

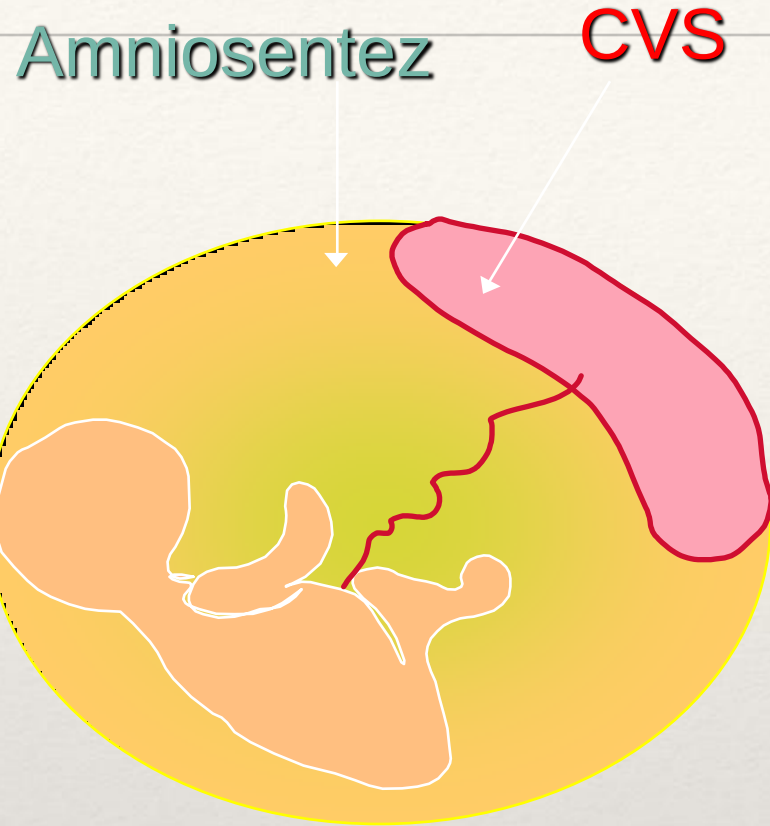


10 Haziran 2016

CVS ve AS Gebelik
Kaybına Yol Açar mı?

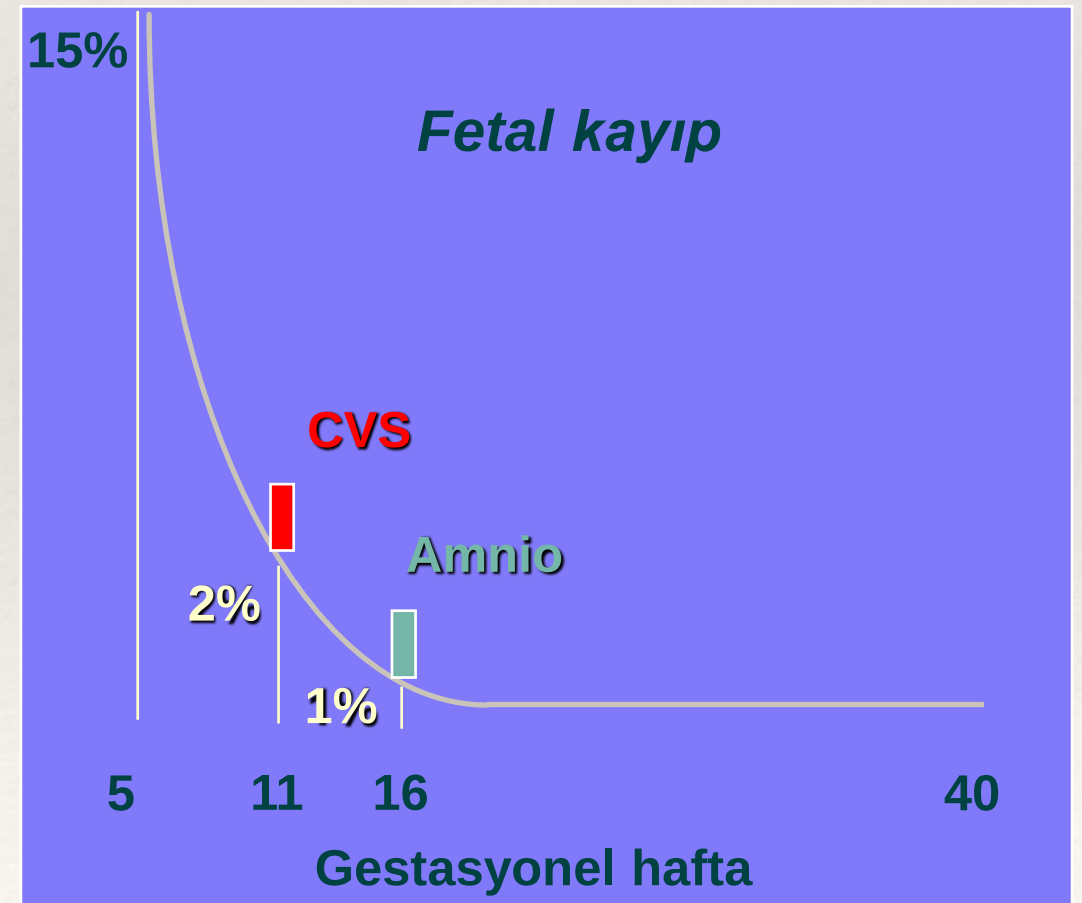
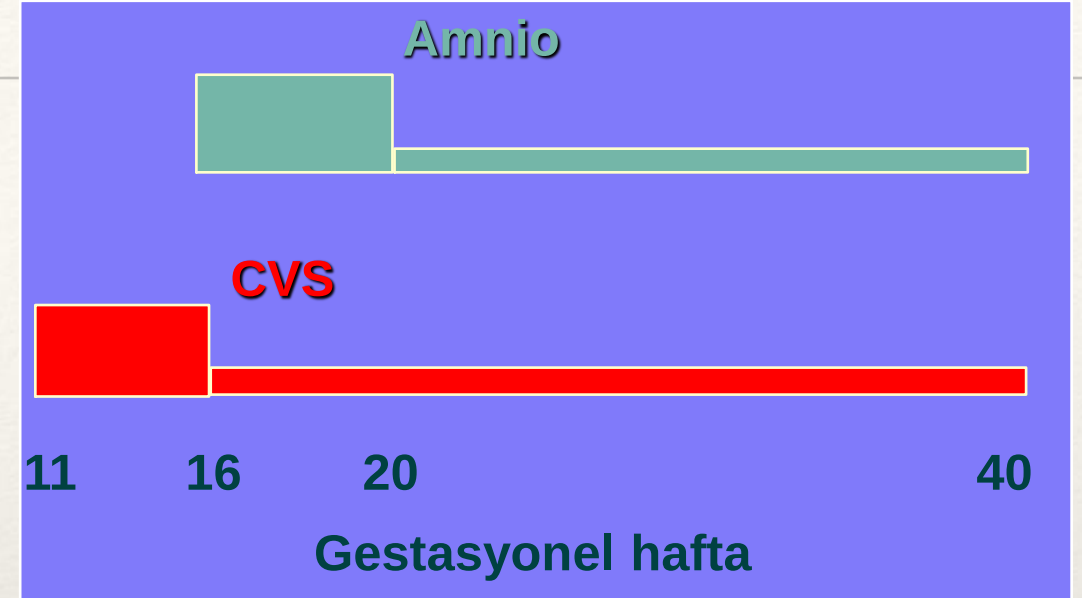
Prof.Dr.
Ali ERGÜN

A/S V CVS



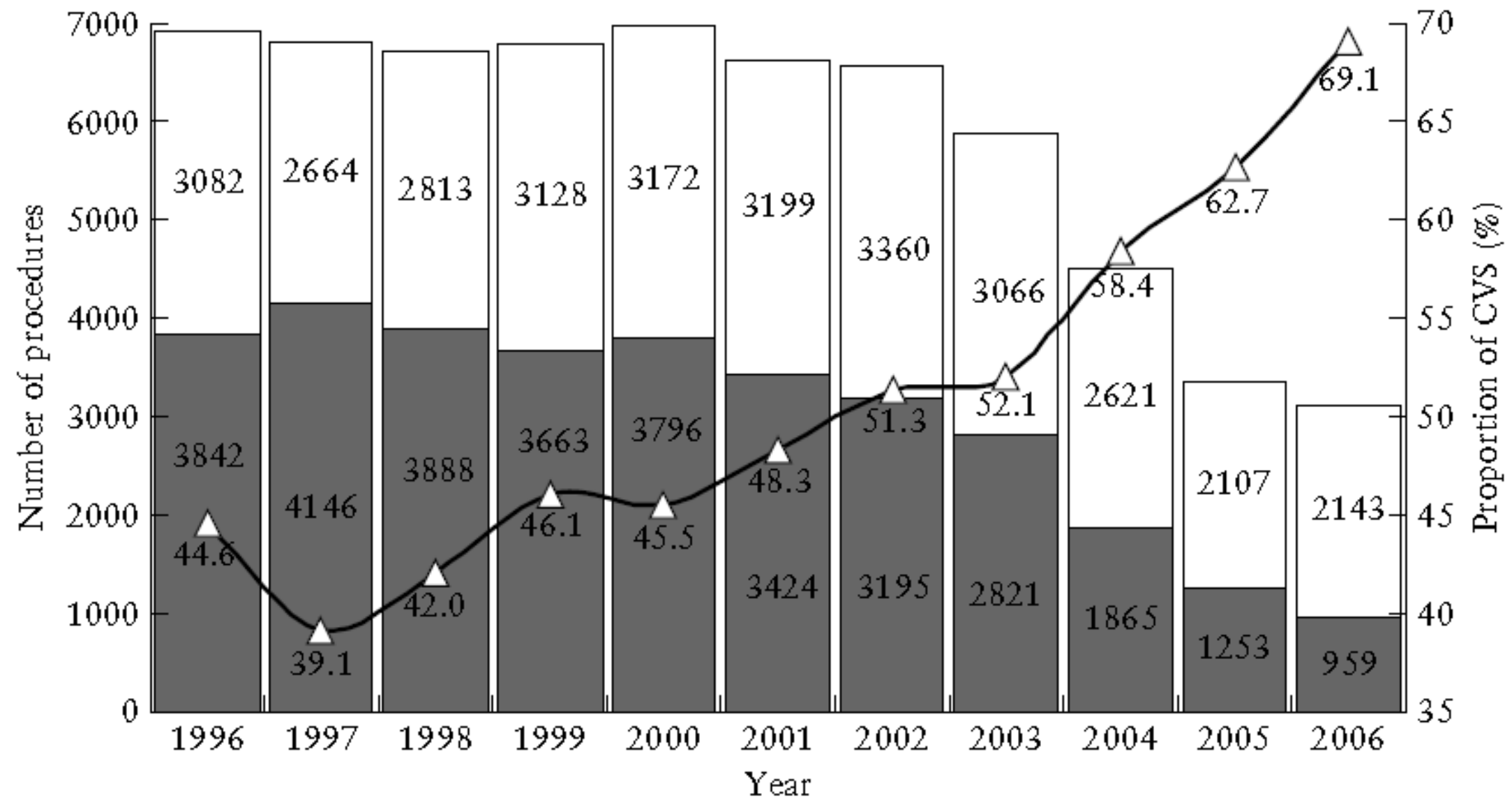
Randomize çalışmalar
Cochrane Library, Issue 4, 2003

2nci trimester amnio – kayıp 1%
TA CVS – amnio ile fark yok
Erken amnio tehlikeli



Prenatal Invaziv Testler

Genel Bilgiler

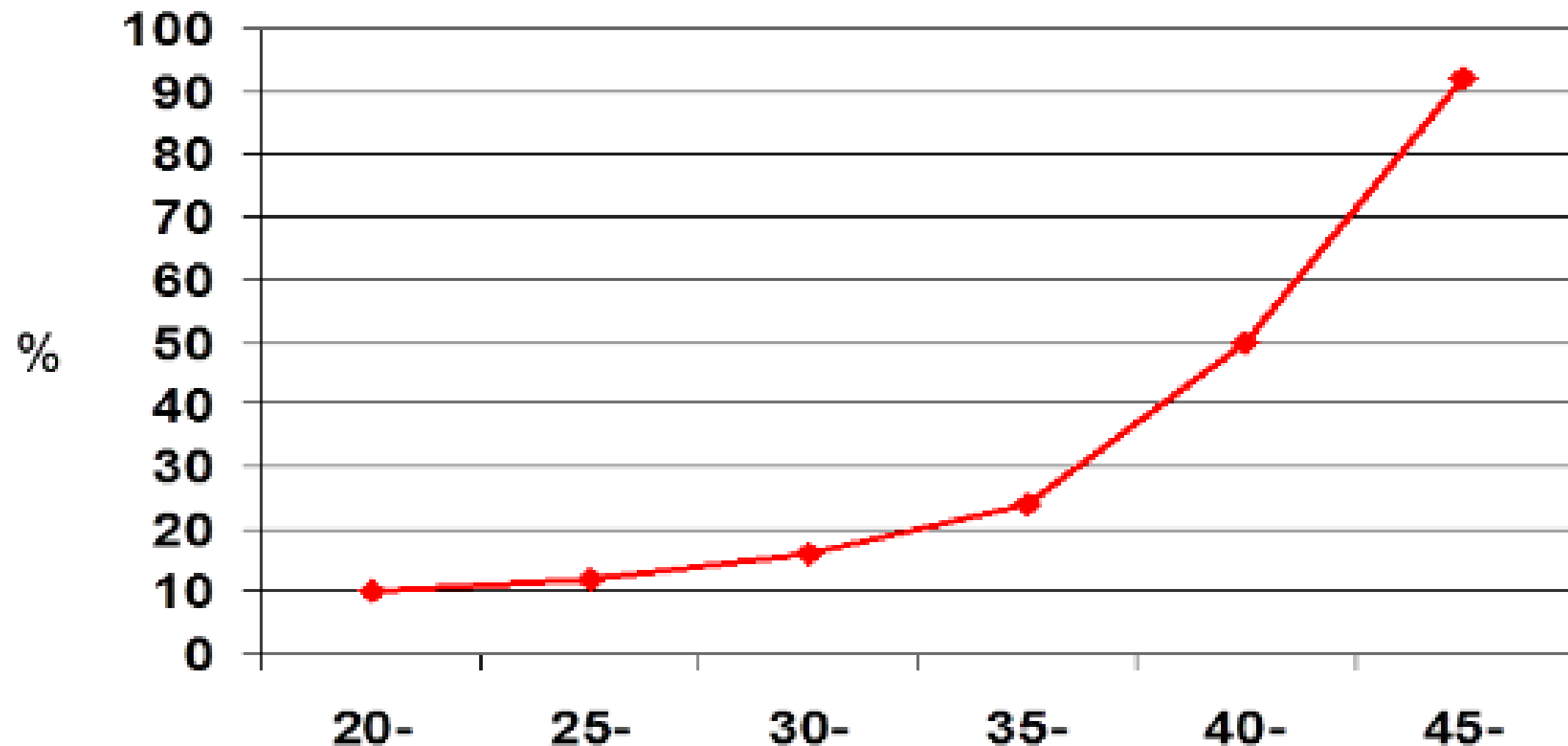


İşleme Bağlı Kayıp Riski

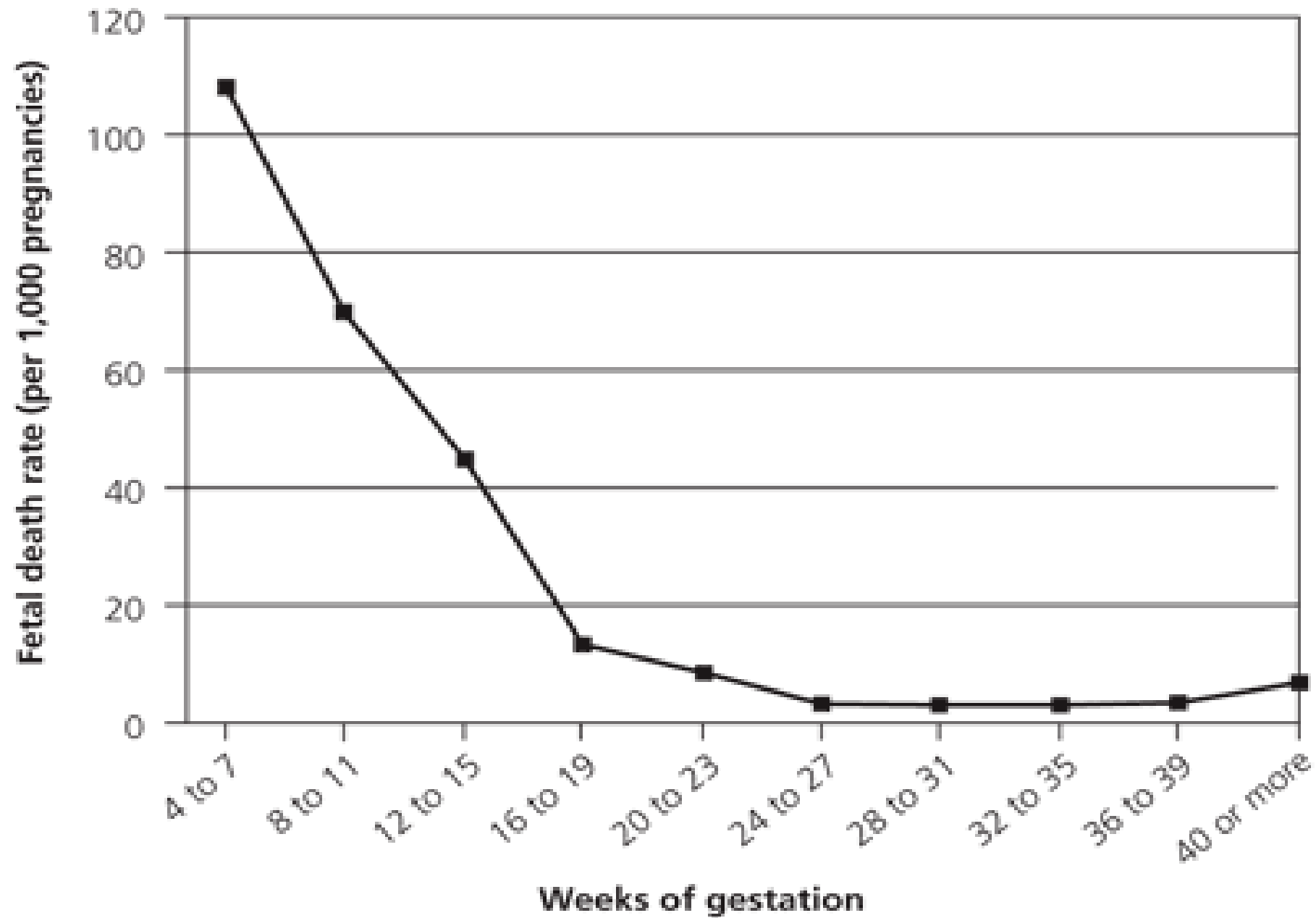
- ❖ Estimated Background Risk
- ❖ İşlem Sonrası Risk

Spontan Kayıp Riski Yaşa Göre

Percentage
of pregnancies
ending in miscarriage

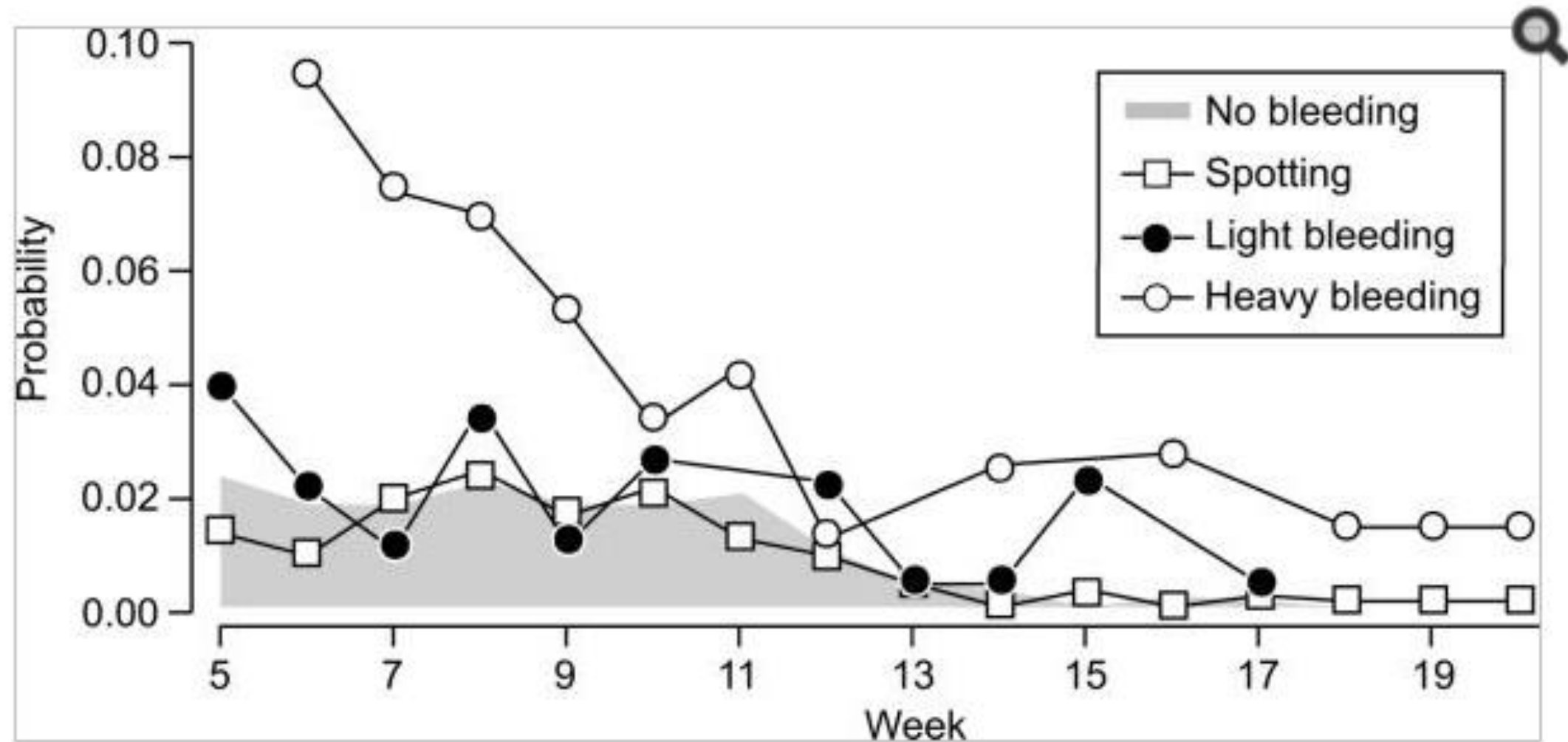


Spontan Kayıp Riski Gebelik Haftasına Göre

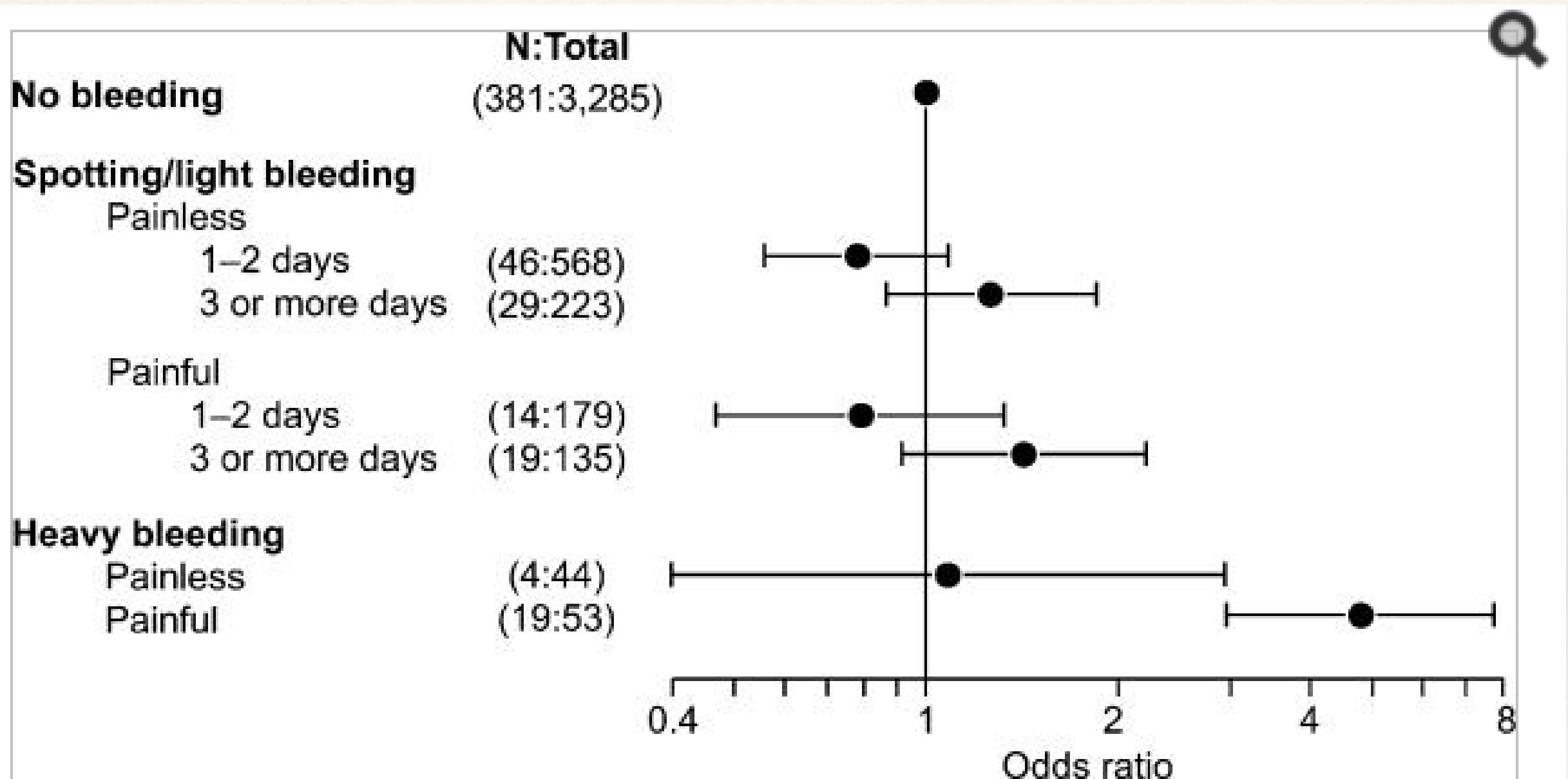


Spontan Kayıp Riski

“Kanama ve Gebelik Haftası”



Spontan Kayıp Riski – Kanama



Fetal Kayıp Riski

Guidelines

- ❖ **UK National Health Service Fetal Anomaly Screening Programme**
 - ❖ Overall kayıp riski
 - ❖ Amniosentez sonrası yaklaşık % 1
 - ❖ CVS sonrası yaklaşık % 1–2%.
- ❖ **Royal College of Obstetricians and Gynaecologists (RCOG)**
 - ❖ Additional Kayıp riski
 - ❖ Amniosentez sonrası yaklaşık % 1
 - ❖ CVS sonrası yaklaşık % 1–2%.

Fetal Kayıp Riski

Guidelines

- ❖ **American College of Obstetricians & Gynecologists**
- ❖ Prosedür ile İlişkili Kayıp Riski
 - ❖ Mid-trimester amniosentez sonrası 1/ 300–500 den daha az.
 - ❖ CVS için oranlar benzer olabilir.
- ❖ **The committee opinion from the Society of Obstetricians and Gynaecologists of Canada**
 - ❖ Amniyosentezi takiben gebelik kayıp riski bireye özgüdür.
 - ❖ Multiple değişkenlere bağlı olarak % 0.19 dan - % 1.53 arasında olabilir.

İnvaziv Test Riski Artırıyor mu?

Ultrasound Obstet Gynecol 2015; 45: 16–26

Published online in Wiley Online Library (wileyonlinelibrary.com). DOI: 10.1002/uog.14636

Procedure-related risk of miscarriage following amniocentesis and chorionic villus sampling: a systematic review and meta-analysis

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2000 – 2014 arası çalışmalar

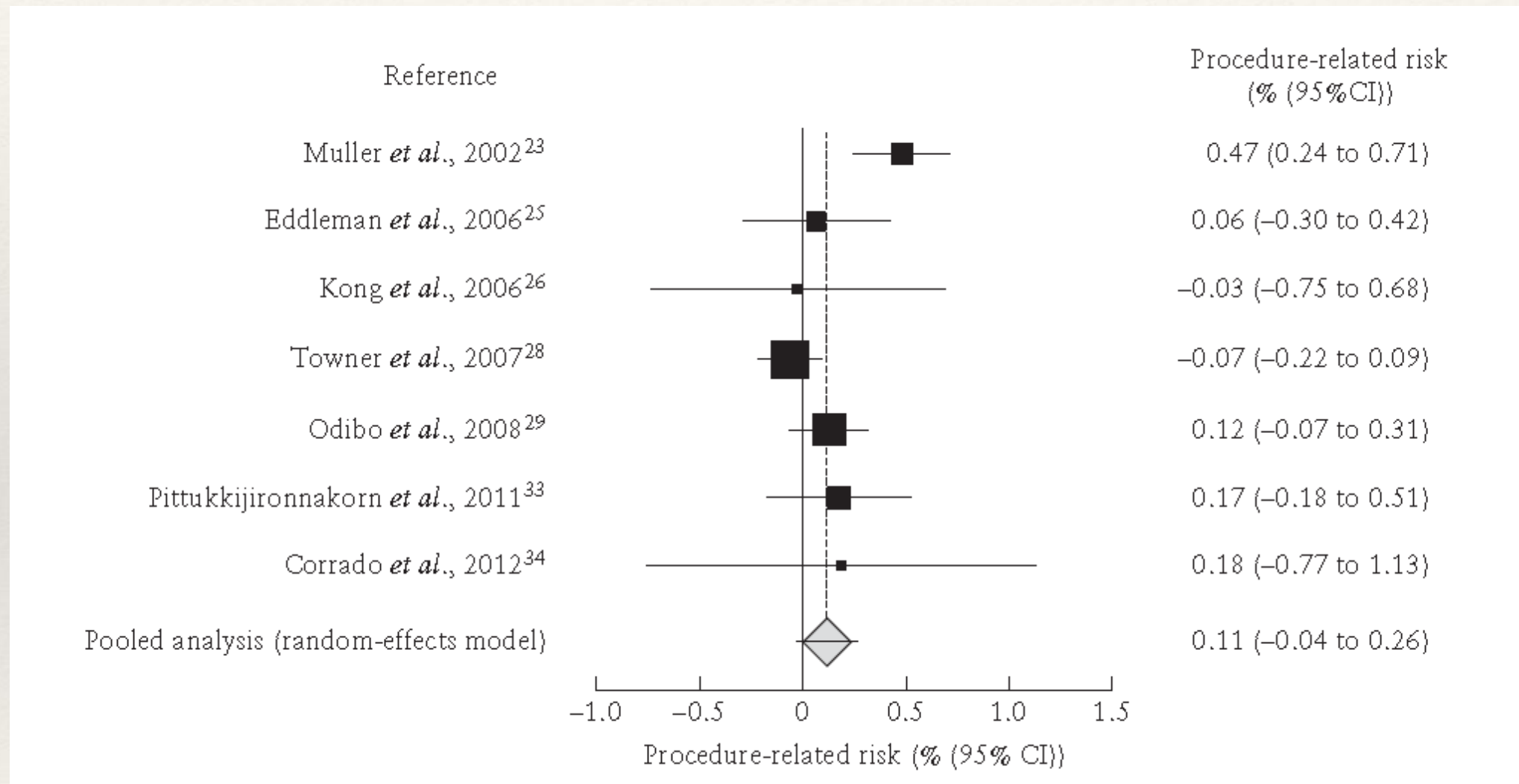
Prosedür ile İlişkili Kayıp Riski

Amniyosentez

Reference	Amniocentesis group		Control group		Procedure-related loss (% (95% CI))	P
	Total (n)	Miscarriage rate (n (%) (95% CI))	Total (n)	Miscarriage rate (n (%) (95% CI))		
Muller, 2002 ²³	3472	31 (0.89 (0.61–1.27))	47 004	197 (0.42 (0.36–0.48))	0.47 (0.24 to 0.71)	0.0003
Eddleman, 2006 ²⁵	3096	31 (1.00 (0.68–1.42))	31 907	300 (0.94 (0.84–1.05))	0.06 (–0.30 to 0.42)	0.6976
Kong, 2006 ²⁶	3468	39 (1.12 (0.80–1.53))	1125	13 (1.16 (0.62–1.97))	–0.03 (–0.75 to 0.68)	0.8727
Towner, 2007 ²⁸	15 005	69 (0.46 (0.36–0.58))	17 045	90 (0.53 (0.42–0.65))	–0.07 (–0.22 to 0.09)	0.4258
Odibo, 2008 ²⁹	11 695	113 (0.97 (0.80–1.16))	39 594	335 (0.85 (0.76–0.94))	0.12 (–0.07 to 0.31)	0.2348
Pitukijronnakorn, 2011 ³³	2990	11 (0.37 (0.18–0.66))	1495	3 (0.20 (0.04–0.59))	0.17 (–0.18 to 0.51)	0.4099
Corrado, 2012 ³⁴	2990	30 (1.00 (0.68–1.43))	487	4 (0.82 (0.22–2.09))	0.18 (–0.77 to 1.13)	0.4720
Pooled analysis (random effects)	42 716	324 (0.81 (0.58–1.08))	138 657	942 (0.67 (0.46–0.91))	0.11 (–0.04 to 0.26)	0.1435
Cochran's Q (P)		46.43 (< 0.0001)		117.15 (< 0.0001)	9.97 (0.1259)	
I ² statistic (% (95% CI))		87.1 (74.5–92.0)		94.9 (92.3–96.3)	39.8 (0.0–73.4)	
Bias (P)		2.9443 (0.3653)		0.2387 (0.9441)	0.6228 (0.6661)	

Prosedür ile İlişkili Kayıp Riski

Amniyosentez



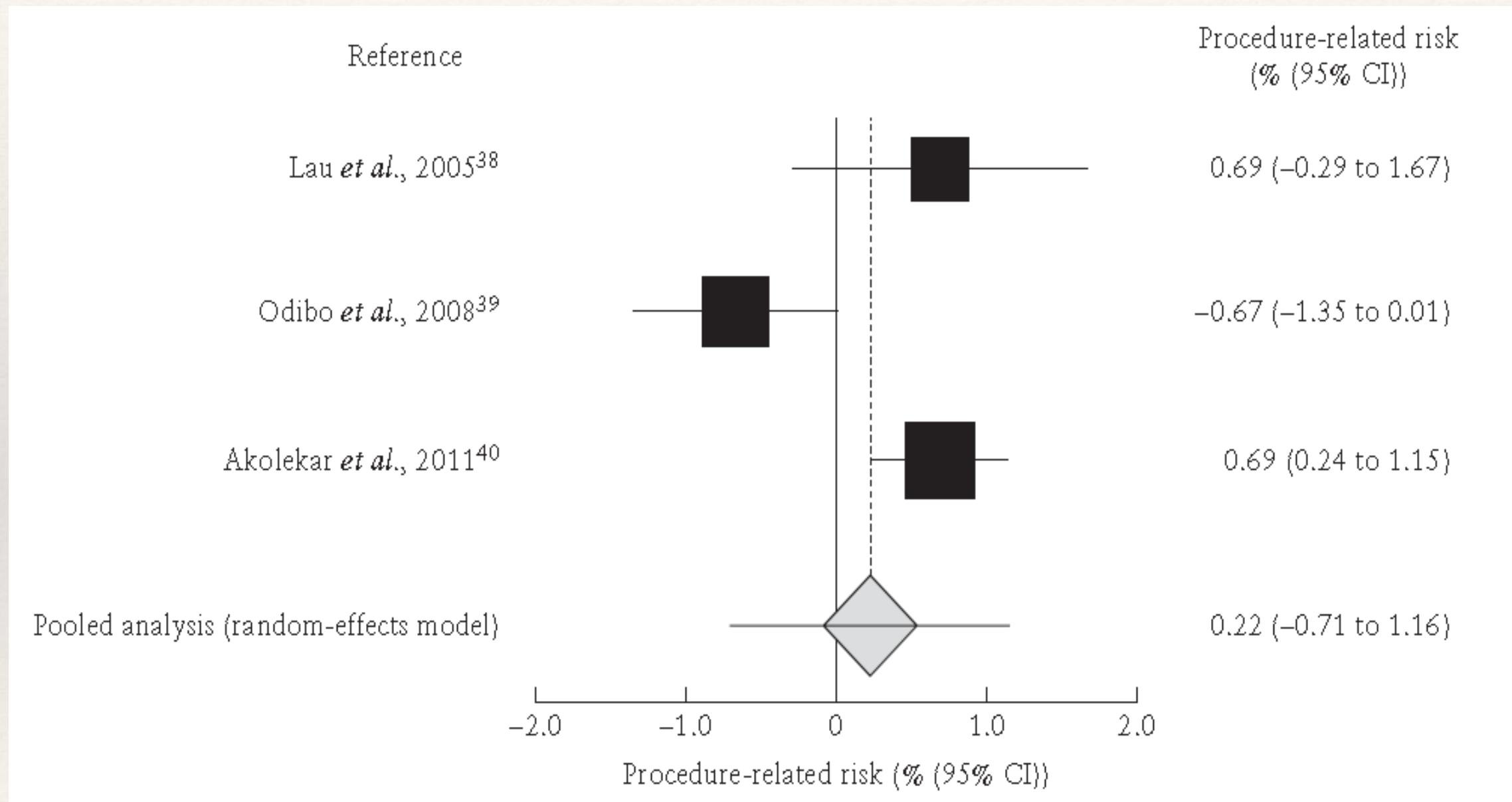
Prosedür ile İlişkili Kayıp Riski

Koriyonik Villus Örneklemesi

Reference	CVS group		Control group		Procedure-related loss (% (95% CI))	P
	Total (n)	Miscarriage rate (n (%) (95% CI))	Total (n)	Miscarriage rate (n (%) (95% CI))		
Lau, 2005 ³⁸	1355	25 (1.85 (1.20–2.71))	1125	13 (1.16 (0.62–1.97))	0.69 (–0.29 to 1.67)	0.2276
Odibo, 2008 ³⁹	5148	138 (2.68 (2.26–3.16))	4803	161 (3.35 (2.86–3.90))	–0.67 (–1.35 to 0.01)	0.0653
Akolekar, 2011 ⁴⁰	2396	44 (1.84 (1.34–2.46))	31460	360 (1.14 (1.03–1.27))	0.69 (0.24 to 1.15)	0.0042
Pooled analysis (random effects)	8899	207 (2.18 (1.61–2.82))	37388	534 (1.79 (0.61–3.58))	0.22 (–0.71 to 1.16)	0.6385
Cochran's Q (P)		6.69 (0.0352)		99.47 (< 0.0001)	10.21 (0.0061)	
I ² statistic (% (95% CI))		70.1 (0.0–89.1)		98.0 (96.9–98.6)	80.4 (0.0–91.9)	
Bias (P)		–5.08 (0.2873)		7.06 (0.6332)	—	

Prosedür ile İlişkili Kayıp Riski

Koriyonik Villus Örneklemesi



Neden Farklı Sonuçlar Var ?

- ❖ **Çalışma dizaynı**
 - ❖ Kontrollü çalışma yetersizliği
- ❖ **Hasta sayısının farklılığı**
 - ❖ (100 – 30.000)
- ❖ **Gebelik kaybının tanımlamasındaki farklılıklar**
 - ❖ 1 hf; 24 hf; 28 hf;
- ❖ **Zaman**
 - ❖ Ekipman, Tecrübe, Teknik.



Diagnostic mid trimester amniocentesis: How safe?

John W. Seeds, MD*

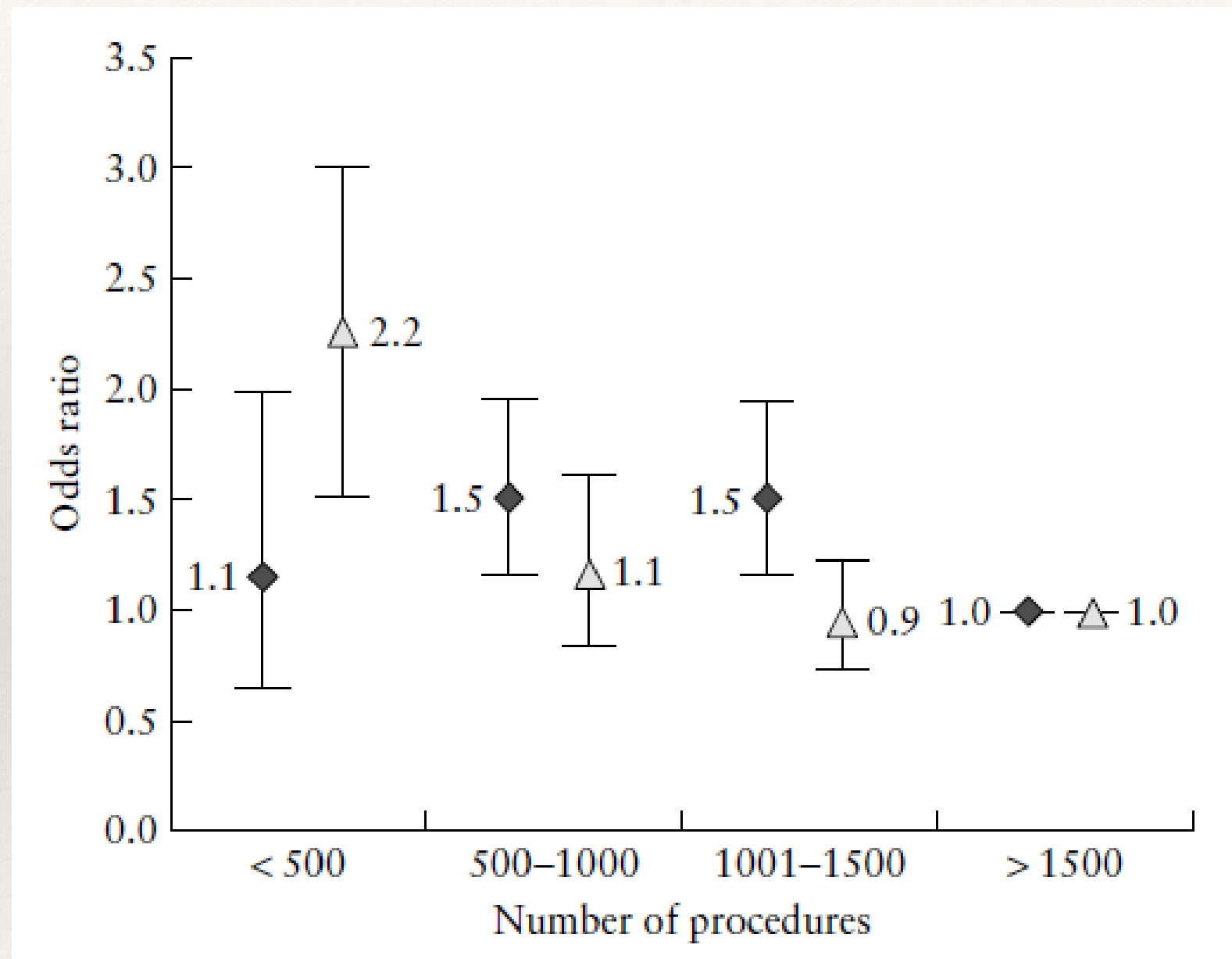
- ❖ 68,119 amniosentez
- ❖ Prosedür ile ilişkili kayıp riski % 0.6
- ❖ Background risk % 1.08
- ❖ Total Kayıp Riski % 1.68
- ❖ USG yardımı yararlı.
- ❖ Transplasental geçiş riski değiştirmiyor.

Neden Farklı Sonuçlar Var ?

- ❖ Sistematik review, Agarwal K, Alfirevic Z, 2012
- ❖ **1990 – 2011 arası**
- ❖ CVS; 9 – 14 hafta
- ❖ A/S; 14 – 22 hafta
- ❖ Randomize çalışmalar olmadığı sürece karar vermek zor
- ❖ CVS ve A/S'de kayıp riski aynı, “background risk + %1”

Fetal Kayıp Riski – Teknik Faktörler

Tecrübe



Fetal Kayıp Riski – Teknik Faktörler

Learning Curve

- ❖ CVS te Learning Curve önemli
- ❖ Prosedür ile ilişkili kayıp riski, 10 hafta üzerinde
 - ❖ Tecrübeli ellerde - Mid-trimester AS ile aynı
 - ❖ Tecrübe az olanlarda – Risk daha yüksek.
- ❖ 13 haftadan önce prenatal tanı isteyen hastalarda en emniyetli prosedür CVS
 - ❖ Erken amniyosentez önerilmemektedir.

Fetal Kayıp Riski – Bireysel Faktörler

Yaş, Kanama, Fetal Kayıp Hikayesi

Table 3. Fetal losses in the three age groups according to the history of bleeding in the current pregnancy or to a history of a previous second trimester miscarriage/termination. Values are given as *n* (%).

Age (years)	Bleeding in the current pregnancy	History of previous fetal loss	Controls
20–34	12/224 (5.4)	4/98 (4.1)	25/1288 (1.9)
35–39	26/410 (6.3)	12/160 (7.5)	66/2280 (2.9)
≥ 40	8/74 (10.8)	8/40 (20)	22/570 (3.9)
Total	46/708 (6.5)	24/298 (8)	113/4024 (2.8)

Fetal Kayıp Riski – Bireysel Faktörler

Yaş, Kanama, Fetal Kayıp Hikayesi

Table 4—Fetal losses for study (amniocentesis) and control groups up to the 28th week, regarding predisposing factors

	Study group (<i>n</i> = 3910)			Control group (<i>n</i> = 5324)			χ^2 <i>p</i> -value
	No.	Losses	%	No.	Losses	%	
Previous bleeding ^a	527	31	5.9	723	28	3.8	<0.04
Previous abortions ^b	258	18	6.9	334	12	3.5	<0.04
No predisposing factors	3125	30	0.96	4267	40	0.93	<0.76
TOPs ^c		214					
Total	3696	79	2.1	5324	80	1.5	<0.06

Fetal Kayıp Riski – Teknik Faktörler

Plaental Lokalizasyon, Transplasental Geçiş

Placental localization	No.	(%)	NoM (%)
Anterior placenta	1012	(48.6)	16 (1.5)
Transpl. needle passage ⁺			
Yes	476	(47.0)	7 (1.5)
No	334	(33.0)	5 (1.5)
Unknown:	202	(20.0)	4 (2.0)
Posterior	928	(44.6)	8 (0.9)
Anterio-posterior	105	(5.0)	3 (2.9)
Fundal placenta	31	(1.5)	1 (3.2)
No registration	7	(0.3)	0

Fetal Kayıp Riski – Teknik Faktörler

İğne İnsersiyon Sayısı

Number of needle insertions	Miscarriages (%)	No. of AMC's
1	22 (1.2)	1807
2	4 (1.8)	228
3	2 (3.8)	53

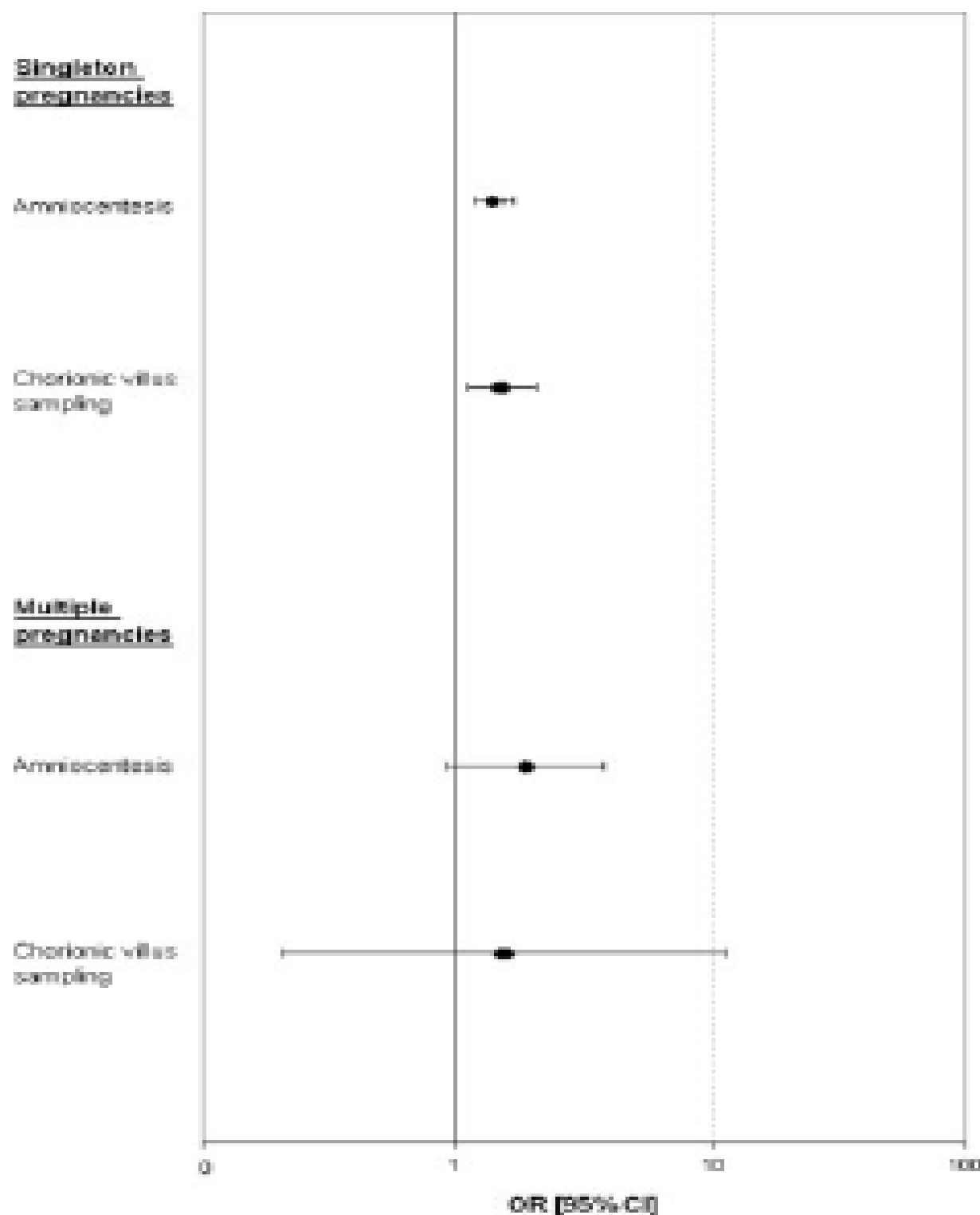
Fetal Kayıp Riski – Bireysel Faktörler

Vücut Kitle İndeksi

Table 4. Fetal Loss by Maternal Body Mass Index Category

	Normal Weight (BMI Lower Than 25)	Overweight (BMI 25.0–29.9)	Class I Obese (BMI 30.0–34.9)	Class II Obese (BMI 35.0–39.9)	Class III Obese (BMI 40.0 or Higher)	<i>p</i>
Amniocentesis (n)	4,777	3,260	1,528	687	527	
All fetal losses	4.2	4.2	4.9	3.8	5.5	.46
Excluding suspected termination	1.3	1.3	2.0	1.2	2.7	.03
CVS (n)	2,755	1,370	553	187	115	
All fetal losses	6.0	6.9	5.8	5.9	10.4	.32
Excluding suspected termination	2.7	3.6	3.3	3.3	4.6	.51

İnvaziv - Riskler



Abruptio Placenta Riski

Tekil Gebeliklerde Risk Yüksek
Çoğul Gebeliklerde Değişmez

	Model	Type of hypertension	OR*	CI (95 %)
AC vs non-exposed	Crude	Gestational HT	0.85	0.74–0.98
		Mild PE	1.06	0.95–1.18
		Severe PE	1.10	0.93–1.30
AC vs non-exposed	Adjusted	Gestational HT	0.77	0.67–0.90
		Mild PE	1.02	0.91–1.14
		Severe PE	0.96	0.80–1.15
CVS vs non-exposed	Crude	Gestational HT	1.20	0.86–1.70
		Mild PE	1.07	0.79–1.45
		Severe PE	1.14	0.72–1.82
CVS vs non-exposed	Adjusted	Gestational HT	1.00	0.70–1.41
		Mild PE	1.07	0.78–1.46
		Severe PE	1.05	0.66–1.69
CVS vs AC	Crude	Gestational HT	1.41	0.99–2.02
		Mild PE	1.01	0.74–1.38
		Severe PE	1.04	0.65–1.67
CVS vs AC	Adjusted	Gestational HT	1.29	0.90–1.84
		Mild PT	1.05	0.77–1.44
		Severe PE	1.10	0.68–1.77

❖ AS ve CVS

❖ **PIH riskini artırmaz.**

Outcome	Non-exposed (N = 47,854)		Amniocentesis (N = 21,748)					Chorionic villus sampling (N = 1984)				
	n	n/1000	n	n/1000	OR		CI adjusted	n	n/1000	OR		CI adjusted
					Crude	Adjusted				Crude	Adjusted	
Normal delivery	26,890	561.9	10,792	496.2	0.77	0.93	0.90–0.97	1085	546.9	0.94	1.06	0.96–1.16
Bleeding in late gestation	1703	35.6	852	39.2	1.10	0.97	0.88–1.06	72	36.3	1.02	0.94	0.73–1.20
Placenta praevia	317	6.6	138	6.3	0.96	0.82	0.65–1.02	11	5.5	0.84	0.70	0.38–1.30
Placental abruption	367	7.7	155	7.1	0.93	0.92	0.74–1.13	14	7.1	0.92	0.88	0.50–1.52
Unspecified bleeding	1096	22.9	597	27.5	1.20	1.04	0.93–1.16	50	25.2	1.10	1.03	0.77–1.39
Postpartum bleeding	4251	88.8	2069	95.1	1.08	0.96	0.90–1.02	161	81.1	0.91	0.88	0.74–1.04
Retained placenta	343	7.2	181	8.3	1.16	1.04	0.85–1.28	13	7.6	1.06	1.27	0.74–2.18
Amnion-related complications	2434	50.9	1382	63.5	1.27	1.15	1.06–1.24	101	50.9	1.00	0.88	0.71–1.09
Oligohydramnios	379	7.9	250	11.5	1.46	1.09	0.91–1.30	9	4.5	0.57	0.44	0.23–0.87
Premature rupture of membranes	583	12.2	302	13.9	1.14	1.13	0.96–1.32	17	8.6	0.70	0.63	0.39–1.04
Delayed delivery	1291	27.0	731	33.6	1.26	1.14	1.03–1.26	61	30.7	1.14	1.10	0.84–1.45
Chorioamnionitis	144	3.0	81	3.7	1.24	1.30	0.96–1.77	10	5.0	1.68	1.60	0.81–3.13
Fever or sepsis in labour	199	4.2	134	6.2	1.48	1.19	0.93–1.51	7	3.5	0.85	0.92	0.42–2.00
Hypotonic uterine dysfunction	5127	107.1	3049	140.2	1.36	1.12	1.06–1.18	198	99.8	0.92	1.10	0.94–1.30
Hypertonic uterine dysfunction	78	1.6	37	1.7	1.04	0.94	0.61–1.46	5	2.5	1.55	1.54	0.60–3.98
Protracted labour	475	9.9	179	8.2	0.83	0.84	0.70–1.02	21	10.6	1.07	1.10	0.69–1.76
Instrumental vaginal delivery	2789	58.3	1607	73.9	1.29	1.11	1.03–1.19	120	60.5	1.04	1.11	0.91–1.36
Elective caesarean	3902	81.5	2204	101.3	1.27	1.09	1.02–1.16	202	101.8	1.28	1.02	0.86–1.21
Emergency caesarean	3789	79.2	1898	87.3	1.11	0.93	0.87–0.99	167	84.2	1.07	0.97	0.82–1.16

35–49 yaş arası

- ❖ AS önemli advers sonuçlar yaratmaz.
- ❖ Ancak, 15. hafta öncesi uygulanırsa minör risk artışı mevcut.

Fetal Kayıp Riski – Bireysel Faktörler

Kanama, Sigara, Abortion Hikayesi, VKİ, Yaş

	n = patients (%)	n = procedure-related fetal losses	significance p<0.05
vaginal bleeding (first trimester)	511 (6.5%)	6 (0.096%)	p= 0.008
maternal smoking > 10 cigarettes/d	439 (5.6%)	3 (0.05%)	p= 0.471
history of ≥ 3 spontaneous abortions	163 (2.1%)	2 (0.03%)	p= 0.119
BMI > 30	748 (9.6%)	2 (0.03%)	p= 0.449
maternal age > 40	832 (10.7%)	8 (0.11%)	p= 0.081

Fetal Kayıp Riski – Bireysel Faktörler

Yaş , Kanama, Abortion Hikayesi, VKİ, Myom, Boyalı AS

	OR (95% CI)	<i>p</i> value
Age (years)		
<34	1.0*	
≥34	2.19 (1.00–4.77)	0.049
Hemorrhage during pregnancy		
No	1.0	
Minor (spotting)	1.96 (1.14–3.38)	0.015
Serious	2.74 (1.23–6.11)	0.014
Number of surgical abortions		
None	1.0	
1–2	1.32 (0.8–2.18)	0.280
≥3	3.44 (1.32–8.94)	0.011
Number of miscarriages in 1st trimester		
None	1.0	
1–2	1.25 (0.75–2.06)	0.391
≥3	1.93 (1.00–3.73)	0.050
Uterine fibroid		
No	1.0	
Yes	2.52 (1.01–6.40)	0.048
Stained amniotic fluid		
No	1.0	
Yes	3.74 (1.05–13.29)	0.041

Amniyosentez Sonrası Kayıp Riski Ne Kadar Devam Eder ?

Weeks after amniocentesis	Fetal loss	
	No.	%
0	2	5.3
1	2	5.3
2	3	7.9
3	4	10.5
4	7	18.4
5	5	13.5
6	4	10.5
7	2	5.3
8	2	5.3
9	2	5.3
10	2	5.3
13	2	5.3
20	1	2.6
23	1	2.6
24	1	2.6
Total	40	100

Amniyosentez Sonrası Kayıp Riski Ne Kadar Devam Eder ?

Table 2. Fetal losses according to time intervals in the three age groups. Values are given as *n* (%).

Time interval	Age 20-34 (<i>n</i> = 1610)	Age 35-39 (<i>n</i> = 2850)	Age ≥ 40 (<i>n</i> = 570)	<i>P</i>
0-15 days	15 (0.9)	20 (0.7)	4 (0.7)	0.687
15 th day-28 weeks	16 (1)	46 (1.6)	15 (2.6)	0.014
28 weeks-term	10 (0.6)	32 (1.1)	10 (1.8)	0.016
Total	41 (2.54)	98 (3.44)	29 (5.1)	0.003

İkiz Gebeliklerde Fetal Kayıp Riski

- ❖ Birçok yayınlanan Guideline Bu konu üzerinde durmamıştır.
- ❖ **The Royal College of Obstetricians and Gynaecologists' guideline**
 - ❖ CVS ve Amniyosentez sonrası kayıp riski tekil gebeliklere göre daha yüksektir.
- ❖ **The Society of Obstetricians and Gynaecologists of Canada guideline**
 - ❖ İkiz gebeliklerde Amniyosentez sonrası 24-28 nci gebelik haftalarından önce kayıp riski % 1 – 4 arasındadır.
- ❖ **The American College of Obstetricians and Gynecologists**
- ❖ **The Royal Australian and New Zealand College of Obstetricians and Gynaecologists guideline**
 - ❖ İkiz gebeliklerde kayıp riskini belirtmemiştir.

Second-trimester amniocentesis vs. chorionic villus sampling for prenatal diagnosis in multiple gestations

A. ANTSAKLIS*, A. P. SOUKA*, G. DASKALAKIS*, Y. KAVALAKIS* and S. MICHALAS*

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Outcome of twin pregnancies that underwent selective feticide

	<i>Amniocentesis</i>	<i>Chorionic villus sampling</i>	<i>P</i>
Pregnancies	9	12	
Miscarriage	0	1/12 (8.3%)	NS
Intrauterine death	0	0	
Delivery ≤ 32 weeks	2/9 (22.2%)	2/11 (18.2%)	NS
Delivery ≤ 35 weeks	2/9 (22.2%)	2/11 (18.2%)	NS
Perinatal mortality ≤ 28 weeks	0	0	NS
Perinatal mortality > 28 weeks	1/9 (11.1%)	0	NS
Neonatal mortality	1/9 (11.1%)	0	NS
Total loss	1/9 (11.1%)	1/12 (8.3%)	NS

İkiz Gebeliklerde Fetal Kayıp Riski

Koriyonik Villus Örneklemesi

Study or subgroup	Transabdominal approach		Transcervical approach		Weight	Risk ratio		Year	Risk ratio	
	Events	Total	Events	Total		M-H, Random, 95% CI			M-H, Random, 95% CI	
De Catte (1996) ¹²	6	48	3	72	60.8%	3.00 (0.79, 11.42)		1996		
De Catte (2000) ¹⁰	4	93	2	55	39.2%	1.18 (0.22, 6.25)		2000		
Total (95% CI)		141		127	100.0%	2.08 (0.73, 5.91)				
Total events	10		5							
Heterogeneity: $\tau^2 = 0.00$; $\chi^2 = 0.73$, $df = 1$ ($P = 0.39$); $I^2 = 0\%$										
Test for overall effect: $Z = 1.38$ ($P = 0.17$)										

0.01 0.1 1 10 100

Favors experimental Favors control

İkiz Gebeliklerde Fetal Kayıp Riski

Amniyosentez

Study or subgroup	Experimental		Control		Weight	Risk ratio		Year	Risk ratio	
	Events	Total	Events	Total		M-H, Random, 95% CI			M-H, Random, 95% CI	
Ghidini (1993) ²⁹	2	101	2	108	8.5%	1.07 (0.15, 7.45)		1993		
Ko (1998) ²⁶	0	128	2	205	3.5%	0.32 (0.02, 6.60)		1998		

İkiz Gebeliklerde AS ve CVS sonrası kayıp riski ile ilgili
RCT ler olmadan karar vermek zor
Background risk + % 1

Test for overall effect: $Z = 2.04$ ($P = 0.04$)

0.01 0.1 1 10 100
Favors experimental Favours control

Figure 7 Pregnancy loss before 24 weeks after amniocentesis. M-H, Mantel-Hanzel.

Komplikasyon riskini Azaltacak Önlem var mı?

Progesteron

Pregnancy complications after sampling^a

	Progesterone (311)	Controls (273)
Miscarriages	4 (1.3%)	3 (1.1%)
Premature rupture of membranes	19 (6.1%)	17 (6.2%)
Premature delivery (<37 weeks)	27 (8.7%)	20 (7.3%)
Intrauterine fetal death	2 (0.6%)	3 (1.1%)

^a $P > 0.05$.

Corrado F et al, European Journal of Obstetrics & Gynecology and
Reproductive Biology 2002.

“Impact of a new national screening policy for Down’s syndrome in Denmark; “population based cohort study”

- ❖ Birinci trimestir tarama (2004-6), sonuçlar (2005-6)
- ❖ 19 merkez, yılda 65 000 gebelik
- ❖ Sonuçlar;
 - Down send. Doğum; 2000-4 *yılda 55-65*, 2005’te *31*, 2006’da *32*
 - Toplam CVS ve AS sayısı; 2000’de *7524*, 2006’da *3510*
 - DR; 2005’te %86, 2006’da %93 ve *FP sırası ile %3.9 ve %3.3*

Ekelund CK, Tabor A Nov 2008, BMJ

Ultrasound Obstet Gynecol 2009; 34: 19–24

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Fetal loss rate after chorionic villus sampling and amniocentesis: an 11-year national registry study

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Risk

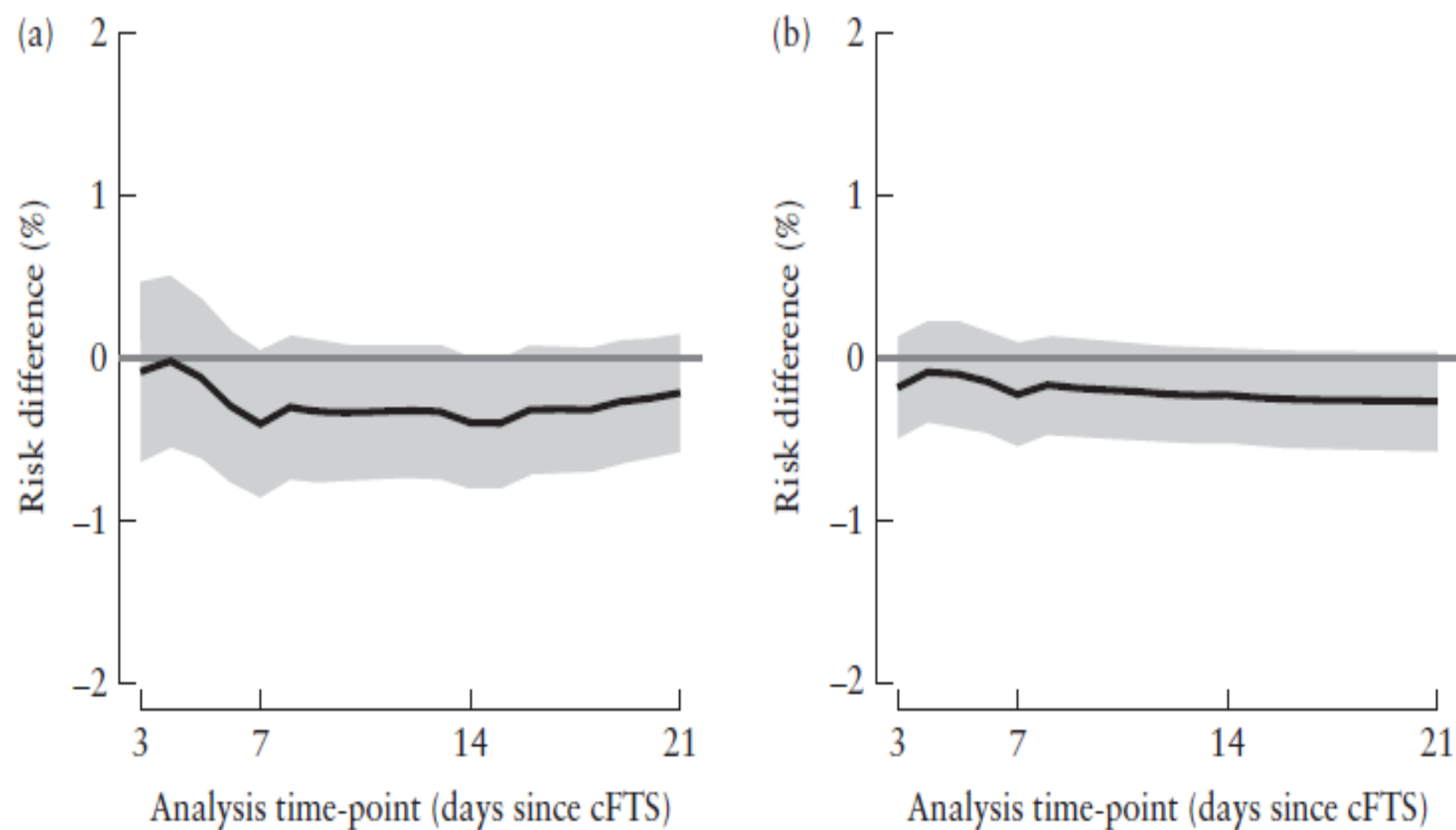


Figure 2 Effect of chorionic villus sampling (CVS) on risk of miscarriage (a) and risk of stillbirth (b) from analysis time-points 3–21 days after combined first-trimester screening (cFTS), showing average risk differences across propensity score strata.

Risk

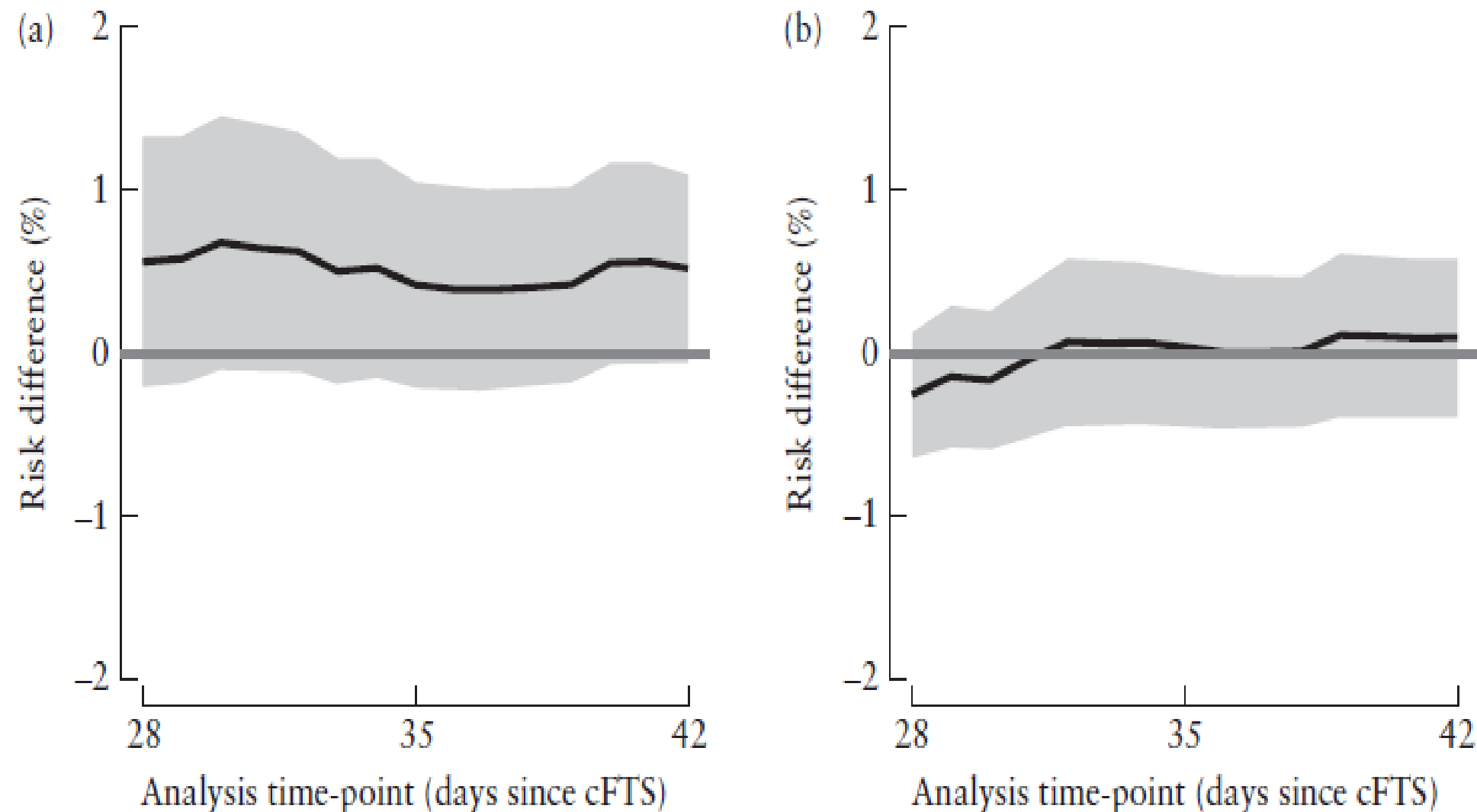
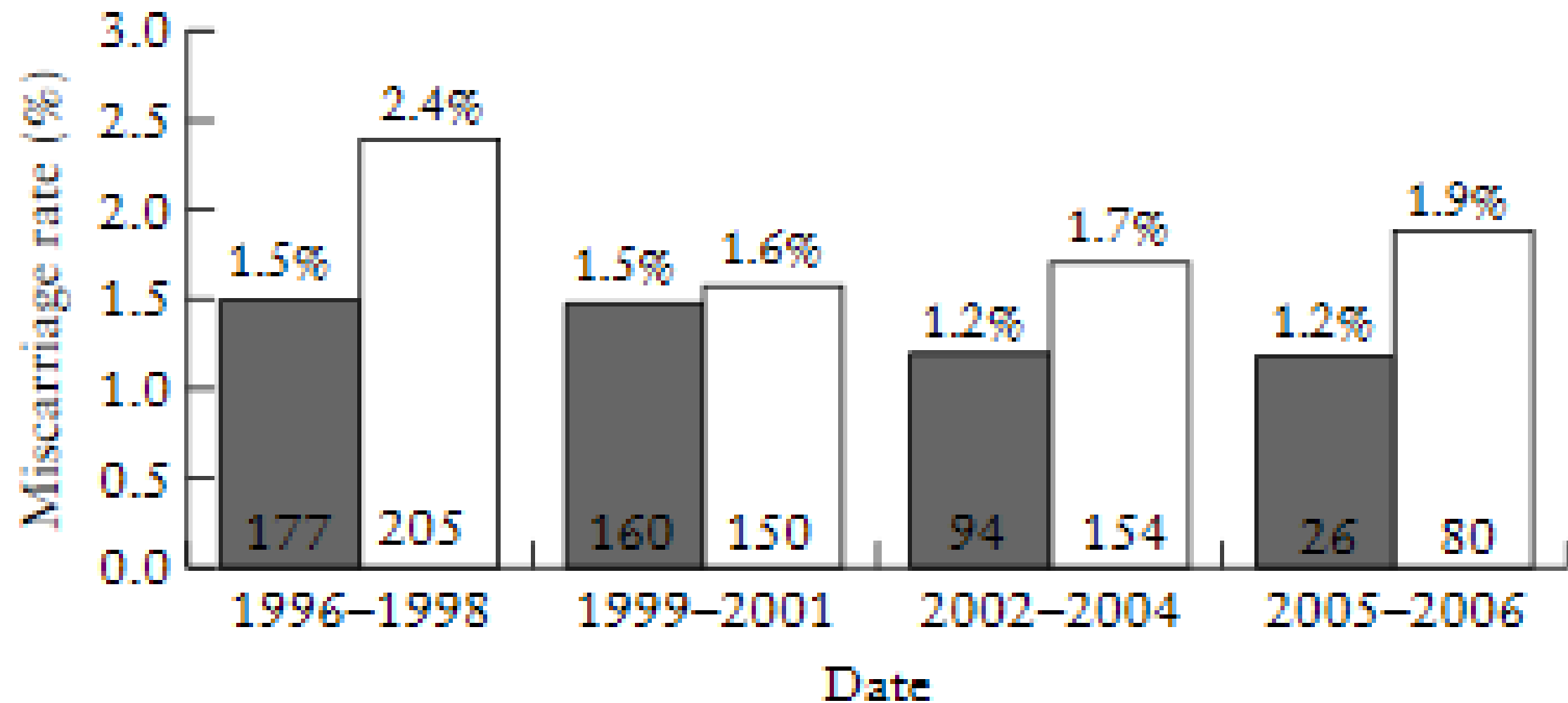


Figure 3 Effect of amniocentesis (AC) on risk of miscarriage (a) and risk of stillbirth (b) from analysis time-points 28–42 days after combined first-trimester screening (cFTS), showing average risk differences across propensity score strata.

Yıllara Göre Kayıp Riski



Maternal Yaşı Göre Risk

Table 4 Postprocedural miscarriage rate per 100 pregnancies following amniocentesis (AC) and transabdominal chorionic villus sampling (CVS) in four time periods, according to maternal age

<i>Time period</i>	<i>Maternal age (years)</i>					
	<i>< 30</i>		<i>30–34</i>		<i>> 34</i>	
	<i>AC</i>	<i>CVS</i>	<i>AC</i>	<i>CVS</i>	<i>AC</i>	<i>CVS</i>
1996–1998	1.1	1.2	1.4	2.2	1.7	2.6
1999–2001	1.3	1.4	1.4	1.5	1.6	1.6
2002–2004	2.2	1.3	1.3	1.0	1.0	1.9
2005–2006	2.3	2.3	0.8	2.0	0.9	1.6
Overall	1.5	1.5	1.3	1.7	1.4	2.0

Kromozomal defektlerin tanısı

16. Hafta Amnio Vs CVS at 9-13 hafta

Randomize Çalışmalar

Çalışma	N	merkez	Fetal kayıp (%)	
			CVS	Amnio
Kanada 1990	2391	11	7.6	7.1
Avrupa 1991	3201	31	13.6	9.0
Danimarka 1992	2069	2	6.3	6.4
Finlandiya 1993	800	1	7.8	8.3
İspanyol 1999	1313	1	2.2	2.8

CVS'te işleme bağlı kayıp oranı, amnio ile aynı (%1)

Sonuçlar

- ❖ Prenatal Tarama ve Tanı stratejilerindeki Gelişim son dekatta prenatal invaziv test uygulamalarını birinci trimestire ve CVS'e doğru kaymasına neden olmuştur.
- ❖ Prenatal İnvaziv Test Kayıp Riski Bireysel olarak farklılıklar göstermektedir.
- ❖ Genel olarak 1/200 olan kayıp riski teknik ve ultrasonografinin kullanımı ile 1/600'lere inmesine rağmen Guideline'lar halen 1/200 oranını söylemektedir.
- ❖ Progesteron kullanımı, İnvaziv prosedür sonrası kayıp riskini azaltmamaktadır.

Sonuçlar

- ❖ Cihaz kalitesindeki iyileşme
- ❖ Deneyim, işlem sayısı
- ❖ İşlem için doğru zamanlama, erken amnio, CVS!
- ❖ Doğru vaka seçimi
 - kanama(uzun süreli, hematoma, ileri yaş)
 - kontrakte uterus, kısa serviks
 - zor vaka(plasental lokasyon, fetal pozisyon, hareketlilik, CVS'te posterior plasenta, anne uyumu)

Teşekkürler