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ABSTRACT

BACKGROUND AND PURPOSE: Fetal lateral ventriculomegaly is a relatively common finding with much debate over its clinical significance. The purpose of this study was to examine the association between ventriculomegaly and asymmetry and concomitant CNS findings as seen in fetal brain MR imaging.

MATERIALS AND METHODS: Fetal brain MR imaging performed for various indications, including ventriculomegaly, with or without additional ultrasound findings, was assessed for possible inclusion. Two hundred seventy-eight cases found to have at least 1 lateral ventricle with a width of \( \geq 10 \) mm were included in the study. Ventriculomegaly was considered mild if the measurement was 10–11.9 mm; moderate if, 12–14.9 mm; and severe if, \( \geq 15 \) mm. Asymmetry was defined as a difference of \( \geq 2 \) mm between the 2 lateral ventricles. Fetal brain MR imaging findings were classified according to severity by predefined categories.

RESULTS: The risk of CNS findings appears to be strongly related to the width of the ventricle (OR, 1.38; 95% CI, 1.08–1.76; \( P = .009 \)). The prevalence of associated CNS abnormalities was significantly higher (\( P = .005 \)) in symmetric ventriculomegaly compared with asymmetric ventriculomegaly (38.8% versus 24.2%, respectively, for all CNS abnormalities and 20% versus 7.1%, respectively, for major CNS abnormalities).

CONCLUSIONS: In this study, we demonstrate that the rate of minor and major findings increased with each millimeter increase in ventricle width and that the presence of symmetric ventricles in mild and moderate ventriculomegaly was a prognostic indicator for CNS abnormalities.

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From the Department of Obstetrics and Gynecology (E.B., R.A., E.K.) and Pediatric Neurology Unit (O.B.-Y.), Sheba Medical Center, Tel-Hashomer, Ramat-Gan, Israel; and Sackler School of Medicine (E.B., O.B.-Y., S.D., R.A., E.K.), Tel Aviv University, Tel Aviv, Israel.

E. Barzilay and O. Bar-Yosef contributed equally to this work.

Please address correspondence to Eran Barzilay, MD, PhD, Department of Obstetrics and Gynecology, Sheba Medical Center, Tel-Hashomer, Ramat-Gan, Israel; e-mail: eran.barzilay@gmail.com

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"Ventriculomegaly" was defined as a lateral ventricle width of ≥10 mm.²–⁴ Lateral ventricle asymmetry was defined as a difference of ≥2 mm between the 2 lateral ventricles. All MR imaging scans during the study period were assessed for possible eligibility, regardless of the reason for the referral. Inclusion criteria for this study were singleton pregnancy and ventriculomegaly (at least 1 lateral ventricle with a width of ≥10 mm). Exclusion criteria were fetal cytomegalovirus or toxoplasma infection and cases with abnormal genetic tests (karyotype or microarray). Two hundred seventy-eight pregnancies were included in the study. The average maternal age was 31.8 years (interquartile range, 29–33 years). Two hundred seventy-eight cases met the inclusion/exclusion criteria.

Fetal MR imaging was performed with a 1.5T system (Optima 1.5T; GE Healthcare, Milwaukee, Wisconsin) as previously published.¹⁷ Single-shot fast spin-echo T2-weighted sequences in 3 orthogonal planes were performed by using the following parameters: section thickness, 3–4 mm; no gap; flexible coil (8-channel cardiac coil); matrix, 320 × 224; TE, 90 ms; and TR, 1298 ms. The FOV was determined by the size of the fetal head: 24 cm for the smaller fetuses and 30 cm for the larger fetuses. T1 fast-spoiled gradient-echo sequences were performed only in the axial plane with a larger FOV (400 mm), with section thickness, 4 mm; gap, 0.5 mm; TR, 160 ms; and TE, 2.3 ms. Measurements of the lateral ventricle widths were obtained on the coronal plane at the level of the ventricles (with good visibility of the choroid plexuses). For this study, ventriculomegaly was considered mild if the measurement was 10–11.9 mm; moderate if, 12–14.9 mm; and severe if, ≥15 mm. Asymmetry was defined as a difference of ≥2 mm between the 2 lateral ventricles.

Fetal brain MR imaging findings were classified according to severity by predefined categories as shown in the Table. Classification of findings into minor or major was determined on the basis of previously published data.¹⁸–²⁷ Statistical analyses were conducted by using the Statistical Package for the Social Sciences (SPSS, Version 23; IBM, Armonk, New York). Q-Q plots were used to assess normality. Continuous variables were described as mean (SD) or median (interquartile range) as appropriate. Continuous variables were compared by using an unpaired t test and a 1-way ANOVA test as appropriate or by using the Mann-Whitney U test and Kruskal Wallis test for nonparametric continuous variables as appropriate. The χ² test was used for comparison of categoric variables. Logistic regression analysis was used to examine the relationship among ventricle width, width difference, and CNS findings. Significance was P < .05.

The study was approved by the local institutional review board of the Sheba Medical Center.

RESULTS
Two hundred seventy-eight pregnancies were included in the study. The average maternal age was 31.8 ± 5.2 years. Maternal age did not differ significantly between cases with or without ventricle asymmetry (32 versus 31.4 years, respectively; P = .693) or among cases of mild, moderate, and severe ventriculomegaly (32, 31.5, and 29.5 years of age, respectively; P = .235). The median gestational age at MR imaging acquisition was 32 weeks (interquartile range, 31–34 weeks). There was no difference in median gestational age between cases with or without ventricle asymmetry (32 versus 32 weeks, respectively; P = .079) or among cases of mild, moderate, and severe ventriculomegaly (32, 32, and 31.5 weeks, respectively, P = .785).

Median ventricle width for the widest ventricle was 11 mm (interquartile range, 11–13 mm), and for the contralateral ventricle, it was 9 mm (interquartile range, 7–11 mm). Median ventricle width difference was 3 mm (interquartile range, 1–4 mm).

Figure 1 depicts the distribution of ventriculomegaly cases according to severity and the presence or absence of asymmetry.
Most cases were defined as mild ventriculomegaly (73%), while moderate and severe ventriculomegaly comprised 22.7% and 4.3% of the cases, respectively. Most cases of ventriculomegaly were asymmetric (71.2%).

Figure 2 depicts the distribution of CNS findings according to the width of the ventricle (the larger of the 2 ventricles). The risk of CNS findings appears to be strongly related to the width of the ventricle. The association between CNS findings and ventricle width was particularly evident in moderate ventriculomegaly, in which each 1-mm increase in ventricle width increases the risk for both minor and major CNS findings. Logistic regression analysis, with each ventricle width along with maternal age, gestational age at the time of MR imaging, and fetal sex as variables, demonstrated that the widths of both ventricles were positively associated with an increased risk for CNS anomalies (OR, 1.38; 95% CI, 1.08–1.76; P = .009, for the maximal ventricle; and OR, 1.21; 95% CI, 1.03–1.44; P = .025, for the contralateral ventricle). Gestational age at the time of MR imaging was found to be negatively associated with CNS anomalies (OR, 0.86; 95% CI, 0.77–0.95; P = .003), possibly indicating that the more complicated cases tended to be diagnosed earlier. Maternal age (P = .75) and fetal sex (P = .304) were not found to be associated with CNS anomalies. The correlation between ventricle width and associated anomalies appears to be strongest within the 12- to 15-mm range because the OR for CNS anomalies increased from 1.38 (95% CI, 1.08–1.76; P = .009) to 2.53 (95% CI, 1.48–4.34; P = .001) when the analysis was restricted to cases with a ventricle width of 12–15 mm.

Most cases in this study were asymmetric ventriculomegaly (71%). We have found more CNS abnormalities in symmetric ventriculomegaly compared with asymmetric ventriculomegaly (38.8% versus 24.2%, respectively) and more major CNS abnormalities in symmetric ventriculomegaly compared with asymmetric ventriculomegaly (20% versus 7.1%, respectively). These differences between symmetric and asymmetric ventriculomegaly were found to be statistically significant (P = .005).

Figure 3 illustrates the distribution of CNS abnormalities according to ventriculomegaly severity and the presence or absence of asymmetry. We found that the differences in the risk of CNS abnormalities between symmetric and asymmetric ventriculomegaly were only apparent in the mild and moderate groups. In cases of severe ventriculomegaly, the rates of minor (25%) and major (50%) findings were identical in symmetric and asymmetric ventriculomegaly. The number of cases in the severe ventriculomegaly group (n = 12) was much smaller than in the moderate (n = 63) and mild (n = 203) groups.

Next, we used regression analysis to explore the individual impact of ventricle width and ventricle asymmetry on the risk of CNS findings. Using maximal ventricle width and width difference as variables, we found that ventricle width was an independent risk factor for CNS findings (OR, 1.56; 95% CI, 1.32–1.85; P < .001), while ventricle width difference was inversely associated with CNS findings (OR, 0.79; 95% CI, 0.68–0.92; P = .003). When average ventricle width was used in the regression model instead of maximal ventricle width, average ventricle width appeared to be independently associated with CNS findings (OR, 1.56; 95% CI, 1.32–1.85; P < .001). However, width difference lost its significant association with CNS findings (OR, 0.99; 95% CI, 0.84–1.16; P = .896).

**DISCUSSION**

In this study we analyzed a large number of MR imaging scans of fetuses diagnosed with fetal ventriculomegaly. We have shown that the risk of minor and major CNS findings is closely related to the width of the lateral ventricle. The association between ventricle width and CNS findings is particularly evident in moderate ventriculomegaly. As seen in Fig 2, the rate of minor and major findings significantly increased with each millime-
Moreover, these findings are contrary to those in previous studies to adequately portray the risk of CNS anomalies, taking into account the increase in ventricle width. Thus, while the prevalent categorization of mild, moderate, and severe ventriculomegaly appears to adequately portray the risk of CNS anomalies, taking into account the ventricle width greatly improves the risk estimation. Moreover, these findings are contrary to those in previous studies that defined mild ventriculomegaly at an atrial width of 10–15 mm.\textsuperscript{2,16}

Another intriguing result of this study is the correlation between ventricle asymmetry and CNS findings in fetal MR imaging. We have found that symmetric ventriculomegaly was associated with more CNS findings than asymmetric ventriculomegaly. This difference was apparent in mild and moderate ventriculomegaly, while the rates of minor and major CNS findings were similar between symmetric and asymmetric severe ventriculomegaly. The apparent advantage of asymmetric ventriculomegaly might be somewhat counterintuitive. However, the presence of asymmetry for a case with a given ventricle width translates to a smaller contralateral ventricle. Thus, in asymmetric mild ventriculomegaly, the contralateral ventricle will be in the normal range, while in moderate ventriculomegaly, most cases will have a contralateral ventricle width in the normal-to-mild ventriculomegaly range. This hypothesis gains support from our regression analysis that demonstrated that the association between ventricle width and CNS findings was eliminated when average ventricle width was used as a variable in the analysis instead of maximal ventricle width, thereby encompassing data on both ventricles.

The relatively low incidence of major anomalies in cases of mild symmetric ventriculomegaly (2.6%) may merit re-evaluation of the cost-effectiveness of fetal MR imaging in these cases. Nevertheless, we believe that given the clinical implications of major CNS anomalies and in light of the prevalence of minor findings in mild asymmetric ventriculomegaly (14.5%), fetal brain MR imaging still plays an important role in the work-up of these cases.

The data portrayed in this study aid in establishing the correlation between lateral ventricle width and CNS anomalies. The high rate of CNS anomalies observed in this study stresses the need for a thorough investigation of fetal ventriculomegaly and especially for meticulous examination in cases of moderate-to-severe ventriculomegaly and bilateral ventriculomegaly.

This study is in accordance with previous studies that found more CNS anomalies in moderate and severe ventriculomegaly compared with mild ventriculomegaly.\textsuperscript{10,11} However, in most of these studies, imaging was based almost exclusively on ultrasound,\textsuperscript{3–11} despite the known advantages of fetal brain MR imaging.\textsuperscript{17} Moreover, none of these studies, including those with fetal MR imaging as an imaging technique,\textsuperscript{15,17} have analyzed the impact of ventricle width on the risk of CNS anomalies.

One of the advantages of this study is that it included a large number of fetal MR imaging scans, enabling us to examine in depth the association between ventriculomegaly and CNS findings. Thus, we could identify differences in the rate of CNS findings not only among mild, moderate, and severe ventriculomegaly but also due to minute increases in ventricle width. Another major advantage is that the study is MR imaging–based, thereby increasing the ability to identify CNS abnormalities compared with neurosonography.\textsuperscript{17,28–31}

This study is limited by the lack of data on neurodevelopmental outcome. Here we examined the association between ventriculomegaly and CNS anomalies in the fetus without assessing the outcome of the child. Nevertheless, because the presence of CNS abnormalities has been shown to be closely related to the neurodevelopmental outcome of the child,\textsuperscript{7} findings in fetal MR imaging can be used as an indication of the outcome. Moreover, to improve the clinical significance of the association between ventriculomegaly and CNS anomalies, we classified CNS anomalies according to severity (Table).

CONCLUSIONS
In this study, we demonstrate that the rate of minor and major findings increases with each millimeter increase in ventricle width and that the presence of symmetric ventricles in mild and moderate ventriculomegaly is a prognostic indicator for CNS anomalies in MR imaging examinations.

REFERENCES

![FIG 3. Distribution of additional CNS findings in cases of fetal ventriculomegaly according to ventriculomegaly severity and the presence or absence of asymmetry. Filled bars represent major CNS findings, dotted bars represent minor CNS findings, and empty bars represent no additional CNS findings.](image-url)