

# PRENATAL SCREENING FOR DOWN SYNDROME, TRISOMY 18 AND OPEN NEURAL TUBE DEFECTS

Prenatal screening estimates the fetal risk of Down syndrome, trisomy 18, and open neural tube defects. The results will assist in determining the need for further diagnostic testing.

The screening tests offered will vary according to the gestational age at the time of presentation, maternal age at the time of delivery, and singleton versus twin gestation.

**Table 1: Tests available through the BC Prenatal Genetic Screening Program**

Test Name	Markers/ Measurements	Possible Timeframe	Best Timeframe
Serum Integrated Prenatal Screen (SIPS) blood test #1	PAPP-A	9–13 <sup>+6</sup> wks	10–11 <sup>+6</sup> wks
SIPS blood test #2	AFP uE3 hCG Inhibin-A	14–20 <sup>+6</sup> wks	15–16 wks
Integrated Prenatal Screen (IPS)	Same as SIPS (blood test #1 & #2) with addition of NT ultrasound <sup>1</sup>	See SIPS for blood tests 11–13 <sup>+6</sup> wks	See SIPS for blood tests 12–13 <sup>+3</sup> wks
Quad blood screen	Same as SIPS blood test #2	14–20 <sup>+6</sup> wks	15–16 wks
Non-Invasive Prenatal Screening (NIPS)	Cell-free DNA in maternal blood	10 weeks and onwards	varies by indication

<sup>1</sup> If an NT ultrasound is performed, NT centers require a separate first trimester dating ultrasound.

## Resources

- BC Prenatal Genetic Screening Program; guideline and related patient teaching resources; [www.bcprenatalscreening.ca](http://www.bcprenatalscreening.ca)
- Perinatal Services BC; T 604-877-2121; E [psbc@phsa.ca](mailto:psbc@phsa.ca); [www.perinatalservicesbc.ca](http://www.perinatalservicesbc.ca)
- Canadian Down Syndrome Society; T (800) 883-5608; E [info@cdss.ca](mailto:info@cdss.ca); [www.cdss.ca](http://www.cdss.ca)
- Down Syndrome Resource Foundation (Burnaby, BC); T 604-444-3773; [www.dsrff.org](http://www.dsrff.org)
- Lower Mainland Down Syndrome Society (Canada); T 604-591-2722; [www.lmdss.com](http://www.lmdss.com)
- Society of Obstetricians and Gynaecologists, Clinical Practice Guidelines (Canada); [www.sogc.org/guidelines](http://www.sogc.org/guidelines)
- Spina Bifida and Hydrocephalus Association of BC; T 604-878-7000; E [info@sbhbc.org](mailto:info@sbhbc.org); [www.sbhbc.org](http://www.sbhbc.org)
- Support Organization For Trisomy 18, 13 and Related Disorders (SOFT; US); [www.trisomy.org](http://www.trisomy.org)

## Genetic counselling services (Medical Genetics)

Victoria: T 250-727-4461 Fax for referrals: 250-727-4295  
Vancouver: T 604-875-2157 Fax for referrals: 604-875-3484

## Questions about prenatal screening in BC

Prenatal Biochemistry Laboratory: T 604-875-2331

**Table 2: Screening options available through the BC Prenatal Genetic Screening Program**

Characteristics of woman	Gestational Age at the First Prenatal Visit		
	≤ 13 <sup>+6</sup> weeks	14–20 <sup>+6</sup> weeks	≥ 21 weeks (no prior screening)
<35 years	• SIPS	• Quad	• Detailed ultrasound
35–39 years	• IPS; or • If NT not available, SIPS	• Quad	• Detailed ultrasound; and • Amnio
40+ years	• IPS; or • If NT not available, SIPS; or • CVS or Amnio without prior screening	• Quad; or • Amnio without prior screening	• Detailed ultrasound; and • Amnio
Personal/family history that increases risk of fetus with Down syndrome, trisomy 18, or trisomy 13	• NIPS; or • CVS or Amnio without prior screening	• NIPS; or • Amnio without prior screening	• Detailed ultrasound; and • NIPS; or • Amnio
Personal/family history that increases risk of fetus with chromosomal abnormality other than Down syndrome, trisomy 18, or trisomy 13	• CVS or Amnio without prior screening	• Amnio without prior screening	• Detailed ultrasound; and • Amnio
Twin gestation	• IPS; or • If NT not available, SIPS; or • If ≥ 35, Amnio without prior screening	• Quad; or • If ≥ 35, Amnio without prior screening	• Detailed ultrasound; and • If ≥ 35, Amnio
Pregnant following In vitro fertilization with intracytoplasmic injection and without preimplantation genetic testing	• IPS; or • If NT not available, SIPS; or • CVS or Amnio without prior screening	• Quad; or • Amnio without prior screening	• Detailed ultrasound; and • Amnio

If the prenatal screen result is **screen positive for Down syndrome** (assuming gestational dating is confirmed) or **trisomy 18**, women should be counselled by their health care provider and offered further testing. See [www.bcprenatalscreening.ca](http://www.bcprenatalscreening.ca) for details.

If the prenatal screen result is **positive for an open neural tube defect**, and dating is confirmed, see Appendix 4 of the guidelines for follow-up of elevated MSAFP.

**Table 3: Eligibility, Screen Cut-Offs, and Performance of Screening Tests <sup>1</sup>**

	Serum Integrated Prenatal Screen (SIPS)	Integrated Prenatal Screen (IPS)	Quad Screen (QUAD)	Non-Invasive Prenatal Screening (NIPS)	
<b>WHO IS ELIGIBLE</b>	All women who present for their first visit at $\leq 13^{+6}$ wks	Women who present for their first prenatal visit at $\leq 13^{+6}$ wks gestation and who: <ul style="list-style-type: none"> <li>• Are 35 years or older at EDD</li> <li>**OR**</li> <li>• Are pregnant with twins</li> <li>**OR**</li> <li>• Are pregnant following In vitro fertilization with intracytoplasmic injection and without preimplantation genetic testing</li> </ul>	All women who present for their first prenatal visit between 14 and $20^{+6}$ wks gestation	<ul style="list-style-type: none"> <li>• Women with a positive (IPS / SIPS / Quad) screen for Down syndrome or trisomy 18</li> <li>**OR**</li> <li>• Women with a previous pregnancy with trisomy 21, 18, or 13</li> <li>**OR**</li> <li>• Women with an increased risk (<math>\geq 1/300</math>) for Down syndrome based on ultrasound marker(s) and serum screen result</li> </ul>	
<b>DOWN SYNDROME</b>	<b>Screen cut-off</b>	<b>1:900</b>	<b>1:200</b>	<b>1:900</b>	
	Detection rate	< 35 yrs: 86% 35 – 39 yrs: 96% $\geq 40$ yrs: 100% <sup>2</sup>	< 35 yrs: 86% 35 – 39 yrs: 98% $\geq 40$ yrs: 97%	< 35 yrs: 83% 35 – 39 yrs: 100% <sup>2</sup> $\geq 40$ yrs: 100% <sup>2</sup>	~98%
	False positive rate	< 35 yrs: 8% 35 – 39 yrs: 21% $\geq 40$ yrs: 44%	< 35 yrs: 7% 35 – 39 yrs: 9% $\geq 40$ yrs: 16%	< 35 yrs: 9% 35 – 39 yrs: 24% $\geq 40$ yrs: 45%	< 0.1%
	Chance a screen negative result is a false negative result	< 0.1%	< 0.1%	< 0.1%	< 0.01%
<b>TRISOMY 18</b>	<b>Screen cut-off</b>	<b>1:300</b>	<b>1:300</b>	<b>1:300</b>	
	Detection rate	88%	90%	75%	~85%
	False positive rate <sup>3</sup>	0.6%	2%	1%	< 0.1%
	Chance a screen negative result is a false negative result	< 0.1%	< 0.1%	< 0.1%	< 0.1% <sup>4</sup>

<sup>1</sup> Performance of screening tests applies to singleton pregnancies. SIPS/IPS/Quad data from Perinatal Services BC. British Columbia Perinatal Data Registry. Years provided: April 1, 2016 to March 31, 2021. Resource type: Tabulated data. NIPS data from Perinatal Services BC based on 229 cases of T21 and 45 cases of Trisomy 18.

<sup>2</sup> The detection rates listed are based on the small cohort of Down syndrome pregnancies in BC. SIPS, IPS and Quad are screening tests so may not have 100% detection rate.

<sup>3</sup> Higher false positive rate of IPS reflects that this test is done in women who are at a higher apriori risk.

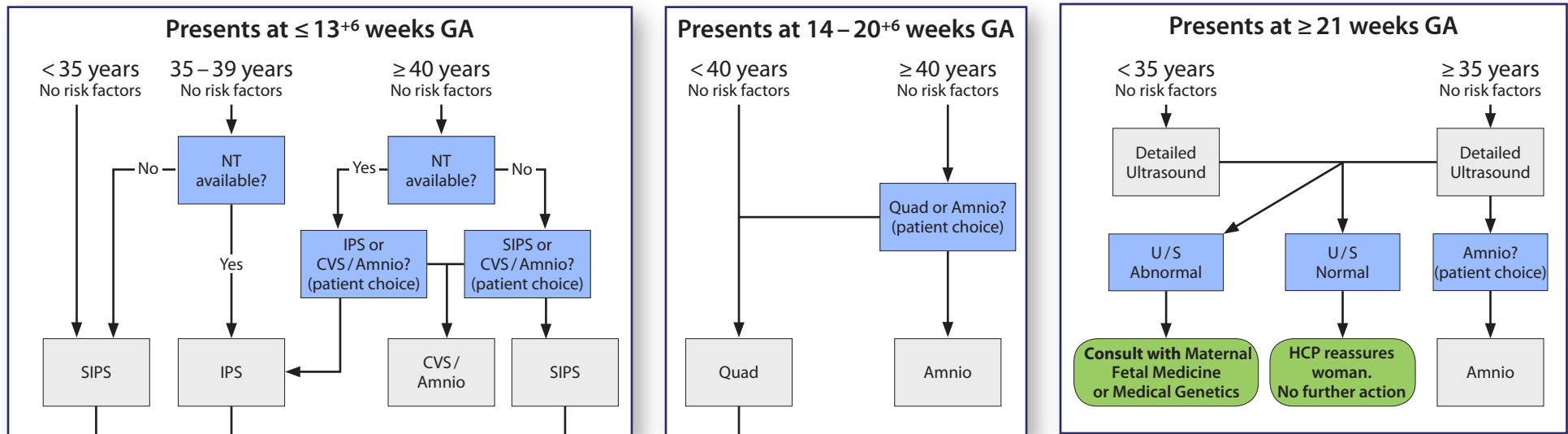
<sup>4</sup> May be higher if ultrasound abnormalities present.

**Table 4: Risk of Down Syndrome and Other Chromosome Abnormalities in Live Births by Maternal Age**

Maternal Age (At Term)	Risk		Maternal Age (At Term)	Risk		Maternal Age (At Term)	Risk	
	Down Syndrome	Total Chromosome Abnormality		Down Syndrome	Total Chromosome Abnormality		Down Syndrome	Total Chromosome Abnormality
25	1 in 1,250	1 in 476	32	1 in 637	1 in 323	39	1 in 125	1 in 81
26	1 in 1,190	1 in 476	33	1 in 535	1 in 286	40	1 in 94	1 in 63
27	1 in 1,111	1 in 455	34	1 in 441	1 in 224	41	1 in 70	1 in 49
28	1 in 1,031	1 in 435	35	1 in 356	1 in 179	42	1 in 52	1 in 39
29	1 in 935	1 in 417	36	1 in 281	1 in 149	43	1 in 40	1 in 31
30	1 in 840	1 in 385	37	1 in 217	1 in 123	44	1 in 30	1 in 21
31	1 in 741	1 in 385	38	1 in 166	1 in 105	$\geq 45$	$\geq 1$ in 24	$\geq 1$ in 19

Source: Hecht CA and Hook EB. 1996

**Woman with Singleton Pregnancy Presents for 1st Prenatal Visit**  
(Excluding women with personal / family history that increases risk of fetus with Down syndrome, trisomy 18, or other chromosomal abnormality, who should be referred to Medical Genetics for counselling of their prenatal screening/diagnostic options)

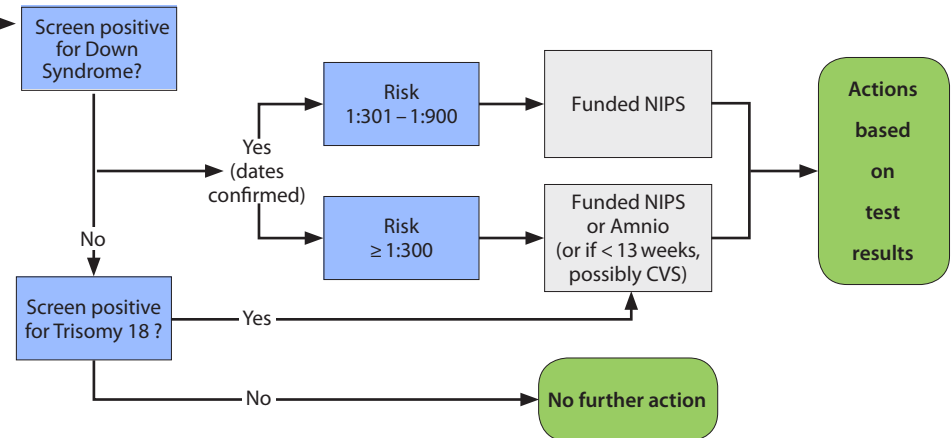


## BC Prenatal Genetic Screening Program

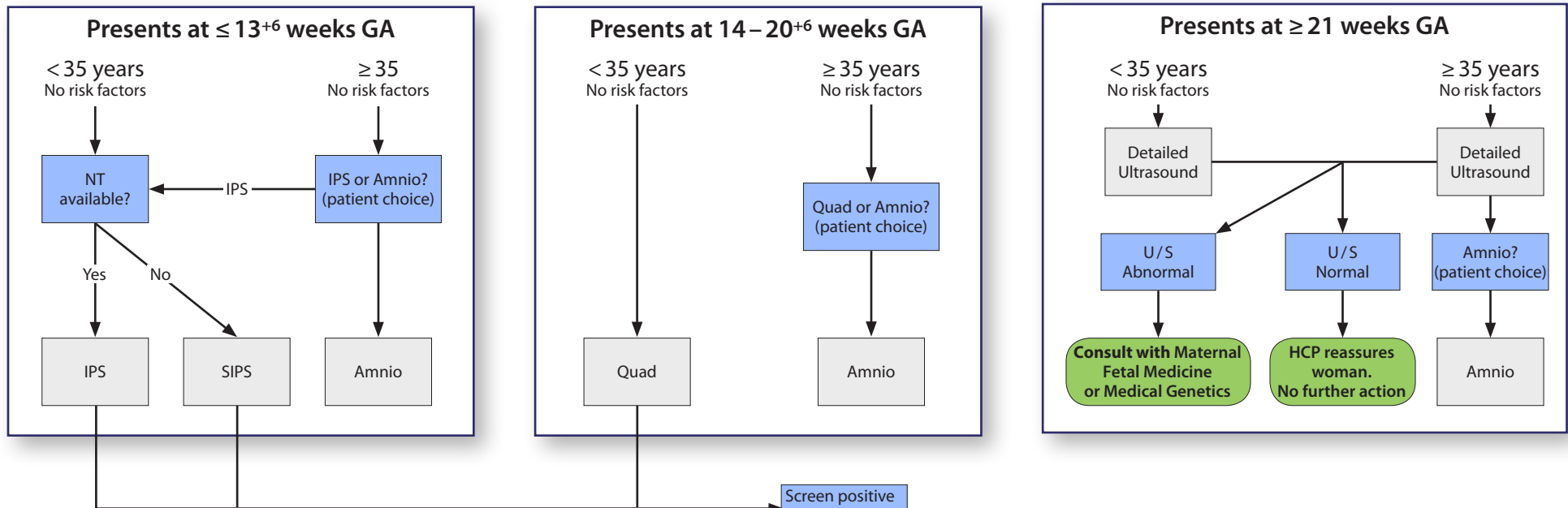
See [www.bcprenatalscreening.ca](http://www.bcprenatalscreening.ca) for more details on amniocentesis and NIPS

**Legend**

- GA = Gestational Age
- IPS = Integrated Prenatal Screen
- NIPS = Non-Invasive Prenatal Screening
- NT = Nuchal Translucency
- SIPS = Serum Integrated Prenatal Screen
- U/S = Ultrasound



**Woman with Twin Pregnancy Presents for 1st Prenatal Visit**  
(Excluding women with personal / family history that increases risk of fetus with Down syndrome, trisomy 18, or other chromosomal abnormality, who should be referred to Medical Genetics for counselling of their prenatal screening/diagnostic options)



## BC Prenatal Genetic Screening Program

See [www.bcprenatalscreening.ca](http://www.bcprenatalscreening.ca) for more details on amniocentesis and NIPS options

\* Screening in higher multiples will remain based on NT alone. If NT is not available and the woman is  $\ge 35$  years old, amniocentesis is an option.

**Legend**

- GA = Gestational Age
- IPS = Integrated Prenatal Screen
- NIPS = Non-Invasive Prenatal Screening
- NT = Nuchal Translucency
- SIPS = Serum Integrated Prenatal Screen
- U/S = Ultrasound

