




## Research paper

# Maximal velocity of fetal pulmonary venous blood flow



Oskar Sylwestrzak<sup>1</sup> , Maciej Słodki<sup>2,3,4</sup> , Maria Respondek-Liberska<sup>3,4</sup> 

<sup>1</sup>Medical Faculty, Medical University of Lodz, Poland

<sup>2</sup>Faculty of Health Sciences, Mazovian State University in Plock, Poland

<sup>3</sup>Department of Prenatal Cardiology, Polish Mother's Memorial Hospital Research Institute, Poland

<sup>4</sup>Department of Diagnoses and Prevention of Fetal Malformations, Medical University of Lodz, Poland

## Abstract

**Introduction:** Doppler imaging of foetal pulmonary veins plays a crucial role in prenatal cardiology centres. Therefore, we established reference ranges for maximal fetal pulmonary venous blood flow velocity for our unit.

**Material and methods:** A database of fetal ultrasound and echocardiographic examinations was analysed retrospectively. Healthy fetuses with no evidence of heart defect or any abnormality at the time of examination were selected as a study group. Fetuses with functional or morphological anomalies were excluded. The obtained data included: gestational age of pregnancy according to last menstrual period, maximal velocity of blood flow through pulmonary veins ( $V_{max}$  for PVs), prenatal cardiological diagnosis, extracardiac anomalies, and extracardiac abnormalities.

**Results:** Ultrasound data were collected at 18–39 weeks of gestation in singleton pregnancies. The study group contained 184 healthy fetuses. Scatter graph and reference ranges for their  $V_{max}$  for PVs during pregnancy were prepared. The regression equation for  $V_{max}$  for PVs as a function of gestational age (GA) in days was:  
 $V_{max}$  for PVs (cm/sec) =  $0.1 \times GA$  (in days) + 5.5 ( $r = 0.45$ , CI 0.95).

**Conclusion:** We presented normal ranges for pulmonary vein Doppler flow for the 18<sup>th</sup>–38<sup>th</sup> week of gestation in fetuses with normal heart anatomy, showing a steady increase towards the term in statistical analysis. However, each case should be approached individually because interpretation of the calculated values might not be very easy or straightforward.

**Key words:** fetal echocardiography, fetal heart, pulmonary veins, maximal velocity.

### Corresponding author:

Oskar Sylwestrzak  
Medical Faculty  
Medical University of Lodz, Poland  
e-mail: [sylwestrzakoskarpatriyk@gmail.com](mailto:sylwestrzakoskarpatriyk@gmail.com)

## Introduction

Colour-coded imaging of fetal pulmonary veins was described for the first time in 1992 by DeVore. Now it is possible to detect the pulmonary veins with their blood flows as early as week 12 of gestation and to trace them in detail as far as the distal end with advancing age [1].

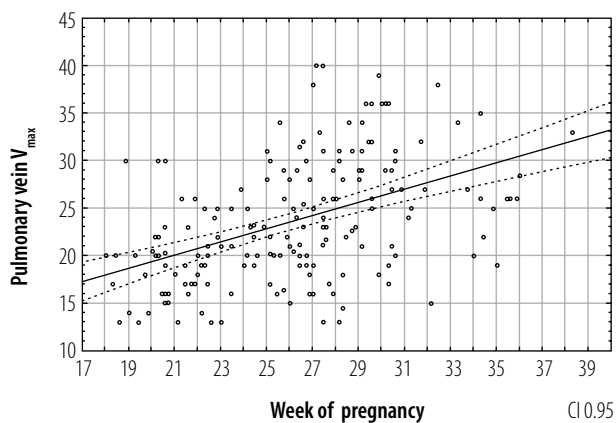
According to Polish Prenatal Cardiology Society and American Ultrasound Institute of Medicine, examination of fetal pulmonary venous blood flow is not compulsory during basic

obstetrical ultrasound, but it is an essential part of advanced examination in prenatal cardiology referential centres [2–3].

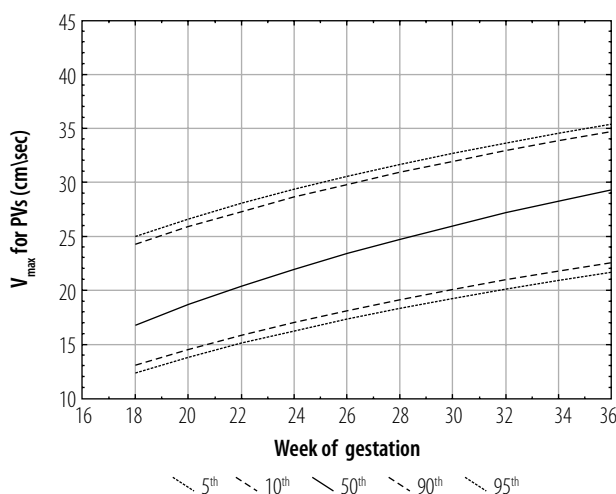
The purpose of this research was to establish reference ranges for maximal fetal pulmonary venous blood flow velocity for our unit.

### Material and methods

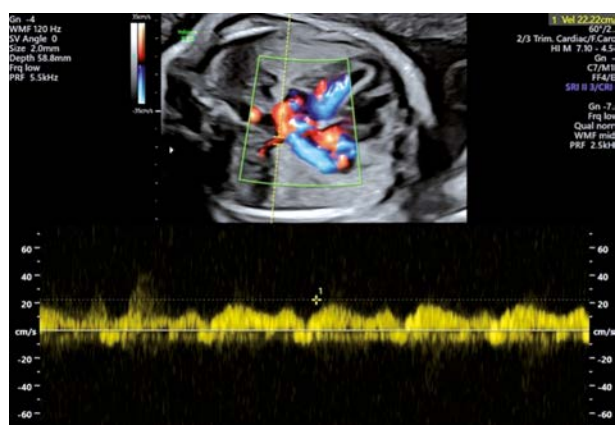
The database of fetal ultrasound and echocardiographic examinations performed by experienced fetal cardiologists in the referential centre of prenatal cardiology was analysed ret-



**Figure 1.** Scatter graph of fetal  $V_{max}$  for PVs in the course of pregnancy with regression line



**Figure 2.** Percentile graph of  $V_{max}$  for PVs between 18<sup>th</sup> and 36<sup>th</sup> week of gestation



**Figure 3.** Measurement of blood flow through pulmonary vein (30 hbd)

respectively. Obtained data included: gestational age of pregnancy according to last menstrual period, maximal velocity of blood flow through pulmonary veins ( $V_{max}$  for PVs), prenatal cardiological diagnosis, extracardiac anomalies, and extracardiac abnormalities. In our centre, visualisation of two PVs is a standard.  $V_{max}$  for PVs was measured for left and pulmo-

nary veins at their entrance to the left atrium separately, using the transducer position to obtain almost zero degree for blood flow. A correction angle was not used. The final result was the mean of these two measurements. Healthy fetuses with no evidence of heart defect or any abnormality at the time of examination were selected as a study group. In our database they had labels of normal heart anatomy, no extracardiac malformations, and no extracardiac anomalies. Fetuses with functional or morphological anomalies, oligo- or polyhydramnios, bright spot, arrhythmias, and umbilical cord around the neck and/or 2 vessel cord were excluded. For statistical analysis Statistica 13.1 software was used.

**Results**

Ultrasound data were collected at 18–39 weeks of gestation in singleton pregnancies. The study group contained 184 healthy fetuses. A scatter graph for their  $V_{max}$  for PVs during pregnancy was prepared using Pearson correlation (CI 0.95) (Figure 1). The regression equation for  $V_{max}$  for PVs as a function of gestational age (GA) in days was:

$$V_{max} \text{ for PVs (cm/sec)} = 0.1 \times \text{GA (in days)} + 5.5 \quad (r = 0.45, \text{ CI } 0.95).$$

The 5<sup>th</sup>, 10<sup>th</sup>, 50<sup>th</sup>, 90<sup>th</sup>, and 95<sup>th</sup> percentiles of  $V_{max}$  for PVs between 18<sup>th</sup> and 35<sup>th</sup> week of gestation were calculated. For consecutive percentiles a logarithmic dispersion line was created, and they were combined into a graph (Figure 2).

**Discussion**

As fetal lung parenchyma develops, so does the pulmonary vasculature. Its development starts by 34 days of gestation. Fetal pulmonary blood flow increases with gestation, from an initial low to almost 50% of the combined ventricular output by term [4]. Only a small amount of blood reaches the fetal pulmonary veins, which is caused by preferential shunting of oxygenated blood from the right to left side of the fetal heart and high vascular resistance in fetal lungs. Nevertheless, visualisation of fetal pulmonary veins connecting to the left atrium plays a crucial role in prenatal cardiology centres (Figure 3).

Many studies show normal fetal pulmonary venous blood flow pattern and velocity [1, 5–7]. Fetal pulmonary venous internal diameters were also evaluated [8]. Considering socio-economic, environmental, and behavioural changes in the contemporary population, it is essential to re-evaluate this data. Normal ranges for pulmonary vein blood flow require continuous elaboration. We estimated ranges of normal  $V_{max}$  for PVs in our region, for our own centre as a reference value for this and subsequent analyses, but it may be used by other obstetrical/cardiac centres. According to Paladini et al., normal  $V_{max}$  for fetal PVs differs from our results marginally [5]. Differences in ranges may be caused by various regions of study or different ultrasound devices used. Pulmonary blood flow velocity and its pattern should always be evaluated during fetal echocardiographic examination in a referential centre. Proper fetal cardiopulmonary circulation yields better outcomes after delivery.

Haemodynamic changes in the lung veins may occur in utero as early as in the second trimester and could explain re-

ports of chronic changes in the lung veins complicating neonatal or postoperative outcome in some heart anomalies [9]. An abnormal pattern may be seen in anomalous pulmonary venous connection or obstructed left atrium and restrictive foramen ovale [10, 11]. In the case of fetal critical aortic stenosis, a double reverse pattern in the pulmonary veins is associated with a poor prognosis [12]. In the IUGR population and in fetuses of diabetic mothers, we can observe higher pulsatility, which is lower during fetal respiratory movements [13–15]. That is why velocity measurement of blood flow through pulmonary veins is of great value.

Mother hyperoxygenation is considered to be a potential therapy in some cases of prenatal congenital heart defects [16–18]. Such a new way of treatment requires the response of pulmonary veins to a high concentration of oxygen. It is used also in our centre. Nonetheless, its precise role should be carefully evaluated [19].

## Conclusions

We present normal ranges for pulmonary vein Doppler flow for the 18<sup>th</sup>–36<sup>th</sup> week of gestation in fetuses with normal heart anatomy, showing a steady increase towards term in statistical analysis. However, the scattered graph of  $V_{max}$  for PVs suggested that for an individual case the proper interpretation of the calculated values might not be very easy or straightforward.

## Conflict of interest

The authors declare no conflict of interest.

## REFERENCES

- Bahlmann F, Galliani R, Schmidt-Fittschen M, Al Naimi A, Reinhard I, Willruth A. Fetal pulmonary venous blood flow velocities in a normal population and new calculated reference values. *Prenat Diagn* 2016; 36: 1033-1040.
- Respondek-Liberska M. [Prenatal USG/ECHO diagnostics. Functional disturbances in cardiovascular system]. 1<sup>st</sup> Ed. PZWL Wydawnictwo Lekarskie, Warszawa 2019.
- Respondek-Liberska M. [Prenatal cardiology for obstetricians and pediatric cardiologists]. 1<sup>st</sup> Ed. Wydawnictwo Czelej, Lublin 2006.
- Morton SU, Brodsky D. Fetal physiology and the transition to extrauterine life. *Clin Perinatol* 2016; 43: 395-407.
- Paladini D, Palmieri S, Celentano E, Guida F, Salviati M, Morra T, et al. Pulmonary venous blood flow in the human fetus. *Ultrasound Obstet Gynecol* 1997; 10: 27-31.
- Hong YM, Choi JY. Pulmonary venous flow from fetal to neonatal period. *Early Hum Dev* 2000; 57: 95-103.
- Laudy JA, Huisman TW, de Ridder MA, Wladimiroff JW. Normal fetal pulmonary venous blood flow velocity. *Ultrasound Obstet Gynecol* 1995; 6: 277-281.
- Dong FQ, Zhang YH, Li ZA, Hou ZZ, He XJ, Guo YZ. Evaluation of normal fetal pulmonary veins from the early second trimester by enhanced-flow (e-flow) echocardiography. *Ultrasound Obstet Gynecol* 2011; 38: 652-657.
- Lenz F, Chaoui R. Changes in pulmonary venous Doppler parameters in fetal cardiac defects. *Ultrasound Obstet Gynecol* 2006; 28: 63-70.
- Respondek-Liberska M, Janiak K, Moll J, Ostrowska K, Czichos E. Prenatal diagnosis of partial anomalous pulmonary venous connection by detection of dilatation of superior vena cava in hypoplastic left heart. *Fetal Diagn Ther* 2002; 17: 298-301.
- Respondek-Liberska M, Sokołowski Ł, Słodki M, Zych-Krekora K, Strzelecka I, Krekora M, et al. Prenatal diagnosis of TAPVC on Monday, delivery of Tuesday and cardiac surgery at Wednesday – a model of perinatal care in 3<sup>rd</sup> trimester in case of fetal/neonatal critical heart defect in tertiary center. *Prenat Cardio* 2016; 6: 37-42.
- Sukegawa S, Yamamoto Y, Sato K, Tanaka S, Tanaka T, Mitsuhashi N. Ultrasound evaluation of fetal critical aortic stenosis using the left atrium area/cardiac area ratio and the Doppler patterns in the pulmonary veins. *J Med Ultrason* 2019; 46: 267-272.
- Bravo-Valenzuela NJ, Zielinsky P, Huhta JC, Acacio GL, Nicoloso LH, Piccoli A, et al. Dynamics of pulmonary venous flow in fetuses with intrauterine growth restriction. *Prenat Diagn* 2015; 35: 249-253.
- Chemello K, Zielinsky P, Nicoloso LH, de Moraes MR. Behavior of pulmonary venous flow during fetal respiratory movements. *Congenital Heart Dis* 2009; 4: 265-268.
- Zielinsky P, Piccoli AL Jr, Teixeira L, Gus EI, Mânica JL, Satler F, et al. Pulmonary vein pulsatility in fetuses of diabetic mothers: prenatal Doppler echocardiographic study. *Arq Bras Cardiol* 2003; 81: 604-607, 600-603.
- Co-Vu J, Lopez-Colon D, Vyas HV, Weiner N, DeGroff C. Maternal hyperoxygenation: a potential therapy for congenital heart disease in the fetuses? A systematic review of the current literature. *Echocardiography* 2017; 34: 1822-1833.
- Zarkowska-Szaniawska A, Janiak K, Forys S, Słodki M, Respondek-Liberska M. Maternal hyperoxygenation test in prediction of fetal lung hypoplasia – preliminary report. *Ginekol Pol* 2011; 82: 834-839.
- Zeng S, Zhou J, Peng Q, Deng W, Zhang M, Zhao Y, et al. Sustained maternal hyperoxygenation improves aortic arch dimensions in fetuses with coarctation. *Sci Rep* 2016; 6: 39304.
- McHugh A, El-Khuffash A, Bussmann N, Doherty A, Franklin O, Breathnach F. Hyperoxygenation in pregnancy exerts a more profound effect on cardiovascular hemodynamics than is observed in the nonpregnant state. *Am J Obstet Gynecol* 2019; 220: 397.e1-397.e8.

### Division of work:

Oskar Sylwestrzak (ORCID: 0000-0001-9325-7304): collection and/or assembly of data, data analysis and interpretation, writing the article, final approval of article

Maciej Słodki (ORCID: 0000-0002-0160-8013): writing the article, critical revision of the article

Maria Respondek-Liberska (ORCID: 0000-0003-0238-2172): research concept and design, data analysis and interpretation, writing the article, critical revision of the article, final approval of article