# NIPT REQUEST FORM

SOGI SCAN

Cell free fetal DNA blood test for Down Syndrome

**Patient Details** 

Name

Address

A MEMBER OF THE I-MED NETWORK

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Clinical Details		
LNMP	EDC	Blood Group
10 week scan and NIPT with pre/post test result genetic counselling and specialist consultation		
12 week NT scan and NIPT with pre/post result genetic counselling and specialist consultation		
NIPT with pre/post test result genetic counselling and specialist consultation for high risk NT scan elsewhere		
NIPT with pre/post test result genetic counselling and specialist consultation for scan finding or for further reassurance		

#### Notes

NIPT blood test does not replace 12 week NT scan.

Invasive testing (CVS/amniocentesis) recommended if:

- 1. Nuchal translucency >3.5mm with microarray testing
- 2. Fetal structural lesion

Fetal echocardiography at a tertiary centre recommended if NT >3.5mm.

12 week scan will effectively screen for open spina bifida.

12 week scan will screen for some major structural anomalies.

Medicare rebates are available for your patient for:

- NIPT specialist consultation and NIPT scan but not for NIPT blood test (sent to USA)
- 10 week scan confirms dates and viability prior to blood test

Arlosa Harmony test as of July 2013 Detection rate:

- · trisomy 21 >99%
- trisomy 18 >98%
- trisomy 13 80% sex chromosome anomalies: depends on anomaly

Up-to-date costs are explained at www.sogiscan.com.au



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