

**UNITED STATES DISTRICT COURT
SOUTHERN DISTRICT OF NEW YORK**

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SAMANTHA MARSH, individually and on	:	
behalf of all others similarly situated,	:	
	:	Case No.
Plaintiff,	:	
v.	:	
	:	
	:	
WANABANA LLC and WANABANA USA LLC,	:	CLASS ACTION COMPLAINT
	:	
Defendants.	:	<u>JURY TRIAL DEMANDED</u>
	:	
	:	
	:	
_____	X	

Plaintiff Samantha Marsh (hereinafter “Plaintiff”), individually and on behalf of all others similarly situated, by her attorneys, alleges the following upon information and belief, except for those allegations pertaining to Plaintiff, which are based on personal knowledge:

NATURE OF THE ACTION

1. This action seeks to remedy the deceptive and misleading business practices of WanaBana LLC and WanaBana USA LLC (hereinafter “Defendants”) with respect to the manufacturing, marketing, and sale of Defendants’ WanaBana, Schnucks, and Weis fruit puree products throughout the state of New York and throughout the country (hereinafter the “Products”):

- WanaBana Apple Cinnamon Fruit Purée Pouch
- Schnucks Apple Sauce 90g pouches with cinnamon
- Weis Cinnamon Apple Sauce 90g

2. Defendants have improperly, deceptively, and misleadingly labeled and marketed their Products to reasonable consumers, like Plaintiff, by omitting and not disclosing to consumers

on their packaging that the Products are contaminated with unsafe levels of lead, which is a powerful neurotoxin that is known to cause cognitive deficits, mental illness, dementia, and hypertension.

3. The Products' contamination is particularly egregious given the potentially severe and irreversible consequences of lead consumption.

4. Defendants specifically list the ingredients in the Products on the labeling; however, Defendants fail to disclose that the Products contain, or are at the risk of containing, lead.

5. A few representative examples of Defendants' lack of disclosure on the Products are depicted below:







6. Lead is a powerful neurotoxin. There is no safe blood level of lead.¹ Lead consumption has been shown to reduce intelligence, and to increase the risk of mental illness, dementia, hypertension, arrhythmia, and breast cancer.²

7. Consumers like the Plaintiff trust manufacturers such as Defendants to sell products that are safe and free from harmful known substances, including lead.

8. Plaintiff and those similarly situated (hereinafter “Class Members”) certainly expect that the food products they purchase will not contain, or risk containing, any knowingly harmful substances that cause disease.

9. Unfortunately for consumers, like Plaintiff, the food Products they purchased contained, or were at risk of containing, lead.

10. The FDA issued a public health alert on October 28, 2023, and Defendants announced a product recall (“Recall”) on October 29, 2023.³

11. Independent testing confirmed and demonstrated the presence of *lead* in the Products.

12. Defendants are using a marketing and advertising campaign that omits from the ingredients lists that the Products contain lead. This omission leads a reasonable consumer to believe they are not purchasing a product with a known neurotoxin when in fact they are purchasing a product contaminated with lead.

¹ CDC – Lead – Tips – Sources of Lead – Folk Medicine, CENTERS FOR DISEASE CONTROL AND PREVENTION (Oct. 15, 2013), <http://www.cdc.gov/nceh/lead/tips/folkmedicine.htm>.

² Maryse F. Bouchard, PhD et al., *Blood Lead Levels and Major Depressive Disorder, Panic Disorder, and Generalized Anxiety Disorder in US Young Adults*, 66 ARCHIVES OF GENERAL PSYCHIATRY 1313, 1317 (Dec 2009); Marc G. Weisskopf et al., *Cumulative Lead Exposure and Prospective Change in Cognition Among Elderly Men*, 160 AMERICAN JOURNAL OF EPIDEMIOLOGY 1184, 1185, 1188, 1190-91 (2004); Olusegun I. Alatise, Gerhard N. Schrauzer, *Lead Exposure: A Contributing Cause of the Current Breast Cancer Epidemic in Nigerian Women*, BIOLOGICAL TRACE ELEMENT RESEARCH 127, 138 (Mar. 3, 2010).

³ *Id.*

13. Defendants' marketing and advertising campaign includes the one place that every consumer looks when purchasing a product – the packaging and labels themselves. As such, a reasonable consumer reviewing Defendants' labels reasonably believes that they are purchasing a product that is safe for oral ingestion and does not contain any harmful neurotoxins. Indeed, consumers expect the ingredient listing on the packaging and labels to accurately disclose the ingredients within the Products. Thus, reasonable consumers would not think that Defendants are omitting that the Products contain, or are at risk of containing, lead.

14. Defendants' advertising and marketing campaign is false, deceptive, and misleading because the Products do contain, or risk containing, lead, which is dangerous to one's health and well-being. Nevertheless, Defendants do not list or mention lead anywhere on the Products' packaging or labeling.

15. Plaintiff and Class Members relied on Defendants' misrepresentations and omissions of the safety of the Products and what is in the Products when they purchased them.

16. Consequently, Plaintiff and Class Members lost the entire benefit of their bargain when what they received was a food product contaminated with a known neurotoxin that is harmful to consumers' health.

17. That is because Defendants Products containing, or at risk of containing lead, a known dangerous substance, have no value.

18. As set forth below, food products, such as Defendants' Products, are in no way safe for human consumption and are entirely worthless.

19. Alternatively, Plaintiff and Class Members paid a price premium for the Products based upon Defendants' marketing and advertising campaign including their false and misleading representations and omission on the Products' labels. Given that Plaintiff and Class Members paid

a premium for the Products, Plaintiff and Class Members suffered an injury in the amount of the premium paid.

20. Accordingly, Defendants' conduct violated and continues to violate, *inter alia*, New York General Business Law §§ 349 and 350. Defendants also breached and continue to breach their warranties regarding the Products.

21. Plaintiff brings this action against Defendants on behalf of herself and Class Members who purchased the Products during the applicable statute of limitations period (the "Class Period").

FACTUAL BACKGROUND

22. Defendants manufacture, market, advertise, and sell food products.

23. Consumers have become increasingly concerned about the effects of ingredients in products that they orally ingest. Companies, such as Defendants, have capitalized on consumers' desire for food products, and indeed, consumers are willing to pay, and have paid, a premium for these products.

24. Consumers lack the meaningful ability to test or independently ascertain or verify whether a product contains unsafe substances, such as lead, especially at the point of sale, and therefore must and do rely on Defendants to truthfully and honestly report what the Products contain or are at risk of containing on the Products' packaging or labels.

25. The Products' packaging does not identify lead. Indeed, lead is not listed in the ingredients section, nor is there any warning about the inclusion (or even potential inclusion) of lead in the Products. This leads reasonable consumers to believe the Products do not contain, and are not at risk of containing, lead.

26. However, the Products contain, or are at risk of containing, lead.

27. The FDA found that Defendants' Products contained extremely high levels of lead when the North Carolina Department of Health and Human Services (NCDHHS) and the North Carolina Department of Agriculture & Consumer Services (NCDA&CS) reported four children with elevated blood lead levels, indicating potential acute lead toxicity. "The NCDHHS investigation identified WanaBana apple cinnamon fruit puree pouches as a potential shared source of exposure."⁴

28. The FDA issued a public health alert on October 28, 2023, and Defendants announced a product recall ("Recall") on October 29, 2023.⁵

29. Independent testing confirmed and demonstrated the presence of lead in the Products.

30. An FDA investigation confirmed that the lead contamination originated from the production facility owned by Austrofoods in Ecuador. FDA investigators collected samples of cinnamon supplied by Negasmart to Austrofoods, and the results showed extremely high levels of lead contamination of up to 5110 parts per million (ppm).⁶

31. "For context, the international standard-setting body, Codex Alimentarius Commission (Codex) is considering adopting a maximum level of 2.5 ppm for lead in bark spices, including cinnamon, in 2024."⁷

32. In other words, the amount of lead found in the cinnamon used in the Products was 2,044 times the maximum level proposed by an international standard-setting body.⁸

⁴ <https://www.fda.gov/food/outbreaks-foodborne-illness/investigation-elevated-lead-levels-cinnamon-applesauce-pouches-november-2023#Time>

⁵ *Id.*

⁶ <https://www.fda.gov/food/outbreaks-foodborne-illness/investigation-elevated-lead-levels-cinnamon-applesauce-pouches-november-2023>

⁷ *Id.*

⁸ <https://www.cnn.com/2023/12/18/health/cinnamon-applesauce-lead-levels/index.html>

33. The FDA has received at least 65 reports of elevated blood lead levels in children under 6 years old as a result of the contaminated Products. “To date, the FDA has included adverse event reports submitted directly to the FDA that note blood lead levels at or above 3.5 micrograms of lead per deciliter of whole blood ($\mu\text{g}/\text{dL}$) within 3 months after consuming recalled product.”⁹

34. The CDC has received at least 125 suspected or confirmed cases of elevated lead levels as a result of the Products.¹⁰

35. Lead is a powerful neurotoxin. There is no safe blood level of lead.¹¹ Lead consumption has been shown to reduce intelligence, and to increase the risk of mental illness, dementia, hypertension, arrhythmia, and breast cancer.¹²

36. This is true even at low levels of lead consumption.¹³ For example, research has shown that an increase of only 0.3 micrograms/deciliter of median blood lead levels is associated with a doubling of the risk for panic disorder.¹⁴ People exposed to low levels of lead lose an average of 1.37 IQ points per 1 microgram/deciliter increase in blood lead concentration.¹⁵ Ingested lead accumulates in the bones and brain and can cause health problems even decades

⁹ <https://www.fda.gov/food/outbreaks-foodborne-illness/investigation-elevated-lead-levels-cinnamon-applesauce-pouches-november-2023#Time>

¹⁰ *Id.*

¹¹ CDC – Lead – Tips – Sources of Lead – Folk Medicine, CENTERS FOR DISEASE CONTROL AND PREVENTION (Oct. 15, 2013), <http://www.cdc.gov/nceh/lead/tips/folkmedicine.htm>.

¹² Maryse F. Bouchard, PhD et al., *Blood Lead Levels and Major Depressive Disorder, Panic Disorder, and Generalized Anxiety Disorder in US Young Adults*, 66 ARCHIVES OF GENERAL PSYCHIATRY 1313, 1317 (Dec 2009); Marc G. Weisskopf et al., *Cumulative Lead Exposure and Prospective Change in Cognition Among Elderly Men*, 160 AMERICAN JOURNAL OF EPIDEMIOLOGY 1184, 1185, 1188, 1190-91 (2004); Olusegun I. Alatise, Gerhard N. Schrauzer, *Lead Exposure: A Contributing Cause of the Current Breast Cancer Epidemic in Nigerian Women*, BIOLOGICAL TRACE ELEMENT RESEARCH 127, 138 (Mar. 3, 2010).

¹³ *Id.*

¹⁴ Bouchard, *supra*, at 1317.

¹⁵ Richard L. Canfield, Ph.D et al., *Intellectual Impairments in Children with Blood Lead Concentrations Below 10 Micrograms per Deciliter*, THE NEW ENGLAND JOURNAL OF MEDICINE 1517, 1521 (April 17, 2003)

later.¹⁶ Chronic low dose exposure to lead is believed to be associated with cognitive decline and dementia in older adults.¹⁷

37. Children are at especially high risk of developing adverse effects from lead exposure due to their developing brains, and because, compared to adults, less lead is stored by the body in bones and teeth and more in the nervous system.¹⁸

38. “Even low levels of lead in blood have been shown to affect a child’s learning capacity, ability to pay attention, and academic achievement. The effects of lead exposure can be permanent.”¹⁹

39. “CDC currently uses a blood lead reference value (BLRV) of 3.5 micrograms per deciliter to identify children with blood lead levels that are higher than most children’s levels. This level is based on the on the 97.5th percentile of the blood lead values among U.S. of children ages 1-5 years from the 2015-2016 and 2017-2018 National Health and Nutrition Examination Survey (NHANES) cycles. Children with blood lead levels at or above the BLRV are among the top 2.5% of U.S. children with the highest blood lead levels.”²⁰

40. Children found to have a blood lead level greater than 3.5 µg/dL should be reported to state and local health departments which may prompt an investigation of the child’s home and environment and regular monitoring.²¹

¹⁶ Marc G. Weisskopf et al., *Cumulative Lead Exposure and Prospective Change in Cognition Among Elderly Men*, 160 AMERICAN JOURNAL OF EPIDEMIOLOGY 1184, 1185, 1188, 1190-91 (2004); Jennifer Weuve et al., *Cumulative Exposure to Lead in Relation to Cognitive Function in Older Women*, 117 ENVIRONMENTAL HEALTH PERSPECTIVES 574, 578 (April 2009).

¹⁷ Marc G. Weisskopf et al., *Cumulative Lead Exposure and Prospective Change in Cognition Among Elderly Men*, 160 AMERICAN JOURNAL OF EPIDEMIOLOGY 1184, 1185, 1188, 1190-91 (2004); Jennifer Weuve et al., *Cumulative Exposure to Lead in Relation to Cognitive Function in Older Women*, 117 ENVIRONMENTAL HEALTH PERSPECTIVES 574, 578 (April 2009); Bouchard, *supra*, at 1318.

¹⁸ <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1314903/>

¹⁹ <https://www.cdc.gov/nceh/lead/docs/lead-levels-in-children-fact-sheet-508.pdf>

²⁰ <https://www.cdc.gov/nceh/lead/prevention/blood-lead-levels.htm>

²¹ <https://www.cdc.gov/nceh/lead/advisory/acclpp/actions-blls.htm>

41. Children found to have blood levels greater than 20 µg/dL are put on more advanced treatments, including abdominal x-ray, bowel decontamination, chelation therapy, or even admission to a hospital.²²

42. Children found to have lead in their blood are recommended to have their levels monitored and potentially enroll in various treatments, including feeding the child a diet high in iron and calcium, x-rays, and chelation therapy to remove lead from their blood.

43. Independent testing confirmed and demonstrated the presence of *lead* in the Products.

44. Defendants are large and sophisticated corporations that have been in the business of producing, manufacturing, selling, and distributing food products for many years, including producing and manufacturing the contaminated Products.

45. Defendants are in the unique and superior position of knowing the ingredients and raw materials used in the manufacturing of their Products and possess unique and superior knowledge regarding the manufacturing process of the Products, the manufacturing process of the ingredients and raw materials the Products contain, and the risks associated with those processes, such as the risk of lead contamination, as well as the ability to test the Products for lead contamination prior to releasing the Products into the stream of commerce.

46. Accordingly, Defendants possess superior knowledge regarding the risks involved in the production and manufacturing of their Products. Such knowledge is not readily available to consumers like Plaintiff and Class Members.

47. Defendants have a duty to provide consumers, like Plaintiff and Class Members, with accurate information about the contents of the Products.

²² *Id.*

48. Therefore, Defendants' false, misleading, and deceptive omissions regarding the Products containing lead is likely to continue to deceive and mislead reasonable consumers and the public, as they have already deceived and misled Plaintiff and the Class Members.

49. Defendants' misrepresentations and omissions were material and intentional because people are concerned with what is in the products that they orally ingest. Consumers such as Plaintiff and the Class Members are influenced by the marketing and advertising campaign, the Products' labels, and the listed ingredients. Defendants know that if they had not omitted that the Products contained lead, then Plaintiff and the Class would not have purchased the Products at all.

50. Through their deceptive advertising and labeling, Defendants have violated, *inter alia*, NY General Business Law § 392-b by: a) putting upon an article of merchandise, bottle, wrapper, package, label, or other thing containing or covering such an article, or with which such an article is intended to be sold, or is sold, a false description or other indication of or respecting the kind of such article or any part thereof; and b) selling or offering for sale an article which, to their knowledge, is falsely described or indicated upon any such package or vessel containing the same, or label thereupon, in any of the particulars specified.

51. Consumers rely on marketing and information in making purchasing decisions.

52. By omitting that the Products include lead on the labels of the Products throughout the Class Period, Defendants know that those omissions are material to consumers since they would not purchase a product with a harmful neurotoxin such as lead.

53. Defendants' deceptive representations and omissions are material in that a reasonable person would attach importance to such information and would be induced to act upon such information in making purchase decisions.

54. Plaintiff and the Class Members reasonably relied to their detriment on Defendants' misleading representations and omissions.

55. Defendants' false, misleading, and deceptive misrepresentations and omissions are likely to continue to deceive and mislead reasonable consumers and the general public, as they have already deceived and misled Plaintiff and the Class Members.

56. In making the false, misleading, and deceptive representations and omissions described herein, Defendants knew and intended that consumers would pay a premium for a product marketed without lead over comparable products not so marketed.

57. As an immediate, direct, and proximate result of Defendants' false, misleading, and deceptive representation and omission, Defendants injured Plaintiff and the Class Members in that they:

- a. Paid a sum of money for Products that were not what Defendants represented;
- b. Paid a premium price for Products that were not what Defendants represented;
- c. Were deprived of the benefit of the bargain because the Products they purchased was different from what Defendants warranted;
- d. Were deprived of the benefit of the bargain because the Products they purchased had less value than what Defendants represented;
- e. They ingested a substance that was of a different quality than what Defendants promised; and
- f. Were denied the benefit of the properties of the Products Defendants promised.

58. Had Defendants not made the false, misleading, and deceptive representations and omissions, Plaintiff and the Class Members would not have been willing to pay the same

amount for the Products they purchased and, consequently, Plaintiff and the Class Members would not have been willing to purchase the Products.

59. Plaintiff and the Class Members paid for Products that do not contain Lead. Since the Products do indeed contain lead, a harmful neurotoxin, the Products Plaintiff and the Class Members received were worth less than the Products for which they paid.

60. Plaintiff and the Class Members all paid money for the Products; however, Plaintiff and the Class Members did not obtain the full value of the advertised Products due to Defendants' misrepresentations and omissions. Plaintiff and the Class Members purchased, purchased more of, and/or paid more for, the Products than they would have had they known the truth about the Products. Consequently, Plaintiff and the Class Members have suffered injury in fact and lost money as a result of Defendants' wrongful conduct.

61. Plaintiff and Class Members read and relied on Defendants' representations about the Products and purchased Defendants' Products based thereon. Had Plaintiff and Class Members known the truth about the Products, i.e., that they contain a harmful neurotoxin (i.e. lead), they would not have been willing to purchase them at any price, or, at minimum would have paid less for them.

JURISDICTION AND VENUE

62. This Court has subject matter jurisdiction under the Class Action Fairness Act, 28 U.S.C. section §1332(d) in that (1) this is a class action involving more than 100 class members; (2) Plaintiff is a citizen of New York, and Defendant Wanabana LLC is a citizen of Florida; Defendant Wanabana USA LLC is a citizen of Florida; and (3) the amount in controversy is in excess of \$5,000,000, exclusive of interests and costs.

63. This Court has personal jurisdiction over Defendants because Defendants conduct and transact business in the state of New York, contract to supply goods within the state of New York, and supply goods within the state of New York.

64. Venue is proper because Plaintiff and many Class Members reside in the Southern District of New York, and throughout the state of New York. A substantial part of the events or omissions giving rise to the Classes' claims occurred in this district.

PARTIES

Plaintiff

65. Plaintiff is a citizen and resident of Dutchess County, New York. During the applicable statute of limitations period, Plaintiff purchased and used Defendants' Products that contained lead, including Products that were subject to the recall. More specifically, during the class period Plaintiff purchased WanaBana Apple Cinnamon Fruit Purée Pouch at a Dollar Tree brick-and-mortar store in Dutchess County, New York during the Class Period at an approximate retail price of \$1.25 per unit.

66. Had Defendants not made the false, misleading, and deceptive representations and omissions regarding the contents of the Products, Plaintiff would not have been willing to purchase the Products. Plaintiff purchased, purchased more of, and/or paid more for, the Products than she would have had she known the truth about the Products. The Products Plaintiff received were worthless because they contain the known harmful neurotoxin, Lead. Alternatively, Plaintiff paid a price premium based on Defendants' false, misleading, and deceptive misrepresentations and omissions. Accordingly, Plaintiff was injured in fact and lost money as a result of Defendants' improper conduct.

Defendants

67. Defendant, Wanabana LLC is a Florida company with its principal place of business in Miami Shores, Florida.

68. Defendant, Wanabana USA LLC is a Florida company with its principal place of business in San Juan, Puerto Rico.

69. Defendants manufacture, market, advertise, and distribute the Products throughout the United States. Defendants created and/or authorized the false, misleading, and deceptive advertisements, packaging, and labeling of their Products.

CLASS ALLEGATIONS

70. Plaintiff brings this matter on behalf of herself and those similarly situated. As detailed at length in this Complaint, Defendants orchestrated deceptive marketing and labeling practices. Defendants' customers were uniformly impacted by and exposed to this misconduct. Accordingly, this Complaint is uniquely situated for class-wide resolution.

71. The Class is defined as all consumers who purchased the Products anywhere in the United States during the Class Period.

72. Plaintiff also seeks certification, to the extent necessary or appropriate, of a subclass of individuals who purchased the Products in the state of New York at any time during the Class Period (the "New York Subclass").

73. The Class and New York Subclass are referred to collectively throughout the Complaint as the Class.

74. The Class is properly brought and should be maintained as a class action under Rule 23(a), satisfying the class action prerequisites of numerosity, commonality, typicality, and adequacy because:

75. Numerosity: Class Members are so numerous that joinder of all members is impracticable. Plaintiff believes that there are thousands of consumers in the Class and the New York Class who are Class Members as described above who have been damaged by Defendants' deceptive and misleading practices.

76. Commonality: The questions of law and fact common to the Class Members which predominate over any questions which may affect individual Class Members include, but are not limited to:

a. Whether Defendants were responsible for the conduct alleged herein which was uniformly directed at all consumers who purchased the Products;

b. Whether Defendants' misconduct set forth in this Complaint demonstrates that Defendants have engaged in unfair, fraudulent, or unlawful business practices with respect to the advertising, marketing, and sale of their Products;

c. Whether Defendants made false and/or misleading statements and omissions to the Class and the public concerning the contents of their Products;

d. Whether Defendants false and misleading statements and omissions concerning their Products were likely to deceive the public; and

e. Whether Plaintiff and the Class are entitled to money damages under the same causes of action as the other Class Members.

77. Typicality: Plaintiff is a member of the Class. Plaintiff's claims are typical of the claims of each Class Member in that every member of the Class was susceptible to the same deceptive, misleading conduct and purchased Defendants' Products. Plaintiff is entitled to relief under the same causes of action as the other Class Members.

78. Adequacy: Plaintiff is an adequate Class representative because her interests do not conflict with the interests of the Class Members she seeks to represent, her consumer fraud claims are common to all members of the Class, she has a strong interest in vindicating her rights, she has retained counsel competent and experienced in complex class action litigation, and counsel intends to vigorously prosecute this action.

79. Predominance: Pursuant to Rule 23(b)(3), common issues of law and fact identified above predominate over any other questions affecting only individual members of the Class. The Class issues fully predominate over any individual issues because no inquiry into individual conduct is necessary; all that is required is a narrow focus on Defendants' deceptive and misleading marketing and labeling practices.

80. Superiority: A class action is superior to the other available methods for the fair and efficient adjudication of this controversy because:

a. The joinder of thousands of individual Class Members is impracticable, cumbersome, unduly burdensome, and a waste of judicial and/or litigation resources;

b. The individual claims of the Class Members may be relatively modest compared with the expense of litigating the claims, thereby making it impracticable, unduly burdensome, and expensive—if not totally impossible—to justify individual actions;

c. When Defendants' liability has been adjudicated, all Class Members' claims can be determined by the Court and administered efficiently in a manner far less burdensome and expensive than if it were attempted through filing, discovery, and trial of all individual cases;

d. This class action will promote orderly, efficient, expeditious, and appropriate adjudication and administration of Class claims;

e. Plaintiff knows of no difficulty to be encountered in the management of this action that would preclude their maintenance as a class action;

f. This class action will assure uniformity of decisions among Class Members;

g. The Class is readily definable and prosecution of this action as a class action will eliminate the possibility of repetitious litigation;

h. Class Members' interests in individually controlling the prosecution of separate actions is outweighed by their interest in efficient resolution by a single class action; and

i. It would be desirable to concentrate in this single venue the litigation of all Class Members who were induced by Defendants' uniform false advertising to purchase their Products.

81. Accordingly, this Class is properly brought and should be maintained as a class action under Rule 23(b)(3) because questions of law or fact common to Class Members predominate over any questions affecting only individual members, and because a class action is superior to other available methods for fairly and efficiently adjudicating this controversy.

CLAIMS

FIRST CAUSE OF ACTION **VIOLATION OF NEW YORK GBL § 349** **(On Behalf of Plaintiff and New York Subclass Members)**

82. Plaintiff repeats and realleges each and every allegation contained in all the foregoing paragraphs as if fully set forth herein.

83. New York General Business Law Section 349 (“GBL § 349”) declares unlawful “[d]eceptive acts or practices in the conduct of any business, trade, or commerce or in the furnishing of any service in this state . . .”

84. The conduct of Defendants alleged herein constitutes recurring, “unlawful” deceptive acts and practices in violation of GBL § 349, and as such, Plaintiff and the New York Subclass Members seek monetary damages against Defendant, enjoining them from inaccurately describing, labeling, marketing, and promoting the Products.

85. There is no adequate remedy at law.

86. Defendants misleadingly, inaccurately, and deceptively advertise and market their Products to consumers.

87. Defendants’ improper consumer-oriented conduct—including failing to disclose that the Products have lead—is misleading in a material way in that it, *inter alia*, induced Plaintiff and the New York Subclass Members to purchase Defendants’ Products and to use the Products when they otherwise would not have. Defendants made the untrue and/or misleading statements and omissions willfully, wantonly, and with reckless disregard for the truth.

88. Plaintiff and the New York Subclass Members have been injured inasmuch as they purchased Products that were mislabeled, unhealthy, and entirely worthless. Accordingly, Plaintiff and the New York Subclass Members received less than what they bargained and paid for.

89. Defendants’ advertising and Products’ packaging and labeling induced Plaintiff and the New York Subclass Members to buy Defendants’ Products.

90. Defendants’ deceptive and misleading practices constitute a deceptive act and practice in the conduct of business in violation of New York General Business Law §349(a) and Plaintiff and the New York Subclass Members have been damaged thereby.

91. As a result of Defendants' recurring, "unlawful" deceptive acts and practices, Plaintiff and the New York Subclass Members are entitled to monetary, statutory, compensatory, treble and punitive damages, restitution, and disgorgement of all moneys obtained by means of Defendants' unlawful conduct, interest, and attorneys' fees and costs.

SECOND CAUSE OF ACTION
VIOLATION OF NEW YORK GBL § 350
(On Behalf of Plaintiff and the New York Subclass Members)

92. Plaintiff repeats and realleges each and every allegation contained in all the foregoing paragraphs as if fully set forth herein.

93. N.Y. Gen. Bus. Law § 350 provides, in part, as follows:

False advertising in the conduct of any business, trade, or commerce or in the furnishing of any service in this state is hereby declared unlawful.

94. N.Y. Gen. Bus. Law § 350a(1) provides, in part, as follows:

The term 'false advertising, including labeling, of a commodity, or of the kind, character, terms or conditions of any employment opportunity if such advertising is misleading in a material respect. In determining whether any advertising is misleading, there shall be taken into account (among other things) not only representations made by statement, word, design, device, sound or any combination thereof, but also the extent to which the advertising fails to reveal facts material in the light of such representations with respect to the commodity or employment to which the advertising relates under the conditions proscribed in said advertisement, or under such conditions as are customary or usual . . .

95. Defendants' labeling and advertisements contain untrue and materially misleading statements and omissions concerning their Products inasmuch as it misrepresents that the Products are safe for use and doesn't list that the Products contain lead.

96. Plaintiff and the New York Subclass Members have been injured inasmuch as they relied upon the labeling, packaging, and advertising and purchased Products that were mislabeled,

unhealthy, and entirely worthless. Accordingly, Plaintiff and the New York Subclass Members received less than what they bargained and paid for.

97. Defendant's advertising, packaging, and Products' labeling induced Plaintiff and the New York Subclass Members to buy Defendants' Products.

98. Defendants made their untrue and/or misleading statements and representations willfully, wantonly, and with reckless disregard for the truth.

99. Defendants' conduct constitutes multiple, separate violations of N.Y. Gen. Bus. Law § 350.

100. Defendants made the material misrepresentations described in this Complaint in their advertising and on the Products' packaging and labeling.

101. Defendants' material misrepresentations were substantially uniform in content, presentation, and impact upon consumers at large. Moreover, all consumers purchasing the Products were and continue to be exposed to Defendants' material misrepresentations.

102. As a result of Defendants' recurring, "unlawful" deceptive acts and practices, Plaintiff and New York Subclass Members are entitled to monetary, statutory, compensatory, treble and punitive damages, restitution, and disgorgement of all moneys obtained by means of Defendants' unlawful conduct, interest, and attorneys' fees and costs.

THIRD CAUSE OF ACTION
BREACH OF EXPRESS WARRANTY
(On Behalf of Plaintiff and All Class Members)

103. Plaintiff repeats and realleges each and every allegation contained in the foregoing paragraphs as if fully set forth herein.

104. Defendants provided Plaintiff and Class Members with an express warranty in the form of written affirmations of fact promising and representing that the Products are safe for use and do not contain lead.

105. Defendants omitted that the Products contain a known neurotoxin from their ingredients labeling. This omission would lead reasonable consumers did not contain a known neurotoxin, when in fact, the Products were contaminated with Lead as stated herein.

106. The above affirmations of fact were not couched as “belief” or “opinion,” and were not “generalized statements of quality not capable of proof or disproof.”

107. These affirmations of fact became part of the basis for the bargain and were material to Plaintiff and Class Members’ transactions.

108. Plaintiff and Class Members reasonably relied upon Defendants’ affirmations of fact and justifiably acted in ignorance of the material facts omitted or concealed when they decided to buy Defendants’ Products.

109. Defendants knowingly breached the express warranties by including lead in the Products sold to Plaintiff and the Class without properly notifying them of their inclusion in the Products.

110. Within a reasonable time after it knew or should have known, Defendants did not change the Products’ labels to include lead in the ingredients list or to otherwise warn consumers that the Products contain, or are at risk of containing, lead.

111. Defendants thereby breached the following state warranty laws:

- a. Code of Ala. § 7-2-313;
- b. Alaska Stat. § 45.02.313;
- c. A.R.S. § 47-2313;

- d. A.C.A. § 4-2-313;
- e. Cal. Comm. Code § 2313;
- f. Colo. Rev. Stat. § 4-2-313;
- g. Conn. Gen. Stat. § 42a-2-313;
- h. 6 Del. C. § 2-313;
- i. D.C. Code § 28:2-313;
- j. Fla. Stat. § 672.313;
- k. O.C.G.A. § 11-2-313;
- l. H.R.S. § 490:2-313;
- m. Idaho Code § 28-2-313;
- n. 810 I.L.C.S. 5/2-313;
- o. Ind. Code § 26-1-2-313;
- p. Iowa Code § 554.2313;
- q. K.S.A. § 84-2-313;
- r. K.R.S. § 355.2-313;
- s. 11 M.R.S. § 2-313;
- t. Md. Commercial Law Code Ann. § 2-313;
- u. 106 Mass. Gen. Laws Ann. § 2-313;
- v. M.C.L.S. § 440.2313;
- w. Minn. Stat. § 336.2-313;
- x. Miss. Code Ann. § 75-2-313;
- y. R.S. Mo. § 400.2-313;
- z. Mont. Code Anno. § 30-2-313;

- aa. Neb. Rev. Stat. § 2-313;
- bb. Nev. Rev. Stat. Ann. § 104.2313;
- cc. R.S.A. 382-A:2-313;
- dd. N.J. Stat. Ann. § 12A:2-313;
- ee. N.M. Stat. Ann. § 55-2-313;
- ff. N.Y. U.C.C. Law § 2-313;
- gg. N.C. Gen. Stat. § 25-2-313;
- hh. N.D. Cent. Code § 41-02-30;
- ii. II. O.R.C. Ann. § 1302.26;
- jj. 12A Okl. St. § 2-313;
- kk. Or. Rev. Stat. § 72-3130;
- ll. 13 Pa. Rev. Stat. § 72-3130;
- mm. R.I. Gen. Laws § 6A-2-313;
- nn. S.C. Code Ann. § 36-2-313;
- oo. S.D. Codified Laws, § 57A-2-313;
- pp. Tenn. Code Ann. § 47-2-313;
- qq. Tex. Bus. & Com. Code § 2.313;
- rr. Utah Code Ann. § 70A-2-313;
- ss. 9A V.S.A. § 2-313;
- tt. Va. Code Ann. § 59.1-504.2;
- uu. Wash. Rev. Code Ann. § 6A.2-313;
- vv. W. Va. Code § 46-2-313;
- ww. Wis. Stat. § 402.313; and

xx. Wyo. Stat. § 34.1-2-313.

112. As a direct and proximate result of Defendants' breach of the express warranties, Plaintiff and Class Members were damaged in the amount of the price they paid for the Products, in an amount to be proven at trial.

JURY DEMAND

Plaintiff demands a trial by jury on all issues.

PRAYER FOR RELIEF

WHEREFORE, Plaintiff, on behalf of herself and the Class, prays for judgment as follows:

- (a) Declaring this action to be a proper class action and certifying Plaintiff as the representative of the Class under Rule 23 of the FRCP;
- (b) An Order requiring Defendants to establish a blood testing program for the Plaintiff and the Class, as well as to establish a medical monitoring protocol for the Plaintiff and the Class to monitor the consumers' individual health and diagnose at an early stage any ailments associated with exposure to lead;
- (c) Awarding monetary damages and treble damages;
- (d) Awarding statutory damages of \$50 per transaction, and treble damages for knowing and willful violations, pursuant to N.Y. GBL § 349;
- (e) Awarding statutory damages of \$500 per transaction pursuant to N.Y. GBL § 350;
- (f) Awarding punitive damages;
- (g) Awarding Plaintiff and Class Members their costs and expenses incurred in this action, including reasonable allowance of fees for Plaintiff's attorneys, experts, and reimbursement of Plaintiff's expenses; and
- (h) Granting such other and further relief as the Court may deem just and proper.

Dated: December 21, 2023

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