# Abstracts

## **Cesarean Section Short- and Long-term Medical Implications**

#### **Apostolos Papageorgiou**

Department of Pediatrics, Obstetrics and Gynecology, Jewish General Hospital, McGill University, Montreal, Canada

The increasing rates of cesarean sections (C/section) are presently of intense medical interest and of public health importance. We are witnessing an extraordinary variation in the number of C/sections not only among countries but also among institutions. They range from 14 to over 65%.

This large discrepancy cannot be explained only on the basis of population differences. Of particular concern are elective C/sections before 39 weeks of gestation.<sup>1</sup>

Many studies indicate a high incidence of neonatal pathologies requiring several days of hospitalization in neonatal intensive care units.<sup>2</sup>

The most important immediate complications are of respiratory nature, including respiratory distress syndrome (RDS), transient tachypnea of newborn (TTN), need for ventilation, and prolonged oxygen requirements. Parental separation and delayed breastfeeding are further complications (Table 1).

Cesarean sections lead also to more frequent long-term risks. They include higher incidence of asthma, food allergies, diabetes, and neurodevelopmental difficulties.

A Norway study indicates a 52% increased risk for asthma following C/section *vs* spontaneous vaginal delivery (hazard ratio = 1.5).<sup>3</sup> The advanced reason is based on the "hygiene hypothesis," which assumes that the initial gut colonization takes place with the "wrong" microbes. The other reason relates to secondary complications following RDS, TTN, and ventilation.

There is also evidence of increased food allergies due to increased sensitization to antigen-specific immunoglobulin E following C/sections.<sup>4</sup>

Finally, neurodevelopment in infants born between 37 and 39 weeks is reported to be affected by an odds ratio of 1.16 (95% confidence interval, 1.12-1.20).<sup>5</sup>

It is also important to mention that high C/section rates do not guarantee low maternal and neonatal mortality, as proven by data from Latin America, where high C/section rates are the norm.

In conclusion, C/sections are on the rise and not always justified. Judicious C/sections undoubtedly have improved perinatal mortality and morbidity. It is now well-established that elective C/sections before 39 weeks of gestation carry unacceptable neonatal risks and the cost of hospitalizations is not negligible. Parents requesting C/ sections should be made aware of the risks and benefits to the fetus and newborn.

			(10,200)	
Outcome	37 (n=834)	38 (n=3,909)	39 (n=6,512)	p trend
Any adverse	15.3%	11.0%	8.0%	< 0.001
RDS+TTN	8.2%	5.5%	3.4%	< 0.001
Ventilation	1.9%	0.9%	0.4%	< 0.001
Hospital≥5d	9.1%	5.7%	3.6%	< 0.001

Table 1: Elective C/section after 37 weeks (13,258)

RDS: Respiratory distress syndrome; TTN: Transient tachypnea of newborn

#### REFERENCES

- 1. Wilmink FA, Hukkelhoven CW, Luncshof S, Mol BW, van der Post JA, Papatsonis DN. Neonatal outcome following elective cesarean section beyond 37 weeks of gestation: a 7 year retrospective analysis of a national registry. Am J Obstet Gynecol 2010 Mar;202(3):e1-e8.
- Hansen AK, Wisborg K, Uldbjerg N, Henriksen TB. Risk of respiratory morbidity in term infants delivered by elective caesarean section: cohort study. BMJ 2008 Jan 12;336(7635):85-87.
- 3. Tollånes MC, Moster D, Daltveit AK, Irgens LM. Cesarean section and risk of severe childhood asthma: a population-based cohort study. J Pediatr 2008 July;153(1):112-116.
- 4. Koplin J, Allen K, Gurrin L. Is caesarean delivery associated with sensitization to food allergens and IgE-mediated food allergy: a systematic review. Ped Allergy Immunol 2008 Dec;19(9):682-687.
- 5. MacKay DF, Smith GC, Dobbie R, Pell JP. Gestational age at delivery and special educational need: retrospective cohort study of 407,593 schoolchildren. PLoS Med 2010 Jun;7(6):e1000289.

## **Clinical and Social Issues of Noninvasive Prenatal Testing**

### <sup>1</sup>Giovanni Monni, <sup>2</sup>Ambra luculano, <sup>3</sup>Rossana Contu

<sup>1-3</sup>Department of Obstetrics and Gynecology, Prenatal and Preimplantation Genetic Diagnosis—Fetal Therapy, Ospedale Pediatrico Microcitemico "A Cao", Cagliari, Italy, Phone: +3907052965546/7, e-mail: prenatalmonni@tiscalicit

Cell-free deoxyribonucleic acid (cf DNA) analysis of maternal blood (noninvasive prenatal testing, NIPT) is the most advanced and efficient prenatal screening test for Trisomy 21, 18, and 13.<sup>1</sup>

The test is currently performed in high-risk population following 1st trimester screening.

Meta-analysis reported a detection rate of 99% for Trisomy 21, 96% for Trisomy 18, and 91% for Trisomy 13, with 0.35% false-positive rate.

Several limitations of the NIPT application, such as maternal obesity, aneuploidy and cancer, low DNA fraction, mosaicisms, and donor oocytes are reported. Patient's acceptability, nondefinitive results, patient's anxiety, and increase top have also been reported (Table 1).

Patients and doctors must also be careful for several aggressive commercial companies creating an increasing economic burden and prenatal policies disparity in access to genetic services.

The most important issue remains the genetic counseling about the accuracy and the efficacy of the NIPT<sup>2,3</sup> and the following necessity to undergo invasive prenatal diagnosis by chorionic villous sampling (CVS) and amniocentesis (Fig. 1).<sup>4</sup>

Since modern society is ever more-internet-web-based, patients must be aware that the easy access to information can prove misleading to their choices which should rather be counseled by a geneticist.

This article describes the clinical aspects as well as the social, economic, and legal issues related to NIPT.

**Table 1:** Prenatal diagnosis: Advantages and disadvantages

0	0		0
	CVS	Amniocentesis	NIPT
Gestational age	++	+	+
Multiple pregnancies	++	+	-
Safety (losses %)	0.1–0.5%	0.1–0.5%	+
Accuracy	++	++	+ only for T21
Days to obtain response	24–48 h	5–10 days	10 days
Repetition frequency	1%	0.5%	5%
Mendelian disease	++	+	-/+
Infections	++	+	-
Women's preference	?	?	?

CVS: Chorionic villous sampling; NIPT: Noninvasive prenatal testing



Fig. 1: Invasive prenatal diagnosis by CVS

#### REFERENCES

- Gil MM, Quezada MS, Revello R, Akolekar R, Nicolaides KH. Analysis of cell-free DNA in maternal blood in screening for fetal aneuploidies: updated meta-analysis. Ultrasound Obstet Gynecol 2015 Mar;45(3):249-266.
- 2. Masala M, Saba L, Zoppi MA, Puddu R, Piccau A, Capponi V, Iuculano A, Monni G, Rosatelli MC. Pitfalls in non-invasive fetal RhD and sex determination due to a vanishing twin. Prenat Diagn 2015 May;35(5):506-508.
- 3. Norton ME, Jacobsson B, Swamy GK, Laurent LC, Ranzini AC, Brar H, Tomlinson MW, Pereira L, Spitz JL, Hollemon D, et al. Cellfree DNA analysis for noninvasive examination of trisomy. N Engl J Med 2015 Apr 23;372(17):1589-1597.
- 4. Monni G, Zoppi MA, Iuculano A, Piras A, Arras M. Invasive or non-invasive prenatal genetic diagnosis? J Perinat Med 2014 Sep;42(5):545-548.

