Amniotic fluid as a vital sign for fetal wellbeing

Abstract

Introduction: Amniotic fluid, once thought to merely provide protection and room for necessary movement and growth for the fetus, is now understood to be a highly complex and dynamic system that is studied as a data point to interpret fetal wellbeing.

Methods: Assessment of amniotic fluid volume is now routine when performing a sonographic evaluation of fetal status and is an important consideration in the assessment and management of perinatal morbidity and mortality. In this review, we will cover the dynamics that affect amniotic fluid volume, review methods for measurement and quantification of volume, review definitions for normative data as related to neonatal outcomes, and provide evidence based guidance on the workup and management options for oligohydramnios and polyhydramnios in singleton and twin pregnancies.

Conclusions: When abnormalities of fluid exist, appropriate workup to uncover the underlying etiology should be initiated as adverse fetal outcomes are sometimes associated with these variations from normalcy.

Keywords: amniotic fluid volume, oligohydramnios, polyhydramnios.

Dynamics

In order to understand how variations in amniotic fluid volume (AFV) can affect the fetus, we must first describe amniotic fluid dynamics. In early pregnancy, the factors that affect AFV are not well known. It is known that the osmolality of amniotic fluid (AF) and maternal plasma are the same, suggesting that the fluid is a transudate of maternal plasma either across the placental surface or through the fetal skin. The fetal skin is non-keratinised until week 22–25, allowing it to act as a membrane through which amniotic fluid can readily pass. Beyond 24 weeks, the surfaces in the mouth and nose can act to exchange fluid, but this is not considered a major source of AFV regulation.

The major pathways affecting AF in the latter half of pregnancy are pulmonary excretion, fetal urine production, fetal swallowing, intramembranous movement between fetal blood and the placenta, and transmembranous movement across the amnion and chorion. A fundamental source of AF is the fetal renal system. This is evidenced by the almost complete lack of AF in fetuses with renal agenesis. Evidence of a functioning renal system first starts around 8–11 weeks when urine is initially observed in the fetal bladder. The dilute urine that is produced is thought to cause the observed drop in osmolality and sodium concentration in the AF that persists until delivery. As pregnancy advances, urinary byproducts are observed at two to three times the concentration found in fetal plasma. Estimates of fetal urine output have been made by 3D measurements of bladder volume and timed intervals and have been found to range between 7 and 70 mL/hour from 24 weeks until delivery in two separate studies.

Akin to renal agenesis being associated with low amniotic fluid, disruption in fetal swallowing is associated with excess AFV. The human fetus demonstrates swallowing around the same time that urine production begins. The ovine model has historically been an accurate approximation of human fetal development. Studies evaluating the volume of AF swallowed by the fetus were done primarily in sheep models, and found to range between 8.75 and 43.2 mL/hr. Ovine studies have also shown that the amount of AF swallowed daily in late gestation is correlated to the AFV, suggesting the fetus can react and attempt to regulate its amniotic fluid surroundings. It is not, however, thought to be a major regulator of AFV, although the fetus is able to modulate swallowing. Primate studies performed by Minei and Suzuki showed initial development of polyhydramnios with esophageal ligation, but the hydramnios had resolved prior to delivery.

Fluid is also excreted from the fetal pulmonary system. It has been demonstrated in fetal sheep by Brace, et al. that half of secreted lung fluid enters the amniotic fluid, and the other half is swallowed upon exit from the trachea. As further evidence of the entrance of secreted lung fluid into amniotic fluid, the phospholipids found in amniotic fluid are produced by pulmonary cells and excreted in fetal urine. The trachea acts as a one-way valve in most situations preventing amniotic fluid from...
entering the lungs. Intra-amniotic injection of contrast media by Liley, et al. resulted in few cases of detectable contrast found in the fetal or neonatal lung. Additionally, meconium staining is not infrequent with a rate in the literature of 7–27% based on the population, whereas meconium aspiration past the trachea is rare, anywhere from less than a percent up to 5% and usually associated with neonatal hypoxia. Aspiration of unimitted amniotic fluid has been reported as a cause of neonatal death confirmed at autopsy.16,17

Intramembranous exchange of fluid describes the resorption of fluid and solutes from the amniotic compartment to the fetal blood via the amnion. This is thought to be regulated at least partially by aquaporins found in the human chorionicamniotic membranes and the placenta, which have been found to be adaptive to abnormal AF levels and may represent a possible therapeutic target for abnormal AF regulation.19-21

As the fluid requirements of the fetus increase with increasing gestation, water flow from the amniotic cavity to the fetal circulation via fetal membranes increases up to 400 mL/day.22 The osmotic gradient between the amniotic fluid and fetal blood also drives fluids and solutes from the amniotic fluid into the fetal blood.23 The inflow of solutes and fluid, although bidirectional, are not necessarily equal.24 Transmembranous movement of amniotic fluid describes the transport of water and solutes from the maternal circulation to the amniotic compartment via the placenta. This has been found to be an insignificant contribution to amniotic fluid dynamics.24

**Methods of quantifying amniotic fluid**

Measurement of amniotic fluid volume can be made directly, indirectly, or estimated sonographically. Direct measurement is done at the time of cesarean or uterine hysterotomy.25 Indirect measurement is done via amniocentesis by dye-dilution techniques. Dye-dilution using para-amino hippurate has been shown to be representative of actual AFV obtained by direct measurement at the time of cesarean delivery.26 Because these techniques to measure amniotic fluid volume are time consuming, invasive, and may require laboratory support, amniotic fluid volumes are usually estimated by ultrasound. Magnetic resonance imaging (MRI) has also been evaluated as a means for estimating AFV; however, this is an impractical approach to everyday screening.

There are four methods of sonographic evaluation of amniotic fluid volume: subjective assessment, 2 x 2 measurement, single deepest pocket (SDP), also known as maximum vertical pocket (MVP), and amniotic fluid index (AFI). Amniotic fluid index, first proposed by Phelan and Rutherford, is the summation of the vertical diameter of the largest pocket in each of the four quadrants with the maternal umbilicus as a central reference point. The transducer should be oriented in the longitudinal plane and there should be a minimum horizontal measurement of one centimeter for each pocket. The single deepest vertical pocket was proposed by Chamberlain and is simply found by identifying the largest pocket of amniotic fluid after a global assessment and selecting the largest vertical measurement with a minimum horizontal measurement of one centimeter. A two-diameter pocket (cm²) is calculated by multiplying the depth and width of the largest single pocket.

The advantages of sonographic estimates are that they are simple to perform, easy to teach to residents, midwives, and nurses, and are reproducible. The disadvantage is that sonograms are two dimensional representations of a complex, three-dimensional structure with limited impact on clinical outcome. The reproducibility of these measurements was shown by Magann, et al. comparing all four measurements in singleton pregnancies with dye-dilution technique across different operator experiences. The accuracy of subjective estimates ranged from 65–70% and the accuracy of the three sonographic estimates were similar, ranging from 59–67%. These were similar across operator experience. Alarmingly, none of the techniques consistently identified abnormal AFVs (oligohydramnios and polyhydramnios). The accuracy in twin pregnancies in a similar study was even more dismal, ranging from 7–29%.

Throughout multiple studies and trials, no single sonographic method has emerged superior to the others. A randomised trial comparing the use of SDP versus AFI during modified biophysical profiles in complicated pregnancies showed no difference in delivery mode, NICU admission, umbilical artery pH, or Apgar score. The only significant finding was that AFI identified more pregnancies as oligohydramnios than SDP. This was confirmed in two meta-analyses comparing the measurement of AFI to SDP. AFI over-diagnosed clinically insignificant oligohydramnios leading to superfluous intervention without a difference in outcome, arguing that the use of SDP for sonographic volume assessment may be better.

**Modification of ultrasound quantification**

The use of color Doppler has been added as a means to identify the umbilical cord in amniotic fluid in an effort to better diagnose oligohydramnios. It has, however, not been shown to aid in identification of pregnancies with adverse outcomes. Using dye-determined AFV as the benchmark, Magann, et al. compared gray scale ultrasonography to Doppler color ultrasonography and found that Doppler not only over-diagnosed oligohydramnios, but labeled 37% of women with normal AFV as having oligohydramnios. The use of color Doppler is not recommended when evaluating amniotic fluid volume. Sahin, et al. attempted to apply the Cavalieri method, a mathematical estimate of volumes, to estimate AFV by ultrasound images. Their results correlated with concurrent AFI measurements, and although this is a promising theory, it is complicated to perform and has not been correlated with pregnancy outcome or directly measured AFV.

3D ultrasound technology has found relevance in the evaluation of the fetus; however, it has not yet found relevance in the assessment of amniotic fluid. There has only been one attempt to assess third trimester AFV with 3D ultrasonography. The authors of that study concluded that 3D volume data sets are reliable for determining AFV; however, AFV was determined subjectively by the sonographer based on five volume acquisitions and compared to the value obtained during the 2D ultrasound; there was no attempt at calculating a volume based on a gold standard. Interestingly, in the first trimester, growth charts have been developed for gestational sac volume and embryonic volume, the ratio of which correlates positively with
gestational age. Clinical applications for this data have not yet been developed, but normal versus abnormal early pregnancies are being investigated to determine if these parameters can help predict early adverse outcomes such as miscarriage.

**Normal volume**

Normality can be defined in two ways. Mathematically, it can represent the majority of the population as expressed by area under the curve in a Gaussian distributed population. Typically, the lower and upper 5% are excluded as ‘abnormal’. Normality can also be defined in terms of outcome. Undesired outcomes are tracked and measurements associated with these outcomes are defined as ‘abnormal’. This second definition is more clinically useful, but much more difficult to define.

Amniotic fluid volume has been defined in both of these manners as well. The 5th and 95th percentiles have been defined for AFI, SDP, the 2-diameter pocket, and dye-directed techniques across gestational ages, as fluid levels have been found to vary significantly throughout pregnancy. Nomograms for amniotic fluid in normal pregnancies have been developed by Queenan, et al. by dye-determined methods; Moore and Cayle by AFI; Brace and Wolfe by dye-dilution and direct measurement; and Magann, et al. by dye-dilution, direct measurement, and sonographic estimate. It is worth noting that although for diagnostic purposes a single set of cutoffs for abnormal amniotic fluid volumes are used throughout a pregnancy, the actual values vary by each week. In the figure below, two studies showed amniotic fluid volume increasing steadily in early gestation and remain relatively stable between 22 and 38 weeks, thereafter declining until delivery. The third nomogram found that amniotic fluid volume continues to increase throughout gestation with a mean volume of approximately 800 mL at term. Table 1 shows typical references for normal ranges of amniotic fluid volume based on sonographic estimates.

A study of over 15,000 patients done by Shanks, et al. showed that AFI less than the 5th percentile versus a cutoff of 5 cm better predicts fetuses at risk for newborn intensive care (NICU) admission, although this study has been criticised for its retrospective design and the variable length of time between the time the measurement was obtained and the time of delivery. Additionally, an investigation of 291 pregnancies with dye-determined AFVs in which the 3rd and 5th percentiles of the AFI were assessed to determine if either of these percentiles was superior to the fixed cut off of < 5 for the AFI to detect the 75 pregnancies with oligohydramnios discovered that both the percentiles (3rd and 5th) and the fixed cut off (< 5) poorly

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**Table 1: Normal values of amniotic fluid estimates.**

| Ultrasound estimate | Oligohydramnios | Normal Values | Polyhydramnios |
|---------------------|-----------------|---------------|----------------|
| AFI                 | < 5 cm          | 5–25 cm       | > 25 cm        |
| SDP                 | < 2 cm          | 2–8 cm        | > 8 cm         |
| 2 diameter pocket   | < 4 cm²         | 4–50 cm²      | > 50 cm²       |

Modified from Magann EF. Oligohydramnios: Amniotic Fluid and the Clinical Relevance of the Sonographically Estimated Amniotic Fluid Volume, *J Ultrasound Med* 2011; 30 (11): 1573–85, and The amniotic fluid index, single deepest pocket, and 2-diameter pocket in normal human pregnancy, *Am J Obstet Gynecol*, 2000; 182 (6): 1581–8.
identified oligohydramnios, but neither was superior to the other.48

**Oligohydramnios**

Oligohydramnios has been defined as an AFV that is less than 200 mL25 or 500 mL.49 By ultrasound techniques, it has been estimated as an SDP less than 2 cm,50 AFI less than 5 cm28,51 or an AFI that is below the 5th percentile for gestational age,35,43 or a subjectively low AFV.52 Borderline fluid has been defined as an AFI between 5–8 cm or 5–10 cm and has been associated with fetal malformations if diagnosed at age 24–34 weeks.52,53

Low amniotic fluid can be due to underproduction, loss, or can be idiopathic. Underproduction can be the result of absent or dysfunctional kidneys, urinary tract obstruction, abnormal placental function, or maternal dehydration. Loss is due to rupture of membranes. Appropriate workup for this abnormal vital sign is the review of maternal history, assessment for evidence of membrane rupture, anatomic evaluation of the renal system and bladder, and assessment of placental function and fetal growth.

Regardless of the etiology, fetuses in pregnancies complicated by oligohydramnios are at increased risk of adverse outcomes in the form of cord accidents. Oligohydramnios/anhydramnios can also result in fetal lung hypoplasia, malformations, and contracture if it is severe enough and persistent.44 Oligohydramnios was retrospectively evaluated in 7582 fetuses in high risk pregnancies with normal anatomy. Perinatal mortality was 11% in these pregnancies versus 0.2% in pregnancies with normal fluid based on SDP. Borderline fluid of SDP 1–2 cm was associated with an increased mortality at 3.7%.50 Morbidity such as cesarean delivery, non-reassuring fetal heart rate patterns, NICU admission, and meconium aspiration are also increased in pregnancies complicated by oligohydramnios.55

There is conflicting data that argues that the most significant risk to the normal fetus with incidental oligohydramnios is iatrogenic prematurity resulting from the subsequent rush to delivery.56 Low AF as determined by dye-dilution along with SGA has been associated with NICU admission (OR 11.1) but no other measures of infant morbidity in another study.57 A retrospective cohort study evaluating the consequences of induction in 206 pregnancies with oligohydramnios as compared to 206 spontaneous labors with normal AFV showed a significant increase in operative deliveries (forceps, vacuum and cesarean) citing non-reassuring fetal status as the driving factor. There was no difference found in neonatal acidosis, Apgar score, NICU admission, or perinatal rate, as well as higher perinatal mortality when all abnormal fluid pregnancies were compared to normal.71 It is notable that none of the above studies evaluated clinically important outcomes. Maternal hydration, therefore, has a limited clinical application and may help in cases where adequate amniotic fluid assists in outcome, such as external cephalic versions or amniocentesis.

Management of oligohydramnios is dependent on the gestational age when discovered. If discovered after 37 weeks gestational age, and rupture of membranes ruled out, induction of labor would not be unreasonable. A retrospective cohort study evaluating the consequences of induction in 206 pregnancies with oligohydramnios as compared to 206 spontaneous labors with normal AFV showed a significant increase in operative deliveries (forceps, vacuum and cesarean) citing non-reassuring fetal status as the driving factor. There was no difference found in neonatal acidosis, Apgar score, NICU admission, or perinatal morbidity and mortality.72 A prospective study looked at outcomes in both oligohydramnios and polyhydramnios as compared to pregnancies with normal AFV. They reported a 56% induction rate with oligohydramnios and 57% cesarean rate, as well as higher perinatal mortality when all abnormal fluid pregnancies were compared to normal.73 Unfortunately, there is no direct comparison in either study of outcomes of inductions with low and normal amniotic fluid.

There is a single randomised trial evaluating induction versus expectant management in 54 patients with oligohydramnios beyond 40 weeks. No difference was found in mode of delivery or neonatal Apgar score or cord blood pH.74 However, in over 500 United States Maternal-Fetal Medicine members surveyed, 92% would recommend induction without documented lung maturity before 39 weeks, and 35% before 37 weeks, even though only one-third of respondents felt induction would decrease
adverse outcomes. If expectant management is chosen, there is no consensus on what fetal monitoring is best, or even if fetal monitoring is necessary; however, the use of Doppler monitoring in pregnancies thought to be at risk of placental insufficiency has shown benefit. Consistent maternal oral hydration can be encouraged.

Prior to term, the discovery of oligohydramnios should initiate a sonographic evaluation if not already undertaken, focusing on the renal system and bladder as well as fetal growth and placental function. Chromosomal analysis can be offered to patients, and premature rupture should be excluded.

The Royal Australian and New Zealand College of Obstetricians and Gynaecologists recommend a pregnancy with oligohydramnios be referred to a specialist Obstetrician. The American College of Obstetrics and Gynecology indicates that oligohydramnios may be an indication for induction of labour. The Society of Obstetricians and Gynaecologists of Canada states that antepartum testing may be beneficial.

Polyhydramnios
Polyhydramnios has been defined as an AFV of greater than 2000 mL. By ultrasound techniques, it has been estimated as an SDP greater than 8 cm, or 25 cm or above the 95th percentile for gestational age. Elevated amniotic fluid can be due to decreased absorption, overproduction, or be idiopathic. Decreased absorption typically results from a failure of fetal swallowing from etiologies such as tracheal atresia, tracheal or bowel obstruction, or neurologic abnormalities such as anencephaly. Chromosomal abnormalities, non-immune hydrops, and diabetes are also recognised reasons for polyhydramnios. Twin-twin transfusion syndrome (TTTS), another origin of hydramnios, will be covered elsewhere. Appropriate workup for this abnormal vital sign is review of the maternal history, possible repeat of 75 g glucose test, anatomic evaluation of the fetus, and assessment of fetal growth.

Older studies of pregnancies with polyhydramnios show an association with macrosomia, premature birth, cesarean delivery, non-reactive non-stress tests, perinatal morbidity, and congenital anomalies. More recent studies are conflicting. Increased rates of congenital malformations are consistently reported; however, perinatal morbidity and mortality vary, possibly due to the retrospective nature of some studies and possibly due to improvements in management of underlying conditions for polyhydramnios.

In a recent study of 788 infants with polyhydramnios, the rate of congenital malformations was 2.3% compared to 0.13% in those with normal AF. Those with no major congenital malformation diagnosis were at risk for the development of respiratory distress and hypoglycemia. One hundred eighteen pregnancies with polyhydramnios as defined by SDP > 8 cm with a normal fetal anatomical scan, 75 g glucose test, and TORCH serology, were studied retrospectively by Abele, et al. Eleven of those pregnancies had postnatal abnormalities identified with the majority being gastrointestinal atresia. Another prospective study looked at 7111 pregnancies over a two-year period starting in 2007. Only 50 patients were found to have polyhydramnios, but out of those 50, perinatal mortality was reported at 42%, with 31% of those being due to congenital anomalies.

Polyhydramnios has been linked to maternal pre-gestational and gestational diabetes and fetal macrosomia. A study of several ultrasound criteria (fetal adipose tissue, asymmetrical macrosomy, cardiac circumference and width, intervillar septum thickness, immature appearance of the placenta, and polyhydramnios) was performed by Perovic, et al. and found to have a sensitivity and specificity of 90.9% and 89.6% respectively. Isolated polyhydramnios has not been found to be a strong independent predictor of gestational diabetes. Known diabetics with polyhydramnios have been found to have an increased hemoglobin A1C indicative of poor diabetic control, however, there was no significant increase in adverse perinatal outcomes in those pregnancies apart from iatrogenic preterm deliveries and elective Cesarean sections not related to fetal macrosomia.

Management of pregnancies with polyhydramnios, if identified early, should include sonographic evaluation for anatomical causes, evaluation for maternal diabetes, TORCH serology testing, and consideration of isoimmunisation as a cause. If polyhydramnios is present with a growth restricted fetus, evaluation for chromosomal abnormalities should be undertaken. If polyhydramnios is a late presentation, all the above should be considered, excluding an anatomical evaluation as this should have already been completed. Antenatal testing should be initiated at 24 weeks gestation, although there is no consensus on what type of testing should be begun, or at what interval. This can be left to the discretion of the physician. There are no recommendations for induction of labor if discovered at term.

The Royal Australian and New Zealand College of Obstetricians and Gynaecologists recommend a pregnancy with polyhydramnios be referred to a specialist Obstetrician. Surprisingly, there are currently no practice management recommendations from the American College of Obstetrics (ACOG), Royal College of Obstetrics and Gynaecology (RCOG) or the Society of Obstetricians and Gynaecologists of Canada (SOGC) beyond the statement that antepartum testing may be beneficial by ACOG and SOGC.

Twins and amniotic fluid
When evaluating the health of the amniotic fluid in twin pregnancies, singleton growth curves currently provide the best predictors of adverse outcomes, and the evaluation of fluid with singleton nomograms is frequently used. This approach appears reasonable since in the only evaluation of AFV in third trimester diamniotic twin pregnancies the AFV of each sac was shown to be similar to that of normal singleton pregnancies. Morin and Lim performed a literature review with the Diagnostic Imaging Committee of the Society of Obstetricians and Gynaecologists of Canada and suggest that the SDP be used in each sac with standard definitions for singleton pregnancies, although they state that there is not enough evidence to suggest that one method is more predictive than the others on adverse pregnancy outcome. The AFI has been used to estimate amniotic fluid volume in twins; however, when using the summated AFI (measurement of all four quadrants as is done in singletons but without taking membrane placement into consideration) the measurement poorly predicts oligohydramnios and polyhydramnios.
Table 2: SDP percentiles in twins across gestation.

| EGA (weeks) | 2.5th | 5th | 10th | 50th | 90th | 95th | 97.5th | Twin sacs assessed |
|-------------|-------|-----|------|------|------|------|--------|-------------------|
| < 17        | 2.02  | 2.11| 2.54 | 3.96 | 6.48 | 6.86 | 6.88   | 22                |
| 17-19       | 2.13  | 2.14| 2.64 | 4.07 | 6.56 | 6.97 | 7.02   | 143               |
| 20-22       | 2.22  | 2.23| 2.79 | 4.22 | 6.60 | 7.04 | 7.21   | 276               |
| 23-25       | 2.23  | 2.28| 2.93 | 4.38 | 6.67 | 7.19 | 7.45   | 322               |
| 26-28       | 2.29  | 2.37| 3.06 | 4.53 | 6.80 | 7.44 | 7.79   | 390               |
| 29-31       | 2.43  | 2.46| 3.20 | 4.67 | 6.91 | 7.64 | 8.09   | 323               |
| 32-34       | 2.44  | 2.51| 3.32 | 4.80 | 6.98 | 7.76 | 8.27   | 283               |
| 35-37       | 2.41  | 2.54| 3.39 | 4.90 | 7.01 | 7.81 | 8.39   | 129               |

Reprinted from: Magann EF, Doherty DA, Ennen CS, Chauhan SP, Shields D, Gjesdal SM, Morrison JC. The ultrasound estimation of amniotic fluid volume in diamniotic twin pregnancies and prediction of peripartum outcomes. Am J Obstet Gynecol 2007; 196 (6): 570. e1-6. With permission from Elsevier.

In a single prospective observational study of 299 diamniotic twin pregnancies, SDP was found to be constant between 17 and 37 weeks with an increase in fetal labor intolerance and neonatal morbidity in twin pregnancies with hydramnios. Alternate fixed cutoffs of < 2.2 cm for a low amniotic fluid volume and > 7.5 cm for a high amniotic fluid volume were suggested in diamniotic twin pregnancies to identify pregnancies at risk for adverse intrapartum and neonatal outcomes.36

Hernandez, et al.37 performed a retrospective review of 1951 twin pregnancies, both dichorionic and monochorionic, with twin-twin transfusion syndrome excluded. Hydramnios was identified in 18% of pregnancies, defined as mild (SDP 8–9.9 cm), moderate (SDP 10–11.9 cm), and severe (SDP ≥ 12). Pregnancy outcomes were reviewed and hydramnios was not associated with preterm delivery, fetal growth restriction, NICU admission, or neonatal death in either twin pregnancy. The incidence of major anomalies became more common in increasing hydramnios in both monochorionic and dichorionic pregnancies with a prevalence of almost 20% in severe hydramnios. Severe hydramnios was significantly associated with stillbirth in monochorionic pregnancies (27% P < .001).

Twin-twin transfusion syndrome carries with it a 3–5 fold increase in mortality and morbidity.38 It is often uncovered with ultrasound evidence of discordant fetal weights and amniotic fluid in monochorionic/diamniotic twins, although it has been documented in monochorionic/monoamniotic pregnancies.39 When diagnosing TTTS by ultrasound, two criteria are required: monochorionic/diamniotic pregnancy and the presence of oligohydramnios in one sac and polyhydramnios in the other.100 It has been shown that if the difference of amniotic fluid in the two sacs fail to meet that criteria, but a subjective difference is noted, TTTS will develop in 15% of those pregnancies.101 The criteria for oligohydramnios and polyhydramnios follow singleton values. Growth discordance and selective growth restriction is sometimes associated with TTTS, but it is not a diagnostic criteria.100,102 Evaluation for TTTS in these pregnancies should begin in the second trimester with weekly surveillance if discordance is detected and every two weeks evaluation for concordant pregnancies.103 Selective intrauterine growth restricted (sIUGR) twins should be monitored with umbilical artery dopplers, keeping in mind that results will vary based on severity of sIUGR.102

Management of abnormal amniotic fluid volume in twin pregnancies does not vary much from singleton pregnancies with the single exception of twin-twin transfusion syndrome. Pregnanies with polyhydramnios-oligohydramnios sequence or those with diagnosed twin-twin transfusion syndrome should be referred to Maternal-Fetal Medicine for evaluation and potential treatment as indicated. The diagnosis of abnormal amniotic fluid in monoamniotic twins does not alter management of the pregnancy, as antenatal testing should begin at viability and delivery planned early.104 The discovery of abnormal fluid volumes may prompt earlier evaluation of causes such as twin-twin transfusion syndrome, IUGR, and congenital anomalies. Dichorionic/diamniotic twin pregnancies should be treated as described above for singleton pregnancies. In pregnancies where polyhydramnios is found in both sacs, or in a single shared amnion, fluid reduction may be necessary for maternal comfort, and preterm labor may be encountered more frequently.105

In summary, amniotic fluid is a highly complex and dynamic system that should be utilised in the interpretation of fetal wellbeing. Routine assessment of amniotic fluid volume has become commonplace with sonography and many options exist for its estimation. Practitioners should be familiar with all of the methods above, but choose one method for which to evaluate their patients, knowing the strengths and limitations of each. When abnormalities of fluid exist, appropriate workup to uncover the underlying etiology should be initiated as adverse fetal outcomes are sometimes associated with these variations from normalcy. Management options should be related to cause, if discovered. The ultimate goal of appropriate timing of delivery while reducing concurrent morbidity and mortality can be challenging, but continued research in this field will no doubt continue to elevate clinical practice to this goal.

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