively, but both diets still elicited increased plasma Lp(a) compared with baseline. The differences between groups were also small at the end of the intervention (<4 mg/dL difference).³⁸

In individuals with low baseline Lp(a) concentrations (approximately 5.5 mg/dL), the consumption of copious saturated fat, cholesterol (derived from egg consumption) and polyunsaturated fat did not influence Lp(a) concentrations.¹³ Carbohydrate intake was low in this trial as well (39% to 46% carbohydrate as a percent of energy). While fat consumption does not appear to influence serum Lp(a) concentrations in the fasting state, a variety of fats may significantly increase postprandial, transient plasma Lp(a) concentrations over the course of 8 hours.³⁹ Investigators found that linoleic, oleic, palmitic, and stearic acid all resulted in significant transient increases in Lp(a) concentrations. While saturated fats, stearic acid and palmitic acid, appeared to have the greatest increase in serum Lp(a) compared with oleic acid and linoleic acid, this differing response did not reach statistical significance.

4.3 | Mechanisms contributing to reduced plasma Lp(a)

The observed reduction in Lp(a) in the present study may be due to decreased hepatic synthesis of apolipoprotein(a) and Apo-B. This may be in part due to decreased expression of the LPA gene. Since the LPA gene is almost exclusively expressed in the liver,⁴⁰ hepatic influences, including the production of *hs*-CRP and inflammatory cytokines, such as IL-6, may upregulate LPA gene expression.⁴¹ Indeed, those with inflammatory conditions may have increased Lp(a) concentrations compared with healthy controls.⁴²

Current data in our plant-based study supports this hypothesis, as reduced hs-CRP and IL-6 was observed. In contrast, previous studies utilizing plant-centered diets to reduce Lp(a) were unsuccessful, as animal products were still substantially consumed.^{35,36} Animal-based foods, including lean meat, can induce a postprandial inflammatory response, including increased hs-CRP and IL-6.43 Pooled data of those consuming non-vegan, plant-based diets have shown reduced hs-CRP and IL-6,⁴⁴ although to a lesser extent compared with the present study (hs-CRP; -0.55 mg/dL vs -2.42 mg/dL, IL-6; -0.25 pg/mL vs -0.64 pg/mL). The elimination of animal products and processed foods completely on a defined, plant-based diet may be a more prudent dietary strategy to avoid potential fluctuations in inflammation. Thus, the fact that there were only minimally processed plant foods consumed during this dietary intervention may account for the observed reduction in serum Lp(a) concentrations that may be associated with reduced LPA gene expression. Further mechanistic research is needed to confirm this hypothesis.

4.4 | Strengths and limitations

The high dietary adherence and provision of all food to subjects supports the conclusion that the intervention likely fully accounted for the observed biochemical changes among the subjects. Furthermore, the study took place in an outpatient clinical setting with established patients providing a real-world example of a standard clinical practice. This study provides a model for the implementation of this intervention across other medical practices. In contrast, a limitation in the design of this study was the lack of a control group and the small sample size. A larger sample size and a control group would be needed to strengthen a causal relationship.

5 | CONCLUSION

A defined, plant-based diet has a favorable impact on Lp(a) and other atherogenic lipoproteins and particles. Lp(a) concentration was previously thought to be only minimally altered by lifestyle interventions. In this study, however, a defined plant-based diet resulted in a substantial reduction in Lp(a) in only 4 weeks. Further investigations are warranted to elucidate the specific mechanisms that contribute to reduced Lp(a) concentrations, which may include alterations in LPA gene expression mediated via hepatic inflammation.

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Conflict of interest

The authors declare no potential conflicts of interest.

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of the article.

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