

	ANN	N	
Survey number (partners):			





# SURVEY FOR PARTNERS

## Please read the following before completing the survey:

#### STUDY DESCRIPTION

NIPT (Non-Invasive Prenatal Testing) is a new technology that tests the risk for Down syndrome. The purpose of this survey is to learn what pregnant women and their partners know about NIPT and to learn about their views regarding its use. This survey is part of a larger study on NIPT, called PEGASUS, see: <a href="http://pegasus-pegase.ca/">http://pegasus-pegase.ca/</a>.

#### **BENEFITS AND RISKS**

Participating in the study will not provide you with any immediate or specific benefit, but it will allow you to contribute to the advancement of knowledge about NIPT and this may help other women who use this test in the future. The survey does not pose any risk and its results will be kept confidential. You are free to withdraw from this study at any time.

#### **CONSENT**

By completing and returning this survey, you consent to participate in this part of the PEGASUS study and authorize Dr. Vardit Ravitsky and her colleagues to analyze the content of the completed survey. Completing this survey can take about 20 minutes.

#### CONFIDENTIALITY

This survey is coded and you do not have to sign your name on it. All information obtained in connection with this survey will be kept confidential. Access to this survey will be restricted to the members of the research team, for the duration of the study. The surveys will be kept in a secure place, under lock and key, for a maximum of 10 years after the project ends. The results of the study may be published, but no identifiable information will ever be disclosed.

#### **CONTACT PERSONS**

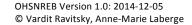
For further information regarding this project, you are welcome at any time to contact Dr. Vardit Ravitsky at (514) 343-6111 extension 3375 or at <a href="mailto:vardit.ravitsky@umontreal.ca">vardit.ravitsky@umontreal.ca</a>.

## **INSTRUCTIONS**

Please answer directly on the survey. If you change your mind, cross out your first mark. When you are finished, please seal it in the attached envelope and hand it in or return it in the pre-addressed envelope.

If you prefer to complete this survey online, you can find it at: <a href="http://nipt.hostedincanadasurveys.ca/s1">http://nipt.hostedincanadasurveys.ca/s1</a>

We thank you for participating.









## PART 1: WHAT DO YOU KNOW ABOUT DOWN SYNDROME (DS)?

1.	Before today, have you ever heard of:
	(PLEASE CHECK ONE ANSWER FOR EACH STATEMENT

(PLEASE CHECK ONE ANSWER FOR EACH STATEMENT)						
		Yes	No			
a.	Down syndrome					
b.	Trisomy 21					
c.	Prenatal screening for Down syndrome					
d.	Noninvasive prenatal testing (NIPT)					
e.	Amniocentesis					

2. The next question is about Down syndrome (DS). Do you think these statements are true or false?

(PLEASE CHECK ONE ANSWER FOR EACH STATEMENT)

		True	False
a.	Other people can tell when a child has DS		
b.	All individuals with DS have some kind of intellectual disability		
c.	People with DS can live until at least their 50's-60's		
d.	Intellectual disability is the only health issue related to DS		
e.	DS can be caused by a woman's unhealthy lifestyle		
f.	A woman has a risk of having a baby with DS <b>only if</b> somebody in her family has DS		

3. The next question is about the different options available to pregnant women who want to know more about the risk of Down syndrome (DS) for their pregnancy. Do you think these statements are true or false?

(PLEASE CHECK ONE ANSWER FOR EACH STATEMENT)

		True	False
a.	All babies with DS show signs of DS on ultrasound		
b.	Current screening can be used to diagnose with certainty a baby with DS		
c.	Amniocentesis can be used to diagnose with certainty a baby with DS		
d.	NIPT can be used to diagnose with certainty a baby with DS		
e.	Amniocentesis can predict the severity of the symptoms of DS		
f.	There is an increased risk of miscarriage (losing the pregnancy) with amniocentesis		
g.	There is no increased risk of miscarriage (losing the pregnancy) with NIPT		

## **INFORMATIONAL SHEET**

Down syndrome (DS) is a genetic condition caused by the presence of an extra chromosome 21 (also called 'trisomy 21') which affects 1 in 770 newborns. Individuals with DS usually share physical features that are characteristic of DS. All have some degree of intellectual disability, which varies from person to person; their development is slower than other kids, but they will eventually learn to walk, talk, and dress themselves. Most children attend their neighborhood schools, some in regular classes and others in special education classes. Some children have more significant needs and require a more specialized program. Many adults with DS are capable of working in the community, but some require a more structured environment. Many will also have other health problems (for example heart defects). 99% of cases of DS are not inherited from the parents; it usually occurs by chance.

There are ways to check during pregnancy if there is a possibility that the baby has DS:

	MATERNAL SERUM SCREENING (MSS or 'current screening')	AMNIOCENTESIS	NIPT	
Description of the procedure	<ul> <li>Checks the level of risk for DS</li> <li>Measures the level of hormones produced by the baby or placenta that end up in the mother's blood</li> <li>Includes one or two blood draws from the mother</li> <li>Where available, an ultrasound is done early in the pregnancy to measure nuchal translucency (level of fluid at the nape of the baby's neck)</li> </ul>	<ul> <li>Medical procedure that can confirm DS during the pregnancy</li> <li>Allows checking the number and appropriate structure of all chromosomes in the baby's cells</li> <li>Requires inserting a thin needle into the uterus – through the mother's abdomen - to extract amniotic fluid (fluid in which the baby floats in the mother's womb)</li> </ul>	<ul> <li>Checks the level of risk for DS</li> <li>Analyses the baby's DNA that is floating in the mother's blood</li> <li>Includes one blood draw from the mother</li> </ul>	
Timing: When in pregnancy	<ul> <li>1<sup>st</sup> blood draw: usually between the 10<sup>th</sup> and 13<sup>th</sup> week of pregnancy</li> <li>2<sup>nd</sup> blood draw: usually between the 15<sup>h</sup> and 16<sup>th</sup> week of pregnancy</li> <li>Results can be available between the 16<sup>th</sup> and 17<sup>th</sup> week of pregnancy</li> </ul>	<ul> <li>Available from the 15<sup>th</sup> week of pregnancy</li> <li>Results can be available between the 17<sup>th</sup> and 19<sup>th</sup> week of pregnancy</li> </ul>	<ul> <li>Available as of the 10<sup>th</sup> week of pregnancy</li> <li>Results can be available between the 11<sup>th</sup>-12<sup>th</sup> week of pregnancy</li> </ul>	
Risk to pregnancy	No increased risk of miscarriage	Risk of miscarriage around 1 in 200 (0.5%)	No increased risk of miscarriage	
Accuracy	Detects between 77% and 88% of DS cases (supported by a lot of evidence)	100% accurate in detecting DS (supported by a lot of evidence)	98% accurate (or more) for DS in women who are considered "high risk based on MSS (supported by some evidence)	
Type of test	Screening	Diagnostic	Screening	
What it detects	<ul> <li>Down syndrome</li> <li>Trisomy 18</li> <li>Neural tube defects (e.g. spina bifida)</li> <li>Possible pregnancy complications</li> </ul>	<ul> <li>Down syndrome</li> <li>Trisomy 13</li> <li>Trisomy 18</li> <li>Other chromosome anomalies</li> <li>Neural tube defects (e.g. spina bifida)</li> <li>Sex of the baby</li> </ul>	<ul> <li>Down syndrome</li> <li>Trisomy 13</li> <li>Trisomy 18</li> <li>Sex of the baby</li> </ul>	
Possible outcome	The test can predict that the pregnancy is at:     → Low risk for DS (less than 1/200 – 1/300) so no further test is recommended     → High risk for DS (higher than 1/200 – 1/300)     • Amniocentesis is offered to check if the baby actually has DS or other abnormalities detectable by chromosome analysis.	<ul> <li>Normal result: the baby does not have DS and has normal chromosomes.</li> <li>Abnormal result: the baby has DS or has another significant chromosome abnormality. In this case, the parents can choose to:         <ul> <li>→ continue the pregnancy</li> <li>→ stop the pregnancy</li> </ul> </li> </ul>	<ul> <li>The test can predict that the pregnancy is at:</li> <li>→ Very low risk for DS - so no further test is recommended</li> <li>→ Very high risk for DS</li> <li>• Amniocentesis is recommended to confirm that the baby has DS</li> </ul>	

Note: NIPT is not yet considered as a first-tier screening test (i.e. that could replace serum screening); the current available evidence supports its use as a second tier screening test – after a positive serum screening and before an amniocentesis.







4.	Information about a baby having Down syndrome (DS) interested in knowing whether your baby has DS? (PLEASE CHECK ONE ANSWER)	) can have differe	ent uses for diffe	rent people. Wh				
	☐ I want to know in advance to prepare for the birth of a baby with DS if baby is diagnosed with DS ☐ I would consider terminating the pregnancy if the baby was diagnosed with DS							
	☐ I'm unsure							
☐ I don't want to know but my partner does								
	Other:							
<u>PAI</u> 5.	RT 2: INFORMED CONSENT  When would be the best time for you to receive the for (PLEASE CHECK ONE ANSWER PER STATEMENT)	ollowing informa	tion:					
	,	Early prenatal appointment ahead of time of NIPT	Same day as blood test for NIPT	When your partner gets NIPT results	Not interested in this information			
a.	What is Down syndrome (DS)							
b.	How good is NIPT in detecting DS							
c.	What the test can and cannot tell							
d.	What is the chance that you can have a baby with DS (according to family history, your partner's age, etc.)							
e.	What are the possible results (high vs. low risk for DS)							
f.	What are the options if the result is positive							
g.	What are the available resources for families who have children with DS							
Oth	er:							
6.	Do you think it is important to get written consent for (PLEASE CHECK ONE ANSWER ONLY)	NIPT considering	there is no risk	of miscarriage?				
	☐ Yes ☐ No ☐ I don't k	now $\square$	Other:					







7.	Please rank in order of importar (1= YOUR FIRST CHOICE, 6 = YOUR LA		would like to d	iscuss NIPT with:		
	Family physician					
[	Obstetrician/Gynecologist					
[	Genetics specialist					
[	Nurse					
[	Midwife					
[	Other:					
8.	How would you like to be inforr	med about NIPT?				
	☐ Brochures ☐	☐ Video	☐ Discu	ssion with prenatal	caregiver	
	☐ Website ☐	☐ Group meetings	☐ Other	r:		
9.	How much time would you wan  (PLEASE CHECK ONE ANSWER ONLY)  I would make a decision d			eartner is offered th	e test.	
	☐ I would want a few days t	o think about it.				
	☐ Not sure.					
10.	What would be the most appro	•	o receive the re	esults of NIPT?		
		In person	By phone	By mail, fax or email	Through a secured website	I don't care
a.	If NIPT result is LOW risk of DS					
b.	If NIPT result is HIGH risk of DS					
Oth	er:					







## **PART 3: HOW SHOULD NIPT BE USED?**

- **11.** George and Melissa are expecting a baby. Melissa is 40 years old and is 10 weeks pregnant. They are meeting her doctor for her first prenatal visit. During their visit, the doctor explains that:
  - Current screening can assess the risk that the baby might have Down syndrome (DS). It can also tell her the risk of neural tube defects (such as spina bifida) and give her information about the risk of complications during pregnancy and labor. However, current screening detects only about 80% of cases of DS and Melissa will only get her results after the 15<sup>th</sup> week of pregnancy.
  - NIPT is available earlier in the pregnancy (around 10 weeks) and will detect 99% of cases of DS. NIPT can also predict the risk that the baby might have a sex chromosome disorder (where the baby has an extra or missing sex chromosome). However, it cannot tell whether the baby might have a neural tube defect or about the risk of pregnancy and labour complications.
  - 11.1. If you were George and you had to choose between current screening and NIPT as a first step, which test would you choose for the pregnancy?
    (PLEASE CHECK ONE ANSWER ONLY)

☐ Current screening
□ NIPT
$\square$ I would not want any testing (please skip the next question and go to question 12)

**11.2.** How did the following reasons influence your decision?

		Does not influence		Somewhat influences		Strongly influences
a.	Current screening estimates the risk that the baby has neural tube defects and NIPT doesn't	1	2	3	4	5
b.	Current screening can indicate the possibility of pregnancy and labor complications and NIPT cannot	1	2	3	4	5
c.	NIPT is much more accurate than current screening in assessing the risk of DS	1	2	3	4	5
d.	Results of NIPT can be available earlier in the pregnancy than the result of current screening	1	2	3	4	5

Other:







**12.** Robert and Carolyn are expecting a baby. Carolyn is 40 years old and is 16 weeks pregnant. They are meeting her doctor for a prenatal visit. The pregnancy is at high risk for Down syndrome (meaning that there is more than 1 in 300 chance that her baby has DS) based on current screening.

They are referred to a genetic specialist, who explains that there are further tests available:

- Amniocentesis is an invasive procedure that is available from the 15<sup>th</sup> week of pregnancy. Because a needle would be inserted into Carolyn's womb, there is an increased risk of miscarriage due to the procedure (0.25 to 0.5%). This test is practically 100% accurate, and makes it possible to check the number of all of the baby's chromosomes.
- NIPT is done earlier in the pregnancy (about 10 weeks). Because it only requires a blood draw, there is no increased risk of miscarriage. However, NIPT cannot check all of the baby's chromosomes. NIPT is a very accurate test (will detect about 99% of cases of DS), but is not as accurate as amniocentesis.
- 12.1. If you were Robert and you had to choose between amniocentesis and NIPT as a diagnostic test after current screening, which test would you choose for the pregnancy?
  (PLEASE CHECK ONE ANSWER ONLY)
  Amniocentesis
  NIPT
  I would not want any testing (please skip the next question and go to question 13)
- **12.2.** How did the following reasons influence your decision?

		Does not influence		Somewhat influences		Strongly influences
a.	Amniocentesis is more accurate than NIPT	1	2	3	4	5
b.	Amniocentesis gives more information about possible chromosome anomalies than NIPT	1	2	3	4	5
c.	With NIPT there is no increased risk of miscarriage	1	2	3	4	5
d.	NIPT tests for the common chromosome disorders (like DS), which is all I need to know	1	2	3	4	5
e.	NIPT is more convenient than amniocentesis. (only requires a blood draw)	1	2	3	4	5

Other:	







13.	Right now, NIPT is not willing to pay for NIPT (PLEASE CHECK ONE ANSW	?	alth care syster	n in most provinces	and territories. Ho	w much would you be
	<b>□</b> 0\$	□ 100\$ to 4	.99\$ <b></b>	<b>1</b> 1000\$ to 4999\$		
	☐ 1\$ to 99\$	☐ 500\$ to 9	99\$	More than 5000\$		
14.	Who do you think sho		NIPT free of ch	arge?		
	☐ All women	Low	risk women on	ly	Other:	
	☐ High risk wome	n only 🔲 Nobo	ody (you should	d pay for the test)		
15.	If NIPT were free of ch (PLEASE CIRCLE ONE ANSW	/ER)	Some impact	u iiripact your decisi	A lot of impact	. <b>:</b> 
	No impact	3	onic impact			!
	No impact	2	3	4	5	
<u>PAR</u> 16.	1 RT 4: DECISION MAK	2  NG AND THE INV	3  OLVEMENT Country of the following the following states and the following states are also as a few sections and the following states are also as a few sections are also as a few sections and the few sections are also as a few sections	F OTHERS	5	for the pregnancy:
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	1  RT 4: DECISION MAKI  Please rank in order o (1 = THE MOST IMPORTA  Partner  Family  Friends and co	2 ING AND THE INV f importance the in NT; 5/6 = THE LEAST II	3  OLVEMENT C  put of the follo	F OTHERS	5	for the pregnancy:
	1  RT 4: DECISION MAKI  Please rank in order o  (1 = THE MOST IMPORTA  Partner  Family  Friends and co  Your own pren	2 ING AND THE INV f importance the in NT; 5/6 = THE LEAST II	3  OLVEMENT C  put of the follo  MPORTANT)	F OTHERS	5	for the pregnancy:







17. How much input do you feel a partner **should have** on a woman's decision about one of these following tests? (PLEASE CIRCLE ONE ANSWER FOR EACH STATEMENT)

		No input		Some input		A lot of input
a.	Current screening	1	2	3	4	5
b.	Amniocentesis	1	2	3	4	5
c.	NIPT	1	2	3	4	5

**18.** Do you feel that your opinion is **actually** taken into account by your **partner** when deciding about one of the following tests?

(PLEASE CIRCLE ONE ANSWER FOR EACH STATEMENT)

		Not at all		Somewhat		Very much
a.	Current screening	1	2	3	4	5
b.	Amniocentesis	1	2	3	4	5
c.	NIPT	1	2	3	4	5

**19.** Do you feel that your opinion as a partner is taken into account by **healthcare professionals** when deciding about one of the following tests?

(PLEASE CIRCLE ONE ANSWER FOR EACH STATEMENT)

		Not at all		Somewhat		Very much
a.	Current screening	1	2	3	4	5
b.	Amniocentesis	1	2	3	4	5
c.	NIPT	1	2	3	4	5

20. If your partner disagreed with your desire to do or not NIPT, what impact would that have on your view? (PLEASE CIRCLE ONE ANSWER ONLY)

No impact		Some impact		A big impact
1	2	3	4	5

Any other comments on the impact a disagreement would have on your decision:







## **PART 5: SOCIAL IMPACT OF NIPT**

**21.** Provincial health care systems cover routine prenatal care. Right now, NIPT is not part of routine prenatal care in most provinces and territories. If NIPT were covered as part of routine prenatal care, which of the following outcomes would be of concern to you?

(PLEASE CIRCLE ONE ANSWER FOR EACH STATEMENT)

	,	Not concerned		Somewhat concerned		Very concerned
a.	Increased pressure on women to use NIPT	1	2	3	4	5
b.	Increased use of NIPT leading to increased pressure to terminate if the baby has Down syndrome (DS)	1	2	3	4	5
c.	Increased availability of NIPT making people less willing to accept children with disabilities	1	2	3	4	5
d.	Decrease of the population of people with DS	1	2	3	4	5
e.	Reduction in resources available for people with DS and their families	1	2	3	4	5
f.	Negative impact on individuals with DS and their families (stigma, discrimination)	1	2	3	4	5

Other:		

 $\textbf{22.} \quad \text{If NIPT became part of routine tests offered during the pregnancy, how likely is it that:} \\$ 

		No pressure		Some pressure		A lot of pressure
a.	your partner would feel pressure to take the test because it would be routine	1	2	3	4	5
b.	<b>you</b> would feel pressure to take the test for the pregnancy because it would be routine	1	2	3	4	5







# **PART 6: FUTURE USES OF NIPT**

**23.** In the future, NIPT may be able to test for many genetic conditions. How interested would you be in NIPT for the following conditions:

		Not interested		Somewhat interested		Very interested
a.	Inherited disorders (such as Tay-Sachs, cystic fibrosis, sickle cell disease, Gaucher disease)	1	2	3	4	5
b.	Paternity testing	1	2	3	4	5
c.	Physical and behavioural attributes (eye colour, intelligence, sexual orientation)	1	2	3	4	5
d.	Predisposition to childhood-onset diseases (autism, leukemia)	1	2	3	4	5
e.	Predisposition to late-onset diseases (heart problems, Alzheimer's disease, cancer)	1	2	3	4	5
f.	Predisposition to mental disorders (schizophrenia, bipolar disease)	1	2	3	4	5

	Other:				
DAD	T 7: ABOUT YOURSELF				
FAN	11 7. ADOUT TOURSLLF	MM/AAAA			
24.	Your birth date (Month/				
25.	Your gender:				
26.	Your country of birth:			-	
27.	What language do you m	nostly speak at home?			
	□English	☐ French	☐ Other:		







	In which pro (PLEASE CHECK		ritory do you <b>curr</b> R <b>ONLY)</b>	rently live?					
□ Alb	erta	□ New Br	unswick	☐ Northwe Territori		☐ Ontario		☐ Saskatch	ewan
□ Bri Co	tish Iumbia	☐ Newfou Labrado	indland and or	☐ Nunavik		☐ Prince-Ed	dward-	☐ Yukon	
☐ Ma	anitoba	□ Nova So	otia	☐ Nunavu	t	☐ Quebec			
	What is you								
☐ Ca	ucasian		☐ African desce	nt	□ North A	merican Abo	original (Fir	st Nation, Inu	it, Metis)
☐ La	tin American		☐ Middle Easter	rn	Other:				_
☐ As	ian								
	What is your		ultural backgroun PLY)	ıd?					
□Bu	ıddhist		☐ Christian		☐ Muslim		☐ No reli	gious affiliati	on
□ Ca	tholic		☐ Jewish		☐ Protest	ant	☐ Other:		
☐ Ca	tholic Ortho	dox	☐ Hindu		☐ Sikh				
	How importa		on / spirituality in						
	Not impo	rtant		Somewh importa			Very	important	
	1		2	3		4		5	
	What is the (PLEASE CHECK		el of school you <i>cc</i>	ompleted?					
	☐ Elementa	ary school	☐ High s	school	Г	☐ University			
	□ Trades/A	pprentices	nip     Colleg	ge	Г	<b>□</b> Other:			







33.	Are you a healthcare profession (PLEASE CHECK ONE ANSWER ONLY)	onal?				
	☐ Yes ———— 3	<b>3.1.</b> If yes, what profession?				
	□No					
34.	What is your relationship statu (PLEASE CHECK ONE ANSWER ONLY)	us?				
	☐ Married	☐ Single	☐ Widowed			
	☐ Common Law	☐ Divorced/separated	☐ Other :			
35.	Do you have children? (PLEASE CHECK ONE ANSWER ONLY)					
	□Yes	□No				
36.	Do you have a child with Down syndrome?  (PLEASE CHECK ONE ANSWER ONLY)					
	□Yes	□No				
37.	Do you have a child with a physical or intellectual disability?  (PLEASE CHECK ONE ANSWER ONLY)					
	□Yes	□No				
38.	Does anyone close to you have (PLEASE CHECK ONE ANSWER ONLY)	e a child with Down syndrome?				
	□Yes	□No				







39.	For this pregnancy, is your partner considered:  (PLEASE CHECK ONE ANSWER ONLY)					
	☐ Low/average risk for D	own syndrom	ne ☐ High risk for Down	n syndrome	☐ Unsure	
40.	For this pregnancy, who provided you information about current screening, amniocentesis and/or NIPT? (PLEASE CHECK ALL THAT APPLY)					
	☐ Family physician☐ Obstetrician/Gynecologist		☐ Genetic Counsellor	□Partner		
			☐ Nurse [			
	☐ Medical geneticist		☐ Midwife			
41.	This pregnancy was conce (PLEASE CHECK ONE ANSWER C					
	□ Naturally		☐ with assisted reproductive to	echnology (ART),	other than IVF	
40. 41.	☐ by <i>in vitro</i> fertilization (IVF)		☐ Other :			
42.	Has your partner had <b>prenatal screening</b> in a <b>previous pregnancy</b> ?  (PLEASE CHECK ONE ANSWER ONLY)					
	☐ Yes ———	=	was the result of the prenatal s	creening?		
	□ No	☐ High risk	for Trisomy (e.g. Down Syndron	ne, trisomy 13, tr	isomy 18)	
	☐ Unsure	☐ High risk	for a Neural Tube Defect (e.g. s	pina bifida)		
		☐ Low or av	verage risk			
		☐ Unsure				
43.	Has your partner had <b>prenatal diagnosis</b> (chorionic villus sampling or amniocentesis) in a <b>previous pregnancy</b> ? (PLEASE CHECK ONE ANSWER ONLY)					
	☐ Yes ———	-	was the result of the prenatal d	liagnosis?		
	□ No	☐ Diagnosis	of a genetic condition (e.g. cys	tic fibrosis, Down	syndrome)	
	☐ Unsure	☐ Normal re	esult (no genetic condition dete	cted)		
		Other:				







# Thank you for completing this survey.

you have any additional comments or thoughts, please write them below.						