

Measurements – what do they tell you?

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Measurements across the trimesters

- Distinguishing between ongoing pregnancy & missed miscarriage
- Accuracy of combined screening risk(s)
- Estimation of expected date of delivery (EDD)
- Exclusion of abnormal findings
- Assessment of growth velocity & 'well being'
- Cervical length
- Relationship between placental site & os



CRL, NT & combined screening

Correct sections for measuring CRL & NT

Correct placement of the callipers

- What difference do they make to:
- dating
- NT risk
- combined screening risk?





UK National Screening Committee



Screening Programmes

Fetal Anomaly

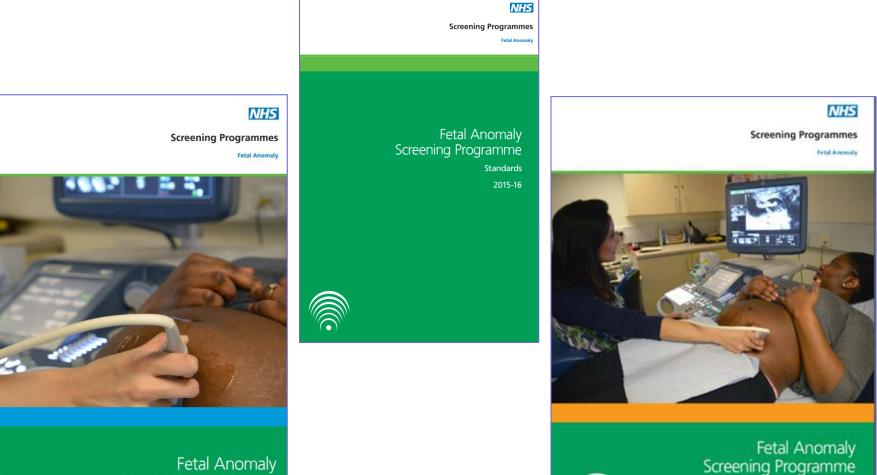
FASP Public Health England

April 2013



Part of Public Health England

Fetal Anomaly Screening Programme Guidance (England)



Screening Programme

April 2015

Handbook for ultrasound practitioners

Screening Programme

Programme handbook

June 2015

Early pregnancy scan - dating

Purpose of dating scan

- confirm viability
- singleton or multiple
- estimate gestational age (CRL or HC)
- detect major structural abnormalities
- e.g. anencephaly















Dating - national guidance (UK)





Ucmissiousia - August 2009 - Valume 17 - Number 3-

Fetal size and dating: charts recommended for clinical obstetric practice

Pam Loughna¹, Lyn Chitty², Tony Evans³ & Trish Chudleigh⁴ ¹Academic Division of Obstetrics and Gynaecology, Nottingham University Hospitals NHS Trust, ²Genetics and Fetal Medicine, Institute of Child Health and University College London Hospitals NHS foundation Trust, London, ³Medical Physics, University of Leeds, Leeds and ⁴The Rosie Hospital, Cambridge, UK

- British Medical Ultrasound Society BMUS.org
- FASP <u>www.gov.uk/topic/population-screening-programmes/fetal-anomaly</u>
- cpd.screening.nhs.uk

CRL - practical issues

- appreciation of 'unflexed' length sagittal not coronal
- rotate or slide past selected section
- better or worse?
- freeze or return to optimal section
- measure using linear callipers
- CRL from 3 separate images +/- 1-3mm



CRL – practical issues





















Fetal Anomaly Screening Programme Guidance - CRL

Table 4 - Recommended criteria for measurement of CRL for pregnancy dating and combined screening (Loughna P et al (2009))

CRL	Detail to be demonstrated
Midline section	 Sagittal section of the fetus with the head in line with the full length of the body
	Echogenic tip of the nose
	Rectangular shape of the palate
	Translucent diencephalon
	 CRL axis should be between 0° and 30° to the horizontal
	Clearly defined crown and rump
Position	 Pocket of fluid, at least equivalent in size to the width of the palate, should be visible between the fetal chin and chest
	 Fetal palate angle should be 30° to 60° relative to the horizontal
	Nasal tip should be level or above the anterior abdominal wall
Magnification	Entire CRL section should fill over 60% of the screen
Calliper placement	Correct calliper placement on outer borders of crown and rump
	Longest length of the fetus should be measured
Image archiving	 The CRL should be measured at least twice and the maximum measurement that meets the criteria should be recorded
	 The image demonstrating the measured CRL which has been reported should be archived





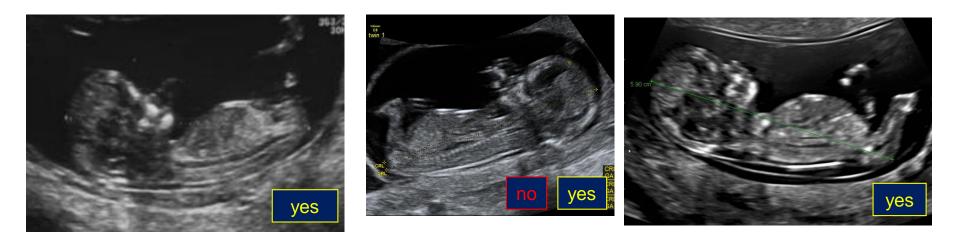


CRL – correct section





CRL – calliper placement





Fetal Anomaly Screening Programme Guidance - CRL



Diagram 4 Examples of scoring CRL images

Image 4 Good



Midline section				Position			Mag	Callipers				
1	2	3	4	5	6	7	8	9	10	11	12	
1	~	~	1	1	1	~	1	1	~	~	~	
12/1	2 compo	nents p	resent									Good

Image 5 Acceptable



Midline section				Position			Mag	Callipers				
1	2	3	4	5	6	7	8	9	10	11	12	
1	*	*	х	1	1	*	1	*	*	*	1	
	2 compo ienceph											Accept-

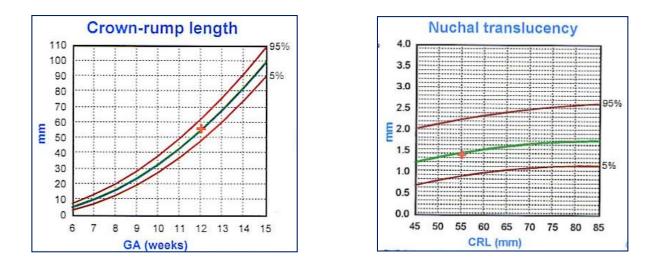
Image 6 Poor



Midline section				Posi	Position			Callipers		Overall		
1	Z	3	4	5	6	7	8	9	10	11	12	
x	*	~	1	*	x	1	1	1	*	х	x	
1) the full length of the body is not present 6) the rump is not clearly defined									Poor			
11) c	allipers	are not o	correctly	placed o	on the rul	mp						

Combined screening





- risk algorithm only for CRL 45.0 84.0mm (11⁺² 14⁺⁰ wks)
- NT normal range 1.0 ~ 2.5mm
- NT increases with CRL (& therefore gestational age)









Fetal Anomaly Screening Programme Guidance - NT

Table 3 - Recommended criteria for measurement of NT for combined screening

NT	Detail to be demonstrated
Midline section	 Horizontal sagittal* section of the fetus extending from crown to upper aspect of the heart which may be supine or prone**
	Head in line with the body with the NT visible along the length of the neck
	Echogenic tip of the nose
	Rectangular shape of the palate
	Translucent diencephalon
	 Frontal process of the maxilla should not be visible
Position	 Pocket of fluid, at least equivalent in size to the width of the palate, should be visible between the fetal chin and chest
	 Angle of the palate relative to the horizontal should be between 30° and 60°
	 Nasal tip should be level with, or above, the anterior chest wall
Magnification	 The section should fill over 60% of the screen
Calliper placement	Callipers should be placed on the upper and lower edges of the NT
	 Widest part of the NT should be measured
Image archiving	 The NT should be measured at least twice and the maximum measurement that meets the criteria should be recorded
	 The image demonstrating the measured NT which has been reported should be archived

* In all criteria the term sagittal describes a midline longitudinal section

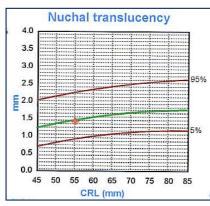
** FASP does not recommend screening for nasal bone absence or hypoplasia, thus allowing measurement of the NT with the fetus in the prone position

NT – correct section & caliper placement







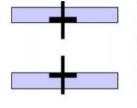




Fetal Anomaly Screening Programme Guidance - NT



Diagram 2 Where to place callipers for the NT measurement





Measurement should be taken with the inner border of the horizontal of the callipers placed ON the line that defines the NT thickness. The crossbar of the calliper should be such that it is hardly visible as it emerges with the white line of the border. It should not be visible in the nuchal fluid.

Diagram 3 Examples of scoring NT images

Image 1 Good



Midline section					Position			Mag	Callipers		Overall	
1	2	3	4	5	6	7	8	9	10	11	12	
1	1	*	*	*	1	1	1	*	*	1	*	
12/12	compo	nents pr	esent									Good

Image 3 Poor

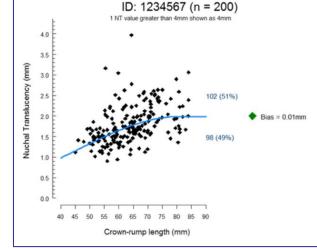


Midline section						Position			Mag	Callipers		Overall
1	2	з	4	5	6	7	8	9	10	11	12	
1	1	~	1	1	х	x	1	x	*	x	x	
7) No 9) Na 11) C	ontal pro o pocket usal tip b Callipers i Videst pi	of fluid elow an are not c	under th terior ch correctly	ne chin est wall placed o	on the sl	kin lines						Poor

National audit of combined screening programme

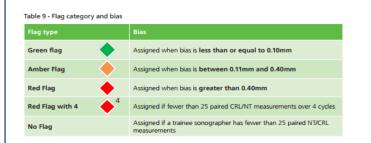
- all providers must be DQASS registered
- All CRL & NT data sets submitted to DQASS, by lab
- annual DQASS audit
 - individual, departmental, regional, national
- 3 monthly departmental review individual NT & CRL images (3 x 2)



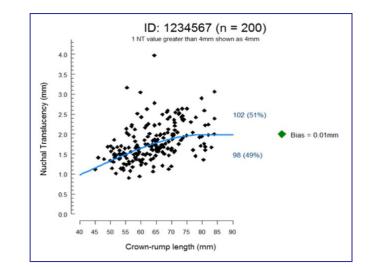


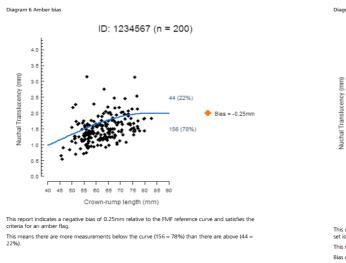


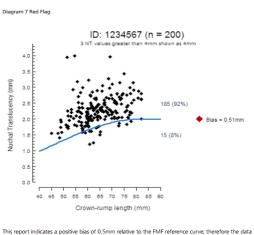
Audit of combined screening programme - flags



- each individual report demonstrates the NT and CRL measurements relative to the FMF reference curve
- bias describes the number of measurements above and below the FMF reference curve
- the bias is either negative in terms of under-measurement (below the FMF reference curve) or positive which refers to over-measurement (above the FMF reference curve)
- the evidence used to develop the flag status was derived from the impact on screening
 performance. For positive biases greater than 0.40mm, the standardised screen positive rate (SPR)
 exceeds 5% and increases the number of pregnancies exposed to the potential risks and anxieties
 associated with a screen positive result which may lead to invasive diagnostic procedures







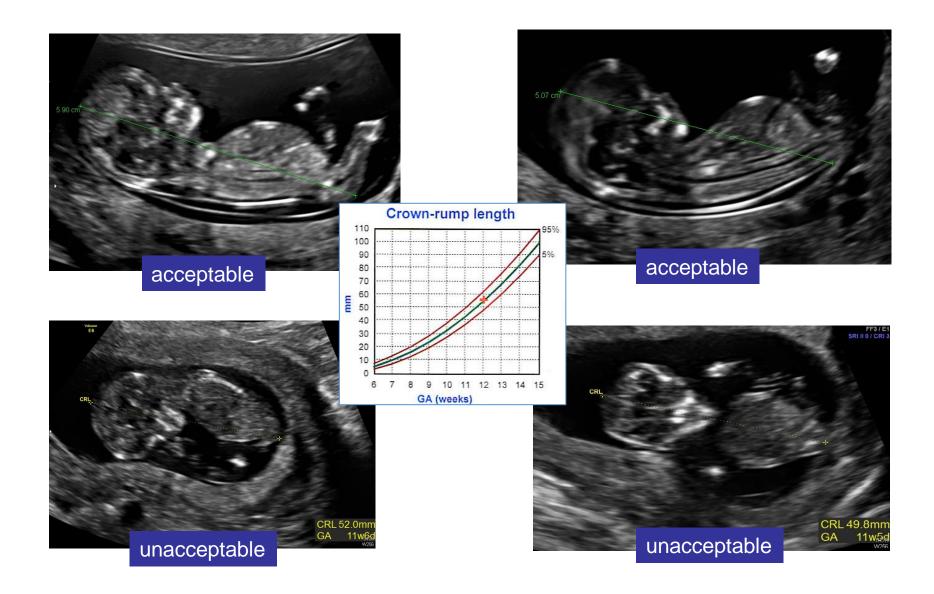
set is assigned a red flag.

This means 92% (185) measurements plotted above the curve, with only 8% (15) plotted below.

Bias can directly impact on the risk calculation women receive.

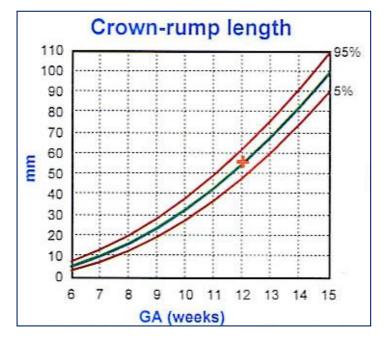
This dataset indicates that there will be an increase in detection rate (91%), however, the

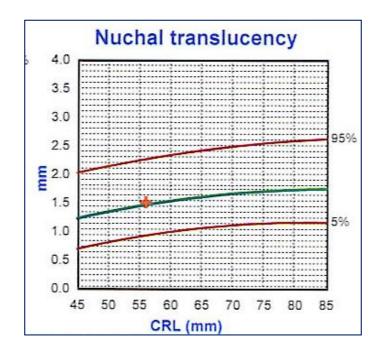
Combined screening – relies on accurate CRL



Impact of CRL in combined screening

- CRL determines gestational age
- CRL determines maternal age (at conception)
- underestimating CRL:
 - → increases MA risk (older at conception)
 - \rightarrow increases risk from NT





Practical implications of poor technique for Tri 21 - CRL



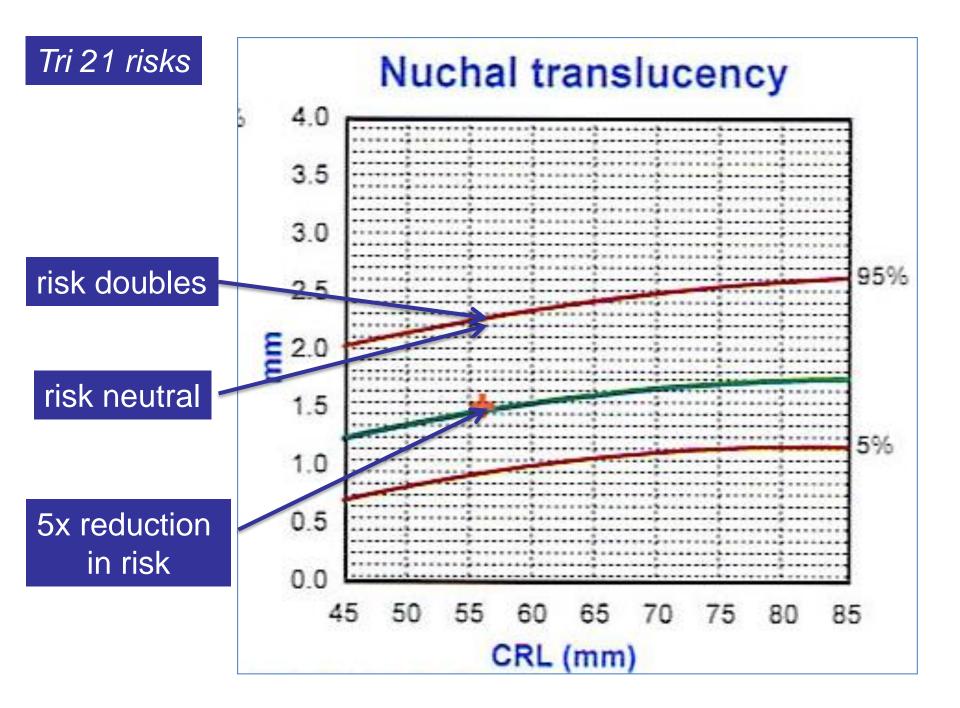


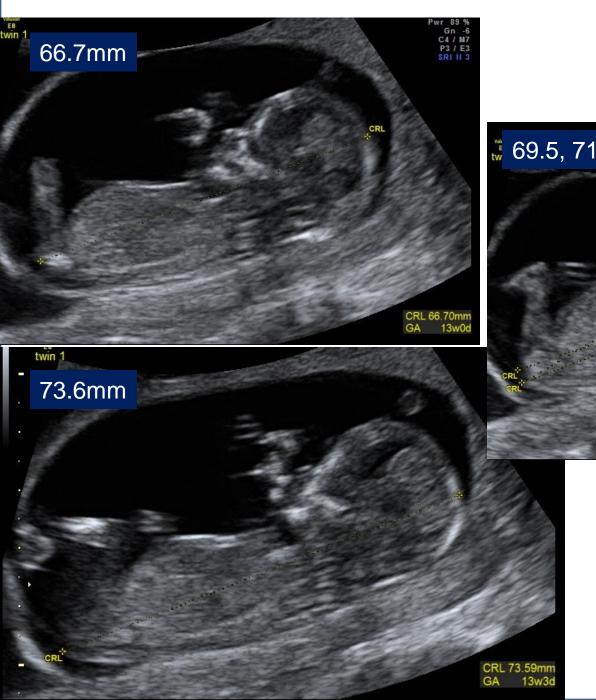
30 yrs, NT 2.4mm, dating by CRL (Tri 21 risks at term)

CRL	GA	Background risk	Adjusted risk
52.8	11+6	1:906	1:182
48.9	11+4	1:905	1:143

NT & Adjusted risk CRL 60mm = 12 wks + 2 days

NT (mm)	25yrs (1:1378)	35yrs (1:425)	40yrs (1:28)
1.5 (50 th c)	7418	2289	679
2.0	3439	1060	315
2.1	2688	823	244
2.2	2005	619	184
2.3 (95 th c)	1082	333	100
2.4	635	133	59
2.5	431	95	40
2.6	309	71	29
2.7	230	71	22
2.8	177	55	17





CRL 71.95mm GA 13w1d CRL 73.36mm GA 13w3d

CRL & Risk (NT 2.0mm)

CRL (mm)	Gest Age (wks + days)	35 yrs Maternal risk	35 yrs Adjusted risk	40yrs Maternal risk	40 yrs Adjusted risk
66.7	12+6	1:383	1:1208	1:112	1:352
69.5	13+0	1:383	1:1297	1:112	1:378
71.9	13+2	1:383	1:1362	1:112	1:397
73.6	13+2	1:383	1:1393	1:112	1:406

Effects of CRL biases on biochemistry MoM values (Tri 21)

CRL Error	Error in GA	Media	an MoM		
(mm)	(days)	PAPP-A	hCG beta	DR	FPR
-5	-2.7	1.17	0.93	80%	1.4%
-4	-2.2	1.13	0.94	81%	1.6%
-3	-1.6	1.10	0.96	82%	1.9%
-2	-1.1	1.06	0.97	83%	2.0%
-1	-0.5	1.03	0.99	84%	2.4%
0	0.0	1.00	1.00	85%	2.7%
1	0.5	0.97	1.01	86%	3.1%
2	1.0	0.94	1.03	86%	3.7%
3	1.6	0.92	1.04	87%	4.1%
4	2.1	0.89	1.06	88%	4.8%
5	2.6	0.87	1.07	89%	5.5%

Note that Down's syndrome presents as biochemical immaturity so effects of errors on PAPP-A and hCG combine

Data provided by Prof D Wright, lead statistician for DQASS, NHS FASP

Effects of NT bias on FPR & DR for Tri 21

Bias (mm)	FPR	DR
-0.4	2%	79%
-0.3	1.8%	80%
-0.2	2.0%	82%
-0.1	2.2%	83%
0	2.6%	85%
0.1	3.1%	86%
0.2	3.7%	87%
0.3	4.6%	88%
0.4	5.7%	90%

Data provided by Prof D Wright, lead statistician for DQASS, NHS FASP



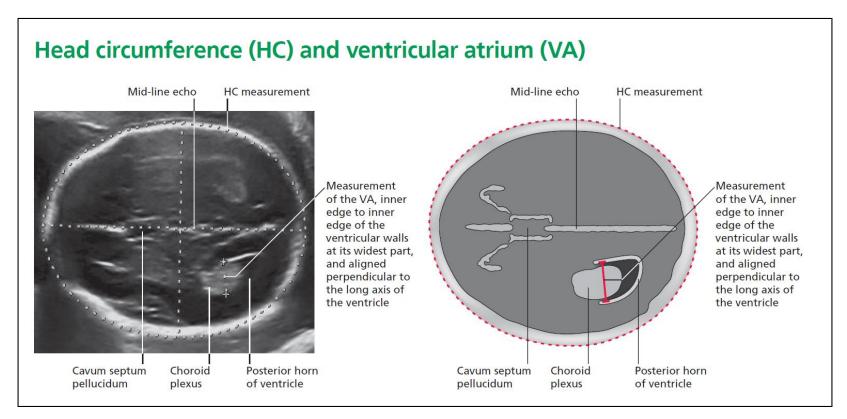


18⁺⁰ - 20⁺⁶ weeks – biometry, HC

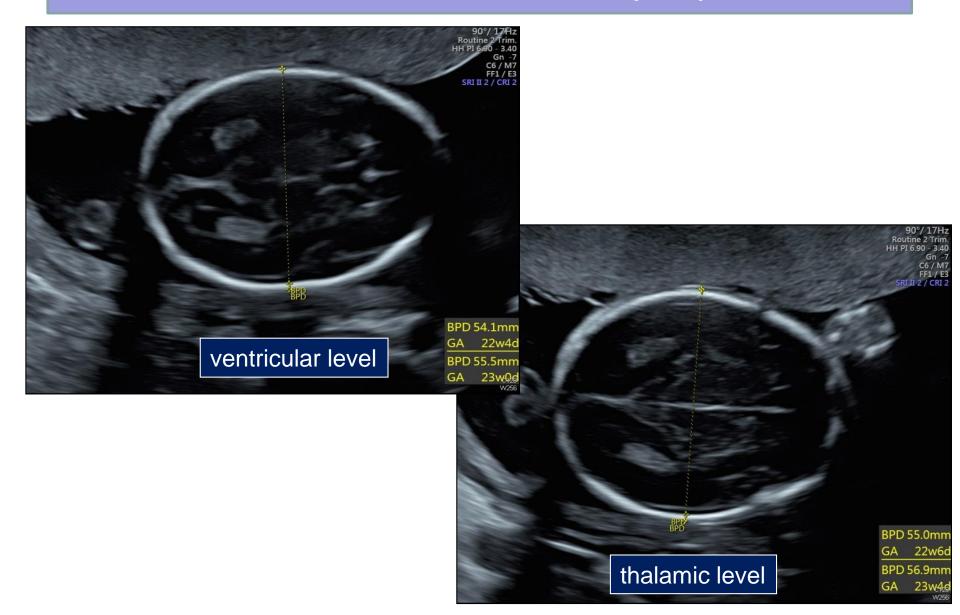


Assess fetal growth velocity, based on EDD assigned at dating/combined screening by measurement of:

HC (Chitty) - ventricular level, with image



Head measurements – level & calliper placement

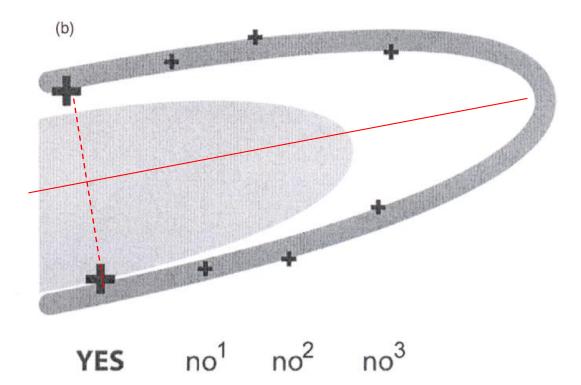


Femur length measurements- calliper placement





Measurement of the posterior horn - correct calliper placement

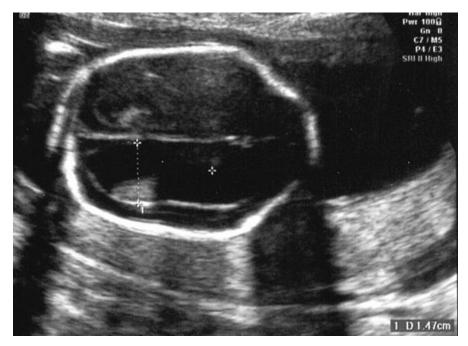


Ref: ISUOG (2007) Sonographic examination of the fetal central nervous system: guidelines for performing the 'basic examination' and 'fetal neurosonogram'. *The International Society of Ultrasound in Obstetrics & Gynaecology.* Vol 29; Pp 109 – 116.

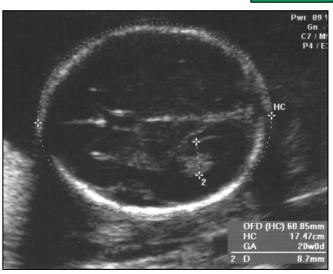
Size & abnormal findings

refer if:

- nuchal fold >6.0mm
- ventriculomegaly (PH>10.0mm)
- RPD (AP pelvis >7.0mm)
- significantly small (<< 5th centile)





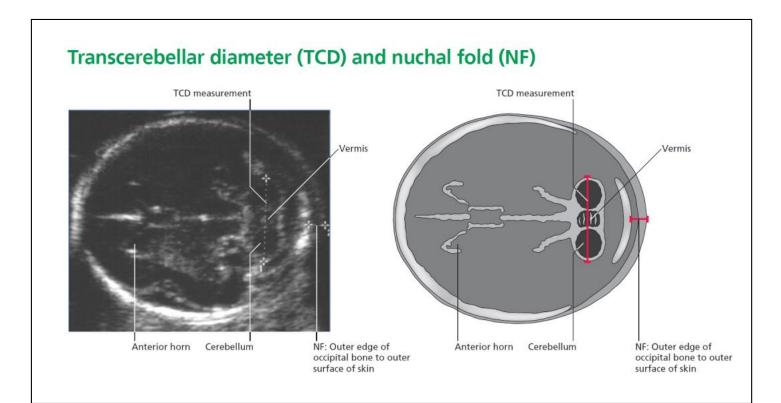




18⁺⁰ - 20⁺⁶ weeks – biometry, TCD

Assess skull, brain & neck

- measure TCD, with image
- assess nuchal fold (<6.0mm)</p>

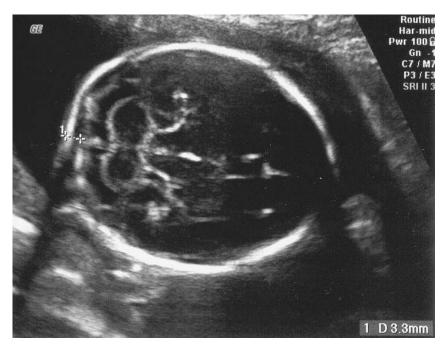


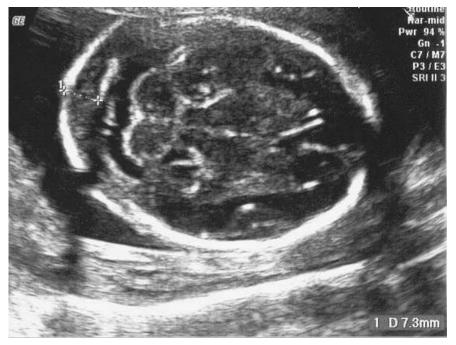
Baterore and the second second

18⁺⁰ - 20⁺⁶ weeks – biometry, nuchal fold

refer if:

- nuchal fold >6.0mm
- ventriculomegaly (PH>10.0mm)
- RPD (AP pelvis >7.0mm)
- significantly small (<< 5th centile)

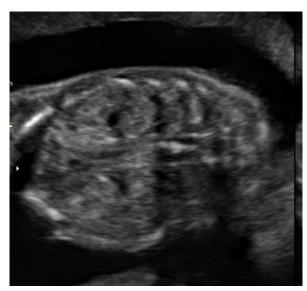




18⁺⁰ - 20⁺⁶ weeks – biometry, renal pelvis

refer if:

- nuchal fold >6.0mm
- ventriculomegaly (PH>10.0mm)
- RPD (AP pelvis >7.0mm)
- significantly small (<< 5th centile)







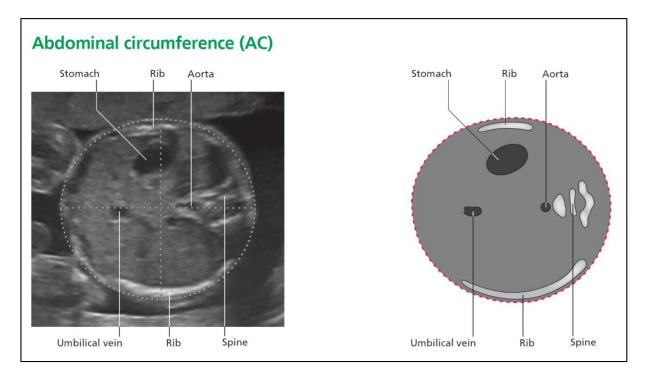


Fetal biometry – abdominal circumference (AC)



Assess fetal growth velocity, based on EDD assigned at dating/combined screening by measurement of:

AC (Chitty) with image

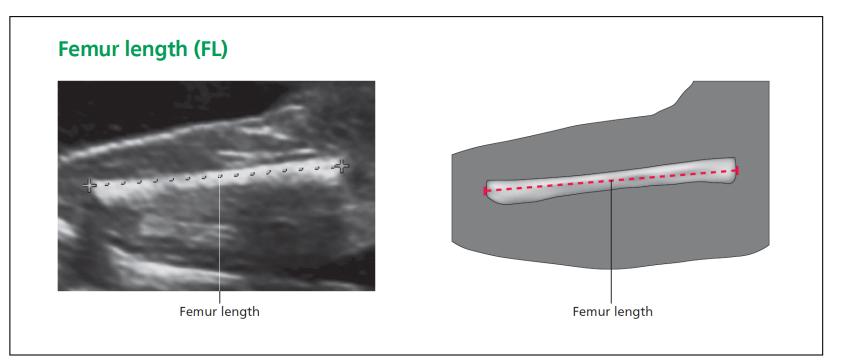


Fetal biometry – femur length (FL)

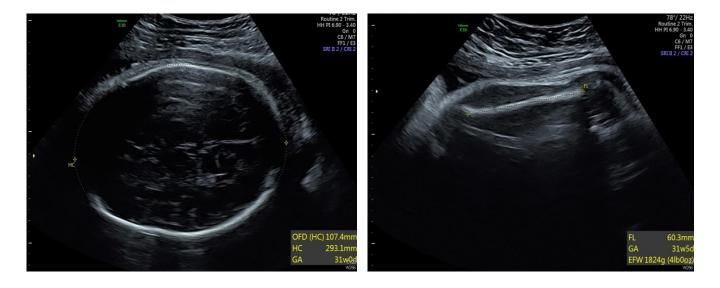


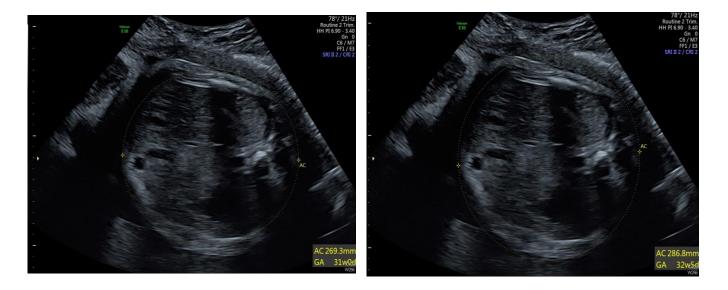
Assess fetal growth velocity, based on EDD assigned at dating/combined screening by measurement of:

FL (Chitty), with image

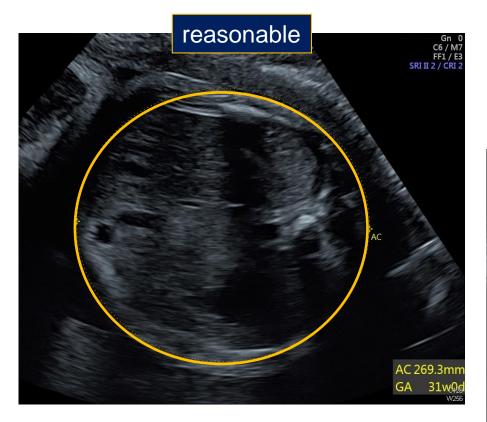


31+ weeks - growth



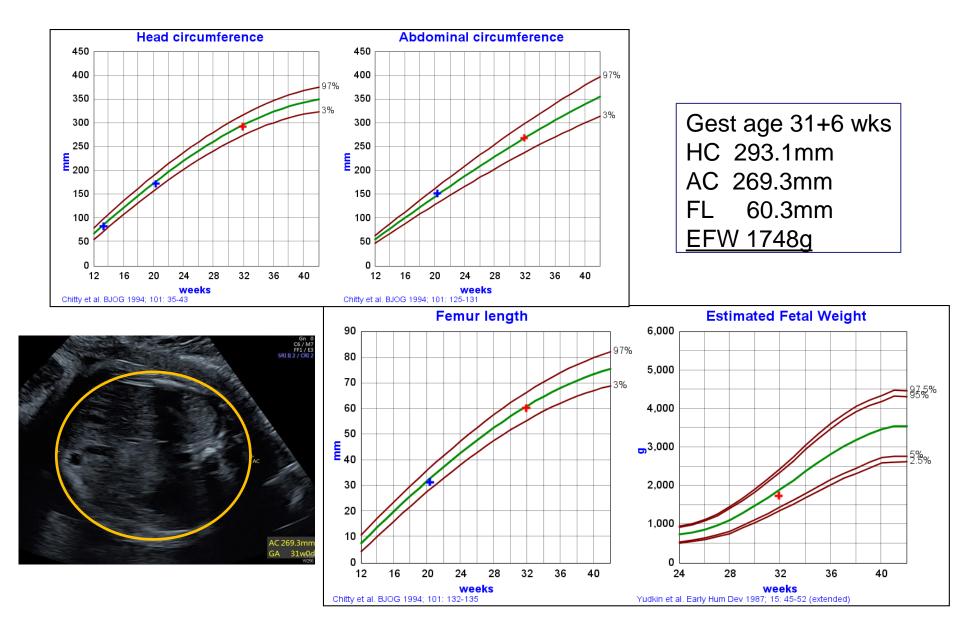


AC measurements

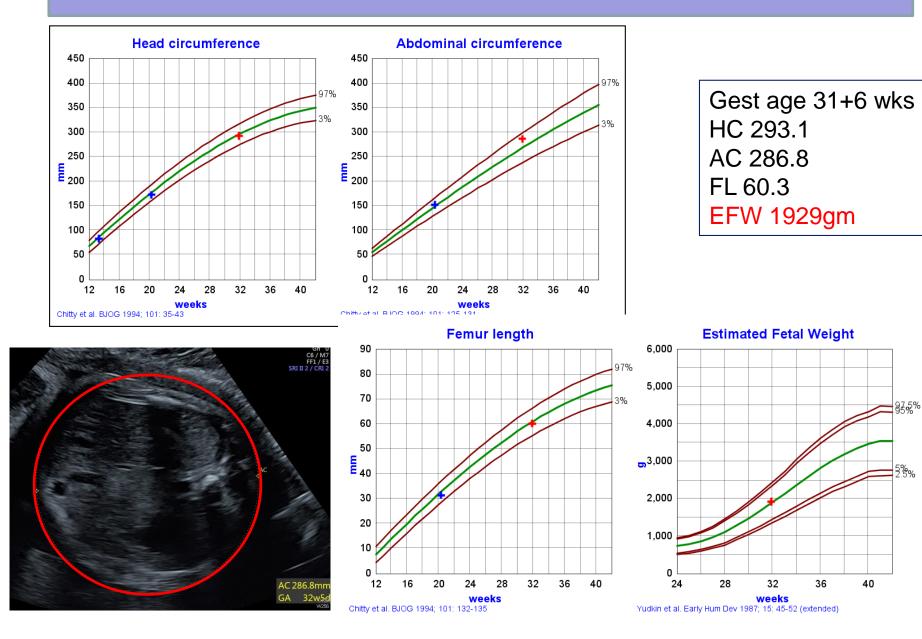




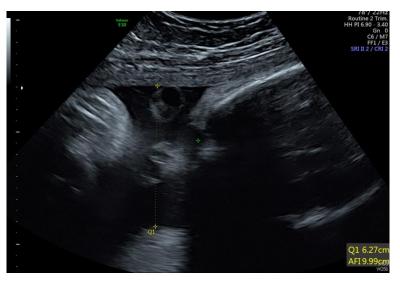
Acceptable measurements including AC



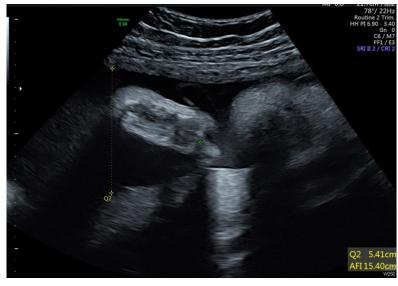
AC overmeasured

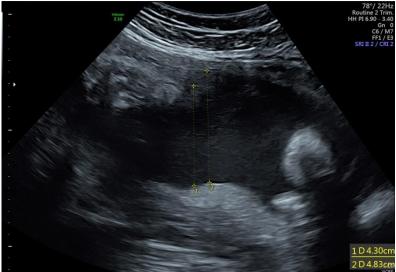


AFI measurements – pool or reverberation



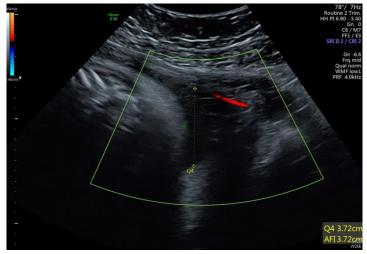




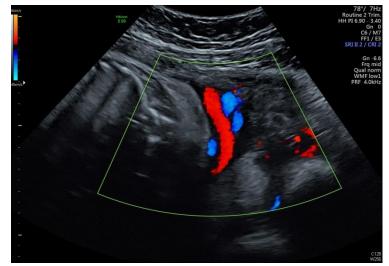


AFI measurements - cord & reverberation









Take home messages



- Accurate fetal biometry is a vital component of every fetal examination
- It is as easy to perform biometry badly as well.
- Selecting the correct section for measurement, & placing the callipers correctly, is as important as other methods of fetal screening
- Be aware of the clinical implications of selecting incorrect sections &/or measuring them incorrectly

Thank you for your attention

