

**Exposure to High Ultra-processed Food and Sodium Intake and  
its effect on Hypertension using the cross sectional study UK  
National Dietary and Nutritional Survey (NDNS) in England  
2008-2019**

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**Dissertation submitted in partial fulfilment of the requirements for the degree  
of Master of Public Health, The University of Liverpool**

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## Dedication

To Julie Andrew and Sophie  
for your loving patience and support

## Acknowledgments

Thanks to Zoe and Martin

Thanks to Paul for the project which didn't quite come together

## Abstract

This study looked at hypertension and exposure to high intake of UPF and high Sodium intake. This examined the assumption that it is the salt content of UPF which contributes to its 'unhealthiness' and to query the idea that 'reformulation' will generate 'healthy UPF'.

## Method

Secondary data analysis of the National Dietary and Nutrition Survey (NDNS). This rolling cross-sectional study provides stratified sample representative of the UK population.

Multivariable logistic regression analysis of high sodium intake and high UPF intake against hypertension was used. Secondary end points included regression of Sodium intake against UPF intake.

## Results

There was an increased odds ratio of hypertension with higher sodium intake, 5.43 (1.44, 20.4)  $p=0.012$ .

There was a lower odds ratio of hypertension with high upf intake. 0.57(0.35, 0.95)  $p=0.032$ .

There was no correlation between UPF intake and sodium intake.

There is a strong correlation between age and BP, as well as age and Sodium intake.

There is a strong age gradient of UPF intake.

## Conclusion

This study shows that high Na intake is associated with hypertension. Reduction of sodium intake may be effective at reducing the overall risk.

UPF intake is also associated with hypertension. This may show reverse causation in this cross-sectional study.

Policy should aim to reduce intake of sodium, longitudinal studies may be more effective at identifying the causal relationship between UPF and BP.

## Keywords

UPF, Sodium, hypertension, NOVA, Nutrition

**Abstract** 226 words

**Dissertation** 9020

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*Table of Abbreviations used*

Abbreviation	Term
NDNS	National Dietary and Nutrition Survey
BP	Blood Pressure
Na	Sodium intake in mg
UPF	Ultra Processed Foods
NCD	Non communicable Disease
CVD	Cardiovascular Disease
CHAMPs	Cheshire and Merseyside public health collaborative
NOVA	NOVA is a classification system, it is not an acronym
IMD	Index of Multiple Deprivation
BMI	Body Mass Index
AIC	Akaike Information Criterion

## Introduction

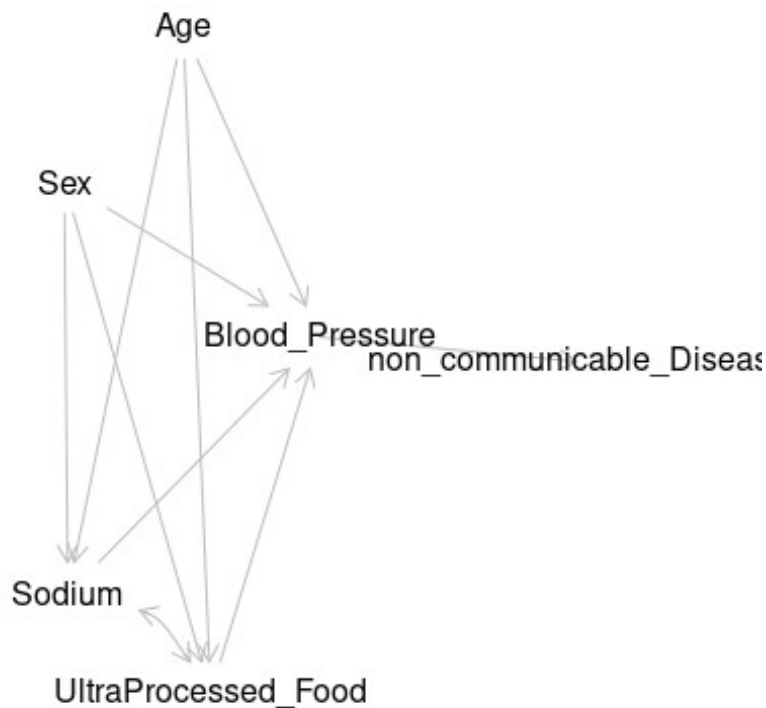
417482 people, 15.4% of the population, have hypertension in Cheshire and Merseyside (1). Cheshire and Merseyside public health collaborative (CHAMPs) have a plan to reduce blood pressure (BP) by 2029 (2). The strategy aims to increase 'awareness'. This is intended to increase individual compliance with testing and treatment of raised BP. This study intends to offer additional opportunities for improving outcomes.

UPF makes up up to 60% diet in UK especially in the North West of England. There is evidence of an association between hypertension and intake of Ultra-processed Foods (UPF) (3) and hypertension and Salt intake (4) from studies of different types in multiple countries. Monteiro et al (5) model a scenario where "halving intake of (NOVA) group 3 foods could result in approximately 22,055 fewer deaths" across the UK in 2030. Local food policies around UPF might be a way of reducing hypertension at a population level. There is potential for significant public health impact.

Marmot (6) identifies the external influences in Cheshire and Merseyside which need to be improved to permit individual action to be effective. Using a better understanding of the role of UPF, and the interaction with sodium, might give a mechanism of action of some of Marmot's categories of influence.

This study assessed prevalence of exposure to sodium and UPF, and hypertension in the data set of the National Dietary and Nutrition Survey (NDNS) (7). The study gives data from the UK from 2008 to 2019. It is stratified to be representative of the population of the UK by sex, age, region and index of multiple deprivation (IMD). BP, UPF and Sodium intake were recorded. Age and sex remain important background factors. This leads to the research question below.

I have used STROBE guidance (8) in producing this report. This study explored this complex web pulling out strands within it, [diagram 1](#) shows a possible arrangement of this.



*The relationships explored in the analysis*

## Research Question

Using PICO (9) approach,

In adults and children across the four home nations of the UK between 2008 and 2019, did exposure to high sodium dietary intake, and or high UPF dietary intake, compared to lower exposure, increase the odds of having a mean systolic blood pressure of over 140mmHg?

This primary question can be split into parts,

For a representative population across the UK What was dietary intake of UPF between 2008 and 2019? What was dietary intake of salt between 2008 and 2019? What was BP between 2008 and 2019? What was the correlation between these?

In addition it may be possible to consider, How did each of these change over that time? Is there evidence of interaction between these? Was UPF or Na most important in these changes?

## Objectives

- 1 Literature Review of UPF and BP, with Sodium
- 2 Descriptive analysis of participants from NDNS with amalgamation of data across the rolling programme.
- 3 Analysis of exposure to UPF and sodium, and prevalence of BP >140mmHg using regression models.
- 4 Discussion of implications of results in relation to limitations of cross- sectional studies, and available data, as well as suggestions for further research

## Epistemology

The epistemological approach of this study is positivist. I use a quantitative approach in a mechanistic and deterministic model. However, I am aware that this model is incomplete. Positivism encourages experimental isolation. The study is isolated from the world through control of the experimental environment.

Real world dietary change requires understanding interaction with social and economic factors, not isolation. Critical realist and social constructionist studies are needed to complement the information from this study. The commercial and social determinants of health are models which have a great deal of impact on exposure to UPF and Na and on dietary effects on BP.

## Positionality

In a positivist paradigm the observer is external to the experiment. Acknowledging the constructivist aspects to this study allows that the observer is closer to the model making my positionality of interest. Jafar (10) argues that understanding the position of the investigator be of interest to understanding this quantitative study.

I bring an attachment to positivist ideals from my biomedical background. As an older physician I am aware of social factors impacting health of participants as Evans and Trotter (11) discuss. I also understand that my perception of the world is from a position of significant privilege. To proceed, I need to be aware of the limitations of the positivist approach. I need to make pragmatic selections to bring some degree of validity to the resulting dataset.

These constructed ideas, social expectations, income, or geography affect food and health 'choices'. They also impact on 'hard' clinical measurements such as BP, through physical position and room temperature as well as by the relationship between the observer and the participant.

This work is primarily to complete requirements for an MPH degree which means that it is influenced by factors around health equity and classic epidemiology as taught on the course. It is produced in collaboration with a research group with a long established reputation in food research in public health, which may steer the results in a conservative direction.

### **University Research Governance and Ethical Review**

The ethics process for the University of Liverpool was followed and confirmation of compliance is attached at [Appendix 2 Ethics Certificate](#)

The storage of the data is in keeping with the research governance agreements of the University and the Data set owners.

## Literature Review

### Rationale

- 1 develop search
- 2 review search and confirm inclusion
- 3 describe literature
- 4 synthesise literature
- 5 critique literature
- 6 explain role of study within context

This literature review was intended as a systematic search, to identify papers with information about UPF, sodium and blood pressure informed by PRISMA (12).

The rationale for this review is to contribute background and to answering the research question;

What is the evidence that in adults and children across the four home nations of the UK between 2008 and 2019, would exposure to high sodium dietary intake, and or high UPF dietary intake, compared to lower exposure, increase the odds of having a mean systolic blood pressure of over 140mmHg?

### Method

Eligible studies were cross-sectional studies, and systematic reviews considering the relationship between the exposure, and outcome in comparable general populations. Papers were excluded where the population was specifically of one type, or had a specified health condition. Another exclusion criterion was where specific foods were considered.

Scopus (13) , Pubmed , Web of Science and Medline(ovid) (14) were searched.

The search strategy is included in [table 1](#) below. The search identified a wide variety of articles, which outlined and augmented the review. Others were identified by reading articles and cross referencing, particularly from the systematic reviews. Colleagues identified further relevant literature.

Table 1: Description of Search

search terms (("ultraprocessed food\*" OR "ultra-processed food\*" OR "ultra processed food\*" OR "NOVA food\*")OR (salt OR sodium OR "sodium intake")) AND ( "blood pressure" OR hypertension OR "cardiovascular disease" OR "cardiovascular risk")AND(cohort OR cross-section\* OR prospective OR meta-analysis OR "systematic review")

Inclusion prospective and observational studies and systematic reviews specifically on hypertension with relevant exposures

Exclusions ("renal cell cancer" OR "gastric cancer" OR "multiple sclerosis" OR dialysis OR DHA OR auto-immunity OR autoimmunity OR diarrhoea OR telomere OR crp OR "c-reactive" OR CKD OR "chronic kidney disease" OR autosomal OR geneti\* OR "Inflammatory bowel disease" OR diabetes OR atherosclerosis OR osteoporosis OR angiotensin\* OR aldosteron\* OR covid-19 OR gestation\* OR stroke OR "birth weight" OR hypertensive OR "immune mechanisms" OR "heart failure"OR taste OR "cognitive decline" OR dementia OR mortality OR validation)

Name	Scopus search	Medline (Ovid)Search	Pubmed search	Web of Science Search	Total
Date of Search	18/6/23	18/6/23	18/6/23	18/6/23	18/6/23
Number of results	240	202	103	48	593

## Selection strategy

Inclusions and exclusions are identified in the table. No time limits, language limits or availability limits were included in the search.

The papers were reviewed by the author only.

The data sought were odds ratios for the effect of UPF or sodium on blood pressure.

There is risk of bias due to the single reviewer approach.

There is little homogeneity of approach to reporting exposure or outcome.



This made it difficult to compare items directly.

### Search results

The search identified 593 papers, 348 after removing duplicates. [Chart 1](#) shows how these were assessed.

These were then reviewed by comparing the titles against the inclusion and exclusion criteria leaving 56. 6 systematic reviews were identified, one which considered both UPF and sodium intake.

## Discussion of literature

### *Table 2: Literature results*

First Author	Type of Study	Subject of Study	Results
Barbosa (15)	systematic review	UPF and Sodium and Hypertension	6 relevant articles
Wang (16)	systematic review and meta analysis	UPF and hypertension	9 articles (odds ratio: 1.23; 95% CI: 1.11, 1.37; P=0.034)
Mambrini (17)	systematic review	UPF and CVD	4/17 UPF and BP
D'Elia (18)	meta-analysis	Salt and CVD	<i>a 2.84% (95% CI 0.51–5.08) reduction in PWV</i>
Leyvraz (19)	meta-analysis	children sodium blood pressure	13/18 studies
Frias (20)	systematic review of cohorts	children and BP	2 studies BP no association systolic BP
Shim (21)	cross-sectional	UPF and Hypertension	
Scaranni (22)	cohort	UPF and Hypertension	
Zhang (23)	cross sectional	UPF and 'CVH'	
Rezende-Alves (24)	cohort	UPF and Hypertension	
Martinez-Peres Monge Harrington (25) Conceicao Asma (26) d'Avila (27) de Deus Mendonca (28) middle aged uni grads	cohort		
	cohort	UPF and (self - reported) hypertension	adjusted HR, 1.21; 95% CI, 1.06, 1.37; P for trend = 0.00
Ming ( <b>mingli?</b> )	cohort	UPF and hypertension	
He (29)	meta-analysis	Salt and	

		hypertension	
Strazzullo (30)	meta-analysis	Salt and CVD	
Graudal (31)	meta-analysis	Salt and CVD	ACM: $HR = 1.16$ , 95% $CI = 1.03-1.30$ ; CVDEs: $HR = 1.12$ , 95% $CI = 1.02-1.24$
Ma (32)	meta-analysis	Salt and CVD	

## UPF and BP

Three meta analyses those of Barbosa et al, Wang et al and Mambrini et al highlight the risk of hypertension with high UPF. Mambrini et al identify that few papers attempt to link UPF with hypertension. Their systematic review identifies Monge, Scaranni, and Mendonca.

Wang provided a systematic review and meta analysis which included these and ivancovsky,

Scaranni used Brazil's ELSA study, in middle aged civil servants

Mendonca in Spain found an affect on hypertension

Shim demonstrated the effect on hypertension in korea

Zhang et al. have demonstrated a reverse gradient between UPF intake and CVH\* using the USA NHANES data. cvh seems to be BMI and diabetes, cholesterol has an opposing affect, and BP seems to zeroed

## UPF and other endpoints

The most extensive work in the most similar line is that of Rauber et al. They have used NDNS to compare nutrients UPF intake and outcomes. Her studies 2018-2021 have looked at the nutrition make up of the UPF diet in the UK, looked at obesity, and at metabolic outcomes. Using sex referenced UPF intake quintiles she has identified some linear associations . ultraprocessed data? She primarily studied NDNS data, great because its comprehensive, and available, not so good as it is cross sectional and has no follow up.

Filou and Srour in France as part of Nutrinet Sante similarly examine the relationship between UPF and CVD

## Salt and CVD

There is more literature on sodium and hypertension.

The systematic reviews identified in the search include Barbosa, and D'Elia. Leyvraz and Frias concentrate on effects in children.

He and Graudal were not identified in the search but were referenced in the other papers.

He identifies reduction in BP with reduction in sodium intake. Graudal reports that sodium reduction can go too far, identifying a 'j'-shaped curve. He directly criticises Graudal's work. The inclusion of papers with too big effect sizes, and too short follow up is to blame.

D'Elia et al look at arterial stiffness pressure wave velocity 'PWV' and show that this increases with salt intake. This arterial stiffness is potentially more sensitive to sodium intake than BP.

*"Eleven studies met the predefined inclusion criteria and provided 14 cohorts with 431 participants and 1–6 weeks intervention time. In the pooled analysis, an average reduction in sodium intake of 89.3mmol/day was associated with a 2.84% (95% CI 0.51–5.08) reduction in PWV."*

D'Elia's results show that BP is less accurately predicted than arterial stiffness. This may be a cause of the equivocal results found by studies looking at BP.

Graudal et al. Studied cohort studies as there were no RCTs of increased sodium intake.

*"Data from 23 cohort studies and 2 follow-up studies of RCTs (n = 274,683) showed that the risks of ACM and CVDEs were.... increased in high sodium vs. usual sodium intake (ACM: HR = 1.16, 95% CI = 1.03-1.30; CVDEs: HR = 1.12, 95% CI = 1.02-1.24)."*

Their findings identify that there might be 'too much' salt reduction possible. They provide an explanation as to how low sodium levels may causes issues.

Straluzzo et al identifies He et al (29) as a source for hypertension and salt.

## Literature review Conclusion

Of the identified papers the majority describe 2/3 elements, UPF or salt intake or BP/ CVD .

They show an odds ratio for cvd with raised sodium intake and hypertension with high UPF intake. The qualification is that BP is an uncertain outcome measurement. Papers often look to CVD outcomes as stronger endpoints.

Where UPF exposure has been studied with hypertension as an endpoint the cross sectional studies have identified a link in adults, but sometimes there is an inverse gradient.

This study aims to identify these two effects within a large representative cross sectional population. The relationship between these two effects can be shown by studying both in the same population. This study also gives the opportunity to consider if there are associated factors.

## Method

### Study Setting and Design

This is a secondary data analysis of data from the National Dietary and Nutritional Survey (NDNS (7)). This analysis intends to analyse the association between sodium intake, UPF intake and BP.

The NDNS was commissioned in collaboration between government departments responsible for health and for food production. Academic partners delivered reports on diet and nutrition across the United Kingdom. The study is designed to be representative across the four home nations, and across age with balanced representation for children. NDNS data are available via the UK national Data service for research purposes.

NDNS is a rolling cross-sectional study, in each year a new cross section of participants is enrolled from the wider population. Questionnaires, food diaries, and nurse assessments are used to gather data. It has been running since 2008. The most recent data is available from 2019.

### Participants, Inclusion and Exclusion

Participants were identified by random selection across postal units. The sample is stratified to ensure a representative sample across the four nations (England, Wales, Scotland, and Northern Ireland) and across regions in England (North, Central/Midlands, South(including London)). The sample is also stratified for age and sex and Index of multiple deprivation (IMD).

For NDNS the intended sample is 1000 per year with 50% adults. Each year the sample is slightly different due to differential uptake. Oversampling is used to control this.

The relationship between salt and systolic blood pressure may be different in individuals with pathologically high BP. Those taking BP controlling medications may have a different relationship to sodium and UPF and so were excluded for analysis.

### Exposure Variables

The participants recorded their food intake as a food diary. Four days of the diary including a weekend day are used for each participant. They record food and portion size as well as where food was eaten.

Based on the food and drink intake reported and with a composition data table, the NDNS team have estimated the daily intake of food by group, and large range of nutrients including sodium.

## Sodium estimation

This analysis used the daily sodium intake in mg from NDNS. This value reflects the expected content of standard foods.

Serum sodium values are available for the early dataset, but not the later one. 24 urinary sodium is a better indicator of dietary sodium but values are not available across the whole time period.

A categorical variable, has been produced with a cut off values at 3000mg, 5000mg and 6000mg. These values are the WHO recommended amount and match values used in Du et al (3).

## UPF

The NOVA classification, developed by Monteiro et al. (33), was used to estimate the intake of UPF. There is no record of NOVA classification in NDNS. The dataset provided by Dr Colombet (personal communication) was used to identify food by NOVA group. This was developed by comparing every food level entry in NDNS against NOVA. A standard methodology describing the approach used has been published by Martinez-Steele et al. (34) .

Next the energy content of the day's food was calculated by NOVA group. This was added to the intake for the other 3 days and the total intake by NOVA group established. The percentage of the total intake of energy was then calculated for each of the 4 Nova categories. Nova group 4 or UPF intake (UPF) is used for this study.

A variable (UPF3) was developed from the mean UPF intake. The central category is the mean with one standard deviation above and below. This effectively identifies 67% in the centre of the distribution. Categories used in other papers eg (35) are low for the UK.

## Outcome Variable

BP is a quality assured mean systolic BP which is reliable across the dataset. It was measured in mmHg using a calibrated automatic sphygmomanometer by a study nurse under specified conditions. These conditions controlled for the effects of exercise, temperature and ill health. The data on all these is in the dataset. Raw BP values are also present in the dataset to allow quality review.

I have created a categorical variable which identifies participants with BP over 140 mmHg to enable logistic regression. This value is identified by Du et al ((3)) and others, though some use lower values such as 120mmHg ( (36) )

## Other Variables

Additional explanatory variables are ones which can also influence BP. They include Age, Sex, and BMI. Age at completion of education, and IMD are also used. These may have effects such as confounding, mediation, and obstruction within the analysis.

Stratification is used in the design and sampling stage to modify for these. Including them in multivariable regression attempts to prevent them confounding the analysis.

## Bias

Selection bias was approached by using random selection of participants using a carefully constructed stratification model. Addresses were selected by postal units to ensure geographic spread of participants. This ensured that whilst random the sample remained representative.

Take up and Drop out bias was approached by ensuring that sample sizing included scope for this to enable comparable sample sizes across annual waves.

Social desirability bias acknowledges that participants remember and record intake framed by their beliefs about the needs of the study, and their beliefs about what is perceived as being healthy. To examine this, in the first wave a double labelled water study was incorporated. This compared reported energy intake with measured values (37).

Finally bias at the analysis stage used weighting to standardise the sample for several variables. Those selected were Age, Sex, region and IMD. Weights are available for different levels of analysis as participants who did not complete the initial interview were not selected for subsequent blood analysis.

Sensitivity analysis looked to see how the variables changed over the survey years after weighting. This is intended to assess changes in sampling over the course of the study.

## Study Size

A sample size calculation for this secondary analysis is available in appendix 1 the initial proposal from OpenEpi (38). This calculated the sample size of 3526, with a ratio of 0.75 unexposed to exposed. An intended power of 80%, at a level of statistical significance of 95% was used. An odds ratio of 1.2 was used based on a meta-analysis by Wang et al (39).

## Statistical Methods

Four data batches of data (2008-2012, 2013-2014, 2015-2016, 2017-2019) were combined. The data was read using 'r-studio' with the processing being carried out using packages (see appendix 3) available from CRAN (40). In particular the package 'survey' (41) was used to manage weighted data. Generated weighting values account for



differences uptake and drop out across the annual cohorts. 'Survey' also accounts for sample stratification.

Data was analysed for correlation by regression. Categorical data was analysed using chi-squared. In all analysis P.values and confidence intervals were calculated and a value of  $p < 0.05$  was taken as the threshold of statistical significance. Multivariable regression models were constructed to manage variables which might have confounding effects on the outcome of the analysis. Sample stratification was also used to reduce potential confounding. Tables of results were produced to best demonstrate the data. For the main results a set of multivariable logistic regression models was developed. Each exposure variable was modelled separately, the final model included both of the exposure variables. AIC was used to understand the relative importance of variables.

## Results

### Participants

Considering participants who opted in and completed questionnaires, the whole NDNS population was 15,655. The median age was 40. Categorising age shows that 22% of the population was between 19 and 35. There were 49% male participants.

After excluding those on medication, the population was 14217 participants.

This table [table 4.1.1](#) shows the participants.

Continuous variables are represented by the median and interquartile range in brackets. Categorical variables give the number of participants and the percentage of the sample in brackets.

*Characteristics of the sample population from National Dietary and Nutrition Study (2008-2019)*

	Whole Populati on	Population with those on BP medication excluded	UPF >63%	Na >5000 mg	hyp >140m mHg
<b>Characteristic</b>	<b>N = 15,655<sup>1</sup></b>	<b>N = 14,217<sup>1</sup></b>	<b>N = 4,793<sub>1</sub></b>	<b>N = 73<sup>1</sup></b>	<b>N = 876<sup>1</sup></b>
Sex					
Male	7,699 (49%)	6,992 (49%)	2,568 (54%)	58 (80%)	505 (58%)
Female	7,956 (51%)	7,225 (51%)	2,225 (46%)	15 (20%)	371 (42%)
Age	40 (22, 58)	37 (20, 54)	23 (12, 42)	31 (22, 39)	60 (48, 70)
agegad3					
(0,18]	3,284 (21%)	3,278 (23%)	1,970 (41%)	4 (5.5%)	12 (1.3%)
(18,35]	3,544 (23%)	3,529 (25%)	1,275 (27%)	44 (61%)	67 (7.7%)
(35,50]	3,355 (21%)	3,241 (23%)	799 (17%)	21 (28%)	177 (20%)
(50,65]	2,912 (19%)	2,475 (17%)	418 (8.7% )	1 (1.8%)	314 (36%)
(65,108]	2,561 (16%)	1,692 (12%)	330 (6.9% )	3 (4.0%)	307 (35%)
educfinh					
Not yet finished	375 (2.9%)	375 (3.2%)	185 (5.1% )	2 (2.4%)	1 (0.2%)
Never went to school	41 (0.3%)	29 (0.2%)	0 (<0.1 %)	0 (0%)	0 (0%)
14 or under	504 (3.9%)	345 (2.9%)	89 (2.5%)	2 (2.4%)	57 (7.2%)

	Whole Populati on	Population with those on BP medication excluded	UPF >63%	Na >5000 mg	hyp >140m mHg
<b>Characteristic</b>	<b>N = 15,655<sup>1</sup></b>	<b>N = 14,217<sup>1</sup></b>	<b>N = 4,793<sub>1</sub></b>	<b>N = 73<sup>1</sup></b>	<b>N = 876<sup>1</sup></b>
			)		
15	1,773 (14%)	1,426 (12%)	472 (13%)	5 (8.3%)	186 (24%)
16	3,483 (27%)	3,160 (27%)	1,180 (33%)	24 (36%)	188 (24%)
17	1,074 (8.3%)	974 (8.3%)	332 (9.2%)	2 (2.5%)	60 (7.6%)
			)		
18	1,588 (12%)	1,484 (13%)	482 (13%)	7 (11%)	78 (9.9%)
19 or over	4,172 (32%)	3,922 (33%)	878 (24%)	25 (38%)	218 (28%)
Unknown	2,645	2,502	1,174	8	89
IMD					
Most deprived	2,977 (19%)	2,748 (19%)	1,139 (24%)	28 (39%)	112 (13%)
2	3,128 (20%)	2,870 (20%)	1,086 (23%)	21 (28%)	169 (19%)
3	2,905 (19%)	2,609 (18%)	850 (18%)	15 (20%)	136 (16%)
4	3,269 (21%)	2,953 (21%)	914 (19%)	5 (7.2%)	210 (24%)
least deprived	3,372 (22%)	3,031 (21%)	804 (17%)	4 (5.2%)	247 (28%)
Unknown	5	5	0		2
region					
England: North	3,684 (24%)	3,313 (23%)	1,231 (26%)	34 (46%)	238 (27%)

	Whole Populati on	Population with those on BP medication excluded	UPF >63%	Na >5000 mg	hyp >140m mHg
<b>Characteristic</b>	<b>N = 15,655<sup>1</sup></b>	<b>N = 14,217<sup>1</sup></b>	<b>N = 4,793<sub>1</sub></b>	<b>N = 73<sup>1</sup></b>	<b>N = 876<sup>1</sup></b>
England: Central/Midland s	2,512 (16%)	2,266 (16%)	834 (17%)	14 (20%)	150 (17%)
England: South(including London)	6,958 (44%)	6,363 (45%)	1,861 (39%)	14 (19%)	329 (37%)
Scotland	1,302 (8.3%)	1,181 (8.3%)	439 (9.2%) )	8 (12%)	78 (8.9%)
Wales	753 (4.8%)	682 (4.8%)	247 (5.2%) )	1 (1.5%)	62 (7.1%)
Northern Ireland	447 (2.9%)	413 (2.9%)	181 (3.8%) )	2 (2.1%)	20 (2.3%)
SurveyYear					
1	1,459 (9.3%)	1,323 (9.3%)	481 (10%)	12 (16%)	100 (11%)
2	1,429 (9.1%)	1,284 (9.0%)	496 (10%)	7 (10%)	83 (9.5%)
3	1,372 (8.8%)	1,246 (8.8%)	472 (9.9%) )	13 (18%)	92 (10%)
4	1,432 (9.1%)	1,291 (9.1%)	495 (10%)	6 (8.4%)	94 (11%)
5	1,485 (9.5%)	1,361 (9.6%)	461 (9.6%) )	6 (8.5%)	86 (9.8%)
6	1,362 (8.7%)	1,234 (8.7%)	473 (9.9%) )	1 (1.7%)	75 (8.6%)
7	1,442	1,312 (9.2%)	421	10	92

	Whole Populati on	Population with those on BP medication excluded	UPF >63%	Na >5000 mg	hyp >140m mHg
Characteristic	<b>N = 15,655<sup>1</sup></b>	<b>N = 14,217<sup>1</sup></b>	<b>N = 4,793<sub>1</sub></b>	<b>N = 73<sup>1</sup></b>	<b>N = 876<sup>1</sup></b>
	(9.2%)		(8.8% )	(14%)	(11%)
8	1,405 (9.0%)	1,276 (9.0%)	378 (7.9% )	5 (6.6%)	75 (8.5%)
9	1,444 (9.2%)	1,305 (9.2%)	362 (7.6% )	5 (6.8%)	81 (9.3%)
10	1,481 (9.5%)	1,360 (9.6%)	375 (7.8% )	1 (1.6%)	98 (11%)
11	1,345 (8.6%)	1,226 (8.6%)	379 (7.9% )	6 (8.1%)	0 (0%)

<sup>1</sup>n (%); Median (IQR)

## Descriptive Data

The study population the median age was 38. The largest age group was 18-35. 49% of the participants were male.

The [table 4.1.1](#) shows the population exposed to UPF >63% of their calories is made up of 4793 from a total of 14217 participants. This compares with 9424 participants with lower exposure. That is an exposure prevalence of 34%.

High UPF is more common in younger males than in the overall population or those not on medication. 37% are 0-16 years old. There is a gradient in deprivation with more exposure in the most deprived group. England's South East has 13% of those with high UPF intake. With 11% in the North West.

The population exposed to Na >5000mg has only 73 out of 14217 participants. An exposure frequency of 0.5%. 61% are 18-35, and 80% male. The least deprived makes up 5.2% of the participants compared with 39% of the most deprived. The north has 46% of those with high Na intake, much the highest.

## Sensitivity to Survey Year

Internal consistency was examined by comparing background data across survey years. This might also be called sensitivity analysis by survey year. General linear regression modelling was used, with wave 1 as a comparator for analysis of the other waves. Age was a variable which the whole sample was weighted to be maintained across the waves. BMI was lower in most years but this was statistically significant only in wave 7 and 8.

Table 4.2.2 shows results for continuous variables.

*Sensitivity of Age and BMI by survey year from National Dietary and Nutrition Study (2008-2019)*

Group	Characteristic	Beta	95% CI <sup>1</sup>	p-value
Age	SurveyYear			0.3
	1	—	—	
	2	0.70	-1.5, 2.9	
	3	-0.21	-2.2, 1.8	
	4	1.8	-0.29, 4.0	
	5	2.0	-0.16, 4.1	
	6	0.14	-1.9, 2.2	
	7	1.6	-0.71, 4.0	
	8	0.94	-1.1, 3.0	
	9	1.5	-0.68, 3.7	
	10	1.5	-0.68, 3.7	
	11	2.0	-0.27, 4.2	
BMI	SurveyYear			<0.001
	1	—	—	



Group	Characteristic	Beta	95% CI <sup>1</sup>	p-value
	2	0.06	-0.56, 0.69	
	3	-0.32	-0.93, 0.29	
	4	0.43	-0.18, 1.0	
	5	-0.35	-0.94, 0.24	
	6	-0.41	-1.0, 0.23	
	7	-2.1	-3.0, -1.3	
	8	-2.1	-3.0, -1.2	
	9	-0.32	-0.93, 0.30	
	10	-0.18	-0.79, 0.43	
	11	-0.35	-0.94, 0.25	

<sup>1</sup>CI = Confidence Interval

The categorical data also had p.values >0.05 for the controlled variables (age, sex, IMD) against annual wave. UK region is part of the weighting, but this sample showed variation with p.value <0.05.

Table 4.2.3 follows.

*Sensitivity of categorical variables to survey year from National Dietary and Nutrition Study (2008-2019)*

Variable	ChiSq <sup>1</sup>	p.value
Sex	0.90	0.53
IMD	0.86	0.71
Age	0.88	0.66

Variable	ChiSq <sup>1</sup>	p.value
BMI	0.78	0.77
Region	2.55	0.00
Sex	1.97	0.53

<sup>1</sup>Chi Squared for categorical data

The exposure variables were compared across annual waves in [Table 4](#) . Most people have sodium exposure less than 3000mg. In year one this was 81%, with only 0.9% exposed to more than 5000mg. By year eleven 92% are reporting less than 3000mg, and 0.5% over 500mg.

UPF exposure was steady with 37%-40% of the participants exposed to 45%-63% throughout the survey. Up to 38.2% of the participants were exposed to levels of more than 63%, the peak being in year 6.

*Exposure by survey year from National Dietary and Nutrition Study (2008-2019)*

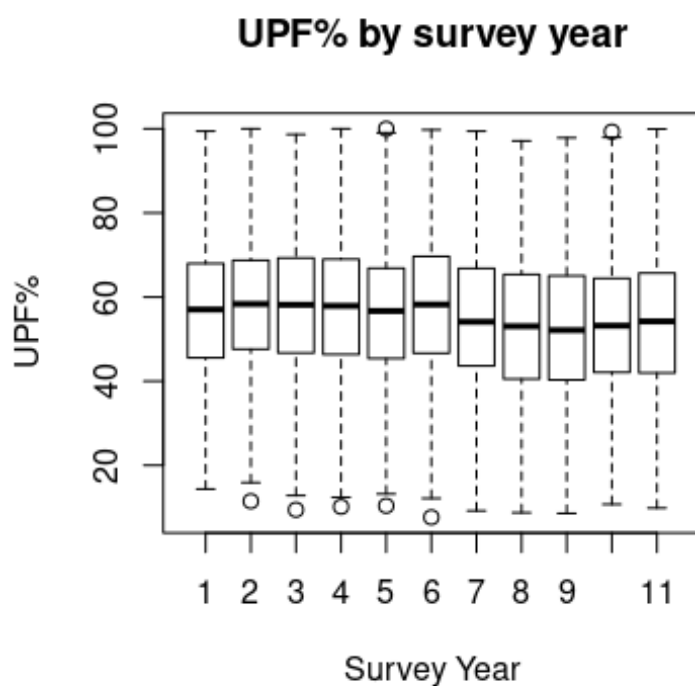
<b>Chara- cteris- tic</b>	<b>1, N = 1,32 3<sup>1</sup></b>	<b>2, N = 1,28 4<sup>1</sup></b>	<b>3, N = 1,24 6<sup>1</sup></b>	<b>4, N = 1,29 1<sup>1</sup></b>	<b>5, N = 1,36 1<sup>1</sup></b>	<b>6, N = 1,23 4<sup>1</sup></b>	<b>7, N = 1,31 2<sup>1</sup></b>	<b>8, N = 1,27 6<sup>1</sup></b>	<b>9, N = 1,30 5<sup>1</sup></b>	<b>10, N = 1,36 0<sup>1</sup></b>	<b>11, N = 1,22 6<sup>1</sup></b>	<b>p- va- lu- e<sup>2</sup></b>
hiNa												< 0. 00 1
(0,1. 5e+0 3]	253 (19 %)	232 (18 %)	240 (19 %)	301 (23 %)	362 (27 %)	330 (27 %)	366 (28 %)	396 (31 %)	410 (31 %)	450 (33 %)	365 (30 %)	
(1.5e +03, 3e+0 3]	820 (62 %)	858 (67 %)	799 (64 %)	837 (65 %)	861 (63 %)	779 (63 %)	821 (63 %)	752 (59 %)	775 (59 %)	794 (58 %)	760 (62 %)	
(3e+ 03,5 e+03 ]	238 (18 %)	187 (15 %)	193 (16 %)	147 (11 %)	131 (9.6 %)	123 (10 %)	114 (8.7 %)	123 (9.6 %)	114 (8.7 %)	114 (8.4 %)	94 (7.7 %)	
(5e+ 03,6 e+03 ]	11 (0.9 %)	0 (<0. 1%)	13 (1.0 %)	4 (0.3 %)	4 (0.3 %)	1 (0.1 %)	7 (0.6 %)	3 (0.2 %)	4 (0.3 %)	0 (0%)	6 (0.5 %)	
(6e+ 03,1 e+04 ]	0 (<0. 1%)	7 (0.5 %)	0 (0% )	2 (0.2 %)	2 (0.1 %)	0 (0% )	3 (0.2 %)	2 (0.1 %)	1 (<0. 1%)	1 (<0. 1%)	0 (0%)	
zPF3												< 0. 00 1
(0,33 ]	90 (6.8 %)	61 (4.8 %)	81 (6.5 %)	86 (6.7 %)	123 (9.1 %)	92 (7.4 %)	111 (8.5 %)	164 (13 %)	159 (12 %)	143 (11 %)	135 (11 %)	
(33,4 5]	226 (17 %)	207 (16 %)	183 (15 %)	207 (16 %)	207 (15 %)	193 (16 %)	258 (20 %)	266 (21 %)	294 (23 %)	301 (22 %)	236 (19 %)	
(45,6 3]	526 (40 %)	520 (40 %)	509 (41 %)	504 (39 %)	569 (42 %)	475 (39 %)	522 (40 %)	467 (37 %)	490 (38 %)	540 (40 %)	476 (39 %)	
(63,8	408	404	353	403	385	378	331	319	298	333	327	

Characteris tic	1, N = 1,32 3 <sup>1</sup>	2, N = 1,28 4 <sup>1</sup>	3, N = 1,24 6 <sup>1</sup>	4, N = 1,29 1 <sup>1</sup>	5, N = 1,36 1 <sup>1</sup>	6, N = 1,23 4 <sup>1</sup>	7, N = 1,31 2 <sup>1</sup>	8, N = 1,27 6 <sup>1</sup>	9, N = 1,30 5 <sup>1</sup>	10, N = 1,36 0 <sup>1</sup>	11, N = 1,22 6 <sup>1</sup>	p- va lu e <sup>2</sup>
0]	(31 %)	(31 %)	(28 %)	(31 %)	(28 %)	(31 %)	(25 %)	(25 %)	(23 %)	(25 %)	(27 %)	
(80,1 00]	73 (5.5 %)	92 (7.2 %)	119 (9.6 %)	92 (7.1 %)	77 (5.6 %)	95 (7.7 %)	90 (6.8 %)	58 (4.6 %)	64 (4.9 %)	42 (3.1 %)	52 (4.2 %)	
Unkn own	0	0	0	0	0	0	1	0	0	0	0	

<sup>1</sup>n (%)

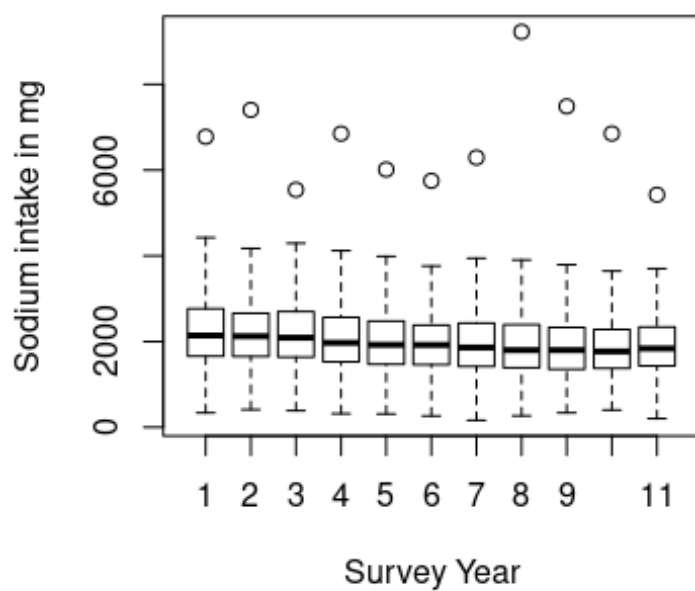
<sup>2</sup>chi-squared test with Rao & Scott's second-order correction

Results were illustrated by plots against survey year. [figure 4](#) showed overlap between the waves for UPF intake. [figure 5](#) showed sodium exposure overlap between waves.



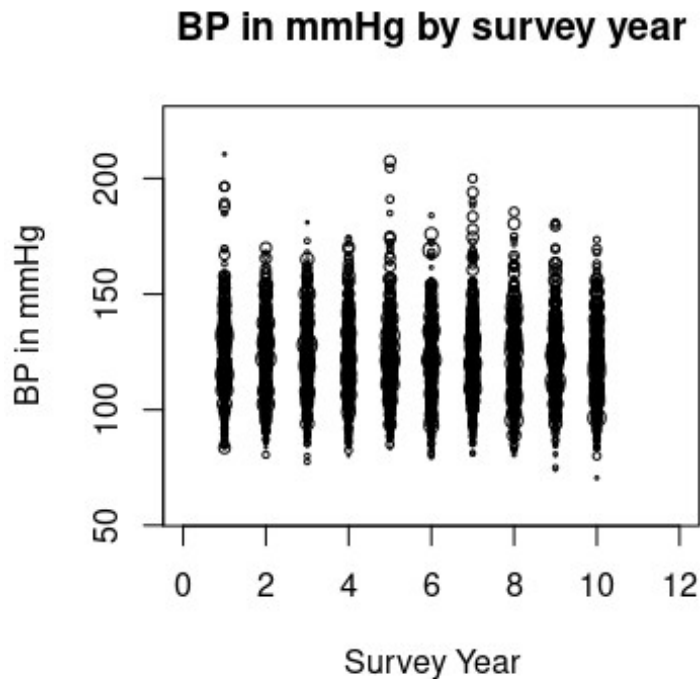
*Energy from UPF% in each annual cohorts NDNS (2008-2019)*

### Sodium Intake in mg by survey year



*Na in mg in each annual cohort NDNS(2008-2019)*

## Outcome variable



*Plot of the BP in mmHg by year from NDNS (2008-2018)*

figure 6 shows that the mean BP is consistent across the waves.

The table 3 shows the population with BP > 140 mmHg is 876 participants. This gives a prevalence of 6%. Men are once again overrepresented. These participants are older than the population median, 44% of these participants are over 65. This group are statistically significantly different from the populations with high Na, or high UPF! There is a reverse gradient with IMD in this population. The most deprived are least represented in this population. The largest proportion are in the least deprived category. The north is second highest in raised BP after the south east. The BP was highest in year one with 125 mmHg, and the lowest 120 mmHg in year 6. BP rose through life to a mean of 134 mm Hg in the over 65 age category.

*BP by survey year from National Dietary and Nutrition Study (2008-2019)*

<b>Char acter istic</b>	<b>1, N = 1,32 3<sup>1</sup></b>	<b>2, N = 1,28 4<sup>1</sup></b>	<b>3, N = 1,24 6<sup>1</sup></b>	<b>4, N = 1,29 1<sup>1</sup></b>	<b>5, N = 1,36 1<sup>1</sup></b>	<b>6, N = 1,23 4<sup>1</sup></b>	<b>7, N = 1,31 2<sup>1</sup></b>	<b>8, N = 1,27 6<sup>1</sup></b>	<b>9, N = 1,30 5<sup>1</sup></b>	<b>10, N = 1,36 0<sup>1</sup></b>	<b>11, N = 1,22 6<sup>1</sup></b>	<b>p- va lu e<sup>2</sup></b>
hyp												
(0,1 40]	630 (87 %)	605 (88 %)	573 (86 %)	569 (86 %)	760 (90 %)	600 (89 %)	650 (89 %)	671 (90 %)	670 (89 %)	744 (89% )	0 (NA %)	
(14 0,3 00]	95 (13 %)	80 (12 %)	90 (14 %)	93 (14 %)	81 (9.6 %)	72 (11 %)	83 (11 %)	75 (10 %)	80 (11 %)	94 (11% )	0 (NA %)	
Unk no wn	598	599	583	629	520	561	579	530	555	522	1,22 6	

<sup>1</sup>n (%)

<sup>2</sup>chi-squared test with Rao & Scott's second-order correction



## Main Results

The main results are the correlation between the exposure and the outcome variables. These are presented using 2\*2 tables with no additional control for confounding. Univariable regression demonstrated the interaction with other explanatory variables. This is complicated by the effect of stratification.

Chi squared table The odds ratio for hypertension in participants exposed to UPF >63% is 0.5. The odds ratio for hypertension in participants exposed to Na >5000mg is 1.45

The regression model for Sodium against BP shows that there is no linear relationship between Sodium and BP in this table [Table 4.4.1](#). UPF compared to Na also shows a zero beta value indicating no linear relationship.

UPF does show a negative relationship with age, which is statistically significant. There is also a negative relationship with Age, again statistically significant.

Age has a relationship with BP with a statistically significant positive gradient. There is also a positive relationship with Na, which is also statistically significant to the 95% level.

*Univariable Regression (NDNS data 2008-2019)*

Characteristic	BP			UPF			Sodium		
	Beta	95% CI <sup>1</sup>	p-value	Beta	95% CI <sup>1</sup>	p-value	Beta	95% CI <sup>1</sup>	p-value
zPF	-0.19	-0.22, -0.15	<0.001						
Na	0.00	0.00, 0.00	<0.001	0.00	0.00, 0.00	<0.001			
Age	0.43	0.41, 0.45	<0.001	-0.25	-0.26, -0.23	<0.001	1.5	0.77, 2.3	<0.001
agegad3									
(0,18]	—	—		—	—		—	—	
(18,35]	11	9.6, 12	<0.001	-9.1	-10, -8.0	<0.001	540	485, 595	<0.001
(35,50]	14	13, 15	<0.001	-13	-14, -12	<0.001	408	363, 454	<0.001
(50,65]	22	20, 23	<0.001	-17	-18, -16	<0.001	250	204, 295	<0.001
(65,108]	27	25, 29	<0.001	-14	-15, -13	<0.001	91	45, 138	<0.001
bmival	1.0	0.92, 1.1	<0.001	-0.29	-0.35, -0.23	<0.001	17	14, 19	<0.001
IMD									
Most deprived	—	—		—	—		—	—	
2	2.2	0.49, 4.0	0.012	-1.6	-3.0, -0.18	0.027	47	-31, 126	0.2
3	2.3	0.69, 4.0	0.005	-3.5	-4.8, -2.1	<0.001	77	2.0, 151	0.044
4	3.1	1.5, 4.8	<0.001	-4.2	-5.6, -2.9	<0.001	86	13, 159	0.021

Characteristic	BP			UPF			Sodium		
	Beta	95% CI <sup>1</sup>	p-value	Beta	95% CI <sup>1</sup>	p-value	Beta	95% CI <sup>1</sup>	p-value
least deprived	3.3	1.5, 5.0	<0.001	-5.3	-6.6, -4.0	<0.001	4.6	-60, 69	0.9

<sup>1</sup>CI = Confidence Interval

Multivariable regression models were constructed. They are regressed against hypertension in patients who are not on BP reducing medication.

The model, “Sodium Only”, includes sodium as the exposure variable. The odds ratio for the group taking between 5000mg and 6000mg per day is statistically significantly different from those taking less than 3000mg per day. There is an odds ratio of 5.07 for this group.

“UPF only” shows a significant difference in odds ratio 0.61 for the group 63-80% with a p.value of 0.049.

The last model, “Sodium and UPF”, shows that when combined the effect remains. The odds ratio for 5000-6000mg of Na remains statistically significant. The odds ratio for UPF also remains. These are both changed from the separate models. The Akaike Inclusion coefficient (AIC) is lower for this combined model indicating it is a better fit for the data also.

Table 4.5.1 follows below.

	Na only			UPF only			Na and UPF		
Characteristic	OR <sub>1</sub>	95% CI <sup>1</sup>	p-value	OR <sub>1</sub>	95% CI <sup>1</sup>	p-value	OR <sub>1</sub>	95% CI <sup>1</sup>	p-value
Sodium Intake mg									
(0,1.5e+03]	—	—					—	—	
(1.5e+03,3e+03]	0.99	0.69, 1.44	>0.9				1.05	0.72, 1.52	0.8
(3e+03,5e+03]	1.25	0.75, 2.09	0.4				1.38	0.82, 2.33	0.2
(5e+03,6e+03]	5.20	1.39, 19.5	0.015				5.57	1.47, 21.2	0.012
(6e+03,1e+04]	0.00	0.00, 0.00	<0.001				0.00	0.00, 0.00	<0.001
Ultraprocessed % Food									
(0,33]				—	—		—	—	
(33,45]				0.86	0.54, 1.36	0.5	0.83	0.52, 1.32	0.4
(45,63]				0.73	0.48, 1.12	0.15	0.70	0.46, 1.08	0.11
(63,80]				0.60	0.36, 0.99	0.046	0.57	0.34, 0.94	0.029
(80,100]				0.74	0.27, 2.05	0.6	0.69	0.24, 1.94	0.5
AIC	3594.65			3598.17			3590.81		

<sup>1</sup>OR = Odds Ratio, CI = Confidence Interval

All models include additional variables Sex, Age, BMI, Education, IMD and Survey Year

### Relative Effect Size calculation

Using the AIC statistic assessing goodness of fit for each model gives another way of understanding the comparative effects between variables. The lowest scored model is the optimal model. The 'best' of these models is that with only sodium included "Na only". The UPF models both being further away from the lowest value.

Of the difference between the lowest scoring model and the highest 70/30 is due to the sodium. the additional effect of UPF is in proportion to this.

## Discussion

### Key Results

The chi squared test shows a statistically significant correlation between UPF and hypertension. This simple model does not allow for confounding or additional variables to be taken into account.

The multivariable logistic regression controls a number of additional variables. Sex and age probably contribute the largest effects. These models show a statistically significant correlation between high Na intake and hypertension, and between high UPF and hypertension. This is present with each variable independently and with both in the model.

There is a reduction in reported intake over time, suggesting either actual reduction or efficacy of message/tax/policy

There is a slight reduction in UPF % over time, this might be reporting, or assessment as there were some changes to understanding and analysis.

The UK remains one of the highest % and so the reduction might be behaviour of a saturated market the USA or Canada might behave similarly. Other European countries are still fighting to retain a different food culture. Countries in the rest of the world are at differing levels of transition. This might be influenced by the degree of 'westernisation'/'internationalization' / or 'capitalist colonialisaton' into local culture.

There is a clear association with high sodium intake and hypertension. This remains an issue with UPF. There appear to be additional interactions as the multivariable regression shows a further effect when high UPF is added to the model.

UPF has a negative association with BP. This would fit with Zhang's finding in 'CVH'. It might also explain the curious absence of reporting in the many cross-sectional studies of BP and UPF. Rauber has published 3-4 papers on NDNS without mentioning salt or BP!

Is the explanation of the finding in cross section due to the nature of the data. Cross-section is a snapshot of exposure and outcome at the same time point. This lends itself to 'reverse causality'. This is where the outcome has lead to a change in activity reducing the exposure. In this case those who are concerned about their BP being more likely to take great care of their diet.

One of the odd findings is that there is no relationship between % UPF and sodium intake. If UPF is 'high in salt' then high UPF should = high sodium, the finding suggests that the nature of the UPF varies. Despite this there is still an affect! so it may not be the salt in the UPF which is responsible for this effect, if so reformulation will not be effective!

The effect of sodium is augmented by that of UPF. The two together is greater than either individually

## Limitations

### The study

This is a cross-sectional study, and so has the limitations of this design. In particular the study measures exposure and outcome at the same time. Causal relationships cannot be understood from this data. Here participants aware of their BP might have chosen to reduce their UPF intake.

Time and age have particular effects in cross sectional studies. Researchers in aging, and learning have identified this clearly, separating the effects of experiential learning, from cohort effects and duration effects. Prospective studies such as Cohort or RCT studies have less of this affect sometimes inverting relationships between variables as a result.

This study was organised by government departments connected with food and farming alongside the Department of Health. The sample was designed to monitor relevant outputs. Funding and commissioning processes affect design structure and might also affect participant engagement and expectation. In particular this might affect social desirability bias. Social desirability and other participant reporting bias may well be significant within dietary diaries. Double labelled water studies on the first wave showed some significant differences between measured energy intake, and reported energy intake with differences between different age groups.

### The data

The result depends on participants recording foods in the same way as time goes on. Exposure of the whole population to a stimulus to change their diet or the recording of their diet may result in systematic changes in results.

A uniform change in the nutrient content of the food or changing the nutritional definitions would affect results. Years 9-11 used a slightly different methodology for identifying foods for analysis as researchers have started to become aware of the need to understand 'processing'. This may account for the apparent lower exposure in the last three years.

These changes would affect the outcome variable less. However BP measurement technology has changed over ten years. BP machines derive their results from the changes in pressure detected in the arm of the participant, the algorithm used by the sphygmomanometer may have changed.

Weighting maintains age, sex, IMD, and government region across the waves. BMI is no different, and educational attainment is also unaffected. There are more vegetarians as time goes on.



The populations do change over time as some of the added variables do show statistical significant changes. In particular the number of vegetarians increases, which perhaps is one indicator of social desirability affecting the study.

In populations with exclusions the careful sample selection and weighting are overcome by the biasing effect of different selections. Selecting for BP, UPF or sodium changes the cohort sex balance, age range and IMD pattern making these samples less representative.

## The analysis

### Interpretation

Participants with high Na >5000mg are more likely to have hypertension. Causation from other studies is likely to be that this high Na intake is contributing to high BP. However from this study causation cannot be identified. Other possibilities are that raised BP stimulates ingestion of Na, that Na ingestion increases hypertension, or that people with hypertension share another characteristic that increases their Na intake.

The results around high UPF intake also do no more than imply correlation. Causation might be the high BP reduces intake of UPF, or that participants with high BP are more aware of dietary UPF content. This can affect the detected directionality of the correlation.

### Age

Age is a particular feature in this outcome. BP is very strongly affected by age. In cross sectional studies Age has several dimensions. Age identifies cohorts of people with particular experiences, it identifies duration of experience, it represents physiologically different states, and it also identifies access to resources financial, material, and experiential. In psychology studies of ageing these separate aspects are understood and some can be controlled for, but not all. Prospective, longitudinal studies, and case matching can help reduce some of the effects. However longitudinal studies have reported similarly equivocal results, identifying a potentially more complex interaction.

### Ideas for further research

I will divide these suggestions into quantitative and qualitative.

#### Quantitative

There is scope for more research based on this data set. Within this same biomedical paradigm there are whole range of variables which can be compared against the clinical and biochemical outcomes. These include measured variables such as BMI, biochemical indicators such as HbA1c and medical diagnostic categories such as Diabetes.

Modelling research has allowed projections to be made using the data from studies such as this as a base for projected models. This can evaluate policy effects.

### *Mixed and Qualitative*

The richness of the quantitative data in this survey calls for its use within an approach allowing more detailed description and in depth assessment with participants. In the study data there are data allowing research around cooking activities, hobbies, and eating activities.

It could also be used as a template for studies smaller in geographical scope, but more in depth as cross over studies collecting both quantitative and qualitative data.

### *Ideas for policy*

Policy is an 'upstream' approach. It can be used to reduce exposure indirectly or directly. Ideas include legislation to reduce UPF use, this might be by pricing, or other approaches. Health promotion policy needs to match policy activity. People who know that UPF is bad, are more likely to accept policy limiting availability.

Reformulation is a policy suggestion, where UPF is further processed to remove the salt. This is frequently discussed. By demonstrating that UPF is not just high in salt this study supports the argument of avoiding further formulation as a policy for reducing BP.

Dietary approaches to improving public health are able to deliver proportionate and universal interventions to populations to reduce the incidence of NCD. When delivered upstream at the policy level they are effective and efficient and minimise cost. These approaches offer significant benefits over actions targeted at individuals.

Dietary and 'awareness' approaches can be used by individuals. These approaches risk the development of a culture of blame of individuals and of sub-groups in society. The commercial and social determinants of health play out a significant role in research, and delivery of public health improvements around food.

This study aims to inform local policy to reduce BP and so Non-Communicable Disease. If UPF and sodium intake increase the risk of hypertension then policy to reduce exposure might deliver change at a population level.

### **Generalisability**

This study used national data. This was stratified across the four home nations. It was stratified for IMD, and sex and to cover adults and children. The study can therefore be generalised to the UK population. The results are comparable with those in Korea, Brazil and USA. These include countries with lower UPF intake, but also similar levels.

## Conclusion

In summary,

In adults and children across the four home nations of the UK between 2008 and 2019, did exposure to high sodium dietary intake, and or high UPF dietary intake, compared to lower exposure, increase the odds of having a mean systolic blood pressure of over 140mmHg?

In answer, the odds ratio for hypertension is increased and statistically significant for high Na compared to lower exposure to Na. The odds ratio for hypertension is statistically significant for high UPF, being lower compared to lower exposure to UPF. The OR of each is altered by combining the two exposure variables in the same regression model indicating some degree of interaction. The AIC also reduces by combining the two variables into the same model.

The first finding matches the findings of other researchers in other countries and in past studies in the UK. The second finding seems to be contrary to other studies. Further research will be needed to understand this result.

Other objectives were also met with a literature review examining the research around Na and UPF and BP, and discussion of the interpretation and generalisability of the study. I will present the findings to support policy development.

## Bibliography

::: {#refs} :::

1. Cardiovascular disease - data - OHID [Internet]. Available from: <https://fingertips.phe.org.uk/profile/cardiovascular/data#page/1/gid/1938133106/pat/223/par/E40000010/ati/221/are/nE54000008/iid/219/age/1/sex/4/cat/-1/ctp/-1/yrr/1/cid/4/tbm/1/page-options/car-do-0>
2. High Blood Pressure & Cardiovascular Disease Prevention | Champs Public Health Collaborative [Internet]. 2020. Available from: <https://champspublichealth.com/blood-pressure/>
3. Du S, Neiman A, Batis C, Wang H, Zhang B, Zhang J, et al. Understanding the patterns and trends of sodium intake, potassium intake, and sodium to potassium ratio and their effect on hypertension in China<sup>1,2,3</sup>. *The American Journal of Clinical Nutrition* [Internet]. 2014 Feb 1;99(2):334–43. Available from: <https://www.sciencedirect.com/science/article/pii/S0002916523049511>
4. Cappuccio FP, Capewell S. Facts, Issues, and Controversies in Salt Reduction for the Prevention of Cardiovascular Disease. 2015;7(1):21.
5. Moreira PVL, Baraldi LG, Moubarac JC, Monteiro CA, Newton A, Capewell S, et al. Comparing Different Policy Scenarios to Reduce the Consumption of Ultra-Processed Foods in UK: Impact on Cardiovascular Disease Mortality Using a Modelling Approach. Hernandez AV, editor. *PLOS ONE* [Internet]. 2015 Feb 13;10(2):e0118353. Available from: <https://dx.plos.org/10.1371/journal.pone.0118353>
6. Marmot M, Allen J, Boyce T, Goldblatt P, Callaghan O. All together fairer: Health equity and the social determinants of health in cheshire and merseyside. London; 2022.
7. University Of Cambridge MEU, NatCen Social Research. NDNS RPNational diet and nutrition SurveysNational diet and nutrition survey years 1-11, 2008-2019. 2022; Available from: <https://beta.ukdataservice.ac.uk/datacatalogue/doi/?id=6533#19>
8. Elm E von, Altman DG, Egger M, Pocock SJ, Gøtzsche PC, Vandenbroucke JP. Strengthening the reporting of observational studies in epidemiology (STROBE) statement: guidelines for reporting observational studies. *BMJ* [Internet]. 2007 Oct 18;335(7624):806–8. Available from: <https://www.bmj.com/content/335/7624/806>

9. Bruce N, Pope D, Stanistreet D. Quantitative methods for health research: A practical interactive guide to epidemiology and statistics. Second edition. Hoboken, NJ: Wiley; 2018.
10. Jafar AJN. What is positionality and should it be expressed in quantitative studies? Emergency Medicine Journal [Internet]. 2018 Jan 11; Available from: <https://emj.bmj.com/lookup/doi/10.1136/emmermed-2017-207158>
11. Evans L, Trotter DRM. Epistemology and Uncertainty in Primary Care: An Exploratory Study. Family Medicine. 41(5):8.
12. Page MJ, Moher D, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. PRISMA 2020 explanation and elaboration: updated guidance and exemplars for reporting systematic reviews. BMJ [Internet]. 2021 Mar 29;n160. Available from: <https://www.bmj.com/lookup/doi/10.1136/bmj.n160>
13. Scopus | the largest database of peer-reviewed literature | elsevier [Internet]. Available from: <https://www.elsevier.com/en-gb/solutions/scopus>
14. Ovid: Search form [Internet]. Available from: <https://ovidsp.dc1.ovid.com/ovid-b/ovidweb.cgi?QS2=434f4e1a73d37e8cf66185f15d893d09240dc867c99b537069b551c0426841bbc6d4c4bcf1cb76dab1f9942ed8716272304cfa1dda7c2540178089af00d2be2da024db246851a260e30a99a35a0b305fdd220d8528bc6964ca701aa2828288873ef69169deed62b35b9a90a5e36c7f2d4813a4f135eda2c83f0a385349cbbb758e4c16cb3ac1291c704a94203038dc0ff849a67807328e8000b370b5f2b1191a500a6c4b22c0f909a786216f3817a25c83da34acd720eb96a1f34f944d57a5030f554c4861ef4c235ecd46bd3b4e5737f17c5e58cdb685c139982761767a57a10b3da88adce86aa245ae3f89ecfe6e23cdeadd102ef459f246894b69d120a6ba37c70f9f2c7d19ae>
15. Barbosa LB, Vasconcelos NBR, Dos Santos EA, Dos Santos TR, Ataíde-Silva T, Ferreira H da S. Ultra-processed food consumption and metabolic syndrome: A cross-sectional study in quilombola communities of alagoas, brazil. International journal for equity in health. 2023;22(1):14.
16. Wang L, Du M, Wang K, Khandpur N, Rossato SL, Drouin-Chartier JP, et al. Association of ultra-processed food consumption with colorectal cancer risk among men and women: Results from three prospective US cohort studies. BMJ [Internet]. 2022 Aug 31 [cited 2022 Dec 1];378:e068921. Available from: <https://www.bmj.com/content/378/bmj-2021-068921>
17. Mambrini SP, Menichetti F, Ravella S, Pellizzari M, De Amicis R, Foppiani A, et al. Ultra-processed food consumption and incidence of obesity and cardiometabolic risk factors in adults: A systematic review of prospective studies. Nutrients. 2023;15(11):2583.

18. D'Elia L, Galletti F, La Fata E, Sabino P, Strazzullo P. [Effect of dietary sodium restriction on arterial stiffness: Systematic review and meta-analysis of the randomized controlled trials](#). *Journal of hypertension*. 2018;36(4):734–43.
19. Leyvraz M, Chatelan A, Costa BR da, Taffé P, Paradis G, Bovet P, et al. [Sodium intake and blood pressure in children and adolescents: A systematic review and meta-analysis of experimental and observational studies](#). *International journal of epidemiology*. 2018;47(6):1796–810.
20. Frías JRG, Cadena LH, Villarreal AB, Piña BGB, Mejía MC, Cerros LAD, et al. [Effect of ultra-processed food intake on metabolic syndrome components and body fat in children and adolescents: A systematic review based on cohort studies](#). *Nutrition (Burbank, Los Angeles County, Calif)*. 2023;111:112038–8.
21. Shim SY, Kim HC, Shim JS. Consumption of ultra-processed food and blood pressure in korean adults. *Korean circulation journal*. 2022;52(1):60–70.
22. Scaranni P de O da S, Cardoso L de O, Chor D, Melo ECP, Matos SMA, Giatti L, et al. Ultra-processed foods, changes in blood pressure and incidence of hypertension: The Brazilian Longitudinal Study of Adult Health (ELSA-Brasil). *Public health nutrition*. 2021;24(11):3352–60.
23. Zhang T, Gan S, Ye M, Meng G, Zhang Q, Liu L, et al. [Association between consumption of ultra-processed foods and hyperuricemia: TCLSIH prospective cohort study](#). *Nutrition, metabolism, and cardiovascular diseases*. 2021;31(7):1993–2003.
24. Rezende-Alves K, Hermsdorff HHM, Miranda AE da S, Lopes ACS, Bressan J, Pimenta AM. Food processing and risk of hypertension: Cohort of Universities of Minas Gerais, Brazil (CUME Project). *Public Health Nutrition [Internet]*. 2021 Sep;24(13):4071–9. Available from: <http://www.cambridge.org/core/journals/public-health-nutrition/article/food-processing-and-risk-of-hypertension-cohort-of-universities-of-minas-gerais-brazil-cume-project/100DC0D407DBEAAA99C701A707061329>
25. Harrington JM, Fitzgerald AP, Kearney PM, McCarthy VJ, Madden J, Browne G, et al. DASH diet score and distribution of blood pressure in middle-aged men and women. *American journal of hypertension [Internet]*. 2013;26(11):1311–20. Available from: <https://www.scopus.com/inward/record.uri?eid=2-s2.0-84933499946&doi=10.1093%2fajh%2fhpt106&partnerID=40&md5=61fc359ae16fbc1c228f0c5591b3ebf7>
26. Asma A, Lokman NAH, Hayati MY, Zainuddin AA. Ultra-processed food classification, their contribution to sodium and added sugar availability, and its relationship with nutritional status among adults in terengganu, malaysia. *IIUM Medical Journal Malaysia [Internet]*. 2019;18(3):49–58. Available from:

<https://www.scopus.com/inward/record.uri?eid=2-s2.0-85096873804&doi=10.31436%2fijm.v18i3.193&partnerID=40&md5=a9643393d9f12cca0b443b39a285e939>

27. D'Avila HF, Kirsten VR. [ENERGY INTAKE FROM ULTRA-PROCESSED FOODS AMONG ADOLESCENTS](#). CONSUMO ENERGETICO PROVENIENTE DE ALIMENTOS ULTRAPROCESSADOS POR ADOLESCENTES. 2017;35(1):54–60.
28. De Deus Mendonça R, Souza Lopes AC, Pimenta AM, Gea A, Martinez-Gonzalez MA, Bes-Rastrollo M. Ultra-processed food consumption and the incidence of hypertension in a mediterranean cohort: The seguimiento universidad de navarra project. American journal of hypertension. 2017;30(4):358–66.
29. He FJ, MacGregor GA. Effect of modest salt reduction on blood pressure: A meta-analysis of randomized trials. Implications for public health. J hum hypertens 2002;16:761–70.
30. Strazzullo P, D'Elia L, Kandala NB, Cappuccio FP. [Salt intake, stroke, and cardiovascular disease: Meta-analysis of prospective studies](#). BMJ. 2009;339(7733):1296–b4567.
31. Graudal N, Jürgens G, Baslund B, Alderman MH. [Compared with usual sodium intake, low- and excessive-sodium diets are associated with increased mortality: A meta-analysis](#). American journal of hypertension. 2014;27(9):1129–37.
32. Ma Y, He FJ, Sun Q, Yuan C, Kieneker LM, Curhan GC, et al. [24-hour urinary sodium and potassium excretion and cardiovascular risk](#). The New England journal of medicine. 2022;386(3):252–63.
33. Monteiro CA, Levy RB, Claro RM, Castro IRR de, Cannon G. A new classification of foods based on the extent and purpose of their processing. Cadernos de Saúde Pública [Internet]. 2010 Nov;26:2039–49. Available from: <http://www.scielo.br/j/csp/a/fQWy8tBbJkMFhGq6gPzsGkb/?lang=en>
34. Martinez-Steele E, Khandpur N, Batis C, Bes-Rastrollo M, Bonaccio M, Cediel G, et al. Best practices for applying the nova food classification system | nature food. Nature Food [Internet]. 2023 Jan 6; Available from: <https://www.nature.com/articles/s43016-023-00779-w>
35. Wang M, Du X, Huang W, Xu Y. Ultra-processed foods consumption increases the risk of hypertension in adults: A systematic review and meta-analysis. American journal of hypertension. 2022;35(10):892–901.
36. Shim SY, Kim HC, Shim JS. [Consumption of Ultra-Processed Food and Blood Pressure in Korean Adults](#). Korean Circ J. 2022 Jan;52(1):60–70.

37. Lennox A, Bluck L, Page P, Pell D, Cole D, Steer T, et al. Misreporting in the National Diet and Nutrition Survey Rolling Programme (NDNS RP): summary of results and their interpretation.
38. OpenEpi. OpenEpi - sample size for unmatched case-control studies [Internet]. 2021. Available from: <https://www.openepi.com/SampleSize/SSCC.htm>
39. Wang M, Du X, Huang W, Xu Y. Ultra-Processed Foods Consumption Increases the Risk of Hypertension in Adults: A Systematic Review and Meta-Analysis. *American Journal of Hypertension* [Internet]. 2022 Oct 1 [cited 2022 Nov 12];35(10):892–901. Available from: <https://doi.org/10.1093/ajh/hpac069>
40. R Core Team. R: A language and environment for statistical computing [Internet]. Vienna, Austria: R Foundation for Statistical Computing; 2022. Available from: <https://www.R-project.org/>
41. Lumley T. Analysis of complex survey samples. *Journal of Statistical Software*. 2004;9(1):1–19.
42. Sjoberg DD, Whiting K, Curry M, Lavery JA, Larmarange J. Reproducible summary tables with the gtsummary package. *The R Journal* [Internet]. 2021;13:570–80. Available from: <https://doi.org/10.32614/RJ-2021-053>



## Appendix

## Appendix 1 Approved Proposal

The approved proposal



## Appendix 2 Ethics Certificate

The ethics cert.

## Appendix 3 Software used

The software used

CRAN

[GT Summary \(42\)](#)