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Assessing Australian women's knowledge and knowledge preferences about long-term health after hypertensive disorders of pregnancy: a survey study

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Assessing Australian women's knowledge and
knowledge preferences about long-term health after
hypertensive disorders of pregnancy: a survey study

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ABSTRACT

Objective(s): To (a) assess women's current knowledge regarding long-term cardiovascular health after hypertensive disorders of pregnancy (b) elicit women's preferred educational content and format regarding health after hypertensive disorders of pregnancy. **Design and setting:** A custom-created online survey exploring Australian women's knowledge about long-term health after hypertensive disorders of pregnancy, distributed through consumer groups and social media. **Participants:** 266 women with (n=174) or without (n=92) a history of hypertensive disorders of pregnancy. **Primary and secondary outcome measures**: 1) Proportion of women identifying long-term health risks after hypertensive disorder of pregnancy using a 10-point risk knowledge score with 0-4 'low', 4.1-7.0 'moderate' and 7.1-10 'high'. 2) Exploration of preferred content, format and distribution of educational material post hypertensive disorder of pregnancy. **Results**: Knowledge scores about health after hypertensive disorder of pregnancy were moderate in groups with and without a history of the disorder. Knowledge was highest regarding risk of recurrent hypertensive disorders in a subsequent pregnancy, 'moderate' for chronic hypertension and heart attack, 'moderate' and 'low' regarding risk of heart disease and 'low' for diabetes and renal disease. Only 36% of all participants were aware that risks start within 10 years after the affected pregnancy. The majority of respondents with a history of hypertensive disorder of pregnancy (76%) preferred receiving information about long-term health 0-6 months postpartum from a healthcare

- 52 provider (80%), key organisations (60%), social media (47%) and 53 brochures/flyers (43%).
- Conclusion(s): Women's knowledge regarding health risks after hypertensive
 disorder of pregnancy was 'moderate', although with important disease-specific
 gaps such as increased risk of diabetes. Most women wanted to be informed
 about their long-term health from a healthcare provider.

STRENGTHS AND LIMITATIONS OF THIS STUDY

- Consumer co-created survey exploring health knowledge after hypertensive pregnancy.
- For the first time survey results include findings from women with a history of gestational hypertension as well as from women without a history of hypertensive disorder of pregnancy.
- Recruitment from groups with potentially greater baseline knowledge may bias results, although substantive knowledge gaps still found.
- Although surveys were available in English, Arabic and Mandarin, there remains potential sub-optimal coverage of culturally and linguistically diverse groups.

INTRODUCTION

Hypertensive disorders of pregnancy (HDP) include chronic hypertension (CH),
preeclampsia (PE) and gestational hypertension (GH) and complicate 5-10% of
pregnancies¹. PE is a multi-system disorder characterised by new-onset
hypertension after 20 weeks' gestation and involvement of one or more other
organ systems and/or the fetus^{2 3}. GH is new-onset hypertension after 20

weeks' gestation without any other complications, and apart from increased risk of progression to PE²³, is not associated with adverse pregnancy outcomes. However, both conditions are associated with long-term cardiovascular sequelae⁴⁵. CH is defined as hypertension that is confirmed before pregnancy or before 20 completed weeks gestation, which may worsen during pregnancy and/or on which preeclampsia may be superimposed². Globally, cardiovascular disease (CVD) is one of the leading causes of death in women,⁶ and for women who have experienced an HDP, it is 2-3 times higher compared with those who did not⁴⁷⁸. This risk of premature death is present within 10 years after the affected pregnancy⁷⁹¹⁰ and remains after adjusting for the presence of other cardiovascular risk factors.

Both Australian and international societies, including the Society of Obstetric Medicine of Australia and New Zealand (SOMANZ) and the International Society for the Study of Hypertension in Pregnancy (ISSHP), recommend that women and healthcare providers (HCP) are provided with information about HDP and later CVD² ³. This includes recommending that women have a clinical review several months postpartum, and regular general practitioner (GP) follow-up to monitor blood pressure, fasting lipids and blood sugar². Adopting a healthy lifestyle with maintenance of an ideal weight and regular aerobic exercise is emphasised² ³. The aims of this study were to (a) explore Australian women's current knowledge on the topic of long-term CVD health after any HDP, not just PE (b) elicit women's preferred educational content and format regarding health after HDP, as a basis for creating tailored information and health advice for women after HDP.

METHOD

A national survey of women with and without a history of HDP was conducted, using a custom-created, face-validated online survey. Ethical approval was provided by the relevant Human Research Ethics Committee (HREC 18/POWH/326, REGIS 2019/PID05668).

Patient and Public involvement

As a validated instrument to assess women's knowledge was unavailable, a survey was custom designed. Initially, women with a history of HDP, comprising nine volunteers from the *Postpartum physiology, psychology and paediatric follow up study (P4 Study)*¹¹ and Australian Action on Preeclampsia (AAPEC), were invited to take part in group interviews which addressed the possible content and design of the survey, tested the survey for face validity and provided feedback for improvement. The topics discussed during the interviews were sourced from findings from a scoping literature review¹² and further complemented by questions specifically exploring the Australian context for women experiencing HDP. Following feedback, the survey was modified. The survey was made available in English, Arabic and Mandarin.

Data collection

The final survey was targeted at women in Australia, 18 years and older with a history of pregnancy in the last 3 years. The online survey, using SurveyMonkey™, was open from July to August 2019. Survey distribution occurred through the P4 study participants, organisations such as AAPEC,

maternity consumer groups as well as via the project's consumer representative and social media (Facebook and Twitter) including multicultural networks in order to reach Arabic and Mandarin speaking communities.

The data collection instrument

The survey for women (Appendix 1) explored demographic details, assessed obstetric history, history of HDP and other medical history including family history. The survey was tailored to women's self-reported HDP history (GH, PE, CH with or without worsening in pregnancy or superimposed PE, no hypertension history), with women given definitions of HDP conditions early in the survey to aid their self-report. Questions focused on knowledge of risk after pregnancy, provision of care and education following birth and what information and education women would like to receive. Women with a history of HDP were asked to classify their risk of various long-term health outcomes as greater, less than or equal to that of a woman with a normotensive pregnancy. Women after normotensive pregnancy were also asked to classify whether they believed women who had had HDP were at greater, lesser, or equal risk. The survey included two 'distractor' conditions not known to have an increased risk after HDP (breast cancer and seizures) to elicit negative answers and ascertain whether women could identify what they were not at increased risk of after HDP as well as what they were at risk of. At survey completion, women were provided with a correct risk profile summary and a link to further information.

Data Analysis

Quantitative survey analysis was undertaken using SPSS Version 25 (SPSS Statistics for Windows, Armonk, NY). Demographic data and responses to individual questions were analysed descriptively. To examine difference in knowledge levels amongst the targeted subgroups, (GH, PE, CH in pregnancy, no hypertension history) responses regarding HDP and future health risks were compared using Chi-squared testing or likelihood ratio for categorical data (as appropriate to subgroup sample size) and one-way ANOVA testing for continuous data. A \bar{p} value of <0.05 was considered statistically significant.

A knowledge score was created for the risk matrix, whereby 1 point was allocated to the correct answer, 0 for the incorrect answer, 0 for 'I do not know' and 0 for no answer/left blank. A mean score for each risk factor was calculated and a scale of 'low', 'moderate' and 'high' knowledge was established. The ranking classifications were divided into three score categories. For individual risk mean scores, 'low knowledge' equated to a mean of 0.00-0.40, 'moderate knowledge' was 0.41-0.70 and 'high knowledge' a mean of 0.71-1.00. An overall mean score out of 10 (as there were 10 conditions) was given for the HDP and non-HDP groups and were classified as 'low' 'moderate' or 'high' using the same mean ranges as were used for the individual conditions. Categorical analysis for proportions of each knowledge group ('high', 'moderate' and 'low') was also conducted to provide a further perspective.

RESULTS

In total, 308 survey responses were received (Figure 1). Forty-two were excluded: 40 for discontinuing the survey and not answering the question

asking about their perception of lower/same/higher risk with regards to 10 health conditions, and two with 'CH only' (no worsening hypertension or superimposed PE in pregnancy). Of 266 included responses, 174 (65%) women had a history of any HDP (HDP group) and 92 (35%) did not (non-HDP group). The HDP group consisted of 15 women with GH only (9%), 143 women with PE only (82%) and 16 women with CH plus superimposed pregnancy hypertension or PE (9%; will be known as CH). Of the HDP group, 123 (71%) had their most recent experience with HDP less than three years prior (32% <1 year prior and 39% 1-3 years prior).

Most respondents were in the 26-35 or 36-45 year age groups (91%), 89% were of Caucasian ethnicity, 90% pursued education beyond secondary school and 96% were in a relationship (Table 1). HDP women were more likely to be Caucasian, to have a history of diabetes, renal problems, be overweight and to have at least one additional cardiovascular risk factor than non-HDP women (Supplementary Table 1), and less likely to be university-educated. Half of all participants were sourced through social media (50%), with most of the remainder (45%) recruited via the P4 study (8% of HDP women, 46% of non-HDP women) and AAPEC (35% of HDP women).

Table 1: Respondent demographics

	GH	PE	СН	Total HDP	Non HDP	Total	Р
	n=15	n=143	n=16	n=174	n=92	n=266	HDP
	n (%)*	n (%)*	n (%)*	n (%)*	n (%)*	n (%)*	vs non- HDP
AGE							
18-25	2 (13)	8 (6)	0 (0)	10 (6)	2 (2)	12 (5)	0.157
26-35	5 (33)	68 (48)	8 (50)	81 (47)	36 (39)	117 (44)	0.246
36-45	8 (53)	60 (42)	8 (50)	76 (44)	50 (54)	126 (47)	0.097

45+	0 (0)	7 (5)	0 (0)	7 (4)	3 (3)	10 (3)	0.753
prefer not to answer	0 (0)	0 (0)	0 (0)	0 (0)	1 (1)	1 (0)	-
TOTAL	15 (100)	143 (101)	16	174 (101)	92 (99)	266 (99)	
			(100)				
ETHNICITY							
Caucasian	14 (93)	138 (97)	13 (81)	165 (95)	71 (77)	236 (89)	<0.001
Asian	1 (7)	3 (2)	1 (6)	5 (3)	18 (20)	23 (9)	<0.001
Other ¹	0 (0)	2 (1)	2 (13)	4 (2)	3 (8)	7 (3)	0.646
TOTAL	15 (100)	143 (100)	16 (100)	174(100)	92 (100)	266 (101)	
HIGHEST EDUCATIONAL A	TTAINMENT						
Secondary School	1 (7)	20 (14)	2 (13)	23 (13)	2 (2)	25 (9)	0.003
Diploma/Trade Certificate	5 (33)	47 (33)	8 (50)	60 (35)	9 (10)	69 (26)	<0.001
University Degree	9 (60)	75 (52)	6 (38)	90 (52)	81 (88)	171 (64)	<0.001
Prefer not to answer	0 (0)	1 (1)	0 (0)	1 (1)	0 (0)	1 (0)	-
TOTAL	15 (100)	143 (100)	16 (101)	174 (101)	92 (100)	266 (99)	
RELATIONSHIP STATUS							
In a relationship	15 (100)	133 (93)	14 (88)	162 (93)	92 (100)	254 (96)	0.004
Not in a relationship	0 (0)	9 (6)	2 (13)	11 (6)	0 (1)	11 (4)	0.001
Prefer not to answer	0 (0)	1 (1)	0 (0)	1 (1)	0 (0)	1 (0)	-
TOTAL	15 (100)	143 (100)	16 (101)	174 (100)	92 (100)	266 (100)	
RECRUITED TO SURVEY V	/IA						
P4 Newsletter	2 (13)	11 (8)	1 (6)	14 (8)	42 (46)	56 (21)	<0.001
AAPEC	1 (7)	57 (40)	3 (19)	61 (35)	1 (1)	62 (23)	<0.001
Social Media	12 (80)	74 (52)	11 (69)	97 (56)	37 (40)	134 (50)	0.016
Other ²	0 (0)	1 (1)	1 (6)	2 (1)	12 (13)	14 (5)	<0.001
TOTAL	15 (100)	143 (101)	16 (100)	174 (100)	92 (100)	266 (99)	

^{*} Percentages may not add to 100% as figures are rounded to whole numbers only

PE = preeclampsia GH = gestational hypertension CH = chronic hypertension, worsening in pregnancy and/or superimposed preeclampsia P4= Postpartum Physiology, Psychology, and Paediatric Study AAPEC= Australian Action on Preeclampsia ACM= Australian College of Midwives.

Table 2: Means of risk factor knowledge of women

	GH	PE	СН	P HDP GH vs	P GH vs PE	P CH vs	HDP	Non- HDP	P HDP vs non- HDP
	n=15	n=14 3	n=16	PE vs CH	VSTL	PE	n=174	n=92	1151
Chronic Hypertension	0.53	0.78	0.81	0.10	0.10	0.49	0.76	0.62	0.023
Diabetes	0.27	0.24	0.31	0.84	0.72	0.30	0.25	0.35	0.115
Renal Disease	0.27	0.54	0.69	0.06	0.044	0.25	0.53	0.21	<0.001
Heart Attack	0.53	0.69	0.75	0.40	0.14	0.23	0.68	0.52	0.014

¹ Other: Indigenous Australian (n=1), Polynesian or Maori (n=2), mixed ethnicity (n=4).

² Other: Friend (n=11), ACM (n=1), Clinic (n=1), Maternity Consumer group other than AAPEC (n=1).

							_		
Repeat HDP	0.87	0.90	0.94	0.81	0.52	0.27	0.90	0.71	<0.001
Stroke	0.47	0.62	0.81	0.14	0.37	0.10	0.63	0.53	0.144
Heart						0.29			
Disease	0.47	0.69	0.75	0.17	0.12		0.68	0.50	0.005
PVD	0.33	0.50	0.50	0.46	0.42	0.98	0.32	0.45	<0.001
Breast						0.12			
Cancer*	0.20	0.52	0.31	0.026	0.013		0.47	0.65	0.004
Seizures*	0.27	0.29	0.13	0.39	0.74	0.10	0.27	0.44	0.009
OVERALL MEAN KNOWLEDGE SCORE (OUT OF 10)	4.2	5.8	6.0	0.09	0.19	0.80	5.6	5.2	0.21

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy, PVD= Peripheral Vascular Disease

The colour coding in addition to the mean scores aims to highlight the areas of 'low' (red), 'moderate' (orange) and 'high' (green) knowledge.

MEAN SCORE	
LOW	0-0.40
MODERATE	0.41-0.70
HIGH	0.71-1

Average knowledge scores are shown in Table 2 and detailed results on which these scores are based on are shown in Supplementary Tables 2-11. Overall knowledge of the 10 conditions in the survey was 'moderate' for both groups (5.6/10 amongst HDP and 5.2/10 amongst non-HDP, p=0.21), with 33% in both groups having "high" overall knowledge, 32% and 40% respectively having "low" overall knowledge (Supplementary Table 12). Women with a history of HDP had 'high' knowledge with regards to HDP recurrence in a subsequent pregnancy (0.90) and CH (0.76); and 'moderate' knowledge regarding HDP's increased chance of conditions such as heart attack (0.68), heart disease (0.68) and stroke (0.63). Women without HDP history had 'high' knowledge (0.71) for HDP recurrence in a subsequent pregnancy. The same group of women had moderate knowledge of chronic hypertension (0.62) and stroke (0.53). Lowest knowledge across both groups was around the risk of future diabetes (0.25

^{*} Breast cancer and seizures are distractors within the survey. These were included despite being conditions that women after HDP are not at greater risk of.

HDP group and 0.35 for non-HDP group). Further 'low' scoring conditions were peripheral vascular disease (PVD) and renal disease. HDP women for most conditions had significantly higher knowledge than the non-HDP group, however the non-HDP group were more likely to correctly identify that there is not an increased risk of the two 'distractor' conditions after HDP.

Table 3 shows knowledge score breakdown by recency of pregnancy. In the subgroup of HDP women who experienced PE (n=143), average knowledge was similar amongst women who experienced HDP within the last three years (5.8/10), compared to those who experienced HDP more than three years ago (5.7/10).

Table 3: Means of risk factor knowledge of women with a history of preeclampsia listed by time elapsed since HDP

	PE n=143		P	PE n=143	3			PE n	=143	
	ALL COMBI	0 mont h- 6 mont h	6 mont h - 12 mont h	1-2 years	2-3 years	>3 year s	Р	UND ER 3	OV ER 3	Р
	NED	n=26	n=19	n=32	n=20	n=46	betw een grou ps	n= 97	n= 46	Under 3 yrs vs over 3 yrs
Chronic Hypertensi on	0.78	0.77	0.89	0.78	0.80	0.72	0.64 8	0.80	0.7	0.272
Diabetes	0.24	0.23	0.42	0.31	0.10	0.20	0.14 4	0.27	0.2	0.350
Renal Disease	0.54	0.50	0.74	0.41	0.65	0.52	0.16 9	0.55	0.5 2	0.784
Heart Attack	0.69	0.58	0.89	0.66	0.70	0.67	0.24 8	0.69	0.6 7	0.841
Repeat HDP	0.90	0.88	0.89	0.94	0.95	0.85	0.67 5	0.92	0.8 5	0.253
Stroke	0.62	0.65	0.68	0.50	0.65	0.65	0.61 9	0.61	0.6 5	0.616

Heart Disease	0.69	0.62	0.79	0.63	0.65	0.76	0.49 8	0.66	0.7 6	0.209
Disease	0.09	0.02	0.79	0.03	0.03	0.70	0.89	0.00	0.5	0.209
PVD	0.50	0.54	0.53	0.53	0.40	0.50	5	0.51	0.5	0.954
Breast							0.24		0.5	
Cancer*	0.52	0.46	0.37	0.63	0.40	0.59	5	0.48	9	0.255
							0.06		0.2	
Seizures*	0.29	0.15	0.16	0.41	0.45	0.26	6	0.30	6	0.641
OVERALL MEAN KNOWLED GE SCORE							0.82			2 222
OUT OF 10	5.8	5.4	6.4	5.8	5.8	5.7	5	5.8	5.7	0.890

The colour coding in addition to the mean scores aims to highlight the areas of 'low' (red), 'moderate' (orange) and 'high' (green) knowledge.

MEA	N SCORE	
	LOW	0-0.40
	MODERATE	0.41-0.70
	HIGH	0.71-1

Of the HDP women, only 32% were aware that the cardiovascular conditions may start manifesting within 10 years after an affected pregnancy, compared with 45% of women in the non-HDP group (p=0.036). About a third in each group (30% HDP, 36% non-HDP) were unsure about timing of risk rise/when health conditions manifest (Supplementary Table 13).

Women with HDP history were asked about their personal experience of risk discussion with healthcare providers (Table 4). The most frequent discussions about future health were regarding HDP in subsequent pregnancies (45%), risk of chronic hypertension (43%), and 'No discussion' (37%). Risk discussions were no more likely to have occurred in women with HDP less than 3 years ago or over 3 years ago.

Table 4: Proportion of conditions discussed when addressing future risk (multiple answers collected) within and over 3 years since last HDP

GH	PE	СН	TOTAL	OVERALL	Р
n=15	n=143	n=16	n=174	TOTAL	< vs > 3
				n=174	yrs since

	<3	>3	<3	>3	<3	>3	<3	>3		HDP
	yrs n=13	yrs n=2	yrs n=97	yrs n=46	yrs n=13	yrs n=3	yrs n=123	yrs n=51	n (%)	
	n (%)	n (%)	n (%)	n (%)	n (%)	n (%)	n (%)	n (%)		
HDP next pregnancy	3 (23)	1 (50)	47 (48)	21 (46)	5 (38)	2 (67)	55 (45)	24 (47)	79 (45)	0.777
Chronic Hypertension	4 (31)	1 (50)	47 (48)	16 (35)	4 (31)	2 (67)	55 (45)	19 (37)	74 (43)	0.365
No discussion	7 (54)	1 (50)	33 (34)	17 (37)	5 (38)	1 (33)	45 (37)	19 (37)	64 (37)	0.934
Lifestyle changes	4 (31)	0 (0)	26 (27)	6 (13)	2 (15)	2 (67)	32 (26)	8 (16)	40 (23)	0.140
Heart Attack	1 (8)	0 (0)	20 (21)	6 (13)	1 (8)	(33)	22 (18)	7 (14)	29 (17)	0.503
Renal Disease	1 (8)	0 (0)	20 (21)	3 (7)	2 (15)	(33)	23 (19)	4 (8)	27 (16)	0.072
Stroke	1 (8)	1 (50)	18 (19)	4 (9)	1 (8)	1 (33)	20 (16)	6 (12)	26 (15)	0.449
Peripheral vascular disease	0 (0)	0 (0)	15 15)	5 (11)	1 (8)	0 (0)	16 (13)	5 (10)	21 (12)	0.555
Cannot remember	1 (8)	0 (0)	4 (4)	2 (4)	1 (8)	0 (0)	6 (5)	2 (4)	8 (5)	0.781

*Table represents frequency of each option; percentages add to over 100% as women were asked to select any/all that applied. PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

When asked about preferences of the timing of a future risk discussion, the majority (76%) of women wanted a discussion 0-6 months postpartum. The topics women most wished to discuss (Table 5) are 'impact on my children from the pregnancy affected by HDP' (73%), 'signs and symptoms of the conditions' (67%), 'when does risk rise' (54%) and 'risk reduction for subsequent pregnancy' (54%). Women's preference for receiving information on long-term health after HDP is via a medical professional (80%), through key organisations (60%) and social media (47%).

Table 5: HDP women's preferences for content and distribution of information/education on future risk after HDP (multiple answers collected) in order of preference

GH	PE	СН	Total HDP
n=15	n=143	n=16	n=174
n (%)*	n (%)*	n (%)*	n (%)*

PREFERENCE OF DISCUSSION TOPICS				
Impact on my children from the pregnancy affected by HDP	11 (73)	115 (80)	10 (63)	136 (73)
Signs and Symptoms of the conditions	12 (80)	101 (71)	11 (69)	124 (67)
Risk reduction for subsequent pregnancy	6 (40)	88 (62)	7 (44)	101 (54)
When does the risk rise	6 (40)	87 (61)	8 (50)	101 (54)
Statistics	6 (40)	86 (60)	6 (38)	98 (53)
Reducing risk behaviours (diet, exercise, smoking cessation)	6 (40)	80 (56)	5 (31)	91 (49)
Where to find information	6 (40)	73 (51)	2 (13)	81 (44)
How to discuss the matter with my Healthcare provider	4 (27)	57 (40)	4 (25)	65 (35)
PREFERENCE OF DISTRIBUTION Medical professionals	11 (73)	117 (82)	12 (75)	140 (80)
Key organisations	8 (53)	87 (61)	10 (63)	105 (60)
Social Media	6 (40)	73 (51)	3 (19)	82 (47)
Brochures/Flyers	6 (40)	64 (45)	5 (31)	75 (43)
, i i i i i i i i i i i i i i i i i i i	` '	` '	` '	42 (24)
Online Videos	3 (20)	35 (24)	4 (25)	42 (24)

*Table represents frequency of each option; percentages add to over 100% as women were asked to select any/all that applied. PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

DISCUSSION

This study found overall, 'moderate' knowledge of health conditions after HDP amongst both HDP and non-HDP women. Amongst women with a history of HDP, highest knowledge was identified with regards to future risk of hypertension and repeat HDP in subsequent pregnancies. Conversely, knowledge of future risk of diabetes was low, as was knowledge of the "distractor" conditions among HDP women particularly. Diabetes as a future risk factor post HDP has previously not been reported on in studies of women's knowledge, and our findings suggest this is an important knowledge gap to address.

Other novel aspects of our study are inclusion of women who had a history of GH as well as those with a history of PE, and assessing knowledge of non-HDP

women's knowledge. Women after HDP had somewhat higher knowledge of most health risks than the non-HDP group, however non-HDP group also had better knowledge of some aspects such as timing of risk increase. However, both groups' knowledge of the early increase in risk was low, adding further concern and reason to address the knowledge gap. When looking at the proportion of participants scoring "high", these were equal between the HDP (33%) and non-HDP groups (33%), whilst proportions scoring "low" were similar enough (32% HDP versus 40% non-HDP) to not to show statistical significance. This further highlights that the HDP group remain underinformed about their increased CVD risk.

A further important finding was that many HDP women were not made aware of future health risks, with 37% of HDP women reporting to have had 'no discussion' about their increased long-term risk. Concerningly, women with more recent HDP were no more likely than women with HDP>3 years ago to report having risks discussed. This finding suggests risk discussions may not have improved in recent years despite updated guidelines emphasising long-term health^{2 3}, and that the extensive evidence regarding long-term implications for women after HDP continues to be lost in the translation of research to practice.

Women's knowledge after GH has not been previously reported as far as we are aware even though GH has similar frequency and similar future CVD risk as PE⁴ ¹³. Although only 9% of our sample were GH, this group had somewhat lower knowledge than the PE and CH groups regarding conditions after HDP

(although mostly not reaching statistical significance). Over half reported receiving no discussion of health risks after GH. This suggests substantive knowledge gaps after GH to address in both women and healthcare providers.

International studies exploring women's knowledge have predominantly reported limited or no knowledge about the link between HDP and CVD¹², though our study found overall, 'moderate' knowledge of health conditions after HDP. The two conditions associated with highest knowledge were repeat HDP and risk of future hypertension. Findings were similar in Traylor et al.'s¹⁴ survey where 146 women post HDP were included (PE n=76, PE with severe features n=41, CH=29). Future hypertension and repeat HDP were correctly identified by women as risk factors, however this knowledge was mainly reflected in the group of women who had experienced PE with severe features. Brown et al. 15 (n=12 women attending postnatal follow-up clinic) also found that women are aware of repeat HDP risks, however despite postnatal risk counseling. perception of hypertension and CVD risk was mainly associated with participants who had a family history of CVD. More recently, Hutchesson et al. 16 surveyed 127 women with PE in the two years prior, finding very high knowledge about future hypertension risk (96%, higher than our post-PE findings) and most were aware of stroke (67%) and CVD (66%) risks (similar to our findings). Over a third of women after PE had 'no discussion' about future risk in our study. Hutchesson et. al¹⁶ reported over one third of their participants remained unaware of increased CVD risks, which is similar to our findings. Similarities may be explained by the fact that major source of PE participants for both, the Hutchesson et. al¹⁶ survey and ours, was the patient support/advocacy group AAPEC. Recruitment from this advocacy group may also explain a higher post-PE knowledge than other studies have reported.

Our study findings resonate with those from similarly targeted women in Canada, Portugal, United Kingdom, the United States of America and a previous Australian study, all conducted between 2013 and 2017¹². Therefore, from a global perspective, these findings reinforce a persisting, and concerning, research to consumer gap. With international guidelines, including ISSHP², specifically targeted to assist HCPs providing care to women on an international scale to better manage and address health after HDP, this practice gap of knowledge transmission to women would be expected to narrow.

Education preferences

357 Content

Women mostly wanted educational materials to address HDP impact on their children, signs and symptoms of conditions they are at higher risk of, the timing of when their risks rise, and how to best reduce risk of recurrent HDP. Similar preferences were expressed by the women included in Seely et al.'s¹⁷ focus group of 20 women after PE, with the key concern being the impact the PE pregnancy may have had on the health of their children.

Format of education and access

Our study identified that women mostly wanted to receive information about long-term health after HDP from medical professionals. Key organisations who are experts on the topic, via social media and through information brochures

were other acceptable avenues of access to information. This is in contrast to Skurnik et al's¹⁸ focus group of 14 women after PE, whose preferences for educational materials about the link between CVD and PE were via pamphlets available in doctor's offices as well as via online communities and topical blogs. However, Hird et al's¹⁹ participants also expressed preference for healthcare providers as their information source, including wanting healthcare providers to guide them towards reliable online/external information sources rather than encounter irrelevant or potentially inaccurate information due to their self-initiated search. Hutchesson et al. ¹⁶ report that high knowledge amongst participants was mainly due to the women's own research rather than receiving all possible, relevant information from their healthcare provider. Overall, existing studies including ours would suggest that although women are very open to the use of online sources or information packs, their healthcare providers are seen as central to closing their knowledge gaps.

Time of risk discussion

An important element to consider when communicating about risk with women who have experienced GH or PE is the timing of these discussions, as situational factors of being a new mother may alter when women are most receptive to follow-up. In our study, three-quarters of the women preferred this to occur in the first six months after birth. As well as being their preference, this also aligns with the potential benefits of early intervention, and would allow for addressing knowledge gaps found in this study around how soon the risk rises after HDP. Addressing future risk early but not immediately is also supported by Brown et al.'s study of women after PE, where participants suggested that six

months postpartum was the timeframe where they felt they had transitioned into a more comfortable stage of parenting and were able to focus more on themselves again¹⁵.

Strengths and limitations

The survey was co-created via a formalised process of seeking input and feedback on the usability, language and content from women who have previously experienced HDP. This creation and face validation process involving consumers gives added value to the survey.

Our knowledge score is both a strength, as it allows for a summary of findings across all the conditions and risks, and a limitation, as assigning cut-points for knowledge ranking is an arbitrary designation. Having included the distractor conditions (breast cancer and seizures) may also have altered the overall score. However, we believe it is important for women to not incorrectly believe they are at increased risk of more conditions than they are, as well as having knowledge of their increased cardiovascular risk. The addition of women with a history of GH as well as women without any history of HDP, is also a strength to add broader perspective on this topic.

Limitations include demographic make-up of respondents, with HDP participants predominantly English speaking and Caucasian (95%) despite the survey being available in Arabic and Mandarin as well as English. The non-HDP group (20% Asian background) had similar background demographics of Australian reproductive-aged women²⁰, and as HDP is more prevalent amongst

the Caucasian population²¹, the sample actually is likely close to representative of Australian HDP and non-HDP women. However, it would have been preferable to also gain insight from more culturally and linguistically diverse groups in order to understand their knowledge base and address their needs within this context.

In the survey, women were asked to select their HDP history which was then used to group them for analysis. Women's diagnosis of HDP is by self-report is a limitation, as some bias may be introduced through inaccurate self-report of diagnosis. The broad geographical range and anonymous nature of the survey precluded any verification of diagnosis. However, women were provided with definitions of the various HDP conditions at the start of the survey to aid them in their self-report. Another limitation is where participants were recruited from, with close to half either drawn from the P4 study (an Australian post-HDP research study) or consumer group AAPEC. Therefore, there may be knowledge bias in the sample (i.e. a more knowledgeable group of participants than the overall HDP or non-HDP population). However, as even this group with potentially greater baseline knowledge had substantive knowledge gaps, our study highlights the need for interventions to improve knowledge of health after HDP.

Implications

Close to two decades worth of data have been collected⁸ since research on the link between HDP and increased CVD risk emerged in the early 2000s, with the first systematic review published in 2007²². It could be expected that this

knowledge, by now, would have been translated into practice and shared with HDP women, however our findings suggest that this is still not the case. This study is valuable from the public health perspective, given the wider context of prevalence and importance of cardiovascular disease in women.

Guidelines such as ISSHP² and SOMANZ³ suggest regular follow-up after HDP as well as counselling women with regards to their individual long-term CVD risk. Although available to the public, these are not designed for women. Compiling suitable information for women would be an important step towards closing the knowledge gap. It is important to establish preferred content, presentation and timing of education for post-HDP health for women as we have in this study, to maximise the chance that women will engage with and benefit from education.

CONCLUSION

This Australian survey of women's knowledge of risks after HDP, found varying knowledge from the targeted groups. Despite 'high' knowledge being demonstrated regarding some risks, overall significant knowledge gaps were identified for certain conditions, particularly diabetes, and for knowledge about the relatively early timing of when health risks increase after HDP. Identifying these gaps are important in planning tailored education for women, and to improve early intervention for modifiable CVD risks in women after HDP. Addressing these women's preferences for content and to have this delivered by their healthcare provider may further lead to enhanced counselling, management and improved women's health trajectories.

469	
470	LIST OF ABBREVIATIONS
471	AAPEC - Australian Action on Preeclampsia
472	CH - chronic hypertension worsening in pregnancy and/or with superimposed
473	preeclampsia
474	CVD - cardiovascular disease
475	GH - gestational hypertension
476	GP - general practitioner
477	HCP - healthcare provider
478	HDP - hypertensive disorder of pregnancy
479	ISSHP - International Society for the Study of Hypertension in Pregnancy
480	PE - preeclampsia
481	PVD - peripheral vascular disease
482	SOMANZ - Society of Obstetric Medicine Australia New Zealand
483	
484	STATEMENTS
485	Ethical approval
486	Ethical approval has been provided by South-Eastern Sydney Local Health
487	District Human Research Ethics Committee (Ref: 18/POWH/326). The
488	ratification for the University of Technology Sydney has also been obtained
489	under ETH18-3061.
490	
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Access to data examined in this study

The authors have full access to all data reported in the study.

Transparency statement and competing interests

The article is the authors' original work, has not received prior publication and is not under consideration for publication elsewhere. All the authors have seen and approved the manuscript being submitted. The manuscript is an honest, accurate, and transparent account of the study being reported, no important aspects of the study have been omitted. We have read and understood BMJ policy on declaration of interests and declare that we have no competing interests. This manuscript presents partial results from Heike Roth's PhD research. The project is supervised by Caroline Homer and Amanda Henry.

Authors contributions

Heike Roth, Amanda Henry and Caroline Homer contributed to the conception and design of the study as well as the distribution of the survey and writing of the manuscript. Heike Roth led the analysis of the survey data, drafting and designed the Tables, Figures and Appendixes and wrote the first draft. Grace LeMarquand was a medical Honours student assisting with pre-survey interviews as well as initial data analysis. Lynne Roberts assisted in the survey development, supported the distribution, the interpretation of the findings and the discussion. Mark Brown contributed to the design of the survey and supported the interpretation of the findings and the discussion. As a maternity consumer, Lisa Hanley has assisted with the survey design and ensured appropriate use of language and content as well as supported the distribution. All authors contributed to drafts and revising of the paper and all approved the final version.

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Authors' information (optional)

This manuscript presents partial results from Heike Roth's PhD research. The project is supervised by Caroline Homer and Amanda Henry.

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TABLES, SUPPLEMENTARY TABLES, APPENDICES AND FIGURES

Table legend

Table 1: Respondent demographics

Table 2: Means of risk factor knowledge of women

Table 3: Means of risk factor knowledge of women with a history of preeclampsia listed by time elapsed since HDP

Table 4: Proportion of conditions discussed when addressing future risk (Multiple answers collected) within and over 3 years since last HDP

Table 5: HDP women's preferences for content and distribution of information/education on future risk after HDP (Multiple answers collected) in order of preference

Supplementary data

Supplementary Table 1: Current CVD risk factors of HDP and non-HDP women (Multiple answers collected) in order of frequency

Supplementary Table 2: Chronic Hypertension

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 3: Diabetes

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 4: Renal Disease

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 5: Heart Attack

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 6: HDP next pregnancy

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 7: Stroke

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 8: Heart Disease

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 9: Peripheral Vascular Disease

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 10: Breast Cancer

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 11: Seizures

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 12: Proportion of participants scoring 'high', 'moderate' and 'low' by type of HDP and non-HDP

Supplementary Table 13: HDP and non-HDP women's answers to timing of rise of risk with signs and symptoms starting to show

Appendices

Appendix 1: Survey for women

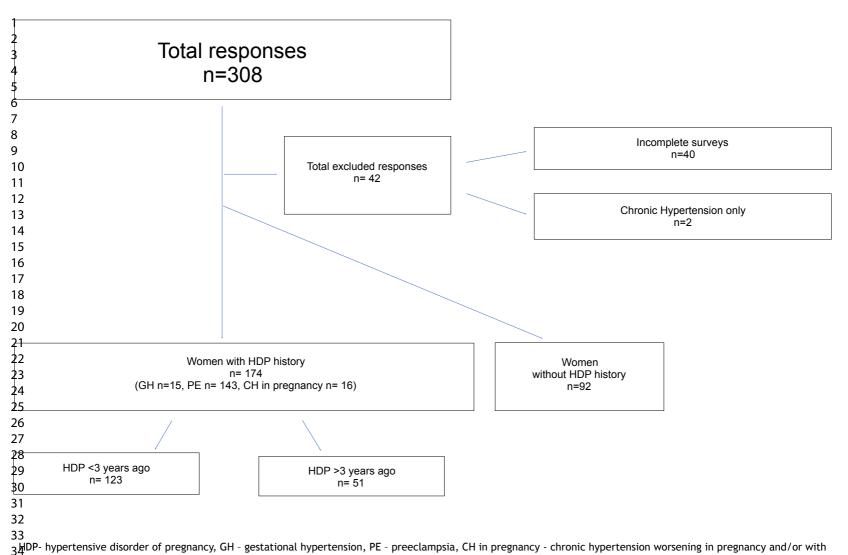
Figure legend

Figure 1: Survey inclusion





Figure 1: Survey Inclusion



35 superimposed preeclampsia

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;	Supplementary Table 1: Current CVD ris	k factors of H	IDP and non	-HDP womer	า (Multiple aı	nswers collec	ted) in orde	r of frequency
								f .

	GH n=15	PE	СН	Total HDP	non-HDP	P HDP vs non HDP	
		n=143	n=16	n=174	n=92		
	n (%)*	n (%)*	n (%)*	n (%)*	n (%)*		
None of the options	6 (40)	52 (36)	2 (13)	60 (34)	57 (62)	<0.001	
Obesity	5 (33)	38 (27)	7 (44)	50 (29)	10 (11)	0.001	
Family History Heart disease	3 (20)	37 (26)	6 (38)	46 (26)	14 (15)	0.037	
Hypertension	3 (20)	30 (21)	12 (75)	35 (20)	0 (0)	<0.001	
Renal problems	0 (0)	21 (15)	2 (13)	23 (13)	0 (0)	<0.001	
Smoking	0 (0)	10 (7)	3 (19)	13 (7)	2 (2)	0.075	
History of cardiovascular event**	0 (0)	1 (1)	0 (0)	1 (1)	0 (0)	0.356	
Diabetes	0 (0)	5 (3)	2 (13)	7 (4)	0 (0)	0.014	
Alcohol consumption	0 (0)	0 (0)	1 (6)	1 (1)	0 (0)	0.356	
High Cholesterol	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	-	

^{*}Table represents frequency of each option; percentages add to over 100% as women were asked to select any/all that applied.

^{**}angina, stroke or heart attack

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

Supplementary Table 2: Chronic Hypertension

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

in the second se	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	1 (1)	0 (0)	1 (1)
Same likelihood	2 (13)	14 (10)	1 (6)	17 (10)
More likely (correct)	8 (53)	111 (78)	13 (81)	132 (76)
I do not know	5 (33)	17 (12)	2 (13)	24 (14)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely (correct)	57 (62)
Same likelihood	22 (24)
More likely	1 (1)
I do not know	12 (13)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

Supplementary Table 3: Diabetes

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	1 (1)	0 (0)	1 (1)
Same likelihood	3 (20)	61 (43)	5 (31)	69 (40)
More likely (correct)	4 (27)	35 (25)	5 (31)	44 (25)
I do not know	8 (53)	44 (31)	6 (38)	58 (33)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n= 92	
	n (%)	
Less likely (correct)	32 (35)	
Same likelihood	39 (42)	
More likely	4 (4)	
I do not know	17 (19)	

Supplementary Table 4: Renal Disease

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

<u> </u>	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	2 (1)	0 (0)	2 (1)
Same likelihood	2 (14)	30 (21)	1 (7)	33 (19)
More likely (correct)	4 (29)	77 (55)	11 (73)	92 (54)
I do not know	8 (57)	32 (23)	3 (20)	43 (25)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

	non-HDP n=92	
	n (%)	
Less likely (correct)	39 (42)	
Same likelihood	31 (34)	
More likely	0 (0)	
I do not know	22 (24)	

Supplementary Table 5: Heart Attack

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

,	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	1 (1)	0 (0)	1 (1)
Same likelihood	1 (7)	20 (14)	1 (7)	22 (13)
More likely (correct)	8 (53)	98 (69)	12 (80)	118 (69)
I do not know	6 (40)	23 (16)	2 (13)	31 (18)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely (correct)	48 (53)
Same likelihood	23 (25)
More likely	0 (0)
I do not know	20 (22)

Supplementary Table 6: HDP next pregnancy

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

3,	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	2 (1)	0 (0)	2 (1)
Same likelihood	0 (0)	9 (6)	0 (0)	9 (5)
More likely (correct)	13 (87)	128 (89)	15 (100)	156 (90)
I do not know	2 (13)	4 (3)	0 (0)	6 (4)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely (correct)	65 (71)
Same likelihood	19 (21)
More likely	0 (0)
I do not know	8 (9)

Supplementary Table 7: Stroke

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

3 /	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	1 (1)	0 (0)	1 (1)
Same likelihood	1 (7)	18 (13)	1 (7)	20 (12)
More likely (correct)	7 (47)	89 (63)	13 (87)	109 (64)
I do not know	7 (47)	33 (23)	1 (7)	41 (24)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely (correct)	49 (53)
Same likelihood	24 (26)
More likely	0 (0)
I do not know	19 (21)

Supplementary Table 8: Heart Disease

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

in the second se	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	1 (1)	0 (0)	1 (1)
Same likelihood	2 (13)	18 (13)	1 (7)	21 (12)
More likely (correct)	7 (47)	99 (70)	12 (80)	118 (67)
I do not know	6 (40)	24 (17)	2 (13)	32 (19)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

	non-HDP n=92	
	n (%)	
Less likely (correct)	46 (50)	
Same likelihood	27 (29)	
More likely	0 (0)	
I do not know	19 (21)	

Supplementary Table 9: Peripheral Vascular Disease

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

3,	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	0 (0)	0 (0)	0 (0)
Same likelihood	2 (13)	20 (14)	1 (7)	23 (13)
More likely (correct)	5 (33)	72 (51)	8 (53)	85 (50)
I do not know	8 (53)	50 (35)	6 (40)	64 (37)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely (correct)	41 (45)
Same likelihood	29 (32)
More likely	0 (0)
I do not know	22 (24)

Supplementary Table 10: Breast Cancer

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

,	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	0 (0)	0 (0)	0 (0)
Same likelihood (correct)	3 (20)	74 (53)	5 (36)	82 (48)
More likely	1 (7)	1 (1)	1 (7)	3 (2)
I do not know	11 (73)	66 (47)	8 (57)	85 (50)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

	non-HDP n=92
	n (%)
Less likely	6 (7)
Same likelihood (correct)	60 (65)
More likely	1 (1)
I do not know	25 (27)

Supplementary Table 11: Seizures

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	0 (0)	0 (0)	0 (0)
Same likelihood (correct)	4 (27)	41 (29)	2 (13)	47 (27)
More likely	1 (7)	41 (29)	4 (27)	46 (27)
I do not know	10 (67)	60 (42)	9 (60)	79 (46)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely	25 (27)
Same likelihood (correct)	40 (44)
More likely	0 (0)
I do not know	27 (29)

Supplementary Table 12: Proportion of participants scoring 'high', 'moderate' and 'low' by type of HDP and non-HDP

	GH n=15 n(%)	PE n=143 n(%)	CH n=16 n(%)	P GH vs PE	P GH vs CH	P PE vs CH	HDP n=174 n(%)	non-HDP n=92 n(%)	P HDP vs non- HDP
High (score 8-10)	4 (27)	48 (34)	5 (31)	0.59	0.78	0.67	57 (33)	30 (33)	0.98
Moderate (score 5-7)	3 (20)	52 (36)	7 (44)	0.21	0.16	0.56	62 (36)	25 (27)	0.16
Low (score 0-4)	8 (53)	43 (30)	4 (25)	0.67	0.11	0.85	55 (32)	37 (40)	0.16
Total	15 (100)	143 (100)	16 (100)	-	-	-	174 (101)	92 (100)	-

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of

The colour coding in addition to the mean scores aims to highlight the areas of 'low' (red), 'moderate' (orange) and 'high' (green) knowledge. to nighting...

LOW
MODERATE
HIGH

Supplementary Table 13: HDP and non-HDP women's answers to timing of rise of risk with signs and symptoms starting to show

	GH	PE	СН	HDP	Non HDP	Р
	n=15	n=143	n=16	n=174	n=92	HDP vs non-HDP
	n (%)*	n (%)*	n (%)*	n (%)*	n (%)*	
<10 years	4 (27)	42 (29)	9 (56)	55 (32)	41 (45)	0.036
10-15 years	3 (20)	28 (20)	2 (13)	33 (19)	15 (16)	0.591
16-20 years	0 (0)	13 (9)	0 (0)	13 (8)	1 (1)	0.013
>20 years	3 (20)	5 (4)	0 (0)	8 (5)	2 (2)	0.302
Not sure/don't know	1 (7)	48 (34)	1 (6)	52 (30)	33 (36)	0.319
I don't think I will get any of these as I maintain a healthy lifestyle	4 (27)	7 (5)	2 (13)	13 (8)	N/A	-
TOTAL	15 (101)	143 (101)	16 (101)	174 (102)	92 (100)	

^{*} Percentages may not add to 100% as figures are rounded to whole numbers only
PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia,
HDP= hypertensive disorder of pregnancy

LONG TERM HEALTH AFTER BLOOD PRESSURE PROBLEMS IN PREGNANCY

You are invited to take part in a survey to gain insight into what women like yourself who have been pregnant before know about women's heart health. We are interested in the views of all women especially women who had high blood pressure (hypertension) or preeclampsia in pregnancy.

You can complete the survey if you are currently pregnant (with no major issues so far this pregnancy) or have been pregnant in the last three (3) years. You may have experienced high blood pressure in pregnancy OR you may have experienced a pregnancy without any serious complications.

The study is being conducted by the University of NSW, University of Technology Sydney and the Sydney Partnership for Health, Education, Research and Enterprise (SPHERE). The study is being undertaken by:

- Dr. Amanda Henry Obstetrician at St George and Royal Hospital for Women, Randwick, Senior Lecturer UNSW and SPHERE member
- Distinguished Professor Caroline Homer UTS, Midwifery Faculty of Health and SPHERE member
- · Dr. Clare Arnott Cardiologist, Royal Price Alfred Hospital
- · Mrs. Heike Roth PhD Candidate at University of Technology, Sydney
- · Mrs. Lynne Roberts Research Midwife at St George Hospital, SESLHD.

This work is occurring as part of Mrs Heike Roth's PhD studies and the NHMRC Fellowship of Dr. Henry. Apart from salary support for Dr. Henry, the study is otherwise unfunded.

If you agree to take part in this survey, it should only take about 15 minutes to complete and will involve answering questions about you, your pregnancy and your understanding of long term health in women who have been diagnosed with blood pressure problems in pregnancy.

Participation in this study is entirely voluntary and if you do not wish to take part it will have no effect on the care you are currently receiving. If you decide to participate, and throughout the survey think you would like to withdraw/not complete the survey, you can simply stop and not submit your answers. If you have already submitted your survey, it will not be possible to withdraw the data you have provided as the surveys are anonymous.

The information you provide will not be identifiable and will be kept securely until destroyed as per the South East Sydney Local Health District's requirements.

The study results will be published in a research thesis, in peer reviewed journals and presented at conferences and other professional forums. No one will be able to identify you from this information.

If you would like to personally receive results, you will have the option to leave your email details. The results will be available one (1) year after conclusion of the survey and your email will not be used for any other purpose.

If you would like any further information about the study or you experience any distress or concern as a result of completing this survey, please contact the Principal Investigator, Dr Amanda Henry on 02 91132315 or via email Amanda.henry1@health.nsw.gov.au. For medical assistance you can consult your General Practitioner. If you would like further information about the topic addressed in this study, you can visit the Australian Heart Foundation on the following link: https://www.heartfoundation.org.au/your-heart/women-and-heart-disease/womens-stories.

If you have any concerns or complaints about the conduct of this study, you should contact the Research Support Office of the South Eastern Sydney Local Health District Human Research Ethics Committee which is nominated to receive complaints from research participants. You should contact them on 02 9382 3587, or email SESLHD-RSO@health.nsw.gov.au and quote HREC 18/156.

* 1. I ackno	wledge that	participation	in the su	ırvey is v	oluntary
--------------	-------------	---------------	-----------	------------	----------

Yes



JT YOU	
-	out about you, your background, and occupation. sk (*) simply mean that they must be answered in order to
What age group are you in?	
18-25	46-55
26-35	56+
36-45	Prefer not to answer
Caucasian Asian	Aboriginal or Torres Strait Islander European
Asian	European
Polynesian or Maori	Prefer not to answer
Other (please specify)	
Other (please specify)	
What is your highest level of forma	
What is your highest level of forma Secondary school	University degree
What is your highest level of forma Secondary school Trade Certificate/Diploma	
What is your highest level of forma Secondary school	University degree
What is your highest level of forma Secondary school Trade Certificate/Diploma	University degree
What is your highest level of forma Secondary school Trade Certificate/Diploma	University degree Prefer not to answer
What is your highest level of forma Secondary school Trade Certificate/Diploma Other (please specify)	University degree Prefer not to answer
What is your highest level of forma Secondary school Trade Certificate/Diploma Other (please specify)	University degree Prefer not to answer
What is your highest level of formation Secondary school Trade Certificate/Diploma Other (please specify) What is your usual occupation/prof	University degree Prefer not to answer fession?
What is your highest level of forma Secondary school Trade Certificate/Diploma Other (please specify)	University degree Prefer not to answer fession?

P4 Newsletter	Maternity Consumer Network	
Australian Action on Preeclampsia (AAPEC)	Maternity Clinics	
Tresillian	Social media (Facebook, Twitter)	
Maternity Choices Australia	Australian College of Midwives	
Other (please specify)		

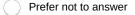
ABOUT YOUR HEALTH

These next questions are about your general health.

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue.

* 8. A	re you	currently	pregnant?
--------	--------	-----------	-----------

Yes



() No

ABOUT YOUR HEALTH Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue. 9. How many weeks pregnant are you? (provide whole numbers only, for example: 24) 10. How many children have you given birth to (20 weeks gestation and over)? * 11. Have you ever had any of the following, whilst pregnant or before or after pregnancy? (select all that apply) High blood pressure Stroke High BMI (overweight) Significant illness Angina Heart attack Diabetes None of the above/ no significant other medical complication Kidney problems Other (please specify) * 12. From the list below, which currently apply to you? (select all that apply) Smoking High cholesterol Obesity High blood pressure Alcohol consumption **Diabetes** Family history of heart disease None of the above Other (please specify) 13. Please provide the details of any prescribed medications you are taking I do not take any prescribed medication I take prescribed medication (please list the medications or leave blank if you prefer not to answer)



PREGNANCY

This section is about your pregnancy history.

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue.

DEFINITIONS OF BLOOD PRESSURE PROBLEMS

Here are some definitions of certain types of blood pressure problems in pregnancy. You may find these useful in order to more easily understand and answer the next questions.

Chronic hypertension: is if you had high blood pressure before falling pregnant, have high blood pressure outside of pregnancy, or were found to already have high blood pressure in the first half of your pregnancy. Chronic hypertension may have no known underlying cause (this is sometimes called "essential" hypertension), or it may be as a result of another underlying condition, such as kidney disease.

Gestational hypertension: is when you might have had high blood pressure for the first time in your pregnancy (after 20 weeks of pregnancy) but were otherwise well (that is, high blood pressure only but no effect on your baby's growth or on your health otherwise).

Preeclampsia: is when you have had high blood pressure in pregnancy (after 20 weeks of pregnancy) and some additional signs or issues in you and/or your baby. For example, you might have had protein in your urine, liver or kidney problems that showed up on blood tests, or there may have been concerns about the growth of your baby while you were pregnant.



PREGNANCY

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue.

We understand that you may find some of the questions difficult to answer as they might remind you of a challenging time in your life. We are grateful for your participation and contribution to improving knowledge on future health for women who had blood pressure problems in pregnancy. You will find some explanations at the end of the survey and a contact, in case you would like to seek further clarification and/or assistance.

* 14. Choose the situation which	best describes your	pregnancy history
----------------------------------	---------------------	-------------------

- At least one pregnancy is/was affected by gestational hypertension
- At least one pregnancy is/was affected by preeclampsia
- I have only been diagnosed with chronic hypertension
- I had chronic hypertension before pregnancy and had/have pregnancies that were complicated further by higher than usual blood pressure
- I had chronic hypertension before pregnancy and had/have pregnancies that were complicated further by preeclampsia
- No pregnancy is/was affected



YOUR HEALTH DURING YOUR PREGNANCY

Questions marked with a green asterisk (*) simply mean that they must be answered in	order to
continue.	

* 15. As someone who has chronic hypertension are you aware of any long term health issues that you are

at risk of? (select all that apply)	
Diabetes	Leukaemia
Kidney disease	Seizures
Breast cancer	Overall mortality risk is higher
Cardiac death	Ischaemic heart disease/heart attack
High blood pressure complications in another pregnancy	I think there are health risks but unsure which conditions I may be at risk of
Stroke	,
Peripheral vascular disease	I do not think that there are increased risks
Other (please specify)	



40.45			after pregnancy	
10-15 years after pregnanc 16-20 years after pregnanc		Not sure/d	o not know	
Other (please specify)	y			

LONG TERM HEALTH RISKS AFTER HIGH BLOOD PRESSURE IN PREGNANCY - Survey for women Chronic Hypertension only -TYPE OF CARE RECEIVED DURING PREGNANCY Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue. 17. If a healthcare provider did speak to you about your future health risks, when did this occur? Before birth 6 months to 1 year Immediately after birth 1 year and over Within first 6 weeks I cannot remember 6 weeks to 6 months * 18. When would be a good time to receive information about long term health risks in your gestational hypertension or preeclampsia experience? 12 months after birth 24 months after birth During pregnancy/at birth * 19. As a result of your pregnancy affected by blood pressure problems, were you referred to any of the below after your baby was born? (tick all that apply) Cardiologist Fitness centre for exercise Renal (kidney) Physician Nutritionist for dietary adjustment General Practitioner I cannot remember Other (please specify)

_ONG	TERM HEALTH	HRISKS A	AFTER I		nen	'KESSUR	E IN PR	(EGNAN(JY - Surv	ey ioi
		_	_	_	_	_	_	_	_	_
	During your pregn blem were you dia			3 years (20 wee	ks pregnan	cy and ov	ver), which	blood pre	essure
		Gestatio Hyperten		Preeclamp	p	No blood pres roblem diagno this pregnan	osed	nnot rememb	er Not a	pplicable
Firs	st Pregnancy									
Sec	cond Pregnancy									
Thi	rd Pregnancy									
Any	comments?									
* 21	How long ago was	r vour mos	t recent r	rognancy	that wa	s affected	hy a bloo	d proceur	nrohlem')
	0-6 months	your mos	i recent p	regriation		2-3 years	by a bloo	a pressure	, problem	·
	6-12 months					more than 3 y	ears ann			
	O 12 months					more triair 5 y	rears ago			
	1-2 years									
	1-2 years									
		ne were vo	ou diaano	sed? (Cho	oose a I	nost accura	ate time f	rame)		
	1-2 years At what point in tin	20-28	28-34	34-37	37-40	40-42	During or after giving	No diagnosis of blood pressure problem this	I cannot remember	Not applicable
* 22.		·		·			During or after	No diagnosis of blood pressure problem		Not applicable
* 22	At what point in tin	20-28	28-34	34-37	37-40	40-42	During or after giving	No diagnosis of blood pressure problem this		
* 22.	At what point in tin	20-28	28-34	34-37	37-40	40-42	During or after giving	No diagnosis of blood pressure problem this		
* 22. A	At what point in tin	20-28	28-34	34-37	37-40	40-42	During or after giving	No diagnosis of blood pressure problem this		
* 22. A	At what point in tin st Pregnancy cond Pregnancy rd Pregnancy	20-28	28-34	34-37	37-40	40-42	During or after giving	No diagnosis of blood pressure problem this		
* 22. A	At what point in tin st Pregnancy cond Pregnancy rd Pregnancy	20-28	28-34	34-37	37-40	40-42	During or after giving	No diagnosis of blood pressure problem this		
* 22. A	At what point in tin st Pregnancy cond Pregnancy rd Pregnancy	20-28	28-34	34-37	37-40	40-42	During or after giving	No diagnosis of blood pressure problem this		
* 22. A	At what point in tin st Pregnancy cond Pregnancy rd Pregnancy	20-28	28-34	34-37	37-40	40-42	During or after giving	No diagnosis of blood pressure problem this		
* 22. A	At what point in tin st Pregnancy cond Pregnancy rd Pregnancy	20-28	28-34	34-37	37-40	40-42	During or after giving	No diagnosis of blood pressure problem this		

REGNANCY						
iestions marked wit ntinue.	th a green aster	isk (*) simply	mean that th	ey must be ans	wered in or	der to
23. Did you have a problems?	olanned induction	n of labour or <i>pl</i>	anned caesar	ean section due	to your bloo	d pressure
,	Yes, planned induction of labour because of blood pressure issues in pregnancy	Yes, planned caesarean section because of blood pressure issues	Yes, planned caesarean for other reasons than blood pressure	No planned induction and no planned caesarean section	I cannot remember	Not applic
First Pregnancy						
Second Pregnancy						
Third Pregnancy						

LONG TERM HEALTH RISKS

This section is about some long term health risks that some women may experience after having had blood pressure problems in pregnancy.

For this section we would like you to think about the long-term health risks of a woman who has been diagnosed with high blood pressure in pregnancy.

Not everyone who experienced blood pressure problems in pregnancy will necessarily have health issues in the future. We would not want you to unnecessarily worry about any of these risks, therefore we will provide you with further information about long-term health after high blood pressure in pregnancy at the end of the survey.

* 24. FOR WOMEN WITH HISTORY OF BLOOD PRESSURE PROBLEMS IN PREGNANCY:

Compare yourself to a woman who has NOT had blood pressure problems in pregnancy.

Do you think you have a lower/same/higher chance of getting the following:

	Less chance than a woman without blood pressure in pregnancy	Same chance as a woman without blood pressure in pregnancy	Higher chance than a woman without blood pressure in pregnancy	Not sure/I do not know
High blood pressure later in life				
Diabetes				
Kidney disease				
Breast cancer				
Heart attack				
High blood pressure in another pregnancy				
Stroke				
Heart disease				
Seizures				
Vascular Disease				

If you are concerned by any of the above potentially affecting you, information is available at the end of the survey where risks are explained.

For this section we would like you to think about the long term health risks of a woman who has been diagnosed with high blood pressure in pregnancy.

Same chance as a

Higher chance than a

* 25. FOR WOMEN WITHOUT HISTORY OF BLOOD PRESSURE PROBLEMS IN PREGNANCY:

Compare yourself to a woman who HAS had blood pressure problems in pregnancy.

Do you think you have a lower/same/higher chance of getting the following:

Less chance than a

	woman with blood pressure in pregnancy	woman with blood pressure in pregnancy	woman with blood pressure in pregnancy	Not sure/I do not know
High blood pressure later in life	\bigcirc		\bigcirc	
Diabetes				
Kidney disease				
Breast cancer				
Heart attack				
High blood pressure in another pregnancy				
Stroke				
Heart disease				
Seizures				
Vascular Disease				
26. How many years a symptoms of the pote	ntial risks may start to	appear?	do you think the vario	ous signs and
10-15 years after preg		O Not sur	re/do not know	
16-20 years after preg	gnancy			
Other (please specify))			

If you are concerned by any of the above potentially affecting you, information is available at the end of the survey where risks are explained.



WOI	men
* 27. How many years after blood pressure problems symptoms of the potential risks may start to appear	
< 10 years after pregnancy	> 20 years after pregnancy
10-15 years after pregnancy	Not sure/do not know
16-20 years after pregnancy	I don't think I will get any of these as I maintain a healthy lifestyle
Other (please specify)	

IN CASE OF DISTRESS

If you experience any distress caused due to the completion of this survey, please contact your GP or the Principal Investigator of this study, Dr Amanda Henry on 02 91132315 or Amanda.henry1@health.nsw.gov.au

For more information on this topic please visit The Australian Heart Foundation on the following link: https://www.heartfoundation.org.au/your-heart/women-and-heart-disease/womens-stories



TYPE OF CARE RECEIVED DURING PREGNANCY WHERE A BLOOD PRESSURE PROBLEM WAS DIAGNOSED

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue.

It is quite likely that some of the following questions may bring back some memories or bring rise to emotions that you find difficult to deal with. Please contact the Principal Investigator, Dr Amanda Henry on 02 91132315 or via email Amanda.henry1@health.nsw.gov.au if you would like to discuss these concerns. For medical assistance you can consult your General Practitioner.

blood pressure problem in pregnancy?	ency Unit of intensive Care Unit as a result of your
Yes	
○ No	
I am not sure	
I cannot remember	
* 29. Have any of your babies been admitted to 'Neona Care Nursery' as a result of your blood pressure prol Yes No I am not sure	atal Intensive Care', 'High Dependancy Unit' or 'Special blem in pregnancy?
30. After your baby was born have you had any of th	e following? (select all that apply)
Blood pressure measurement in hospital	Consultation with a renal (kidney) specialist
Blood pressure measurement with my GP	Consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with a consultati
Consultation with an obstetrician	specialises in complications of pregnancy like high blood pressure)
	I cannot remember
Other (please specify)	

Increased risk of high blood pressure	Risk of hypertensive disease in your next pregnancy
Increased risk of kidney problems	I was told to eat a healthy diet, do some exercise and
Increased risk of stroke	normally No risks were discussed
Increased risk of heart attack	
Increased risk of vascular disease	I cannot remember
Other (please specify)	
0	

TYPE OF CARE RECEIVED DURING PREGNANCY WHEN A BLOOD PRESSURE PROBLEM WAS DIAGNOSED

Questi continu		risk (*) simply mean	that they must be answered in	order to	
32. I	32. If a healthcare provider did speak to you about your future health risks, when did this occur?				
	Before birth		6 months to 1 year		
	Immediately after birth		1 year and over		
	Within first 6 weeks		I cannot remember		
	6 weeks to 6 months				
* 33. \	When would be a good time to re	eceive information abo	out long term health risks in your	gestational	
	ertension or preeclampsia exper		ý		
	During pregnancy/at birth	12 months after birth	24 months after birth		
* 34. As a result of your pregnancy affected by blood pressure problems, were you referred to any of the below after your baby was born? (tick all that apply)					
	Cardiologist		Fitness centre for exercise		
	Renal (kidney) Physician		Nutritionist for dietary adjustment		
	General Practitioner		I cannot remember		
	Other (please specify)				

EDUCATION

In this section we would like to find out about your preferred ways of getting information about long term health after gestational hypertension and preeclampsia.

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue

ontin	ontinue.				
	After experiencing gestational hypertension or preeclampsia what do you want to know about your long n health? (select all that apply)				
	Risk reduction for subsequent pregnancies				
	Reducing risk behaviours (eg. diet, exercise, smoking cessation)				
	Statistics (eg. increased risk)				
	At what point does the risk increase				
	Signs and Symptoms				
	Where to find information				
	How to discuss the matter with my healthcare provider				
	Impact on my children from the pregnancy affected by blood pressure problems				
	Other (please specify)				
* 36.	How do you want to receive the information? (select all that apply).				
	Key organisations addressing heart health (e.g. The Australian Heart Foundation or Her Heart)				
	Social Media channels (e.g. Instagram, Facebook, Twitter)				
	Brochures/Flyers				
	Medical Professionals				
	Podcasts/Media				
	Online videos				
	Other (please specify)				



POST-SURVEY OPTIONS AND SUMMARY OF RISK PROFILE

Some further things we would like you to consider:

38. We would like to have your opinion on what you think would be appropriate education material to improve women's knowledge. We would also like to know how the education could best be distributed to women. Would you like to participate in a **focus group** (one off - 2hrs max at St George Hospital, Sydney) or a **telephone interview** (one off 30 mins max)? By involving 'consumers', meaning women like you, the education package will have added value.

If you are interested in participate in either, you can leave your details here.

Name	
Email Address	
Phone Number	

39. Please leave your email address to **receive results** from this study (in approx. 1 year). Your details will not be used for any other purpose.

Email Address	

IN CASE OF DISTRESS

If you experience any distress caused due to the completion of this survey, please contact your GP or the Principal Investigator of this study, Dr Amanda Henry on 02 91132315 or Amanda.henry1@health.nsw.gov.au

For more information on this topic please visit The Australian Heart Foundation on the following link: https://www.heartfoundation.org.au/your-heart/women-and-heart-disease/womens-stories



RISK PROFILE - LONG TERM RISKS AFTER BLOOD PRESSURE PROBLEMS IN PREGNANCY EXPLAINED

You may like to take a screenshot of the risk profile so you can refer back to it whenever you need to.

RISK PROFILE

Although most women will experience good long-term health after having high blood pressure in pregnancy, there are, unfortunately, some long term health risks associated with having had high blood pressure in pregnancy.

Women who have had high blood pressure during pregnancy are about 3 to 4 times more likely to develop chronic hypertension than women who did not have a blood pressure problems in pregnancy. They are also about twice as likely to get diabetes in later life, even if they did not have diabetes during pregnancy.

Blood pressure diseases are also more likely to happen in the next pregnancy to women who have already had a previous blood pressure problems in pregnancy compared to women who have not. Therefore, if they have had a pregnancy with blood pressure problems, it is important to be seen early in their next pregnancy. There are treatments that can decrease the chance of recurring problems.

Women are also more likely to get various forms of cardiovascular disease (heart disease, stroke, vascular disease) if they have had gestational hypertension, preeclampsia and/or chronic hypertension. All of these cardiovascular problems are about twice as likely to eventually happen to a woman who has had blood pressure problems in pregnancy compared to a woman who has not. This still means most women will not have heart disease or diabetes after having high blood pressure in pregnancy, especially if they can avoid risk factors like smoking or excessive weight gain, and maintain a healthy diet and exercise.

Kidney problems are about 5 to 10 times more common after preeclampsia in particular. Although the relative risk of developing kidney problems is substantially higher after preeclampsia, the absolute risk of long-term kidney disease is still low. Unless the woman already had a kidney problem, well over 90% of women after preeclampsia and gestational hypertension will not have a kidney problem.

Fortunately, although seizures may occur as a result of preeclampsia during pregnancy, women have no higher long term risk of seizures compared to women who did not have a complicated pregnancy. There is no increased risk of getting cancer (e.g. breast cancer, leukaemia) after having high blood pressure in pregnancy.

For all the long term health risks, these start to go up within 10 years after an affected pregnancy and are ongoing after that. Therefore, it is recommended that women attend regular blood pressure checks with their GP and discuss any changes they can make to improve their general health. For more general information about heart health and managing health risks, please visit the National Heart Foundation website: https://www.heartfoundation.org.au/your-heart/know-your-risks

THANK YOU FOR YOUR PARTICIPATION

STROBE Statement— Assessing Australian women's knowledge about cardiovascular risk after hypertensive disorders of pregnancy and exploring information content and format preferences

	Item No	Recommendation	Page No
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or	1
		the abstract	
		(b) Provide in the abstract an informative and balanced summary of what	2
		was done and what was found	
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported	3-4
Objectives	3	State specific objectives, including any prespecified hypotheses	2
Methods		1	1
Study design	4	Present key elements of study design early in the paper	2&
study design		Tresent ney croments of study design early in the puper	5-6
Setting	5	Describe the setting, locations, and relevant dates, including periods of	2&6
~~·········	J	recruitment, exposure, follow-up, and data collection	
Participants	6	(a) Cohort study—Give the eligibility criteria, and the sources and	5-6
.		methods of selection of participants. Describe methods of follow-up	
		Case-control study—Give the eligibility criteria, and the sources and	
		methods of case ascertainment and control selection. Give the rationale	
		for the choice of cases and controls	
		Cross-sectional study—Give the eligibility criteria, and the sources and	
		methods of selection of participants	
		(b) Cohort study—For matched studies, give matching criteria and	-
		number of exposed and unexposed	
		Case-control study—For matched studies, give matching criteria and the	
		number of controls per case	
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders,	-
		and effect modifiers. Give diagnostic criteria, if applicable	
Data sources/	8*	For each variable of interest, give sources of data and details of methods	-
measurement		of assessment (measurement). Describe comparability of assessment	
		methods if there is more than one group	
Bias	9	Describe any efforts to address potential sources of bias	16
Study size	10	Explain how the study size was arrived at	5-6
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If	7
		applicable, describe which groupings were chosen and why	
Statistical methods	12	(a) Describe all statistical methods, including those used to control for	7
		confounding	
		(b) Describe any methods used to examine subgroups and interactions	
		(c) Explain how missing data were addressed	
		(d) Cohort study—If applicable, explain how loss to follow-up was	
		addressed	
		Case-control study—If applicable, explain how matching of cases and	
		controls was addressed	
		Cross-sectional study—If applicable, describe analytical methods taking	
		account of sampling strategy	

(e) Describe any sensitivity analyses

Continued on next page

Results Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers potentially	8
1 articipants	13	eligible, examined for eligibility, confirmed eligible, included in the study,	0
		completing follow-up, and analysed	
		(b) Give reasons for non-participation at each stage	F: 1
D ' ' '	1 4 1/4	(c) Consider use of a flow diagram	Fig 1
Descriptive	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and	8-9
data		information on exposures and potential confounders	Tab1
		(b) Indicate number of participants with missing data for each variable of interest	8
		(c) <i>Cohort study</i> —Summarise follow-up time (eg, average and total amount)	
Outcome data	15*	Cohort study—Report numbers of outcome events or summary measures over time	
		Case-control study—Report numbers in each exposure category, or summary	
		measures of exposure	
		Cross-sectional study—Report numbers of outcome events or summary measures	
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and	
		their precision (eg, 95% confidence interval). Make clear which confounders were	
		adjusted for and why they were included	
		(b) Report category boundaries when continuous variables were categorized	
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a	
		meaningful time period	
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and	
		sensitivity analyses	
Discussion			
Key results	18	Summarise key results with reference to study objectives	9-10
•			Table
			3&4
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or	15-
		imprecision. Discuss both direction and magnitude of any potential bias	16
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations,	10-
. · ·		multiplicity of analyses, results from similar studies, and other relevant evidence	15
Generalisability	21	Discuss the generalisability (external validity) of the study results	17
Other informati		2.2000 in Solicians (citerian variaty) of the study feeding	1 1
		Cive the course of funding and the role of the funders for the present stride and if	10
Funding	22	Give the source of funding and the role of the funders for the present study and, if	19

^{*}Give information separately for cases and controls in case-control studies and, if applicable, for exposed and unexposed groups in cohort and cross-sectional studies.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.

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Assessing Australian women's knowledge and knowledge preferences about long-term health after hypertensive disorders of pregnancy: a survey study

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Assessing Australian women's knowledge and knowledge
preferences about long-term health after hypertensive
disorders of pregnancy: a survey study

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- **Keywords:** Cardiovascular risk, women, preeclampsia, gestational hypertension,
- 25 long-term cardiovascular health, preventive health

ABSTRACT

Objective(s): To (a) assess women's current knowledge regarding long-term cardiovascular health after hypertensive disorders of pregnancy (b) elicit women's preferred educational content and format regarding health after hypertensive disorders of pregnancy. Design and setting: A custom-created online survey exploring Australian women's knowledge about long-term health after hypertensive disorders of pregnancy, distributed through consumer groups and social media. **Participants:** 266 women with (n=174) or without (n=92) a history of hypertensive disorders of pregnancy. **Primary and secondary outcome measures**: 1) Proportion of women identifying long-term health risks after hypertensive disorder of pregnancy using a 10-point risk knowledge score with 0-4 'low', 4.1-7.0 'moderate' and 7.1-10 'high'. 2) Exploration of preferred content, format and distribution of educational material post hypertensive disorder of pregnancy. **Results**: Knowledge scores about health after hypertensive disorder of pregnancy were moderate in groups with and without a history of the disorder. Knowledge was highest regarding risk of recurrent hypertensive disorders in a subsequent pregnancy, 'moderate' for chronic hypertension and heart attack, 'moderate' and 'low' regarding risk of heart disease and 'low' for diabetes and renal disease. Only 36% of all participants were aware that risks start within 10 years after the affected pregnancy. The majority of respondents with a history of

hypertensive disorder of pregnancy (76%) preferred receiving information about long-

term health 0-6 months postpartum from a healthcare provider (80%), key organisations (60%), social media (47%) and brochures/flyers (43%).

Conclusion(s): Women's knowledge regarding health risks after hypertensive disorder of pregnancy was 'moderate', although with important disease-specific gaps such as increased risk of diabetes. Most women wanted to be informed about their long-term health from a healthcare provider.

STRENGTHS AND LIMITATIONS OF THIS STUDY

- Consumer co-created survey exploring health knowledge after hypertensive
 pregnancy.
- For the first time survey results include findings from women with a history of gestational hypertension as well as from women without a history of hypertensive disorder of pregnancy.
 - Recruitment from groups with potentially greater baseline knowledge may bias results, although substantive knowledge gaps still found.
 - Although surveys were available in English, Arabic and Mandarin, there remains
 potential sub-optimal coverage of culturally and linguistically diverse groups.

INTRODUCTION

Hypertensive disorders of pregnancy (HDP) include chronic hypertension (CH), preeclampsia (PE) and gestational hypertension (GH) and complicate 5-10% of pregnancies¹. PE is a multi-system disorder characterised by new-onset hypertension after 20 weeks' gestation and involvement of one or more other organ systems and/or the fetus² ³. GH is new-onset hypertension after 20 weeks' gestation without any other complications. Apart from GH itself being considered an adverse pregnancy outcome

and it carrying an increased risk of progression to PE²³, is not associated with adverse pregnancy outcomes. However, both conditions are associated with long-term cardiovascular and other chronic disease sequelae⁴⁵. CH is defined as hypertension that is confirmed before pregnancy or before 20 completed weeks gestation, which may worsen during pregnancy and/or on which preeclampsia may be superimposed². Globally, cardiovascular disease (CVD) is one of the leading causes of death in women,⁶ and for women who have experienced an HDP, it is 2-3 times higher compared with those who did not⁴⁷⁸. This risk of premature death is present within 10 years after the affected pregnancy⁷⁹¹⁰ and remains after adjusting for the presence of other cardiovascular risk factors. There is also an increasing body of recent research linking PE and GH with other major chronic diseases including chronic kidney disease, end-stage kidney disease, and Type 2 diabetes mellitus¹¹⁻¹⁴.

Both Australian and international societies, including the Society of Obstetric Medicine of Australia and New Zealand (SOMANZ) and the International Society for the Study of Hypertension in Pregnancy (ISSHP), recommend that women and healthcare providers (HCP) are provided with information about HDP and later CVD²³. This includes recommending that women have a clinical review several months postpartum, and regular general practitioner (GP) follow-up to monitor blood pressure, fasting lipids and blood sugar². Adopting a healthy lifestyle with maintenance of an ideal weight and regular aerobic exercise is emphasised²³. The aims of this study were to (a) explore Australian women's current knowledge on the topic of long-term CVD health after any HDP, not just PE and (b) elicit women's preferred educational content and format regarding health after HDP, as a basis for creating tailored information and health advice for women after HDP.

METHOD

A national survey of women with and without a history of HDP was conducted, using a custom-created, face-validated online survey. Ethical approval was provided by the relevant Human Research Ethics Committee (HREC 18/POWH/326, REGIS 2019/PID05668).

Patient and Public involvement

As a validated instrument to assess women's knowledge was unavailable, a survey was custom designed. Initially, women with a history of HDP, comprising nine volunteers from the Postpartum physiology, psychology and paediatric follow up study (P4 Study)¹⁵ and Australian Action on Preeclampsia (AAPEC), were invited to take part in group interviews which addressed the possible content and design of the survey, tested the survey for face validity and provided feedback for improvement. The topics discussed during the interviews were sourced from findings from a scoping literature review¹⁶ and further complemented by questions specifically exploring the Australian context for women experiencing HDP. Nine women participated in the facevalidation process and commented on content, language, flow, survey structure including length, whether the introduction and the risk profile proposed for the end of the survey were informative as well as using appropriate language. Following feedback and integration of suggestions from the women, the survey was modified until consensus over a final version was achieved among study investigators, including the consumer representative (LH). The survey was made available in English, Arabic and Mandarin.

Data collection

The final survey was targeted at women in Australia, 18 years and older with a history of pregnancy in the last 3 years. Women who were currently pregnant were requested to only complete the survey if they had no major issues in their current pregnancy. Women not currently pregnant with pregnancy in the preceding three years were eligible either if they had experienced HDP (CH, GH or PE) or a pregnancy without any serious complications. The online survey, using SurveyMonkeyTM, was open from July to August 2019. Survey distribution occurred through the P4 study participants, organisations such as AAPEC, maternity consumer groups as well as via the project's consumer representative and social media (Facebook and Twitter) including multicultural networks in order to reach Arabic and Mandarin speaking communities. A targeted convenience sample was selected. Prior to acknowledging their voluntary participation at the commencement of the survey, women were presented with an introductory letter outlining the details of the study (Appendix 1). Commencement of the survey was then taken as consent to participate.

The data collection instrument

The survey for women (Appendix 1) explored demographic details, assessed obstetric history, history of HDP and other medical history including family history. The survey was tailored to women's self-reported HDP history (GH, PE, CH with or without worsening in pregnancy or superimposed PE, no hypertension history), with women given definitions of HDP conditions early in the survey to aid their self-report.

Questions focused on knowledge of risk after pregnancy, provision of care and education following birth and what information and education women would like to receive. Women with a history of GH, PE or CH were asked to classify their perceived

risk (based on their own lived HDP sub-type) of experiencing various long-term health outcomes as greater, less than or equal to that of a woman with a normotensive pregnancy. Women who experienced a normotensive pregnancy were also asked to classify whether they believed women who had had HDP were at greater, lesser, or equal risk. The survey included two 'distractor' conditions not known to have an increased risk after HDP (breast cancer and seizures) to elicit negative answers and ascertain whether women could identify what they were not at increased risk of after HDP as well as what they were at risk of. At survey completion, women were provided with a correct risk profile summary and a link to further information.

Data Analysis

Quantitative survey analysis was undertaken using SPSS Version 25 (SPSS Statistics for Windows, Armonk, NY). Demographic data and responses to individual questions were analysed descriptively. To examine difference in knowledge levels amongst the targeted subgroups, (GH, PE, CH in pregnancy, no hypertension history) responses regarding HDP and future health risks were compared using Chi-squared testing or likelihood ratio for categorical data (as appropriate to subgroup sample size) and one-way ANOVA testing for continuous data. A *p* value of <0.05 was considered statistically significant.

A knowledge score was created for the risk matrix whereby 1 point was allocated to the correct answer, 0 for the incorrect answer, 0 for 'I do not know' and 0 for no answer/left blank. A mean knowledge score for each condition/health outcome was calculated and a scale of 'low', 'moderate' and 'high' knowledge was established. The ranking classifications were chosen based on the data distribution and were divided

into three score categories. For each individual condition/health outcome's mean score, 'low knowledge' equated to a mean of 0.00-0.40, 'moderate knowledge' was 0.41-0.70 and 'high knowledge' a mean of 0.71-1.00. An overall mean score out of 10 (as there were 10 conditions) was calculated for the HDP and non-HDP groups (i.e. the HDP group's knowledge regarding their long-term health risks and the non-HDP group's knowledge regarding the long-term health risks of HDP women). This overall score was classified as 'low' 'moderate' or 'high' using the same mean ranges as were used for the individual conditions. Categorical analysis for proportions of each knowledge group ('high', 'moderate' and 'low') was also conducted to provide a further perspective.

RESULTS

In total, 308 survey responses were received (Figure 1). Forty-two were excluded: 40 for discontinuing the survey and not answering the question asking about their perception of lower/same/higher risk with regards to 10 health conditions, and two with 'CH only' (no worsening hypertension or superimposed PE in pregnancy) who were excluded due to small numbers. Of 266 included responses, 174 (65%) women had a lived experience of any HDP (will be known as HDP for reporting purposes) and 92 (35%) did not (will be known as non-HDP for reporting purposes). The HDP group consisted of 15 women with GH only (9%), 143 women with PE only (82%) and 16 women with CH plus superimposed pregnancy hypertension or PE (9%; will be known as CH). Of the HDP group, 123 (71%) had their most recent experience with HDP less than three years prior (32% <1 year prior and 39% 1-3 years prior).

Most respondents were in the 26-35 or 36-45 year age groups (91%), 89% were of Caucasian ethnicity, 90% pursued education beyond secondary school and 96% were in a relationship (Table 1). HDP women were more likely to be Caucasian, to have a history of diabetes, renal problems, be overweight and to have at least one additional cardiovascular risk factor than non-HDP women (Supplementary Table 1), and less likely to be university-educated. Half of all participants were sourced through social media (50%), with most of the remainder (45%) recruited via the P4 study (8% of HDP women, 46% of non-HDP women) and AAPEC (35% of HDP women).

Table 1: Respondent demographics

	Total	GH	PE	СН	Total HDP	Total Non HDP	P value HDP vs non
	N(%)	%	%	%	N (%)	N(%)	HDP
Total N	266	15	43	16	174	92	
AGE							
18-25	12 (5)	13	6	-	10 (6)	2 (2)	0.16
26-35	117 (44)	33	48	50	81 (47)	36 (39)	0.25
36-45	126 (47)	53	42	50	76 (44)	50 (54)	0.10
45+	10 (3)	-	5)	7 (4)	3 (3)	0.75
Prefer not to answer	1 (0)	-	-	-	-	1 (1)	-
ETHNICITY							
Caucasian	236 (89)	93	97	81	165 (95)	71 (77)	<0.001
Asian	23 (9)	7	2	6	5 (3)	18 (20)	<0.001
Other ¹	7 (3)	-	1	13	4 (2)	3 (8)	0.65
HIGHEST EDUCATIONAL ATT	AINMENT						
Secondary School	25 (9)	7	14	13	23 (13)	2 (2)	0.003
Diploma/Trade*	69 (26)	33	33	50	60 (35)	9 (10)	<0.001
University Degree	171 (64)	60	52	38	90 (52)	81 (88)	<0.001
Prefer not to answer	1 (0)	-	1	-	1 (1)	-	-
RELATIONSHIP STATUS							
In a relationship	254 (96)	100	93	88	162 (93)	92 (100)	0.001
Not in a relationship	11 (4)	-	6	13	11 (6)	0 (1)	0.001
Prefer not to answer	1 (0)	-	1	-	1 (1)	-	-
RECRUITED TO SURVEY VIA	4						
P4 Newsletter	56 (21)	13	8	6	14 (8)	42 (46)	<0.001
AAPEC	62 (23)	7	40	19	61 (35)	1 (1)	<0.001
Social Media	134 (50)	80	52	69	97 (56)	37 (40)	0.02
Other ²	14 (5)	-	1	6	2 (1)	12 (13)	<0.001

^{*} Diploma or Trade certificate

¹ Other: Indigenous Australian (n=1), Polynesian or Maori (n=2), mixed ethnicity (n=4).

² Other: Friend (n=11), ACM (n=1), Clinic (n=1), Maternity Consumer group other than AAPEC (n=1).

PE = preeclampsia GH = gestational hypertension CH = chronic hypertension, worsening in pregnancy and/or superimposed preeclampsia P4= Postpartum Physiology, Psychology, and Paediatric Study AAPEC= Australian Action on Preeclampsia ACM= Australian College of Midwives.

Table 2: Means of risk factor knowledge of women listed by type of HDP

	GH n=15	PE n=143	CH n=16	HDP n=174	Non HDP n=92	P Value HDP vs non HDP
Chronic Hypertension	0.53	0.78	0.81	0.76	0.62	0.02
Diabetes	0.27	0.24	0.31	0.25	0.35	0.12
Renal Disease	0.27	0.54	0.69	0.53	0.21	<0.001
Heart Attack	0.53	0.69	0.75	0.68	0.52	0.01
Repeat HDP	0.87	0.90	0.94	0.90	0.71	<0.001
Stroke	0.47	0.62	0.81	0.63	0.53	0.14
Heart Disease	0.47	0.69	0.75	0.68	0.50	0.005
PVD	0.33	0.50	0.50	0.32	0.45	<0.001
Breast Cancer*	0.20	0.52	0.31	0.47	0.65	0.004
Seizures*	0.27	0.29	0.13	0.27	0.44	0.01
OVERALL MEAN KNOWLEDGE SCORE (OUT OF 10)	4.2	5.8	6.0	5.6	5.2	0.21

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy, PVD= Peripheral Vascular Disease * Breast cancer and seizures are distractors within the survey. These were included despite being conditions that women after HDP are not at greater risk of.

MEAN SCORE					
LOW	0-0.40				
MODERATE	0.41-0.70				
HIGH	0.71-1				

Average knowledge scores are shown in Table 2 and detailed results on which these scores are based are shown in Supplementary Tables 2-11. Overall knowledge of the 10 conditions in the survey was 'moderate' for both groups (5.6/10 amongst HDP and 5.2/10 amongst non-HDP, p=0.21), with 33% in both groups having "high" overall knowledge and 32% and 40% respectively having "low" overall knowledge (Supplementary Table 12). Women with a history of HDP had 'high' knowledge with regards to recurrence of HDP in a subsequent pregnancy (0.90) and risk of future CH

(0.76). The same group had 'moderate' knowledge regarding increased chance of conditions such as heart attack (0.68), heart disease (0.68) and stroke (0.63). Women without HDP history had 'high' knowledge (0.71) for HDP recurrence in a subsequent pregnancy. The same group of women had moderate knowledge of chronic hypertension (0.62) and stroke (0.53). Lowest knowledge across both groups was around the risk of future diabetes (0.25 HDP group and 0.35 for non-HDP group). Further 'low' scoring conditions were peripheral vascular disease (PVD) and renal disease. For most conditions HDP women had significantly higher knowledge than the non-HDP group. However, the non-HDP group were more likely to correctly identify that the risk of the two 'distractor' conditions, seizures or breast cancer, were equal for both groups.

Supplementary Table 13 shows knowledge score breakdown by time since pregnancy. In the subgroup of HDP women who experienced PE (n=143), average knowledge was similar amongst women who experienced HDP within the last three years (5.8/10), compared to those who experienced HDP more than three years ago (5.7/10). Of the HDP women, only 32% were aware that the cardiovascular conditions may start manifesting within 10 years after an affected pregnancy, compared with 45% of women in the non-HDP group (p=0.036). About a third in each group (30% HDP, 36% non-HDP) were unsure about timing of risk rise/when health conditions manifest (Supplementary Table 14).

Women with HDP history were asked about their personal experience of risk discussion with healthcare providers (Table 3 represents summary of collective HDP data, Supplementary Table 15 provides all findings by HDP sub-group). The most

frequent discussions about future health were regarding HDP in subsequent pregnancies (45%), risk of chronic hypertension (43%), and 'No discussion' (37%). Risk discussions were no more likely to have occurred in women with HDP less than 3 years ago or over 3 years ago. There were also no statistically significant differences found between HDP subgroups about whether future risks were discussed, or what types of risk were discussed.

Table 3: Proportion of conditions discussed when addressing future risk (multiple answers collected) within and over 3 years since last HDP (summary of collective HDP data)*

	n =	otal = 174 (%)	Overall Total n = 174 N (%)	P value < 3 yrs vs >3 yrs	
	<3yrs	>3yrs			
HDP next pregnancy	55 (45)	24 (47)	79 (45)	0.78	
Chronic Hypertension	55 (45)	19 (37)	74 (43)	0.37	
No discussion	45 (37)	19 (37)	64 (37)	0.93	
Lifestyle changes	32 (26)	8 (16)	40 (23)	0.14	
Heart Attack	22 (18)	7 (14)	29 (17)	0.50	
Renal Disease	23 (19)	4 (8)	27 (16)	0.07	
Stroke	20 (16)	6 (12)	26 (15)	0.45	
Peripheral vascular disease	16 (13)	5 (10)	21 (12)	0.56	
Cannot remember	6 (5)	2 (4)	8 (5)	0.78	

*Table represents frequency of each option; percentages add to over 100% as women were asked to select any/all that applied. PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

When asked about preferences of the timing of a future risk discussion, the majority (76%) of women wanted a discussion 0-6 months postpartum. The topics most women with HDP wished to discuss (Table 4) are 'impact on my children from the pregnancy affected by HDP' (73%), 'signs and symptoms of the conditions' (67%), 'when does risk rise' (54%) and 'risk reduction for subsequent pregnancy' (54%). HDP women's preference for receiving information on long-term health after HDP is via a medical professional (80%), through key organisations such as The Australian Heart Foundation (60%) and social media (47%).

Table 4: HDP women's preferences for content and distribution of information/education on future risk after HDP (multiple answers collected) in order of preference

	GH	PE	СН	Total HDP
	%	%	%	N (%)
Total N	15	143	16	174
Preference of Discussion Topics*				
Impact on my children from the pregnancy affected by HDP	73	80	63	136 (73)
Signs and Symptoms of the conditions	80	71	69	124 (67)
Risk reduction for subsequent pregnancy	40	62	44	101 (54)
When does the risk rise	40	61	50	101 (54)
Statistics	40	60	38	98 (53)
Reducing risk behaviours (diet, exercise, smoking cessation)	40	56	31	91 (49)
Where to find information	40	51	13	81 (44)
How to discuss the matter with my healthcare provider	27	40	25	65 (35)
Preference of Distribution*				
Medical professionals	73	82	75	140 (80)
Key organisations	53	61	63	105 (60)
Social Media	40	51	19	82 (47)
Brochures/Flyers	40	45	31	75 (43)
Online Videos	20	24	25	42 (24)
Podcast/Media	13	23	25	39 (22)

*Table represents frequency of each option; percentages add to over 100% as women were asked to select any/all that applied. PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

DISCUSSION

This study found overall, 'moderate' knowledge of health conditions after HDP amongst both HDP and non-HDP women. Amongst women with a history of HDP, highest knowledge was identified with regards to future risk of hypertension and repeat HDP in subsequent pregnancies. Conversely, knowledge of future risk of diabetes was low, as was knowledge of the "distractor" conditions among HDP women particularly. Diabetes as a future risk factor post HDP has previously not been reported on in

studies of women's knowledge, and our findings suggest this is an important knowledge gap to address.

Other novel aspects of our study are inclusion of women who had a history of GH as well as those with a history of PE, and assessing knowledge of non-HDP women's knowledge. Women after HDP had somewhat higher knowledge of most health risks than the non-HDP group, however non-HDP group also had better knowledge of some aspects such as timing of risk increase. However, both groups' knowledge of the early increase in risk was low, adding further concern and reason to address the knowledge gap. When looking at the proportion of participants scoring "high", these were equal between the HDP (33%) and non-HDP groups (33%), whilst proportions scoring "low" were similar enough (32% HDP versus 40% non-HDP) to not to show statistical significance. Our scoping review in 2019¹⁶ identified that post HDP, women have insufficient knowledge of their long-term risks. By including non-HDP women we wanted to explore whether knowledge was similar between the groups, which if so would suggest HDP women are not receiving tailored, targeted information and/or any information received is not translated into knowledge of personal risk after HDP. Given women after HDP were not markedly more knowledgeable about their health risks than unaffected women, the research-to-practice translational gap is further highlighted and suggests women with a lived experience of HDP remain underinformed about their increased CVD risk.

A further important finding was that many HDP women were not made aware of future health risks, with 37% of HDP women reporting to have had 'no discussion' about their increased long-term risk. Women with more recent HDP were no more likely than

women with HDP>3 years ago to report having risks discussed, which is concerning. This finding suggests risk discussions may not have improved in recent years despite updated guidelines emphasising long-term health^{2 3}, and that the extensive evidence regarding long-term implications for women after HDP continues to be lost in the translation of research to practice. We are exploring reasons for this (e.g. lack of evidence base in guidelines, lack of provider knowledge of guidelines, siloed healthcare with insufficient handover from maternity care team to primary care) in our broader work.

Women's knowledge after GH has not been previously reported as far as we are aware even though GH has similar frequency and similar future CVD risk as PE⁴ ¹⁷. Although only 9% of our sample were GH, this group had somewhat lower knowledge than the PE and CH groups regarding conditions after HDP (although mostly not reaching statistical significance). Over half reported receiving no discussion of health risks after GH. Despite the small number of women with a history of GH (n=15) contributing to the study, this suggests potential substantive knowledge gaps after GH to address in both women and healthcare providers.

International studies exploring women's knowledge have predominantly reported limited or no knowledge about the link between HDP and CVD¹⁶, though our study found overall, 'moderate' knowledge of health conditions after HDP. The two conditions associated with highest knowledge were repeat HDP and risk of future hypertension. Findings were similar in Traylor et al.'s¹⁸ survey conducted in the United States of America (USA), where 146 women post HDP were included (PE n=76, PE with severe features n=41, CH=29). Future hypertension and repeat HDP were

correctly identified by women as risk factors, however this knowledge was mainly reflected in the group of women who had experienced PE with severe features. In the United Kingdom (UK), Brown et al. 19 (n=12 women attending postnatal follow-up clinic) also found that women are aware of repeat HDP risks, however despite postnatal risk counseling, perception of hypertension and CVD risk was mainly associated with participants who had a family history of CVD. More recently in Australia, Hutchesson et al. 20 surveyed 127 women with PE in the two years prior, finding very high knowledge about future hypertension risk (96%, higher than our post-PE findings) and most were aware of stroke (67%) and CVD (66%) risks (similar to our findings). Over a third of women after PE had 'no discussion' about future risk in our study. Hutchesson et. al²⁰ reported over one third of their participants remained unaware of increased CVD risks, which is similar to our findings. Similarities may be explained by the fact that major source of PE participants for both, the Hutchesson et. al²⁰ survey and ours was the patient support/advocacy group AAPEC. Recruitment from this advocacy group may also explain a higher post-PE knowledge than other studies have reported.

Our study findings resonate with those from similarly targeted women in Canada, Portugal, UK, the USA and a previous Australian study, all conducted between 2013 and 2017¹⁶. Therefore, from a global perspective, these findings reinforce a persistent and concerning, research to consumer gap. With international guidelines, including ISSHP², specifically targeted to assist HCPs providing care to women on an international scale to better manage and address health after HDP, this practice gap of knowledge transmission to women would be expected to narrow.

Education preferences

Content

Women mostly wanted educational materials to address HDP impact on their children, signs and symptoms of conditions they are at higher risk of, the timing of when their risks rise, and how to best reduce risk of recurrent HDP. Similar preferences were expressed by the women included in Seely et al.'s²¹ focus group of 20 women after PE, with the key concern being the impact the PE pregnancy may have had on the health of their children. More recently, a UK-based study²², involving women with a history of HDP and healthcare providers (HCP), identified research priorities regarding HDP. The top-ranking priority identified was the long-term physical and mental health consequences of HDP for the woman, baby and family. Other 'uncertainties' expressed by participants regarding their lived experience of HDP included topics such as diagnosis and management in pregnancy, prevention of future complications, short and long-term consequences of HDP for the woman and the baby, prevention of recurrent HDP as well as educational needs of HCPs and support for women and their families. Our study, with focus on women in Australia, suggests that similar uncertainties may benefit from being addressed, hence validating the importance of our findings.

Format of education and access

Our study identified that women mostly wanted to receive information about long-term health after HDP from medical professionals. Key organisations who are experts on the topic, via social media and through information brochures were other acceptable avenues of access to information. This is in contrast to Skurnik et al's²³ focus group of 14 women after PE, whose preferences for educational materials about the link between CVD and PE were via pamphlets available in doctor's offices as well as via

online communities and topical blogs. However, Hird et al's²⁴ participants also expressed preference for healthcare providers as their information source, including wanting healthcare providers to guide them towards reliable online/external information sources rather than encounter irrelevant or potentially inaccurate information due to their self-initiated search. Hutchesson et al.²⁰ report that high knowledge amongst participants was mainly due to the women's own research rather than receiving all possible, relevant information from their healthcare provider. Overall, existing studies including ours would suggest that although women are very open to the use of online sources or information packs, their healthcare providers are seen as central to closing their knowledge gaps.

Time of risk discussion

An important element to consider when communicating about risk with women who have experienced GH or PE is the timing of these discussions, as situational factors of being a new mother may alter when women are most receptive to follow-up. In our study, three-quarters of the women preferred this to occur in the first six months after birth. As well as being their preference, this also aligns with the potential benefits of early intervention and would allow for addressing knowledge gaps found in this study around how soon the risk rises after HDP. Addressing future risk early but not immediately is also supported by Brown et al.'s study of women after PE, where participants suggested that six months postpartum was the timeframe where they felt they had transitioned into a more comfortable stage of parenting and were able to focus more on themselves again¹⁹.

Strengths and limitations

The survey was co-created via a formalised process of seeking input and feedback on the usability, language and content from women who have previously experienced HDP. Although face-validation is a subjective process, involving consumers with a history of HDP gives added value to the survey.

Our knowledge score is both a strength, as it allows for a summary of findings across all the conditions and risks, and a limitation, as assigning cut-points for knowledge ranking is an arbitrary designation. Having included the distractor conditions (breast cancer and seizures) may also have altered the overall score. Whilst women are more likely to experience seizures during a pregnancy complicated by HDP compared to non-HDP women, the long-term risk of seizures is similar for both groups. Similarly, the association of HDP and future increased risk of cancer (including breast cancer) has been examined in a systematic review and meta-analysis, however proven not to be associated with increased future risks after HDP²⁵. Distractor inclusion may well have lowered overall knowledge score, for example women believing that after HDP they are at more risk of ongoing seizures since this is a risk during PE-affected pregnancy. However, we believe inclusion of distractors and assessment of women's response to them is valid, as it is important for women to not incorrectly believe they are at increased risk of more conditions than they are, as well as having knowledge of their increased cardiovascular risk. The addition of women with a history of GH as well as women without any history of HDP, is also a strength to add broader perspective on this topic.

Limitations include demographic make-up of respondents, with HDP participants predominantly English speaking and Caucasian (95%) despite the survey being

available in Arabic and Mandarin as well as English. The non-HDP group (20% Asian background) had similar background demographics of Australian reproductive-aged women²⁶, and as HDP is more prevalent amongst the Caucasian population²⁷, the sample in the context of ethnic background actually is proportionally likely close to representative of Australian HDP and non-HDP women. However, it would have been preferable to also gain insight from more culturally and linguistically diverse groups in order to understand their knowledge base and address their needs within this context.

In the survey, women were asked to select their HDP history which was then used to group them for analysis. Women's diagnosis of HDP is by self-report is a limitation, as some bias may be introduced through inaccurate self-report of diagnosis. The broad geographical range and anonymous nature of the survey precluded any verification of diagnosis. However, women were provided with definitions of the various HDP conditions at the start of the survey to aid them in their self-report. Another limitation is where participants were recruited from, with close to half either drawn from the P4 study (an Australian post-HDP research study) or consumer group AAPEC. Therefore, there may be knowledge bias in the sample (i.e. a more knowledgeable group of participants than the overall HDP or non-HDP population). The women's level of active engagement in pursuing further information on their long-term risks as well as their level of motivation to participate in this study, further contributes to knowledge bias. The number of respondents in all included HDP subgroups are a small proportion of the total number of women experiencing HDP, which suggests volunteer bias and this affects generalisability. However, non-representative, specialised samples of women can be noted within most research addressing women's knowledge on long-term health after HDP¹⁶. As even this group with potentially greater baseline knowledge had substantive knowledge gaps, our study highlights the need for interventions to improve knowledge of health after HDP.

Implications

Close to two decades worth of data have been collected⁸ since research on the link between HDP and increased CVD risk emerged in the early 2000s, with the first systematic review published in 2007²⁵. It could be expected that this knowledge, by now, would have been translated into practice and shared with HDP women, however our findings suggest that this is still not the case. This study is valuable from the public health perspective, given the wider context of prevalence and importance of cardiovascular disease in women. Findings from this study and the broader study it is embedded in, will contribute towards the development, application and evaluation of educational materials for women and healthcare providers. These future projects will address persistent knowledge-to-practice-gaps regarding improving women's cardiovascular health after HDP. Given the prevalence and impact of both HDP and CVD, this is valuable for women's health, and public health more broadly.

well as counselling women with regards to their individual long-term CVD risk.

Although available to the public, these are not designed for women. Compiling suitable information for women would be an important step towards closing the knowledge gap. It is important to establish preferred content, presentation and timing of education for post-HDP health for women as we have in this study, to maximise the chance that

Guidelines such as ISSHP² and SOMANZ³ suggest regular follow-up after HDP as

women will engage with and benefit from education.

CONCLUSION

This Australian survey of women's knowledge of risks after HDP, found varying knowledge from the targeted groups. Despite 'high' knowledge being demonstrated regarding some risks, overall significant knowledge gaps were identified for certain conditions, particularly diabetes, and for knowledge about the relatively early timing of when health risks increase after HDP. Identifying these gaps are important in planning tailored education for women, and to improve early intervention for modifiable CVD risks in women after HDP. Addressing these women's preferences for content and to have this delivered by their healthcare provider may further lead to enhanced counselling, management and improved women's health trajectories.

LIST OF ABBREVIATIONS

- 508 AAPEC Australian Action on Preeclampsia
- 509 CH chronic hypertension worsening in pregnancy and/or with superimposed
- 510 preeclampsia
- 511 CVD cardiovascular disease
- 512 GH gestational hypertension
- 513 GP general practitioner
- 514 HCP healthcare provider
- 515 HDP hypertensive disorder of pregnancy
- 516 ISSHP International Society for the Study of Hypertension in Pregnancy
- 517 PE preeclampsia
- 518 PVD peripheral vascular disease
- 519 SOMANZ Society of Obstetric Medicine Australia New Zealand

STATEMENTS

Ethical approval

Ethical approval has been provided by South-Eastern Sydney Local Health District

Human Research Ethics Committee (Ref: 18/POWH/326). The ratification for the

University of Technology Sydney has also been obtained under ETH18-3061.

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Transparency statement and competing interests

The article is the authors' original work, has not received prior publication and is not under consideration for publication elsewhere. All the authors have seen and approved the manuscript being submitted. The manuscript is an honest, accurate, and transparent account of the study being reported, no important aspects of the study have been omitted. We have read and understood BMJ policy on declaration of interests and declare that we have no competing interests. This manuscript presents partial results from Heike Roth's PhD research. The project is supervised by Caroline Homer and Amanda Henry.

Authors contributions

Heike Roth, Amanda Henry and Caroline Homer contributed to the conception and design of the study as well as the distribution of the survey and writing of the manuscript. Heike Roth led the analysis of the survey data, drafting and designed the Tables, Figures and Appendixes and wrote the first draft. Grace LeMarquand was a medical Honours student assisting with pre-survey interviews as well as initial data analysis. Lynne Roberts assisted in the survey development, supported the distribution, the interpretation of the findings and the discussion. Mark Brown contributed to the design of the survey and supported the interpretation of the

findings and the discussion. As a maternity consumer, Lisa Hanley has assisted with the survey design and ensured appropriate use of language and content as well as supported the distribution. All authors contributed to drafts and revising of the paper and all approved the final version.

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Authors' information (optional)

This manuscript presents partial results from Heike Roth's PhD research. The project is supervised by Caroline Homer and Amanda Henry.

Data availability

Data are available upon reasonable request.

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TABLES, SUPPLEMENTARY TABLES, APPENDICES AND FIGURES

Table legend

Table 1: Respondent demographics

Table 2: Means of risk factor knowledge of women listed by type of HDP

Table 3: Proportion of conditions discussed when addressing future risk (multiple answers collected) within and over 3 years since last HDP (summary of collective HDP data)*

Table 4: HDP women's preferences for content and distribution of information/education on future risk after HDP (multiple answers collected) in order of preference

Supplementary data

Supplementary Table 1: Current CVD risk factors of HDP and non-HDP women (Multiple answers collected) in order of frequency

Supplementary Table 2: Chronic Hypertension

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 3: Diabetes

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 4: Renal Disease

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 5: Heart Attack

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 6: HDP next pregnancy

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 7: Stroke

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 8: Heart Disease

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 9: Peripheral Vascular Disease

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 10: Breast Cancer

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 11: Seizures

- a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")
- b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

Supplementary Table 12: Proportion of participants scoring 'high', 'moderate' and 'low' by type of HDP and non-HDP

Supplementary Table 13: Means of risk factor knowledge of women with a history of preeclampsia listed by time elapsed since HDP

Supplementary Table 14: HDP and non-HDP women's answers to timing of rise of risk with signs and symptoms starting to show

Supplementary Table 15: Proportion of conditions discussed by HDP sub-type when addressing future risk (multiple answers collected) within and over 3 years since last HDP (complements Table 3 which illustrates a summary of this data)

Appendices

Appendix 1: Survey for women

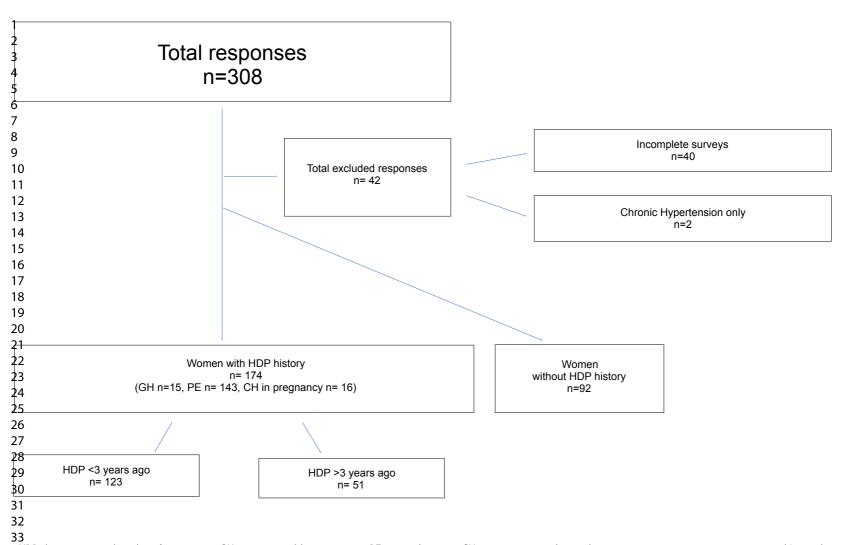
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Figure 1: Survey inclusion





Figure 1: Survey Inclusion



34DP- hypertensive disorder of pregnancy, GH - gestational hypertension, PE - preeclampsia, CH in pregnancy - chronic hypertension worsening in pregnancy and/or with 35 superimposed preeclampsia

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Supplementary Table 1: Current CVD risk factors of HDP and non-HDP women (Multiple answers collected) in order of frequence	Supplementar	v Table 1: Current CVD risk fac	tors of HDP and non-HDP wom	en (Multiple answers collected	d) in order of frequency
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	GH	PE	СН	Total HDP	non-HDP	P HDP vs non HDP
	n=15	n=143	n=16	n=174	n=92	
	n (%)*	n (%)*	n (%)*	n (%)*	n (%)*	
None of the options	6 (40)	52 (36)	2 (13)	60 (34)	57 (62)	<0.001
Obesity	5 (33)	38 (27)	7 (44)	50 (29)	10 (11)	0.001
Family History Heart disease	3 (20)	37 (26)	6 (38)	46 (26)	14 (15)	0.037
Hypertension	3 (20)	30 (21)	12 (75)	35 (20)	0 (0)	<0.001
Renal problems	0 (0)	21 (15)	2 (13)	23 (13)	0 (0)	<0.001
Smoking	0 (0)	10 (7)	3 (19)	13 (7)	2 (2)	0.075
History of cardiovascular event**	0 (0)	1 (1)	0 (0)	1 (1)	0 (0)	0.356
Diabetes	0 (0)	5 (3)	2 (13)	7 (4)	0 (0)	0.014
Alcohol consumption	0 (0)	0 (0)	1 (6)	1 (1)	0 (0)	0.356
High Cholesterol	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	-

^{*}Table represents frequency of each option; percentages add to over 100% as women were asked to select any/all that applied.

^{**}angina, stroke or heart attack

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

Supplementary Table 2: Chronic Hypertension

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

, and the second	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	1 (1)	0 (0)	1 (1)
Same likelihood	2 (13)	14 (10)	1 (6)	17 (10)
More likely (correct)	8 (53)	111 (78)	13 (81)	132 (76)
I do not know	5 (33)	17 (12)	2 (13)	24 (14)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92	
	n (%)	
Less likely (correct)	57 (62)	
Same likelihood	22 (24)	
More likely	1 (1)	
I do not know	12 (13)	

Supplementary Table 3: Diabetes

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

3,	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	1 (1)	0 (0)	1 (1)
Same likelihood	3 (20)	61 (43)	5 (31)	69 (40)
More likely (correct)	4 (27)	35 (25)	5 (31)	44 (25)
I do not know	8 (53)	44 (31)	6 (38)	58 (33)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n= 92	
	n (%)	
Less likely (correct)	32 (35)	
Same likelihood	39 (42)	
More likely	4 (4)	
I do not know	17 (19)	

Supplementary Table 4: Renal Disease

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	2 (1)	0 (0)	2 (1)
Same likelihood	2 (14)	30 (21)	1 (7)	33 (19)
More likely (correct)	4 (29)	77 (55)	11 (73)	92 (54)
I do not know	8 (57)	32 (23)	3 (20)	43 (25)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92	
	n (%)	
Less likely (correct)	39 (42)	
Same likelihood	31 (34)	
More likely	0 (0)	
I do not know	22 (24)	

Supplementary Table 5: Heart Attack

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

3,	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	1 (1)	0 (0)	1 (1)
Same likelihood	1 (7)	20 (14)	1 (7)	22 (13)
More likely (correct)	8 (53)	98 (69)	12 (80)	118 (69)
I do not know	6 (40)	23 (16)	2 (13)	31 (18)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely (correct)	48 (53)
Same likelihood	23 (25)
More likely	0 (0)
I do not know	20 (22)

Supplementary Table 6: HDP next pregnancy

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	2 (1)	0 (0)	2 (1)
Same likelihood	0 (0)	9 (6)	0 (0)	9 (5)
More likely (correct)	13 (87)	128 (89)	15 (100)	156 (90)
I do not know	2 (13)	4 (3)	0 (0)	6 (4)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely (correct)	65 (71)
Same likelihood	19 (21)
More likely	0 (0)
I do not know	8 (9)

Supplementary Table 7: Stroke

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174	
	n (%)	n (%)	n (%)	n (%)	
Less likely	0 (0)	1 (1)	0 (0)	1 (1)	
Same likelihood	1 (7)	18 (13)	1 (7)	20 (12)	
More likely (correct)	7 (47)	89 (63)	13 (87)	109 (64)	
I do not know	7 (47)	33 (23)	1 (7)	41 (24)	

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely (correct)	49 (53)
Same likelihood	24 (26)
More likely	0 (0)
I do not know	19 (21)

Supplementary Table 8: Heart Disease

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174	
	n (%)	n (%)	n (%)	n (%)	
Less likely	0 (0)	1 (1)	0 (0)	1 (1)	
Same likelihood	2 (13)	18 (13)	1 (7)	21 (12)	
More likely (correct)	7 (47)	99 (70)	12 (80)	118 (67)	
I do not know	6 (40)	24 (17)	2 (13)	32 (19)	

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the following")

	non-HDP n=92
	n (%)
Less likely (correct)	46 (50)
Same likelihood	27 (29)
More likely	0 (0)
I do not know	19 (21)

Supplementary Table 9: Peripheral Vascular Disease

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

,	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174
	n (%)	n (%)	n (%)	n (%)
Less likely	0 (0)	0 (0)	0 (0)	0 (0)
Same likelihood	2 (13)	20 (14)	1 (7)	23 (13)
More likely (correct)	5 (33)	72 (51)	8 (53)	85 (50)
I do not know	8 (53)	50 (35)	6 (40)	64 (37)

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely (correct)	41 (45)
Same likelihood	29 (32)
More likely	0 (0)
I do not know	22 (24)

Supplementary Table 10: Breast Cancer

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174	
	n (%)	n (%)	n (%)	n (%)	
Less likely	0 (0)	0 (0)	0 (0)	0 (0)	
Same likelihood (correct)	3 (20)	74 (53)	5 (36)	82 (48)	
More likely	1 (7)	1 (1)	1 (7)	3 (2)	
I do not know	11 (73)	66 (47)	8 (57)	85 (50)	

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely	6 (7)
Same likelihood (correct)	60 (65)
More likely	1 (1)
I do not know	25 (27)

Supplementary Table 11: Seizures

a) Women with HDP likelihood of risk compared to non-HDP ("Do you think you have a lower/same/higher chance of getting the following")

, and the same of	GH pregnancy n=15	PE pregnancy n=143	CH pregnancy n=16	Total n=174	
	n (%)	n (%)	n (%)	n (%)	
Less likely	0 (0)	0 (0)	0 (0)	0 (0)	
Same likelihood (correct)	4 (27)	41 (29)	2 (13)	47 (27)	
More likely	1 (7)	41 (29)	4 (27)	46 (27)	
I do not know	10 (67)	60 (42)	9 (60)	79 (46)	

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of pregnancy

b) Women without HDP likelihood of risk compared to HDP ("Do you think you have a lower/same/higher chance of getting the

following")

	non-HDP n=92
	n (%)
Less likely	25 (27)
Same likelihood (correct)	40 (44)
More likely	0 (0)
I do not know	27 (29)

Supplementary Table 12: Proportion of participants scoring 'high', 'moderate' and 'low' by type of HDP and non-HDP

	GH n=15 n(%)	PE n=143 n(%)	CH n=16 n(%)	P GH vs PE	P GH vs CH	P PE vs CH	HDP n=174 n(%)	non-HDP n=92 n(%)	P HDP vs non- HDP
High (score 8-10)	4 (27)	48 (34)	5 (31)	0.59	0.78	0.67	57 (33)	30 (33)	0.98
Moderate (score 5-7)	3 (20)	52 (36)	7 (44)	0.21	0.16	0.56	62 (36)	25 (27)	0.16
Low (score 0-4)	8 (53)	43 (30)	4 (25)	0.67	0.11	0.85	55 (32)	37 (40)	0.16
Total	15 (100)	143 (100)	16 (100)	-	-	-	174 (101)	92 (100)	1

PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia, HDP= hypertensive disorder of

The colour coding in addition to the mean scores aims to highlight the areas of 'low' (red), 'moderate' (orange) and 'high' (green) knowledge. \$ TO ringring...

LOW
MODERATE
HIGH

Supplementary Table 13: Means of risk factor knowledge of women with a history of preeclampsia listed by time elapsed since HDP

preeclampsia	PE n=143	y	PE n=143						PE n	=143	
	ALL		0 - 6 month	6 - 12 month	1-2 years	2-3 years	>3 years	Р	< 3	> 3	Р
			n=26	n=19	n=32	n=20	n=46		n= 97	n= 46	< 3 yrs vs >3 yrs
Chronic Hypertension	0.78		0.77	0.89	0.78	0.80	0.72	0.65	0.80	0.72	0.27
Diabetes	0.24		0.23	0.42	0.31	0.10	0.20	0.14	0.27	0.20	0.35
Renal Disease	0.54		0.50	0.74	0.41	0.65	0.52	0.17	0.55	0.52	0.78
Heart Attack	0.69		0.58	0.89	0.66	0.70	0.67	0.25	0.69	0.67	0.84
Repeat HDP	0.90		0.88	0.89	0.94	0.95	0.85	0.68	0.92	0.85	0.25
Stroke	0.62		0.65	0.68	0.50	0.65	0.65	0.62	0.61	0.65	0.62
Heart Disease	0.69		0.62	0.79	0.63	0.65	0.76	0.50	0.66	0.76	0.21
PVD	0.50		0.54	0.53	0.53	0.40	0.50	0.90	0.51	0.50	0.95
Breast Cancer*	0.52		0.46	0.37	0.63	0.40	0.59	0.25	0.48	0.59	0.26
Seizures*	0.29		0.15	0.16	0.41	0.45	0.26	0.07	0.30	0.26	0.64
OVERALL MEAN KNOWLEDGE SCORE OUT OF 10	5.8		5.4	6.4	5.8	5.8	5.7	0.83	5.8	5.7	0.89

The colour coding in addition to the mean scores aims to highlight the areas of 'low' (red), 'moderate' (orange) and 'high' (green) knowledge.

MEAN SCORE			
L	-OW	0-0.40	
N	MODERATE	0.41-0.70	
H	HIGH	0.71-1	

Supplementary Table 14: HDP and non-HDP women's answers to timing of rise of risk with signs and symptoms starting to show

	GH	PE	CH	HDP	Non HDP	Р
_	n=15	n=143	n=16	n=174	n=92	HDP vs non-HDP
	n (%)*	n (%)*	n (%)*	n (%)*	n (%)*	
<10 years	4 (27)	42 (29)	9 (56)	55 (32)	41 (45)	0.036
10-15 years	3 (20)	28 (20)	2 (13)	33 (19)	15 (16)	0.591
16-20 years	0 (0)	13 (9)	0 (0)	13 (8)	1 (1)	0.013
>20 years	3 (20)	5 (4)	0 (0)	8 (5)	2 (2)	0.302
Not sure/don't know	1 (7)	48 (34)	1 (6)	52 (30)	33 (36)	0.319
I don't think I will get any of these as I maintain a healthy lifestyle	4 (27)	7 (5)	2 (13)	13 (8)	N/A	-
TOTAL	15 (101)	143 (101)	16 (101)	174 (102)	92 (100)	

^{*} Percentages may not add to 100% as figures are rounded to whole numbers only
PE = preeclampsia GH = gestational hypertension CH= chronic hypertension worsening in pregnancy and/or with superimposed preeclampsia,
HDP= hypertensive disorder of pregnancy

Supplementary Table 15: Proportion of conditions discussed by HDP sub-type when addressing future risk (multiple answers collected) within and over 3 years since last HDP (complements Table 3 which illustrates a summary of this data)

	(SH	P	E	CI	1	То	tal	Overall	P value
	n=15 N (%)					n = 16		n = 174		< 3 yrs
			N (%)		N (%)		N (%)		n = 174	vs
	<3yrs	>3yrs	<3yrs	>3yrs	<3yrs	>3yrs	<3yrs	>3yrs	N (%)	>3 yrs
HDP next pregnancy	3 (23)	1 (50)	47 (48)	21 (46)	5 (38)	2 (67)	55 (45)	24 (47)	79 (45)	0.78
Chronic Hypertension	4 (31)	1 (50)	47 (48)	16 (35)	4 (31)	2 (67)	55 (45)	19 (37)	74 (43)	0.37
No discussion	7 (54)	1 (50)	33 (34)	17 (37)	5 (38)	1 (33)	45 (37)	19 (37)	64 (37)	0.93
Lifestyle changes	4 (31)		26 (27)	6 (13)	2 (15)	2 (67)	32 (26)	8 (16)	40 (23)	0.14
Heart Attack	1 (8)	U,	20 (21)	6 (13)	1 (8)	1 (33)	22 (18)	7 (14)	29 (17)	0.50
Renal Disease	1 (8)	-	20 (21)	3 (7)	2 (15)	1 (33)	23 (19)	4 (8)	27 (16)	0.07
Stroke	1 (8)	1 (50)	18 (19)	4 (9)	1 (8)	1 (33)	20 (16)	6 (12)	26 (15)	0.45
Peripheral vascular disease	-	-	15 15)	5 (11)	1 (8)	-	16 (13)	5 (10)	21 (12)	0.56
Cannot remember	1 (8)	-	4 (4)	2 (4)	1 (8)	-	6 (5)	2 (4)	8 (5)	0.78

LONG TERM HEALTH RISKS AFTER HIGH BLOOD PRESSURE IN PREGNANCY - Survey for women

LONG TERM HEALTH AFTER BLOOD PRESSURE PROBLEMS IN PREGNANCY

You are invited to take part in a survey to gain insight into what women like yourself who have been pregnant before know about women's heart health. We are interested in the views of all women especially women who had high blood pressure (hypertension) or preeclampsia in pregnancy.

You can complete the survey if you are currently pregnant (with no major issues so far this pregnancy) or have been pregnant in the last three (3) years. You may have experienced high blood pressure in pregnancy OR you may have experienced a pregnancy without any serious complications.

The study is being conducted by the University of NSW, University of Technology Sydney and the Sydney Partnership for Health, Education, Research and Enterprise (SPHERE). The study is being undertaken by:

- Dr. Amanda Henry Obstetrician at St George and Royal Hospital for Women, Randwick, Senior Lecturer UNSW and SPHERE member
- Distinguished Professor Caroline Homer UTS, Midwifery Faculty of Health and SPHERE member
- · Dr. Clare Arnott Cardiologist, Royal Price Alfred Hospital
- Mrs. Heike Roth PhD Candidate at University of Technology, Sydney
- · Mrs. Lynne Roberts Research Midwife at St George Hospital, SESLHD.

This work is occurring as part of Mrs Heike Roth's PhD studies and the NHMRC Fellowship of Dr. Henry. Apart from salary support for Dr. Henry, the study is otherwise unfunded.

If you agree to take part in this survey, it should only take about 15 minutes to complete and will involve answering questions about you, your pregnancy and your understanding of long term health in women who have been diagnosed with blood pressure problems in pregnancy.

Participation in this study is entirely voluntary and if you do not wish to take part it will have no effect on the care you are currently receiving. If you decide to participate, and throughout the survey think you would like to withdraw/not complete the survey, you can simply stop and not submit your answers. If you have already submitted your survey, it will not be possible to withdraw the data you have provided as the surveys are anonymous.

The information you provide will not be identifiable and will be kept securely until destroyed as per the South East Sydney Local Health District's requirements.

The study results will be published in a research thesis, in peer reviewed journals and presented at conferences and other professional forums. No one will be able to identify you from this information.

If you would like to personally receive results, you will have the option to leave your email details. The results will be available one (1) year after conclusion of the survey and your email will not be used for any other purpose.

If you would like any further information about the study or you experience any distress or concern as a result of completing this survey, please contact the Principal Investigator, Dr Amanda Henry on 02 91132315 or via email Amanda.henry1@health.nsw.gov.au. For medical assistance you can consult your General Practitioner. If you would like further information about the topic addressed in this study, you can visit the Australian Heart Foundation on the following link: https://www.heartfoundation.org.au/your-heart/women-and-heart-disease/womens-stories.

If you have any concerns or complaints about the conduct of this study, you should contact the Research Support Office of the South Eastern Sydney Local Health District Human Research Ethics Committee which is nominated to receive complaints from research participants. You should contact them on 02 9382 3587, or email SESLHD-RSO@health.nsw.gov.au and quote HREC 18/156.

* 1. I	acknowledge	that partic	cipation in	the sur	vey is '	voluntary
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Yes

OUT YOU	
	out about you, your background, and occupation. isk (*) simply mean that they must be answered in order to
What age group are you in?	
18-25	46-55
26-35	56+
36-45	Prefer not to answer
What ethnic group do you identify	with? (Please select one answer)
Caucasian	Aboriginal or Torres Strait Islander
Asian	European
Polynesian or Maori	Prefer not to answer
Other (please specify)	
-	
. What is your highest level of forma	al education?
What is your highest level of formation Secondary school	al education? University degree
Secondary school	University degree
Secondary school Trade Certificate/Diploma	University degree
Secondary school Trade Certificate/Diploma	University degree
Secondary school Trade Certificate/Diploma	University degree Prefer not to answer
Secondary school Trade Certificate/Diploma Other (please specify)	University degree Prefer not to answer
Secondary school Trade Certificate/Diploma Other (please specify)	University degree Prefer not to answer
Secondary school Trade Certificate/Diploma Other (please specify) What is your usual occupation/pro	University degree Prefer not to answer fession?
Secondary school Trade Certificate/Diploma Other (please specify)	University degree Prefer not to answer fession?

P4 Newsletter	Maternity Consumer Network	
Australian Action on Preeclampsia (AAPEC)	Maternity Clinics	
Tresillian	Social media (Facebook, Twitter)	
Maternity Choices Australia	Australian College of Midwives	
Other (please specify)		

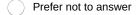
ABOUT YOUR HEALTH

These next questions are about your general health.

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue.

* 8. A	re you	currently	/ pregna	ınt?
--------	--------	-----------	----------	------

Yes



() No

	women
ABOUT YOUR HEALTH	
Questions marked with a green asterisk continue.	c (*) simply mean that they must be answered in order to
9. How many weeks pregnant are you?	(provide whole numbers only, for example: 24)
10. How many children have you given	birth to (20 weeks gestation and over)?
* 11. Have you ever had any of the follow apply)	ving, whilst pregnant or before or after pregnancy? (select all that
High blood pressure	Stroke
High BMI (overweight)	Significant illness
Angina	Heart attack
Diabetes	None of the above/ no significant other medical complication
Kidney problems	
Other (please specify)	
* 12. From the list below, which currently	
Smoking	High cholesterol
Obesity	High blood pressure
Alcohol consumption	Diabetes
Family history of heart disease	None of the above
Other (please specify)	
12 Diago provide the details of any ar	accribed medications you are taking
13. Please provide the details of any properties.I do not take any prescribed medication	escribed medications you are taking
	e medications or leave blank if you prefer not to answer)
Trave hieserinen menicanon (hiease list file	inedications of leave blank if you prefer hot to answer)



PREGNANCY

This section is about your pregnancy history.

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue.

DEFINITIONS OF BLOOD PRESSURE PROBLEMS

Here are some definitions of certain types of blood pressure problems in pregnancy. You may find these useful in order to more easily understand and answer the next questions.

Chronic hypertension: is if you had high blood pressure before falling pregnant, have high blood pressure outside of pregnancy, or were found to already have high blood pressure in the first half of your pregnancy. Chronic hypertension may have no known underlying cause (this is sometimes called "essential" hypertension), or it may be as a result of another underlying condition, such as kidney disease.

Gestational hypertension: is when you might have had high blood pressure for the first time in your pregnancy (after 20 weeks of pregnancy) but were otherwise well (that is, high blood pressure only but no effect on your baby's growth or on your health otherwise).

Preeclampsia: is when you have had high blood pressure in pregnancy (after 20 weeks of pregnancy) and some additional signs or issues in you and/or your baby. For example, you might have had protein in your urine, liver or kidney problems that showed up on blood tests, or there may have been concerns about the growth of your baby while you were pregnant.



PREGNANCY

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue.

We understand that you may find some of the questions difficult to answer as they might remind you of a challenging time in your life. We are grateful for your participation and contribution to improving knowledge on future health for women who had blood pressure problems in pregnancy. You will find some explanations at the end of the survey and a contact, in case you would like to seek further clarification and/or assistance.

* 14.	Choose	the situation	n which b	est descri	bes vour	pregnancy	history

At least one pregnancy is/was affected by gestational
hypertension

- At least one pregnancy is/was affected by preeclampsia
- I have only been diagnosed with chronic hypertension
- I had chronic hypertension before pregnancy and had/have pregnancies that were complicated further by higher than usual blood pressure
- I had chronic hypertension before pregnancy and had/have pregnancies that were complicated further by preeclampsia
- No pregnancy is/was affected



YOUR HEALTH DURING YOUR PREGNANCY

Questions marked with a green asterisk (*) simply	\prime mean that they must be answered in	order to
continue.		

* 15. As someone who has chronic hypertension are y at risk of? (select all that apply)	you aware of any long term health issues that you are
Diabetes	Leukaemia
Kidney disease	Seizures
Breast cancer	Overall mortality risk is higher
Cardiac death	Ischaemic heart disease/heart attack
High blood pressure complications in another pregnancy	I think there are health risks but unsure which conditions I
Stroke	may be at risk of I do not think that there are increased risks
Peripheral vascular disease	T do not think that there are increased lisks
Other (please specify)	



< 10 years after pregnancy		> 20 years a	fter pregnancy	
10-15 years after pregnanc		Not sure/do	not know	
16-20 years after pregnancy Other (please specify)	<i>(</i>			

LONG TERM HEALTH RISKS A	AFTER HIGH BLOOD PRES women	SSURE IN PREGNANCY - Surve	ey for
Chronic Hypertension only -TYF	PE OF CARE RECEIVED D	URING PREGNANCY	
Questions marked with a green a continue.	sterisk (*) simply mean that t	hey must be answered in order to	
		nealth risks, when did this occur?	
Before birth	6 mor	nths to 1 year	
Immediately after birth	1 year	r and over	
Within first 6 weeks	I cann	not remember	
6 weeks to 6 months			
hypertension or preeclampsia ex	xperience?	ng term health risks in your gestation	al
During pregnancy/at birth	12 months after birth	24 months after birth	
* 19. As a result of your pregnancy below after your baby was born? Cardiologist	(tick all that apply)	oblems, were you referred to any of t	:he
Renal (kidney) Physician	Nutriti	ionist for dietary adjustment	
General Practitioner	I cann	not remember	
Other (please specify)			

-	agnosed wit								
	Gestatio Hyperten		Preeclamp	pro	blood press blem diagno his pregnan	osed	not rememb	er Not a _l	pplicable
First Pregnancy								(
Second Pregnancy								(
Third Pregnancy								(
ny comments?									
0-6 months 6-12 months 1-2 years 2. At what point in til	me were yo	u diagno	sed? (Ch	n	-3 years nore than 3 y ost accura				
6-12 months				oose a m	ost accura	ate time f During or after	No diagnosis of blood pressure problem		
6-12 months 1-2 years	me were yo 20-28 weeks	u diagno 28-34 weeks	sed? (Cha 34-37 weeks	n	ore than 3 y	ate time f	No diagnosis of blood pressure	I cannot remember	
6-12 months 1-2 years 2. At what point in til	20-28	28-34	34-37	oose a m	ost accura	ate time for the state of the s	No diagnosis of blood pressure problem this		
6-12 months 1-2 years 2. At what point in till First Pregnancy	20-28	28-34	34-37	oose a m	ost accura	ate time for the state of the s	No diagnosis of blood pressure problem this		
6-12 months 1-2 years	20-28	28-34	34-37	oose a m	ost accura	ate time for the state of the s	No diagnosis of blood pressure problem this		
6-12 months 1-2 years 2. At what point in till First Pregnancy Second Pregnancy	20-28	28-34	34-37	oose a m	ost accura	ate time for the state of the s	No diagnosis of blood pressure problem this		Not applica

Problems? Yes, planned induction of alabour because of blood pressure issues in pregnancy Second Pregnancy Any Comments? Yes, planned induction of a blood pressure issues of blood pressure issues in pregnancy Third Pregnancy Any Comments?	PREGNANCY Questions marked with ontinue. * 23. Did you have a place.	n a green astei	wor risk (*) simply	men mean that th	ey must be ans	swered in or	der to
Second Pregnancy	problems?	induction of labour because of blood pressure issues	caesarean section because of blood	caesarean for other reasons than blood	induction and no planned caesarean		Not applicabl
Third Pregnancy O O O O O O O O O O O O O O O O O O O	First Pregnancy						
Any Comments?	Second Pregnancy						
	Third Pregnancy						

LONG TERM HEALTH RISKS

This section is about some long term health risks that some women may experience after having had blood pressure problems in pregnancy.

For this section we would like you to think about the long-term health risks of a woman who has been diagnosed with high blood pressure in pregnancy.

Not everyone who experienced blood pressure problems in pregnancy will necessarily have health issues in the future. We would not want you to unnecessarily worry about any of these risks, therefore we will provide you with further information about long-term health after high blood pressure in pregnancy at the end of the survey.

* 24. FOR WOMEN WITH HISTORY OF BLOOD PRESSURE PROBLEMS IN PREGNANCY:

Compare yourself to a woman who has NOT had blood pressure problems in pregnancy.

Do you think you have a lower/same/higher chance of getting the following:

	Less chance than a woman without blood pressure in pregnancy	Same chance as a woman without blood pressure in pregnancy	Higher chance than a woman without blood pressure in pregnancy	Not sure/I do not know
High blood pressure later in life	\circ	\circ		0
Diabetes				
Kidney disease				
Breast cancer				
Heart attack				
High blood pressure in another pregnancy				
Stroke				
Heart disease				
Seizures				
Vascular Disease				

If you are concerned by any of the above potentially affecting you, information is available at the end of the survey where risks are explained.

For this section we would like you to think about the long term health risks of a woman who has been diagnosed with high blood pressure in pregnancy.

Same chance as a

Higher chance than a

* 25. FOR WOMEN WITHOUT HISTORY OF BLOOD PRESSURE PROBLEMS IN PREGNANCY:

Compare yourself to a woman who HAS had blood pressure problems in pregnancy.

Do you think you have a lower/same/higher chance of getting the following:

Less chance than a

	woman with blood pressure in pregnancy	woman with blood pressure in pregnancy	woman with blood pressure in pregnancy	Not sure/I do not know
High blood pressure later in life				
Diabetes				
Kidney disease				
Breast cancer				
Heart attack				
High blood pressure in another pregnancy				\bigcirc
Stroke				
Heart disease				
Seizures				
Vascular Disease				
26. How many years a symptoms of the pote < 10 years after pregr	ntial risks may start to	appear?	ears after pregnancy	ous signs and
10-15 years after preg		Not sur	re/do not know	
16-20 years after preg				
Other (please specify))			

If you are concerned by any of the above potentially affecting you, information is available at the end of the survey where risks are explained.



	women
* 27. How many years after blood pressure pr symptoms of the potential risks may start to	roblems in pregnancy do you think the various signs and appear?
< 10 years after pregnancy	> 20 years after pregnancy
10-15 years after pregnancy	Not sure/do not know
16-20 years after pregnancy	I don't think I will get any of these as I maintain a healthy lifestyle
Other (please specify)	

IN CASE OF DISTRESS

If you experience any distress caused due to the completion of this survey, please contact your GP or the Principal Investigator of this study, Dr Amanda Henry on 02 91132315 or Amanda.henry1@health.nsw.gov.au

For more information on this topic please visit The Australian Heart Foundation on the following link: https://www.heartfoundation.org.au/your-heart/women-and-heart-disease/womens-stories



TYPE OF CARE RECEIVED DURING PREGNANCY WHERE A BLOOD PRESSURE PROBLEM WAS DIAGNOSED

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue.

It is quite likely that some of the following questions may bring back some memories or bring rise to emotions that you find difficult to deal with. Please contact the Principal Investigator, Dr Amanda Henry on 02 91132315 or via email Amanda.henry1@health.nsw.gov.au if you would like to discuss these concerns. For medical assistance you can consult your General Practitioner.

blood pressure problem in pregnancy?	ency Unit of intensive Care Unit as a result of your
Yes	
○ No	
I am not sure	
I cannot remember	
* 29. Have any of your babies been admitted to 'Neona Care Nursery' as a result of your blood pressure prol Yes No I am not sure	atal Intensive Care', 'High Dependancy Unit' or 'Special blem in pregnancy?
30. After your baby was born have you had any of th	e following? (select all that apply)
Blood pressure measurement in hospital	Consultation with a renal (kidney) specialist
Blood pressure measurement with my GP	Consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with an obstetric medicine specialist (doctor when the consultation with a consultati
Consultation with an obstetrician	specialises in complications of pregnancy like high blood pressure)
	I cannot remember
Other (please specify)	

Increased risk of high blood pressure	e Risk of hypertensive disease in your next pregnancy
Increased risk of kidney problems	I was told to eat a healthy diet, do some exercise and I
Increased risk of stroke	normally
Increased risk of heart attack	No risks were discussed
Increased risk of vascular disease	I cannot remember
Other (please specify)	
O.	

TYPE OF CARE RECEIVED DURING PREGNANCY WHEN A BLOOD PRESSURE PROBLEM WAS DIAGNOSED

If a healthcare provider did s	peak to you about your future	health risks, when did this occur?
Before birth	6 mo	nths to 1 year
Immediately after birth	1 yea	ar and over
Within first 6 weeks	I can	not remember
6 weeks to 6 months		
During pregnancy/at birth	12 months after birth	24 months after birth
	y affected by blood pressure p	roblems, were you referred to any of th
As a result of your pregnancy ow after your baby was born? Cardiologist		ss centre for exercise
ow after your baby was born?	Fitne	
ow after your baby was born?	Fitne	ss centre for exercise
ow after your baby was born? Cardiologist Renal (kidney) Physician	Fitne	ss centre for exercise tionist for dietary adjustment
ow after your baby was born? Cardiologist Renal (kidney) Physician General Practitioner	Fitne	ss centre for exercise tionist for dietary adjustment

EDUCATION

In this section we would like to find out about your preferred ways of getting information about long term health after gestational hypertension and preeclampsia.

Questions marked with a green asterisk (*) simply mean that they must be answered in order to continue.

Olitili	uc.
	After experiencing gestational hypertension or preeclampsia what do you want to know about your long n health? (select all that apply)
	Risk reduction for subsequent pregnancies
	Reducing risk behaviours (eg. diet, exercise, smoking cessation)
	Statistics (eg. increased risk)
	At what point does the risk increase
	Signs and Symptoms
	Where to find information
	How to discuss the matter with my healthcare provider
	Impact on my children from the pregnancy affected by blood pressure problems
	Other (please specify)
* 36.	How do you want to receive the information? (select all that apply).
	Key organisations addressing heart health (e.g. The Australian Heart Foundation or Her Heart)
	Social Media channels (e.g. Instagram, Facebook, Twitter)
	Brochures/Flyers
	Medical Professionals
	Podcasts/Media
	Online videos
	Other (please specify)



POST-SURVEY OPTIONS AND SUMMARY OF RISK PROFILE

Some further things we would like you to consider:

38. We would like to have your opinion on what you think would be appropriate education material to improve women's knowledge. We would also like to know how the education could best be distributed to women. Would you like to participate in a **focus group** (one off - 2hrs max at St George Hospital, Sydney) or a **telephone interview** (one off 30 mins max)? By involving 'consumers', meaning women like you, the education package will have added value.

If you are interested in participate in either, you can leave your details here.

Name	
Email Address	
Phone Number	

39. Please leave your email address to **receive results** from this study (in approx. 1 year). Your details will not be used for any other purpose.

IN CASE OF DISTRESS

If you experience any distress caused due to the completion of this survey, please contact your GP or the Principal Investigator of this study, Dr Amanda Henry on 02 91132315 or Amanda.henry1@health.nsw.gov.au

For more information on this topic please visit The Australian Heart Foundation on the following link: https://www.heartfoundation.org.au/your-heart/women-and-heart-disease/womens-stories



RISK PROFILE - LONG TERM RISKS AFTER BLOOD PRESSURE PROBLEMS IN PREGNANCY EXPLAINED

You may like to take a screenshot of the risk profile so you can refer back to it whenever you need to.

RISK PROFILE

Although most women will experience good long-term health after having high blood pressure in pregnancy, there are, unfortunately, some long term health risks associated with having had high blood pressure in pregnancy.

Women who have had high blood pressure during pregnancy are about 3 to 4 times more likely to develop chronic hypertension than women who did not have a blood pressure problems in pregnancy. They are also about twice as likely to get diabetes in later life, even if they did not have diabetes during pregnancy.

Blood pressure diseases are also more likely to happen in the next pregnancy to women who have already had a previous blood pressure problems in pregnancy compared to women who have not. Therefore, if they have had a pregnancy with blood pressure problems, it is important to be seen early in their next pregnancy. There are treatments that can decrease the chance of recurring problems.

Women are also more likely to get various forms of cardiovascular disease (heart disease, stroke, vascular disease) if they have had gestational hypertension, preeclampsia and/or chronic hypertension. All of these cardiovascular problems are about twice as likely to eventually happen to a woman who has had blood pressure problems in pregnancy compared to a woman who has not. This still means most women will not have heart disease or diabetes after having high blood pressure in pregnancy, especially if they can avoid risk factors like smoking or excessive weight gain, and maintain a healthy diet and exercise.

Kidney problems are about 5 to 10 times more common after preeclampsia in particular. Although the relative risk of developing kidney problems is substantially higher after preeclampsia, the absolute risk of long-term kidney disease is still low. Unless the woman already had a kidney problem, well over 90% of women after preeclampsia and gestational hypertension will not have a kidney problem.

Fortunately, although seizures may occur as a result of preeclampsia during pregnancy, women have no higher long term risk of seizures compared to women who did not have a complicated pregnancy. There is no increased risk of getting cancer (e.g. breast cancer, leukaemia) after having high blood pressure in pregnancy.

For all the long term health risks, these start to go up within 10 years after an affected pregnancy and are ongoing after that. Therefore, it is recommended that women attend regular blood pressure checks with their GP and discuss any changes they can make to improve their general health. For more general information about heart health and managing health risks, please visit the National Heart Foundation website: https://www.heartfoundation.org.au/your-heart/know-your-risks

THANK YOU FOR YOUR PARTICIPATION

STROBE Statement—

Assessing Australian women's knowledge and knowledge preferences about long-term health after hypertensive disorders of pregnancy: a survey study

	Item No	Recommendation	Page No
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title	1
		or the abstract	
		(b) Provide in the abstract an informative and balanced summary of	2
		what was done and what was found	
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation	3-5
		being reported	
Objectives	3	State specific objectives, including any prespecified hypotheses	2
Methods			
Study design	4	Present key elements of study design early in the paper	2&
			5-8
Setting	5	Describe the setting, locations, and relevant dates, including periods of	2&6
		recruitment, exposure, follow-up, and data collection	
Participants	6	(a) Cohort study—Give the eligibility criteria, and the sources and	5-6
		methods of selection of participants. Describe methods of follow-up	
		Case-control study—Give the eligibility criteria, and the sources and	
		methods of case ascertainment and control selection. Give the rationale	
		for the choice of cases and controls	
		Cross-sectional study—Give the eligibility criteria, and the sources and	
		methods of selection of participants	
		(b) Cohort study—For matched studies, give matching criteria and	-
		number of exposed and unexposed	
		Case-control study—For matched studies, give matching criteria and	
		the number of controls per case	
Variables	7	Clearly define all outcomes, exposures, predictors, potential	-
		confounders, and effect modifiers. Give diagnostic criteria, if applicable	
Data sources/	8*	For each variable of interest, give sources of data and details of	-
measurement		methods of assessment (measurement). Describe comparability of	
		assessment methods if there is more than one group	
Bias	9	Describe any efforts to address potential sources of bias	2& 21 22
Study size	10	Explain how the study size was arrived at	5-6
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If	&Fig1 7-8
		applicable, describe which groupings were chosen and why	
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	7-8
		(b) Describe any methods used to examine subgroups and interactions	
		(c) Explain how missing data were addressed	
		(d) Cohort study—If applicable, explain how loss to follow-up was	
		addressed	

Case-control study—If applicable, explain how matching of cases and	
controls was addressed	
Cross-sectional study—If applicable, describe analytical methods	
taking account of sampling strategy	
(e) Describe any sensitivity analyses	

Continued on next page

Results			
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers potentially	5&6
		eligible, examined for eligibility, confirmed eligible, included in the study,	
		completing follow-up, and analysed	
		(b) Give reasons for non-participation at each stage	
		(c) Consider use of a flow diagram	Fig 1
Descriptive	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and	8-9
data		information on exposures and potential confounders	Tab1
		(b) Indicate number of participants with missing data for each variable of interest	8
		(c) Cohort study—Summarise follow-up time (eg, average and total amount)	
Outcome data	15*	Cohort study—Report numbers of outcome events or summary measures over time	
		Case-control study—Report numbers in each exposure category, or summary measures of exposure	
		Cross-sectional study—Report numbers of outcome events or summary measures	
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates	
		and their precision (eg, 95% confidence interval). Make clear which confounders	
		were adjusted for and why they were included	
		(b) Report category boundaries when continuous variables were categorized	
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a	
		meaningful time period	
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and	
		sensitivity analyses	
Discussion			
Key results	18	Summarise key results with reference to study objectives	8-14
	Tables		
			2-4
Limitations	ons 19 Discuss limitations of the study, taking into account sources of potential b		20-22
		imprecision. Discuss both direction and magnitude of any potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations,	14-19
		multiplicity of analyses, results from similar studies, and other relevant evidence	
Generalisability	21	Discuss the generalisability (external validity) of the study results	22
Other informati	on		
Funding	22	Give the source of funding and the role of the funders for the present study and, if	24
		applicable, for the original study on which the present article is based	

^{*}Give information separately for cases and controls in case-control studies and, if applicable, for exposed and unexposed groups in cohort and cross-sectional studies.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.