



ISUOG VIRTUAL INTERNATIONAL SYMPOSIUM 2021

**State-of-the-art Ultrasound Imaging
in Obstetrics and Gynecology**

17-18 April 2021

REGISTER NOW ►

Using ultrasound together with other technologies to improve the lifelong health of women and babies

- Two streams of scientific content over two days, delivered through the ISUOG virtual platform which will exceed your expectations
- A mixture of lectures and practical, interactive training, including scan demonstrations, pattern recognition sessions and case report discussion
- Leading international and local experts in obstetrics, gynecology and imaging
- Live program delivered from 7:30 - 18:30 Calgary, Canada time (Mountain Daylight Time)
- Content available on Demand, at a time, pace and location to suit you until 17 May 2021
- All non-member registration fees include a 12-month ISUOG basic membership

Provisional program

Sessions will run simultaneously, providing two streams of content both days.

Highlights include:

- Obstetrics: the first trimester, beyond the routine mid-trimester fetal ultrasound scan, screening to improve pregnancy outcomes, fetal growth and health, ultrasound in labor, and more
- Gynecology: ectopic pregnancy, miscarriage, endometriosis, menopause, ovarian tumors, tubal and uterine pathology, and more
- Special sessions include advanced imaging/MRI, fetal therapy and COVID

The symposium will be co-chaired by:

*Jo-Ann Johnson (Canada),
Denise Pugash (Canada)*

Symposium Advisory Group

*Shabnam Bobdiwala (UK)
George Condous (Australia)
Karen Fung-Kee-Fung
(Canada)
Jon Hyett (Australia)
Simon Meagher (Australia)
Liona Poon (Hong Kong)
Angela Ranzini (USA)
Magdalena Sanz Cortes (USA)*

Who should attend?

This interactive course is designed for Maternal Fetal Medicine (MFMs), OB-GYNs, Radiologists, Sonographers, Geneticists, Researchers, Trainees/Residents and other maternity care providers. The program will appeal to a wide global audience, with a focus on North American educational needs.

See you ONLINE in 2021! For more information, please visit:
isuog.org/event/17th-isuog-international-symposium.html





Editorial

Pro forma for ultrasound reporting in suspected abnormally invasive placenta (AIP): an international consensus

Z. ALFIREVIC*, A.-W. TANG*,
S. L. COLLINS†, S. C. ROBSON‡ and
J. PALACIOS-JARAQUEMADA§, on behalf of
the Ad-hoc International AIP Expert Group

*Department of Women's and Children's Health, University of Liverpool, Liverpool, UK; †John Radcliffe Hospital, Oxford, UK; ‡University of Newcastle, Newcastle, UK; §University of Buenos Aires, Buenos Aires, Argentina

*Correspondence. (e-mail: zarko@liv.ac.uk)

Accurate antenatal diagnosis of an abnormally invasive placenta (AIP), allowing multidisciplinary management at the time of delivery, has been shown to improve maternal and fetal outcomes^{1–3}. AIP can be predicted as early as in the first trimester, by identifying cases of suspected Cesarean scar pregnancy (CSP), as there is evidence that CSP in the first trimester and AIP in the second and third trimesters may represent different stages of a similar pathology⁴. Grayscale ultrasonography, with or without color Doppler and performed both transabdominally and transvaginally, has been used widely for antenatal screening and diagnosis of AIP. Many signs have been suggested, with reports varying as to their sensitivity and specificity⁵. Most of these 'signs' are poorly defined and, consequently, it is difficult to assess which are the most robust. To address this, the European Working Group on AIP (EW-AIP) have produced a consensus proposal to standardize the ultrasound descriptions used to define each sign, published in this issue of the Journal⁶.

We assembled an international group of experts in the field with the specific aim of reaching an agreement regarding a standardized means of reporting ultrasound assessment of suspected AIP. If adopted by sonographers, clinicians and researchers worldwide, such a pro forma may facilitate better communication, and better evaluation of our diagnostic performance, in cases of suspected AIP.

The group of international experts comprised an e-mail discussion group ($n=50$) led by Jose Palacios Jaraquemada, members of the EW-AIP ($n=19$) and members of the ISUOG (International Society of Ultrasound in Obstetrics and Gynecology) Clinical Standards Committee ($n=7$). Each expert was asked to participate in a survey which involved completion of an online questionnaire to indicate what they believed should be included in the pro forma for reporting ultrasound assessment of suspected AIP.

The online questionnaire, created using Survey Monkey, included risk factors known to be associated with AIP and all commonly reported ultrasound signs and definitions related to the diagnosis of AIP^{5–11}. Ultrasound signs were divided into three subgroups according to modality: grayscale ultrasound, color Doppler and three-dimensional (3D) power Doppler. Each ultrasound sign in each subgroup had between one and six associated definitions reported in the published literature.

To each selected demographic characteristic and ultrasound sign we assigned three options: (i) definitely include in report; (ii) include optionally in report and (iii) do not include in report. The definitions for each ultrasound sign were also assigned three options: (i) include; (ii) do not include and (iii) unsure. Participants were also asked whether clinical interpretation and relevance of the ultrasound findings should be included in the report. Options for preferred method of reporting clinical interpretation included: (i) give probability of clinically significant AIP, (ii) state whether manual removal of placenta should be attempted, and (iii) give free text description to provide guidance to the local team. There was the opportunity to provide free text comments for each section. A reminder to complete the questionnaire was sent out after 2 weeks, and we allowed 4 weeks for a response.

All demographic characteristics and ultrasound signs for which >50% respondents selected 'definitely include in report' were incorporated into the standardized report, while those for which >50% respondents selected 'do not include in report' were excluded. For each ultrasound sign, the definitions for which >50% of respondents selected either 'include' or 'unsure' were kept for further evaluation. A second questionnaire was created for such items requiring further evaluation, in which respondents could specify first and second choice for definition of the ultrasound sign, and included additional suggestions from the free text comments, such as assessment for suspected parametrial involvement. For confirmation, we distributed a third and final round of the survey, with three domains, addressing: demographic and risk factors, ultrasound signs and clinical interpretation. At this round, consensus was sought from all participants that the ultrasound signs previously agreed on should be defined using the standardized descriptors proposed by the EW-AIP⁶.

There were 42 respondents in the first round of the survey (response rate, 55%). For all of the demographic characteristics, placental location and grayscale ultrasound parameters, and for all but one color Doppler parameter, >50% of respondents chose 'definitely include in report'. Only seven respondents thought that 3D power Doppler volumes should definitely be included and thus this criterion was excluded. All

but one respondent agreed that clinical interpretation should be included in the standardized report. The preferred option for reporting clinical relevance was as probability of clinically significant AIP. There were also suggestions to include reporting of assessment of parametrial involvement and the extent of AIP ('focal' or 'diffuse'), which were included as options in subsequent rounds.

In the second round, 28 (37% of the original group) responded. All demographic characteristics suggested additionally in this round were selected by a clear majority as to be definitely included in the report. In the third and final round (response rate, 42% of the original group ($n=32$)), agreement was obtained from all respondents regarding the description of ultrasound signs, and the majority (87%) preferred categorization of clinical relevance for significance of AIP into high, intermediate or low risk.

We propose that, when performing an ultrasound scan for suspected AIP, this set of ultrasound signs should always be reported. (See ultrasound report on next page, also supplied in pdf format online.) Using these standardized descriptors and this reporting pro forma should facilitate a systematic approach to the assessment of this rare condition. This should allow better comparison between diagnostic centers and enable prospective multicenter evaluation of the diagnostic performance of each sign, or combination of signs, for prediction of AIP, thus reducing the risk of serious perinatal complications and maternal morbidity.

We propose that, at present, 3D color Doppler should not be included in the standardized reporting of suspected AIP. However, centers with experience in this ultrasound modality should continue to evaluate it and report it optionally. This pro forma can also be further adapted for local use to include other findings, such as location of cord insertion and placenta, to assist in planning the operative technique for management of AIP.

The consensus to include clinical interpretation of ultrasound findings was somewhat unexpected. The panel felt that such interpretation is important in order to allow better planning of the intrapartum management within a multidisciplinary team. The current proposal is to have three different levels of suspected risk. It is anticipated that cases of low risk will receive standard care. How best to incorporate the other two levels of risk into clinical pathways

will depend on local circumstances, maternal view on future fertility and surgical and critical-care expertise.

Members of the Ad-hoc International AIP Expert Group

The Ad-hoc International AIP Expert Group comprised: Zarko Alfrevic, Michael A. Belfort, Amar Bhide, Thorsten Braun, Pavel Calda, Giuseppe Cali, Gihad Chalouhi, Kinga Chalubinski, Frederic Chantraine, Min Min Chou, Sally Collins, Ekaterini Domali, Ana Espinosa, Tullio Ghi, Markus Gonser, Jorge Hamer, Irene Hoesli, Andrew D. Hull, Eric Jauniaux, Boon H. Lim, Phillip Lim, Shigeki Matsubara, Felipe Moretti, Donal O'Brien, Per Olofsson, Jose Palacios-Jaraquemada, Pedro Pinto, Steve Robson, Loïc Sentilhes, Jin-Chung Shih, Alexandros Sotiriadis, Vedran Stefanovic, Minna Tikkanen, Ilan Timor-Tritsch, Boris Tutschek, Heleen van Beekhuizen, Oliver Vasilj, Constantin von Kaisenberg and Katharina von Weizsaecker.

REFERENCES

1. Eller AG, Bennett MA, Sharshiner M, Masheter C, Soisson AP, Dodson M, Silver RM. Maternal morbidity in cases of placenta accreta managed by a multidisciplinary care team compared with standard obstetric care. *Obstet Gynecol* 2011; 117: 331–337.
2. Tikkanen M, Paavonen J, Loukovaara M, Stefanovic V. Antenatal diagnosis of placenta accreta leads to reduced blood loss. *Acta Obstet Gynecol Scand* 2011; 90: 1140–1146.
3. Fitzpatrick KE, Sellers S, Spark P, Kurinczuk JJ, Brocklehurst P, Knight M. The management and outcomes of placenta accreta, increta, and percreta in the UK: a population-based descriptive study. *BJOG* 2014; 121: 62–70.
4. Timor-Tritsch IE, Monteagudo A, Cali G, Vintzileos A, Viscarello R, Al-Khan A, Zamudio S, Mayberry P, Cordoba MM, Dar P. Cesarean scar pregnancy is a precursor of morbidly adherent placenta. *Ultrasound Obstet Gynecol* 2014; 44: 346–353.
5. D'Antonio F, Iacovella C, Bhide A. Prenatal identification of invasive placentation using ultrasound: systematic review and meta-analysis. *Ultrasound Obstet Gynecol* 2013; 42: 509–517.
6. Collins SL, Ashcroft A, Braun T, Calda P, Langhoff-Roos J, Morel O, Stefanovic V, Tutschek B, Chantraine F, on behalf of the European Working Group on Abnormally Invasive Placenta (EW-AIP). Proposal for standardized ultrasound descriptors of abnormally invasive placenta (AIP). *Ultrasound Obstet Gynecol* 2016; 47: 271–275.
7. Cali G, Giambanco L, Puccio G, Forlani F. Morbidly adherent placenta: evaluation of ultrasound diagnostic criteria and differentiation of placenta accreta from percreta. *Ultrasound Obstet Gynecol* 2013; 41: 406–412.
8. Chalubinski KM, Pils S, Klein K, Seemann R, Speiser P, Langer M, Ott J. Prenatal sonography can predict degree of placental invasion. *Ultrasound Obstet Gynecol* 2013; 42: 518–524.
9. Chou MM, Ho ES, Lee YH. Prenatal diagnosis of placenta previa accreta by transabdominal color Doppler ultrasound. *Ultrasound Obstet Gynecol* 2000; 15: 28–35.
10. Comstock CH, Bronsteen RA. The antenatal diagnosis of placenta accreta. *BJOG* 2014; 121: 171–181.
11. Shih JC, Palacios Jaraquemada JM, Su YN, Shyu MK, Lin CH, Lin SY, Lee CN. Role of three-dimensional power Doppler in the antenatal diagnosis of placenta accreta: comparison with gray-scale and color Doppler techniques. *Ultrasound Obstet Gynecol* 2009; 33: 193–203.

A sample pro forma is shown on next page



An electronic version of the pro forma is provided in the online version of this article.

SUSPECTED ABNORMALLY INVASIVE PLACENTA (AIP)**Ultrasound report****Demographics and Risk Factors**

Date: __/__/____

Gestational age: __ weeks __ days

Parity ☐

Mode of conception:

Spontaneous ☐IVF ☐Number of previous CS ☐Number of classical CS ☐Number of previous surgical evacuations (including TOP) ☐

Was Cesarean scar pregnancy suspected/diagnosed in first trimester?

Yes ☐No ☐Not known ☐

Previous uterine surgery (e.g. myomectomy, endometrial ablation)

Yes ☐No ☐Not known ☐

History of AIP

Yes ☐No ☐Not known ☐**Placenta previa on ultrasound**

If yes: Anterior placenta previa

< 2 cm from internal os ☐Covering internal os ☐

Posterior placenta previa

< 2 cm from internal os ☐Covering internal os ☐**Ultrasound Signs**

Cervical length (without funnel or placental tissue)	mm		
Grayscale ultrasound parameters and definition	Yes	No	Unsure
Loss of 'clear zone' - Loss, or irregularity, of hypoechoic plane in myometrium underneath placental bed ('clear zone')			
Myometrial thinning - Thinning of myometrium overlying placenta to <1mm or undetectable			
Abnormal placental lacunae - Presence of numerous lacunae including some that are large and irregular, often containing turbulent flow visible on grayscale imaging			
Bladder wall interruption - Loss or interruption of bright bladder wall (hyperechoic band or 'line' between uterine serosa and bladder lumen)			
Placental bulge - Deviation of uterine serosa away from expected plane, caused by abnormal bulge of placental tissue into neighboring organ, typically bladder; uterine serosa appears intact but outline shape is distorted			
Focal exophytic mass - Placental tissue seen breaking through uterine serosa and extending beyond it; most often seen inside filled urinary bladder			
Color Doppler ultrasound parameters and definition	Yes	No	Unsure
Uterovesical hypervascularity - Striking amount of color Doppler signal seen between myometrium and posterior wall of bladder; this sign probably indicates numerous, closely packed, tortuous vessels in that region (demonstrating multidirectional flow and aliasing artifact)			
Subplacental hypervascularity - Striking amount of color Doppler signal seen in placental bed; this sign probably indicates numerous, closely packed, tortuous vessels in that region (demonstrating multidirectional flow and aliasing artifact)			
Bridging vessels - Vessels appearing to extend from placenta, across myometrium and beyond serosa into bladder or other organs; often running perpendicular to myometrium			
Placental lacunae feeder vessels - Vessels with high-velocity blood flow leading from myometrium into placental lacunae, causing turbulence upon entry			
Parametrial involvement	Yes	No	Unsure
- Suspicion of invasion into parametrium			

Clinical Significance of Ultrasound Findings

Probability of clinically significant AIP

High ☐Intermediate ☐Low ☐

Extent of AIP

Focal ☐Diffuse ☐