Lecture - Placenta Development

Introduction

This lecture is an introduction to the development and functions of the placenta.

The placenta (Greek, plakuos = flat cake) named on the basis of this organ's appearance. The placenta a mateno-fetal organ which begins developing at implantation of the blastocyst and is delivered with the fetus at birth. Only recently have we begun to understand the many different functions this organ carries out in addition to its role in embryonic nutrition.

If you have time after working through this online lecture page, have a look at the Medicine practical class, 7 pages the students work through in 2 hours.

The placenta and placental blood at birth has recently been seen as a new source for stem cells in bone marrow replacement therapy in many diseases. The topic of stem cells will be covered later in this current course.

Lecture Objectives

- Understanding of placental villi development
- Understanding of placental structure
- Understanding of placental functions
Brief understanding of placental abnormalities

Lecture Resources

References

Movies

Nutrition

- Histiotrophic nutrition - describes early placental development and the form of initial transfer of nutrition from maternal to embryo.
- Haemotrophic nutrition - describes the later blood-borne nutrition.

Fetal Membranes

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Villi Stages
Primary villi

**Week 2** - first stage of chorionic villi development, trophoblastic shell cells (syncitiotrophoblasts and cytotrophoblasts) form finger-like extensions into maternal decidua.

Secondary villi

**Week 3** - second stage of chorionic villi development, extraembryonic mesoderm grows into villi, covers entire surface of chorionic sac.

Tertiary villi

**Week 4** - third stage of chorionic villi development, mesenchyme differentiates into blood vessels and cells, forms arteriocapillary network, fuse with placental vessels, developing in connecting stalk. Basal region will form chorionic plate.

**Chorionoic Villi Location**

Historic - Semi-diagramatic sagittal section of human uterus containing an embryo. Note the villi locations and the names for different maternal decidual regions.
Originally cover entire chorionic surface and become restricted to decidua basalis region forming 2 regions:

**Week 4 - Carnegie stage 11**

- **Frondosum** - "leafy" where villi are mainly located
- **Capsularis** - smooth chorion, where villi are absent or not abundant

**Week 5 - Carnegie stage 14**
Villi Terms

- **primary villi** - week 2, first stage of chorionic villi development, trophoblastic shell cells (syncitiotrophoblasts and cytotrophoblasts) form finger-like extensions into maternal decidua.

- **secondary villi** - week 3, second stage of chorionic villi development, extraembryonic mesoderm grows into villi, covers
• tertiary villi - third stage of chorionic villi development, mesenchyme differentiates into blood vessels and cells, forms arteriocapillary network, fuse with placental vessels, developing in connecting stalk.
• stem villi - or anchoring villi, cytotrophoblast cells attached to maternal tissue.
• branched villi - or terminal villi, grow from sides of stem villi, region of main exchange, surrounded by maternal blood in intervillous spaces.
• terminal villi - not active outgrowths caused by proliferation of the trophoblast. Passive protrusions induced by capillary coiling due to growth of the fetal capillaries within the mature intermediate villi (third trimester).
• chorionic plate - region of membrane at the base of the villi through which placental arteries and vein passes.

Placenta anchoring villi

Villi first trimester

Placenta at Birth

• Placenta (Greek, plakuos = flat cake)
  • embryonic/maternal organ
  • villous chorion/decidua basalis
  • continuous with amniotic and chorionic sacks

Dimensions

• at birth - discoid up to 20cm diameter and 3 cm thick (term) and weighs 500-600 gm
• Shapes - accessory placenta, bidiscoid, diffuse, horseshoe
• maternal and embryonic surface, both delivered at parturition
  ○ retention may cause uterine hemorrhage
## Fetal Surface
- umbilical cord attachment
  - cord 1-2 cm diameter, 30-90cm long
- covered with amniotic membrane and attached to chorionic plate
- umbilical vessels branch into chorionic vessels which anastomose

## Maternal Surface
- **Cotyledons**
  - form cobblestone appearance, originally placental septa formed grooves.
  - covered with maternal decidua basalis

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Fetus in utero, between fifth and sixth months
CT term placental blood vessels showing branching pattern of placental artery.

**Placental Classification**

Classification of placenta is on the basis of histological (microscopic) structural organization and layers between fetal and maternal circulation, giving 3 main groups:

- **Haemochorial** - placenta where the chorion comes in direct contact with maternal blood (human)
- **Endotheliochorial** - maternal endometrial blood vessels are bare to their endothelium and these comes in contact with the chorion. (dogs, cats)
- **Epitheliochorial** - maternal epithelium of the uterus comes in contact with the chorion.considered as primitive (pigs, cows)
The presence of these three differing types of placenta have also been used to describe the pattern mammalian evolution. See also Placental Layers

**Placental Types**

- Discoid in humans, mice, insectivores, rabbits, rats, and monkeys.
- Zonary in dogs, cats, bears and seals.
- Cotyledenedary in cows, deer, goat, and giraffe.
- Diffuse in horses, pigs, camels, lemurs, opossums, kangaroos, and whales

**Links:** Comparative Placentation
Chorionic Villi Trimester Development

- **Trimester 1 and 2** - immature intermediate villi, developmental steps towards the stem villi.
- **Trimester 3** - mature intermediate villi develop during the last trimester, produce numerous terminal villi.
  - Terminal villi are not active outgrowths caused by trophoblast proliferation, passive protrusions induced by capillary coiling due to excessive longitudinal growth of the fetal capillaries within the mature intermediate villi.
  - Capillary bed arrangement in the terminal villi can vary from simple U-like loops to branched network, due to capillary elongation and sprouting.(Data from PMID 2327595)

First trimester spiral artery trophoblast plug.[1]

Placental Cord Blood Vessels
**Week 6** Umbilicus containing initial placental cord.

**Week 8** Embryo showing the placental blood vessels within the ventral body wall.

- **Arteries** - paired, carry deoxygenated blood (from dorsal aorta) and waste products to the placental villi
- **Veins** - paired initially then **only left** at end of embryonic period, carry oxygenated blood to the embryo (sinus venosus)

- initially in the connecting stalk (then umbilical cord) anastomose in chorion
  - extend maternally - toward chorionic villi
  - extend embryonically - to the sinus venosus and dorsal aorta
Placental Function

- 4 layers separate maternal and fetal blood: syncitiotrophoblast, cytotrophoblast, villi connective tissue and fetal capillary endothelium
- 3 main functions: metabolism, transport and endocrine

Placental Metabolism

Synthesizes: glycogen, cholesterol, fatty acids

- provides nutrient and energy

Placental Transport

gases and nutrition

- oxygen, carbon dioxide, carbon monoxide
- water, glucose, vitamins
- hormones, mainly steroid not protein
- electrolytes
- maternal antibodies
- waste products - urea, uric acid, bilirubin
- drugs and their metabolites (fetal drug addiction)
- infectious agents (cytomegalovirus, rubella, measles, microorganisms)

Placental Endocrine

- Human chorionic gonadotrophin (hCG) - like leutenizing hormone, supports corpus luteum
- Human chorionic somatomommotropin (hCS) (or placental lactogen) - hormone level increases in maternal blood through pregnancy, decreases maternal insulin sensitivity (raising maternal blood glucose
levels and decreasing maternal glucose utilization) aiding fetal nutrition ("anti-insulin" function)

- **Human chorionic thyrotropin (hCT)** - Peptide placental hormone, similar to anterior pituitary released thyroid stimulating hormone (TSH), which along with human chorionic gonadotrophin (hCG) is thought to act on maternal thyroid. There is little recent research published on this hormone, its level and activities.

- **Human chorionic corticotropin (hCACTH)** - placental hormone thought to have corticotropin (ACTH)-like activity, increasing maternal cortisol levels.

- **Steroid Hormones**
  - progestins - progesterone, support of the endometrium and suppress uterine smooth muscle contractility.
  - estrogens - estriol, stimulate growth of the myometrium and mammary gland development.
  - both hormones support maternal endometrium

- **Relaxin** - Humans high levels early in pregnancy than at birth promotes angiogenesis probably plays a role in development of the uterus/placenta than in the birth process

## Fetal Placenta

Trophoblast cells are the major source of placental hormones.

**Placental growth hormone** (PGH) is mainly expressed in the syncytiotrophoblast cells (PGH differs from pituitary derived growth hormone by 13 amino acids).
extravillous cytotrophoblast - arise from anchoring villi invade the uterine spiral arteries, generating fibrinoid material and endovascular trophoblastic cells. syncytiotrophoblast

**Fetal Blood Vessels** At least 2 phases of development during pregnancy driven by vascular endothelial growth factor (VEGF):

1. Initially cytotrophoblasts are the cellular stimulus to vasculogenesis and angiogenesis.
2. Later Hofbauer (lacental villi macrophages of mesenchymal origin) and stromal cells take over the stimulation of blood vessel development.

**Placenta Human chorionic gonadotrophin** (hCG) After implantation cells within the developing placenta (syncitiotrophoblasts) synthesize and secrete Human chorionic gonadotrophin (hCG) into the maternal bloodstream. The main function of serum hCG is to maintain the corpus luteum in the maternal ovary and therefore maintain the early pregnancy, that is block the menstrual cycle. Later the placenta itself supports the pregnancy.

**Maternal Placenta**

<table>
<thead>
<tr>
<th>Uterine and placental vasculature</th>
<th>Placenta spiral artery conversion</th>
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**Fibrinoid** - said to exist as 2 forms of extracellular matrix:
1. Fibrin-type fibrinoid is a maternal blood-clot product which replaces degenerative syncytiotrophoblast.

2. Matrix-type fibrinoid is secreted by invasive extravillous trophoblast cells.

Fibrinoid layer (Nitabuch's layer) is thought to act to prevent excessively deep implantation.

**Decidualization** - process of endometrial stromal cells (fibroblast-like) change in morphology (polygonal cells) and protein expression and secretion (specific decidual proteins: prolactin, insulin-like growth factor binding protein-1, tissue factor, interleukin-15, and VEGF).

1. Estrogen and progesterone - receptive phase, luminal and glandular epithelial cells change in preparation for blastocyst implantation.
2. Human Chorionic gonadotropin - luminal epithelium endoreplication leading to epithelial plaque formation.
3. Human Chorionic gonadotropin - trophoblast invasion and decidualization of human stromal fibroblasts.

**Artery Dilatation** - due to extravillous trophoblast cells invading uterine wall and maternal spiral arteries replacing both smooth muscle with fibrinoid material and part of vessel endothelium. There is also a proliferation of maternal blood vessels.

**Other changes**

- Endoreplication - rounds of nuclear DNA replication without intervening cell or nuclear division (mitosis).
- Cytokines - of maternal origin also act on placental development.
- Natural Killer (NK) cells - 30% of all the decidual cells towards the end of the first trimester of pregnancy. These lymphocytes are present in the maternal decidua in large numbers (70%, normal circulating blood lymphocytes 15%) close to the extravillous trophoblast cells. Have a cytolytic potential against virus-infected and tumor-transformed cells.
Placental Abnormalities

- **Placenta Accreta** - abnormal adherence, with absence of decidua basalis. The incidence of placenta accreta also significantly increases in women with previous cesarean section compared to those without a prior surgical delivery.

- **Placenta Increta** - occurs when the placenta attaches deep into the uterine wall and penetrates into the uterine muscle, but does not penetrate the uterine serosa. Placenta increta accounts for approximately 15-17% of all cases.

- **Placenta Percreta** - placental villi penetrate myometrium and through to uterine serosa.

- **Placenta Previa** - In this placenatal abnormality, the placenta overlies internal os of uterus, essentially covering the birth canal. This condition occurs in approximately 1 in 200 to 250 pregnancies. In the third trimester and at term, abnormal bleeding can require cesarian delivery and can also lead to Abruptio Placenta. Ultrasound screening programs during 1st and early 2nd trimester pregnancies now include placential localization. Diagnosis can also be made by transvaginal ultrasound.
- **Vasa Previa** - (vasa praevia) placental abnormality where the fetal vessels lie within the membranes close too or crossing the inner cervical os (opening). This occurs normally in 1:2500-5000 pregnancies and leads to complications similar too those for Placenta Previa. Type II is defined as the condition where the fetal vessels are found crossing over the internal os connecting either a bilobed placenta or a succenturiate lobe with the main placental mass. Some recent evidence of successful in utero laser ablation of type II vasa previa at 22.5 weeks of gestation.

- **Abruptio Placenta** - a retroplacental blood clot formation, abnormal hemorrhage prior to delivery.

- **Chronic Intervillositis** - (massive chronicintervillositis, chronic histiocytic intervillositis) Rare placental abnormality and pathology defined by inflammatory placental lesions, mainly in the intervillous space (IVS), with a maternal infiltrate of mononuclear cells (monocytes, lymphocytes, histiocytes) and intervillous fibrinoid deposition.

- **Hydatidiform mole** - placental tumor with no embryo development. Several forms of hydatidiform mole: partial mole, complete mole and persistent gestational trophoblastic tumor. Many of these tumours arise from a haploid sperm fertilizing an egg without a female pronucleus (the alternative form, an embryo without sperm contribution, is called parthenogenesis). The tumour has a "grape-like" placental appearance without enclosed embryo formation. Following a first molar pregnancy, there is approximately a 1% risk of a second molar pregnancy.

- **Placenta Abnormalities**
There are few abnormalities associated with umbilical cord development, other than abnormally short or long cords, which in most cases do not cause difficulties. In some cases though, long cords can wrap around limbs or the fetus neck, which can then restrict blood flow or lead to tissue or nerve damage, and therefore affect development.
• **Cord knotting** - can also occur (1%) in most cases these knots have no effect, in some cases of severe knotting this can prevent the passage of placental blood.

• **Cord torsion** - Rare event where even without knot formation can also affect placental blood flow, even leading to fetal demise.

**Placental Infections**

• Several infective agents may cross into the placenta from the maternal circulation, as well as enter the embryo/fetal circulation. The variety of bacterial infections that can occur during pregnancy is as variable as the potential developmental effects, from virtually insignificant to a major developmental, abortive or fatal in outcome.

• Pregnant women have an increased susceptibility to malaria infection. Malarial infection of the placenta by sequestration of the infected red
blood cells leading to low birth weight and other effects. There are four types of malaria caused by the protozoan parasite Plasmodium falciparum (main), Plasmodium vivax, Plasmodium ovale, Plasmodium malariae). This condition is common in regions where malaria is endemic with women carrying their first pregnancy (primigravida).

**Placental Pathology**

**MH** - content in this section is not examinable.

- Chronic Villitis - can occur following placental infection leading to maternal inflammation of the villous stroma, often with associated intervillositis. The inflammation can lead to disruption of blood flow and necrotic cell death.
- Massive Chronic Intervillositis (MCI) - maternal blood-filled space is filled with CD68-positive histiocytes and an increase in fibrin, occurring more commonly in the first trimester.
- Meconium Myonecrosis - prolonged meconium exposure leads to toxic death of myocytes of placental vessels (umbilical cord or chorionic plate).
- Neuroblastoma - a fetal malignancy that leads to an enlarged placenta, with tumor cells in the fetal circulation and rarely in the chorionic villi.
- Thrombophilias - (protein C or S deficiency, factor V Leiden, sickle cell disease, antiphospholipid antibody) can generate an increased fibrin/fibrinoid deposition in the maternal or intervillosous space, this can trap and kill villi.

**Additional Images**

<table>
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<tr>
<th>Virtual Slides</th>
<th>Human Circumvallate Placenta</th>
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<td>Placental Villi</td>
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Historic - Fetal Membranes Images

- Isolated villi from chorion frondosum of a human embryo of eight weeks
- Section of chorion of human embryo of one month (9 mm)
- Histology of the placaenta - diagrammatic representations of the minute structure of the placaenta

External Links

*External Links Notice* - The dynamic nature of the internet may mean that some of these listed links may no longer function. If the link no longer works search the web with the link text or name. Links to any external commercial sites are provided for *information purposes only* and should never be considered an endorsement. UNSW *Embryology* is provided as an *educational resource* with no clinical information or commercial affiliation.

- [Comparative Placentation](#)
- [University of Ottawa - Histology - Placenta](#)
- Virtual Microscopy Histology

Placenta Terms
Labs: 1 Fertility and IVF | 2 ES Cells to Genome Editing | 3 Preimplantation and Early Implantation | 4 Reproductive Technology Revolution | 5 Cardiac and Vascular Development | 6 CRISPR-Cas9 | 7 Somitogenesis and Vertebral Malformation | 8 Organogenesis | 9 Genetic Disorders | 10 Melanocytes | 11 Stem Cells | 12

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Student Projects: 1 Cortex | 2 Kidney | 3 Heart | 4 Eye | 5 Lung | 6 Cerebellum