Placental Pathology in Early-Onset and Late-Onset Fetal Growth Restriction

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Key Words
Placental pathology · Growth restriction · Pre-eclampsia

Abstract
Several histopathological features are found more frequently in placentas from pregnancies complicated by fetal growth restriction (FGR), including villous infarction, maternal vascular changes and villous morphological alterations, although around one quarter of placentas associated with FGR lack any morphological abnormality on routine examination. Since similar changes may also affect clinically uncomplicated pregnancies, the positive predictive value of such findings for pathological FGR in an unselected case remains low. However, the pattern of placental pathologies varies with clinical subgroup. The combination of placental bed and parenchymal lesions in FGR with abnormal uterine artery Doppler velocimetry is essentially identical to preterm pre-eclampsia (PET), and there is an association between FGR with abnormal umbilical artery Doppler findings and lesions of fetal stem arteries and terminal villous hypovascularity. Conversely, placentas from pregnancies complicated by PET or FGR presenting at or near term have a significantly lower frequency of histological abnormalities compared to early-onset disease and absence of a distinctive biochemical profile. The histological placental findings in FGR are therefore varied, from morphologically unremarkable through to severe uteroplacental vasculopathy, with no single pathological feature associated with high sensitivity or specificity. Severe early-onset FGR, overlapping with severe early-onset PET, is mainly associated with features of impaired maternal uteroplacental perfusion secondary to defective extravillous trophoblast invasion, and its consequences. Late-onset FGR probably represents a more heterogeneous group with less characteristic pathological changes. Future research using histopathological assessment of aggregated data from multiple studies into larger datasets with centralised pathology review will allow delineation of distinctive clinicopathological associations and further understanding of pathophysiology.

Introduction
There has been increasing interest in defining patterns of placental pathology associated with intrauterine fetal growth restriction (FGR), with progressive clinical refinement of subgroups of fetuses that are small for gestational age (SGA), with and without other features of FGR,
such as abnormal Doppler ultrasound assessment and maternal serum markers such as placental growth factor and fms-like tyrosine kinase-1. Here, we review the available data on documented histopathological features of the placenta associated with early versus late-onset FGR, their significance, and their relationship to findings in pre-eclampsia (PET). We also infer possible pathogenetic implications of these findings and speculate on future work to better define, understand and manage these conditions.

**General Placental Histopathological Features Associated with FGR**

There are numerous publications describing a range of morphological features in placentas from pregnancies complicated by FGR (or more specifically SGA, since the vast majority of studies, especially early studies, defined the ‘FGR’ group simply based on a fetal size below a given centile value rather than definitive features of abnormal fetal growth) [1]. Despite this large descriptive literature, no reliable and pathognomonic features for either FGR/SGA or specific aetiologies are documented, but several lesions are more common compared to controls in the majority of studies, and hence appear associated with the clinical phenotype of FGR/SGA. However, comparison of findings between different studies, and more detailed analysis of the data, is often not straightforward, due to several reasons. There is biological variation in the disease process due to several complex underlying mechanisms resulting in a final common ‘phenotype’ of FGR, and individual patient responses may differ in response to a given underlying cause. There is variation in placental morphology due to the fact that FGR occurs in parallel with many features of growth and development of the fetoplacental unit, and there may be geographical variation between different areas of the same placenta in relation to sampling adequacy. In addition, studies may have used different sampling protocols, with associated bias and errors, and the extent of such regional variations in placental histological changes remains undefined. Studies have used different definitions and thresholds for recognising changes as ‘lesions’, many of which are based on subjective assessment by an observer, such that variation in interpretation between studies is impossible to control. However, the greatest problem is that the study populations examined are relatively poorly defined and potentially heterogeneous. For example, any arbitrary fixed proportion of a population, such as the smallest 5% of fetuses, will include a mixture of pathologically growth-restricted fetuses and those who are constitutively small, with many studies defining inclusion based on fetal size rather than growth. However, the progressive refinement of clinical subgrouping for inclusion, through use of Doppler studies and plasma biomarkers, has facilitated an improved understanding of FGR pathology, and emerging well-defined subgroups of placental abnormalities are now becoming increasingly recognised.

Despite these potential difficulties in interpretation, there are patterns of placental histological changes that are well reported and common to most studies of the pathology of FGR. First, it should be noted that, overall, approximately one quarter of placentas associated with FGR/SGA (however defined) lack any morphological abnormality on routine macroscopic and histological examination [1]. In those cases with lesions present, the most frequent macroscopic abnormality that occurs with increased frequency in FGR placentas is patchy placental infarction, being present in around 25% of term FGR versus 10% of controls [2–14]. This finding illustrates a common theme when interpreting placental findings in FGR, that the frequency of several ‘lesions’ is increased, but since such lesions are only present in a minority of affected cases and are also found in some normal controls, and since non-FGR is far more frequent than FGR in the population, the positive predictive value of such findings for pathological FGR in any given unselected case will be very low.

Similarly, several microscopic abnormalities have been reported more frequently in FGR placentas, none of which are present in the majority of affected cases in any study, and almost all of which may be encountered in other non-FGR pregnancies, including [2–7, 9–25]: villous infarcts (an example from a placenta in a case of FGR is shown in figure 1a, b), placental abruption/retroplacental haemorrhage, villous morphological abnormalities suggestive of reduced utero-placental and/or fetoplacental flow (‘hypoxic’ lesion) such as syncytiotrophoblast ‘knots’, excess cytrophoblast cells, thickened trophoblastic basement membrane, villous fibrosis, hypovascular terminal villi, reduced villous volume, reduced intervillous space, and non-specific inflammatory lesions [villitis of unknown aetiology (VUE)]. In addition, histological features indicating defective remodelling of spiral arteries into utero-placental vessels may be identified, such as inadequate ‘physiological change’, fibrinoid necrosis and acute atherosis (a dense perivascular lymphocytic infiltrate with intimal arterial foamy macrophages; see fig. 2a, b) [26–28].

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Association of Placental Pathologies with Clinical Subgroups

The use of well-defined specific clinical subgroups of definite pathological FGR should allow determination of any specific patterns of placental abnormalities, and hence, suggest pathophysiological pathways. The introduction of uterine artery (UtA) Doppler studies identified FGR cases with evidence of maternal uterine malperfusion, with placental bed biopsies in these cases demonstrating defective spiral artery remodelling, including inadequate physiological change and fibrinoid necrosis/atherosclerosis [27, 29]. Ferrazzi et al. [30] studied placentas from women with FGR and reported that ‘extensive hy-

Fig. 1. Placental infarct, secondary to maternal vascular underperfusion. At low magnification (a), the infarct can be seen as an area with severe reduction of the intervillous space and aggregation of villi. In the area surrounding the infarct the intervillous space is increased and there are several very small villi. At higher magnification (b), villous aggregation can be better appreciated, and the syncytiotrophoblastic show various stages of degeneration, with pyknosis, karyorrhexis and replacement by fibrin-like eosinophilic amorphous (hyaline) material.

Fig. 2. Lesions of acute atherosis at low (a) and high (b) magnification. The arterial walls are irregular, necrotic and replaced with fibrin-like eosinophilic material. In later stages, there are greater numbers of lipid-laden macrophages (lipophages) and there is a perivascular mononuclear inflammatory infiltrate. These lesions occur in vessels with inadequate extravillous trophoblast invasion.
poxic villous damage’ (more than 30% of terminal villi with one or more of the following: syncytiotrophoblast knotting, excess cytotrophoblast cells, thickened trophoblastic basement membrane, villous fibrosis, hypovascularity), non-peripheral infarcts involving more than 5% of placental parenchyma, and placental abruption were found only in those FGR cases with abnormal UtA velocimetry, regardless of the presence or absence of pregnancy-induced hypertension (PIH). UtA velocimetry is therefore a marker for defective remodelling of spiral arteries with consequent placental malperfusion and associated impaired fetal growth.

**Abnormal UtA Doppler Is Associated with a Common Set of Placental Ischaemic Lesions in FGR and Preterm PET**

The combination of placental bed and parenchymal lesions described in FGR with abnormal UtA velocimetry is essentially identical to lesions described in preterm PET, particularly when subsets of preterm PET and preterm FGR are compared [12, 15, 26, 28–37]. Indeed, abnormally raised UtA resistance is associated with defective trophoblast migration on placental bed biopsies independent of the presence or absence of the clinical syndrome of PIH [38].

Whilst placental findings in preterm PET and FGR are therefore similar, defining the extent of similarity and its dependence on abnormal UtA Doppler would be best achieved by a quantitative meta-analysis, but this is difficult due to the reasons described in the Introduction. Nevertheless, two studies [12, 35] were performed by the same group on the same study population (delivering between 22 and 32 weeks’ gestation); including 48 cases with FGR (<10th centile); 65% had PET and in these cases there were significantly greater frequencies of villous fibrosis and hypovascularity. Other lesions occurred at comparable frequency between groups. Table 1 shows a detailed comparison of the pathological findings reported in the two studies. Subsequent data suggests that these features may be associated with reduced end-diastolic flow (EDF) on umbilical artery (UA) Doppler studies [39]. The findings imply a common pathophysiological pathway involving impaired uteroplacental underperfusion in which PET may become apparent in a subset of FGR with placental bed and placental changes.

However, whilst this association is established, the relationship is complex. Regression modelling showed that the abnormal placental histological findings accounted for only 34% of variation in fetal growth, indicating that histological measures of ‘severity’ of placental changes do not correlate well with FGR severity [12]. For cases with PET, the predictive value was only around 10%, with histological ‘severity’ of placental lesions poorly associated with clinical severity of disease [35]. Thus, whilst there is a pattern of placental histological lesions that indicates failure of trophoblast migration, there are no features which are reliably associated with the clinical severity of either PET or FGR.

**UA Doppler Abnormalities**

Abnormal UA Doppler indices, particularly absent EDF or reduced EDF, are associated with fetal hypoxia and increased perinatal mortality [30, 40–45]. Such UA abnormalities are progressive and usually accompanied by abnormal UtA velocimetry, which typically develops well before the UA abnormality [45–47]. Within the group of non-hypertensive FGR with abnormal UtA Doppler, UA Doppler abnormalities preferentially identify cases with a high risk of perinatal mortality or morbidity (34% perinatal mortality with abnormal UA versus no cases of perinatal mortality with normal UA) [30]. Therefore, we can assess whether specific placental lesions are associated with abnormal UA Doppler indices. In the above study, there was no significant difference between groups, but many changes examined reflected abnormal uteroplacental flow rather than aspects of fetoplacental flow, such as fetal stem artery abnormalities, which were not specifically examined. Other studies examining pathology of abnormal UA Doppler have reported an association between FGR and lesions of the fetal stem arteries particularly, including vessel wall thickening/hypertrophy, apparent reduction of vessel lumina and lumenal herniation [17, 39, 48, 49]. There appears to be a relationship between the UA pulsatility index and both the proportion of abnormal fetal stem arteries [49] and stem vessel wall thickness [50, 51] (see fig. 3). Such stem artery lesions are reported in 90% of cases with absent EDF (versus 35% of FGR with present EDF [39]). Interestingly, in cases of reduced EDF, other lesions appear significantly more frequent than in FGR with either normal or absent EDF, including terminal villous fibrosis, hypovascularity, thin-walled stem arteries and haemorrhagic endovasculosis, but the pathological significance of these lesions specifically in relation to flow dynamics is uncertain [39].

The association of UA Doppler abnormalities with small, fibrotic and hypovascular terminal villi was initially proposed to be due to a primary defect in terminal villus development [21, 52–54]. However, it is difficult to explain the coexistence of other changes found in these cas-
Table 1. Comparison of pathological features in placentas from cases of preterm symmetrical FGR and preterm PET from two comparable studies [12, 35]

<table>
<thead>
<tr>
<th>Pathological Feature</th>
<th>Preterm symmetrical FGR, %</th>
<th>Preterm PET, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of cases</td>
<td>48</td>
<td>76</td>
</tr>
<tr>
<td>Acute inflammation: amnion*</td>
<td>8.0</td>
<td>0</td>
</tr>
<tr>
<td>Acute inflammation: fetal vessels</td>
<td>10.4</td>
<td>not reported</td>
</tr>
<tr>
<td>Chronic villitis</td>
<td>12.6</td>
<td>20.0</td>
</tr>
<tr>
<td>Chronic intervillitis</td>
<td>Not reported</td>
<td>not reported</td>
</tr>
<tr>
<td>Maternal vascular thrombosis: multiple occlusive lesions</td>
<td>33.3</td>
<td>29.0</td>
</tr>
<tr>
<td>Maternal vascular thrombosis: single occlusive lesion</td>
<td>14.6</td>
<td>not reported</td>
</tr>
<tr>
<td>Maternal vascular thrombosis: non-occlusive</td>
<td>6.3</td>
<td>not reported</td>
</tr>
<tr>
<td>Chronic maternal vasculitis: &gt;1 vessel</td>
<td>Not reported</td>
<td>10.0</td>
</tr>
<tr>
<td>Chronic maternal vasculitis: 1 vessel only</td>
<td>Not reported</td>
<td>20.0</td>
</tr>
<tr>
<td>Absence of spiral artery physiological changes</td>
<td>Not reported</td>
<td>30.0</td>
</tr>
</tbody>
</table>

| Fibrinoid necrosis and/or atherosis of spiral arteries:   |
|-----------------------------------------------------------|----------------------------|----------------|
| multiple arteries                                         | 10.4                       | 24.0           |
| 1 artery only                                             | 12.5                       |                |

| Villous fibrosis: severe*                                  | 27.1                       | 82.0           |
| Villous fibrosis: mild                                     | 39.6                       | not reported   |
| Villous hypovascularity: severe*                           | 25.0                       | 72.0           |
| Villous hypovascularity: mild                              | 39.6                       | not reported   |
| Villous infarction: multiple                               | 37.5                       | 45.0           |
| Villous infarction: single                                 | 20.8                       | 24.0 (‘focal’) |
| Villous infarction location: peripheral 10%                | 10.4                       | 11.0           |
| Villous infarction location: central 90%                   | 29.2                       | 36.0           |
| Villous infarction location: peripheral & central          | 16.7                       | 18.0           |
| Syncytiotrophoblast knotting: severely increased           | 39.0                       | 40.0           |
| Syncytiotrophoblast knotting: mildly increased             | 43.8                       | not reported   |
| Proliferation of cytotrophoblast cells: severe            | 36.9                       | 42.0           |
| Proliferation of cytotrophoblast cells: mild              | 27.1                       | not reported   |
| Abruptio: frank                                            | 10.4                       | 12.0           |
| Abruptio: consistent with                                  | 31.3                       | 37.0           |
| Perivillous fibrin deposition: severely increased          | 45.8                       | 43.0           |
| Perivillous fibrin deposition: mildly increased            | 41.7                       | not reported   |
| Fetal vascular thrombosis                                  | not reported               | 4.0            |
| ‘Haemorrhagic endovasculitis’                              | not reported               | 12.0           |
| Avascular terminal villi: multifocal                       | 18.8                       | 20.0           |
| Avascular terminal villi: focal                           | 29.2                       | 25.0           |

| Nucleated erythrocytes: numerous                          | 17.5                       | 76.0           |
| Nucleated erythrocytes: present, not numerous             | 56.3                       |                |

* Parameters with a significant difference between preterm FGR and preterm PET groups, with respective percentages in bold. Percentages are italicised if significantly different from control population used in original study. In the preterm symmetrical FGR study [12], the control population was preterm placentas from pregnancies without FGR (340 cases). In the preterm PET study [35], the control population (353 cases) was a combination of spontaneous premature rupture of membranes (192 cases) and premature labour with intact membranes (161 cases).
es as consequences of such a primary abnormality, and several other lines of evidence suggest that this is erroneous. Changes in the stem villous arteries are strongly suggestive of a response to chronic vasoconstriction [45], and develop in parallel with a progressive UA Doppler abnormality [45, 55], and similar downstream changes in terminal villi are present in the territory of an upstream fetal vessel with occlusive thrombus [56, 57]. Importantly, in most cases there is also defective uteroplacental perfusion with associated UTA Doppler studies preceding the UA Doppler changes [46, 47]. Furthermore, experimental evidence in sheep demonstrates a rapid increase in UA resistance upon reducing maternal placental perfusion [58, 59] and there is experimental evidence of rapid and reversible fetal vasoconstriction in the human placenta in response to hypoxia [60], in an analogous manner to the lung, involving inhibition of potassium channels [61]. Similar to the pulmonary circulation, hypoxia-induced and K⁺ channel-mediated depolarisation of vascular smooth muscle cells activates voltage-gated L-type gated Ca²⁺ channels, which provides cytosolic Ca²⁺ for activating the contractile apparatus [62]. Such hypoxic vasoconstriction allows autoregulation to optimise fetomaternal flow matching, but when widespread results in increased global flow resistance. However, experimental demonstrations of possible fetal hypoxic vasoconstriction in the placenta have been made with oxygen tensions that are higher than those in the placenta in vivo [60–62], and to the best of our knowledge, no studies have reported the effect of hypoxia at levels approximating those in the placenta, on preparations including the entire fetal subchorionic vasculature. There have been reports studying placental levels of hypoxia on isolated chorionic plate arteries and veins with diameters of 268 ± 13 μm (arteries) and 266 ± 19 μm (veins) at 2% oxygen [63, 64], showing increased chorionic vein vasodilation in response to nitric oxide, but no convincing hypoxic chorionic artery vasoconstriction. However, the isolated vessels used in these studies were too large to represent stem villus vessels, whose diameter is typically in the range of 10–100 μm in both normal and FGR pregnancies [54]. Regulation of stem venous tone may also occur to maintain water balance [45, 65–67]. To adequately test these hypotheses, one would need an experimental system that can maintain placental cotyledons at physiological levels of oxygenation and also truly representative levels of hypoxia, while allowing for independent variation of intervillous perfusion pressure and for measurement of vascular tone across all the different vessel types (arterial, venous, different calibres). To date, such a system has not been reported.

More recently, the similarity of the changes in the stem villus arteries between severe FGR and preterm PET was further demonstrated by the discovery that in both clinical situations the smooth muscle cells in the stem villus arteries show markedly reduced expression of cystathionine-lyase, responsible for synthesis of the potent vasodilator hydrogen sulphide (H₂S). Its reduced expression in the fetal villous resistance vessels supports the view that there is vasoconstriction in the fetal stem villi in association with abnormal umbilical Doppler profile [68]. Interestingly, cystathionine-lyase is not only downregulated in cases with abnormal UA Doppler profile, but shows increased expression in PET with normal UA Doppler findings compared to non-PET controls, although expression in both groups appeared heterogeneous by Western blotting. In addition, the PET cases with normal UA Doppler findings were preterm cases (delivered at 29.5 ± 1.5 weeks), while the control cases were delivered at term (39 ± 0.6 weeks) [68].

Other Placental Histological Lesions Associated with FGR

In addition to the typical uteroplacental and fetoplacental ‘vascular’ lesions described above, several studies have reported an excess frequency of VUE in FGR, with widely varying rates ranging from 8 to 90% [5, 11, 13, 16,
These studies report a generally low positive immune reaction, with consequently increased risk of VUE recurrence [73–77]. VUE is approximately twice as common and more diffuse in multigravidity [78], and it has a higher than expected concordance in twin pregnancies [79]. The pathological features of VUE are well reviewed elsewhere [80], although there is considerable variation amongst ‘expert’ pathologists in recognising this histological lesion [81]. Nevertheless, there is a consistently increased frequency of VUE in FGR, and also in placentas from cases of PIH and PET, with and without associated FGR, despite the majority of cases being associated with normal pregnancy outcome [82, 83]. The precise mechanism by which patchy inflammation may lead to impaired fetal growth remains undetermined.

Pathological Features

Multiple studies have reported that placentas from PET presenting at term have a significantly lower frequency of histological lesions associated with impaired maternal uteroplacental perfusion compared to early-onset disease. In an early study of 158 PET placentas there was an inverse relationship between gestation at delivery and the frequency of non-peripheral infarcts, villous hypermaturity and decidual arteriopathy [85] and another study of 37 PET placentas confirmed a significantly reduced frequency of non-peripheral infarcts in cases presenting near term [86]. A large study (with >900 PET cases and >7,000 controls) also reported an inverse relationship between the frequency of placental lesions of maternal underperfusion (villous infarction, villous agglutination, increased syncytial knots, increased interstitial fibrin, distal villous hypoplasia, persistent vascularisation of basal plate arteries, mural hypertrophy of decidual arterioles, acute atherosis), and gestational age. Despite this inverse relationship, even in PET delivered at or beyond 37 weeks’ gestation, the frequency of maternal underperfusion lesions was relatively high at around 30% [87]. In another series of >1,200 PET placentas, a comparable figure of 26% was reported [88]; in this study, the authors also showed an association of low placental weight (<10th centile) with preterm PET but a converse association of relatively increased placental weight with term PET. What has not been established is any volumetric or proportional extent of placental lesions and how these vary with gestational age at delivery in PET or FGR, since this would require histological study of the entire placenta. In addition, none of these studies reported blinding of the reporting pathologist to the clinical information raising the possibility of reporting bias in retrospective series.

UtA Doppler Profiles

UtA Doppler abnormalities in the late second trimester are associated with a greater risk of both subsequent PET and FGR, identifying approximately 40% of PET and 20% of FGR cases overall, but with the sensitivity for both inversely related to the gestational age at delivery [89, 90]. These studies report a generally low positive predictive value in screening populations, since the majority (70–90%) of women with UtA Doppler abnormality on routine screening will not develop PET or FGR [90]. There is no specific study examining histological findings from placentas associated with second trimester UtA Doppler abnormalities and normal versus abnormal fetal outcomes, to determine the relative associations of...
vascular haemodynamic abnormalities versus clinical disease and histological features. When performed in the first trimester, UtA Doppler screening demonstrates a similar inverse relationship between sensitivity and gestational age at delivery, with screening abnormalities significantly associated with preterm PET but not PET at term [91, 92].

Abnormal UtA Doppler studies in early pregnancy are therefore more sensitive for identifying women at risk of early-onset preterm PET/FGR, but the majority with an abnormal screen do not develop disease. This indicates that even when UtA flow is abnormal at around the normal time of completion of the second wave of trophoblast invasion, most women appear to have compensatory mechanisms for subsequent uteroplacental flow maintenance. A considerable number of cases of PET develop later in gestation, at or around term [87, 88, 90, 93], and these are predominantly those without significant FGR, who are also much less likely to be predicted by early UtA Doppler screening. (For example, 93% sensitivity for PET with FGR <32 weeks, versus 14% for PET without FGR delivered at or after 38 weeks, and 50% for PET with FGR delivered at or after 38 weeks.) These findings either represent that a ‘normal’ UtA Doppler profile by population data may not be normal for an individual, or that there is a different pathological process leading to at least a subset of term PET, or a combination of explanations. Studies of large cohorts of women with serial UtA Doppler assessments could theoretically address this issue but most current studies, for pragmatic reasons, are based on cross-sectional data from a dynamic developmental process requiring cut-offs derived from population distributions. Blinded standardised histological placental studies in these cohorts with subsequent correlation to Doppler profile evolution patterns as well as to outcome, would allow assessment of associated pathological features.

Abnormal Maternal Serum Biochemistry and Placental Pathology

There have been several reports of maternal serum biochemical (MSB) differences between early-onset and late-onset PET, which largely reflect the differences noted previously on placental pathological findings and UtA Doppler studies. For example, analysis of serum levels of a number of anti-angiogenic factors, including soluble fms-like tyrosine kinase-1, soluble endoglin and soluble vascular endothelial growth factor receptor-1, and pro-angiogenic factors such as placental growth factor shows an association between preterm PET and high levels of anti-angiogenic factors together with low levels of placental growth factor [94–100]. Those with an ‘angiogenic’ profile are associated with earlier preterm delivery and adverse outcomes, while non-angiogenic cases are not [100].

The subset of early-onset PET with FGR can therefore be largely defined by the presence of uteroplacental vascular lesions in the placenta, UtA Doppler abnormalities and an anti-angiogenic MSB profile. However, conversely, term PET without FGR demonstrates a general paucity of uteroplacental vascular lesions and absence of a distinctive MSB profile. Within this group there may be some who adapt after an early tendency for uteroplacental underperfusion, and another subset with a similar underlying pathophysiology to more severe cases, but who develop the clinical syndrome more slowly.

Placental Pathology of FGR at Term (‘Late-Onset’ FGR)

In keeping with the findings discussed above for PET, whilst placentas from FGR fetuses delivered at term have significantly increased frequencies of uteroplacental vascular lesions (especially infarcts) compared to normal controls, the incidence of such lesions is much lower than in preterm FGR, and many cases are histologically unremarkable [11, 39, 101]. Furthermore, it has been reported that compared to normal term pregnancies, placentas from FGR at term may have an increased incidence of other villous lesions including fibrosis, hypovascularity and avascularity, suggestive of fetal thrombotic events [102], although these changes had not been described in previous studies. In addition, in a minority of cases there is an association with other, non-vascular, pathologies such as VUE [19, 103].

FGR at term may therefore include a group of pregnancies with a similar uteroplacental perfusion defect to that seen with early-onset FGR, but with a less severe phenotype, or other changes which may suggest growth restriction mediated via other mechanisms. Interestingly, the majority of this group does not appear to overlap clinically with PET at term, in which the majority are not associated with FGR, despite the fact that both PET/PIH and FGR at term may show similar ‘non-specific’ placental findings such as VUE.
Conclusions and Future Directions

The histological placental findings in FGR are varied, from morphologically unremarkable through to severe uteroplacental vasculopathy, with no single pathological feature associated with high sensitivity or specificity. This is in part because the condition (FGR) is likely heterogeneous, but also because the extent and distribution of disease in the placenta is uneven and there are technical and practical limitations to the methods of routine study of this organ for use in large series. Nevertheless, it is clear that those with severe early-onset FGR, overlapping with severe early-onset PET, are mainly associated with both clinical and histological features of impaired maternal uteroplacental perfusion secondary to defective extravillous trophoblast invasion, and its consequences. Their histopathological findings generally reflect abnormal uteroplacental flow with secondary chronic fetal vascular vasoconstriction and distal villous changes, reflected in abnormal UtA and UA Doppler profiles. Cases of later onset FGR probably represent a more heterogeneous group, are delivered closer to term, and overlap much less with PET at term. These cases may aetiollogically represent both a subset of the group with typically more severe disease, in which, for unknown reasons, the effect on the fetus (FGR) and the mother (PET) develops more slowly and/or later and/or to a lesser extent, and those with different aetologies or mechanisms.

There is room for greater understanding of the mechanisms of FGR/PET based on integration of clinical and pathological data. The parameters that are assessed clinically, biochemically, sonographically and pathologically are subject to substantial variation within a population, only some of which is related to the disease. It is important that future prospective studies of cohorts undergoing serial assessments for FGR have placental examination included into study protocols.

Pathological examination of the placenta remains subject to technological as well as cost and time limitations. It is challenging to interpret the significance of lesions in an organ with large, but variable and imperfectly defined functional reserve. However, current practice can be improved by reporting pathological features in a standardised manner based on published criteria such as maternal vascular underperfusion [104], fetal vascular obstructive lesions [37] and amniotic fluid infection [105], with detailed descriptive and morphometric assessment for study purposes. Furthermore, it is now technically possible for studies using histopathological assessment to upload high-resolution whole-slide images of all material used in a study to a central repository, in a manner analogous to microarray and sequencing data that is deposited in NCBI’s GEO [106, 107]. This suggestion is not new [108], but with the increasing availability and reducing cost of whole slide imaging scanners and low-cost data storage facilities, it is becoming more feasible to implement and would greatly facilitate the aggregation of data from multiple centres into larger studies with centralised pathology review.

References


Placenta in Early and Late FGR


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