The reproductive significance of human Fallopian tube cilia

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Effective tubal transport of ova, sperm and embryos is a prerequisite for successful spontaneous pregnancy. Although there is much yet to be discovered about the mechanisms involved, it is evident that tubal transit is a far more complicated process than initially thought. Propulsion of gametes and embryos is achieved by complex interaction between muscle contractions, ciliary activity and the flow of tubal secretions. Evidence is accumulating of the important and possibly pre-eminent role of ciliary motion in this process; and this review describes current knowledge about ciliary activity and its physiological regulation. There is also a description of the effects on ciliary function of cigarette smoking and various pathological states, including endometriosis and microbial infection, with consideration given as to how altered ciliary activity may impact upon fertility.

Key words: chlamydia/cilia/ectopic pregnancy/Fallopian tube/tubal infertility

Introduction

With the progress in IVF, the contribution of the Fallopian tube towards successful reproduction has been comparatively overlooked. It is clear from the success of IVF, which of course bypasses tubal transport, that exposure to the tubal milieu is not an absolute requisite for fertilization or implantation to occur. Thus, the Fallopian tube is often now thought of as little more than a mere conduit. However, in fertilization in vivo, the Fallopian tube plays an essential role in gamete transport, fertilization and the early development of the embryo.

It is becoming increasingly evident that the mechanism of tubal transport is much more complex than first thought and can be affected by a wide range of factors and conditions that may impair fertility. It is the purpose of this article to review the recent advances in knowledge about tubal transport predominantly within the human Fallopian tube, with particular reference to the role of the mucosal cilia, the physiological control of their action and how this can be disrupted by various pathological states.

Structure of the human Fallopian tube epithelium

The tubal mucosa is arranged into longitudinal folds, which become increasingly convoluted towards the distal end of the tube (Figure 1). By light microscopy, the Fallopian tube is seen to have a single layer of cuboidal or columnar epithelium which often appears pseudostatified because of crowding of cells of different height. The two major cell types are the ciliated and secretory cells. Ciliated cells are found predominantly on the apex of the mucosal folds (Ferenzy et al., 1972; Patek, 1974). At the base of each cilium is a basal body from which the cilium originates. These develop from centrioles which have migrated towards the luminal surface of the cell, where they are aligned perpendicular to the cell surface giving a refractile appearance on electron microscopy (Hagiwara et al., 2000). Cilia are about 10 μm long and 0.25 μm in diameter (Satir, 1992). There is a progressive decrease in the proportion of ciliated cells from over 50% in the fimbria to less than 35% in the isthmus (Patek et al., 1972a; Amso et al., 1994; Crow et al., 1994).

Cyclical variation in human tubal morphology

The first description of a distinct oviductal cycle in women was made in 1928 (Novak and Everett, 1928). In common with the rest of the female reproductive tract, the Fallopian tube undergoes cyclical changes under the influence of estrogen and progesterone (Critoph and Dennis, 1977a). Cells are low in height during the menstrual phase of the cycle, increasing during the proliferative phase to reach their maximal height in the periovulatory period. At this time, both secretory and ciliated cells are of equal size, with the secretory cells forming domes between the tufts of cilia (Figure 2). Around the time of ovulation, the secretory cells reach peak activity and discharge their contents into the lumen of the tube, consequently reducing in height relative to the ciliated cells (Pauerstein and Eddy, 1979). This results in greater prominence of the cilia and may enable them to move particulate material or viscous secretions more effectively. Subsequently, in the luteal phase, both cell types reduce in height and there is partial deciliation (Patek et al., 1972b; Verhage et al., 1979).
Ovarian hormones affect tubal epithelial structure and expression of cilia. Receptors for estradiol (E\textsubscript{2}) and progesterone have been identified within the Fallopian tube epithelium, and the expression of these receptors varies according to the stage of the ovarian cycle (Pollow et al., 1981). Tubal E\textsubscript{2} receptors are maximal at midcycle, whereas progesterone receptors are present throughout the cycle, even into the late luteal phase (Amso et al., 1994). Estrogen stimulates epithelial cell hypertrophy, secretion and ciliogenesis, whilst atrophy and deciliation are associated with high levels of serum progesterone (Verhage et al., 1979; Donnez et al., 1985).

**Structure of the cilia**

The cilia possess a central bundle of microtubules, called the axoneme, in which nine outer doublet microtubules surround a central pair of single microtubules (Figure 3). This gives a characteristic ‘9 + 2’ arrangement when viewed in cross-section. Each doublet microtubule consists of A and B tubules, or subfibres, each approximately 200 Å in diameter. Rows of dynein arms reach out from the A tubule to the B tubule of the neighbouring doublet.

The basis for axonemal movement is the sliding of doublet microtubules relative to one another. This requires hydrolysis of ATP, utilizing ATPase located in the outer dynein arms. Local regulating factors include calcium ions, cyclic adenosine monophosphate and polypeptides (Huitorel, 1988). The successive active formation and breakage of cross-bridges results in the effective ‘walking’ of the outer dynein arms of the A tubule of one doublet along the B tubule of the adjacent doublet towards its base (Sale and Satir, 1977). Once the dynein arm has become transiently attached to the B tubule, it returns to its original tilt angle, pushing this doublet in a base-to-tip direction (Figure 4). Since microtubule sliding occurs in one direction only, this means that when the doublets on one side of the axoneme are actively sliding, the doublets on the other side are involved in passive movement in the opposite direction. This is termed the switch point hypothesis (Satir and Matsuoka, 1989).

The resultant sliding motion is converted into bending by the selective activation and deactivation of protein cross-links, termed radial spokes. In straight regions of the axoneme, the radial spokes are functionally detached from the central sheath, facilitating the sliding of microtubule doublets. However, in bent regions of the cilium, the spokes remain attached to the inner sheath, converting active interdoublet sliding into local bending (Warner and Satir, 1974).

Analysis of serial cross-sections of the cilia reveals consistent structural asymmetries within the axoneme. The outer dynein arm is absent from one doublet for greater than 90% of the axonemal length. This particular doublet is always orientated in the plane of ciliary beat, opposite to the direction of the effective stroke. Thus, it appears to be structural components within the axoneme which determine the direction of ciliary beat (Hoops and Witman, 1983). Cilia often beat out of phase with one another, producing metachronal waves similar to the pattern of a ‘field of wheat’ (Satir, 1992). However, around the time of ovulation, the strokes of cilia are synchronized and oriented towards the uterus in humans (Gaddum-Rosse et al., 1973).
Cyclical changes in ciliary beat

There have been conflicting reports on the changes in ciliary beat frequency (CBF) throughout the ovarian cycle. In rabbits, the rate of ciliary beat in the oviducts increases by 20% on the 2nd and 3rd days after copulation, corresponding to the time of movement of the ova down the tubes (Borell et al., 1957). Critoph and Dennis detected a significant increase in the CBF of the human isthmus and ampulla after ovulation (Critoph and Dennis, 1977b). In contrast, Westrom found no cyclical or anatomical variation in beat frequencies (Westrom et al., 1977). Recent investigations using analogue contrast enhancement have shown an increase in CBF of the fimbrial section of the tube during the secretory phase (Lyons et al., 2002a). These results agree with those of Critoph and Dennis in confirming a variation in CBF according to the ovarian cycle.

Methods used to measure CBF

The cilia beat at such high frequencies that direct observation allows only the recognition of dramatic changes in CBF and therefore makes this assessment of limited value (Rusznak et al., 1994). Several techniques for measuring CBF have been developed, but all have various limitations such as poor spatial resolution, vibration artefacts from the immersion fluid and inaccuracy when the cilia are in asynchronous rhythm (Naitoh and Kaneko, 1973; Verdugo et al., 1980a; Kennedy and Duckett, 1981). Recent advances in measurement of CBF involve the use of digital technology. The use of digital cameras allows improvement of the signal/noise ratio, enhancement of the image resolution and direct storage into computer memory (Sanderson, 2000).

Considering the numerous and varied techniques employed to measure CBF, it is not surprising that there is no consensus as to the baseline rate of CBF in the human Fallopian tube. A wide range of frequencies from 5 to 20 Hz has been reported in the literature (Westrom et al., 1977; Paltieli et al., 1995). It is important to note that results obtained using the different techniques cannot be directly compared (Chilvers and O’Callaghan, 2000).
As a result, the isthmic lumen opens and allows transport of the early embryo to the uterus (Jansen, 1978). This delay at the AIJ is believed to be important in exposing the embryo to developmental factors and nutrients within the tubal fluid and in postponing entry into the uterus to the time of maximal endometrial receptivity. The concept that the fertilized ovum is delivered to the uterine cavity at an appropriate time for implantation implies the presence of regulatory mechanisms governing tubal transport. Three possible mechanisms have been suggested: endocrine regulation, mainly by ovarian steroids; neuronal regulation through the autonomic nervous system and paracrine regulation by the embryo itself.

Sperm transport

Because pregnancy has been shown to result from intercourse which occurs up to 5 days before ovulation (Wilcox et al., 1995), human sperm must be stored at some site in the female genital tract. Although motile sperm have been recovered from the human cervix up to 5 days after insemination (Gould et al., 1984), it is not known whether these sperm could reach the Fallopian tube so long after deposition or indeed whether they might have re-entered the cervix from the uterus. The available evidence suggests the human Fallopian tube itself as the likely candidate for a sperm storage site as the tubal epithelium provides a favourable environment for sperm. Anatomical sperm reservoirs within the tubal isthmus have been detected in a variety of mammalian species (Yanagimachi and Chang, 1963; Hunter, 1981; Hunter and Nichol, 1983), but a distinct tubal reservoir has not been located in humans (Williams et al., 1993).

Nevertheless, motile human sperm have been shown to bind by their heads to the ciliated apical areas of the tubal epithelium in vitro, and the density of sperm is greater in the isthmus than the ampulla (Baillie et al., 1997). Species-specific carbohydrate moieties are involved in the binding process in animal studies, but no carbohydrate performing such a function has yet been identified in humans. Recently, however, it has been proposed that human sperm–isthmus attachment results from binding of the amino acid sequence Arg-Gly-Asp of an as-yet unidentified sperm protein to integrins on the tubal epithelium (Reeve et al., 2003). Sperm binding in the human endosalpinx appears to be intermittent and less tight than in other mammalian species studied (Pacey et al., 1995a, b). Despite this, when associated with the convoluted architecture and thick tenacious mucus of the isthmic endosalpinx, these factors may be sufficient to act as a functional sperm reservoir and slow sperm progression through the tube.

Sperm–endosalpingeal contact preserves the viability of sperm. Incubation of human sperm with cultured tubal epithelium preserves viability for longer than culture in medium alone, as does incubation with vesicles prepared from the apical membrane of the human endosalpinx (Kervancioglu et al., 1994; Murray and Smith, 1997). This indicates that the epithelium can enhance sperm survival by direct contact rather than by secretions. Retention of sperm within the tubal isthmus could therefore prolong the availability of viable sperm and increase the chance of successful fertilization.

There is evidence from animal studies that the incidence of polyspermy is increased if sperm is freely available in the tubal ampulla. If porcine sperm is injected directly into the ampulla, or the isthmus is excised or the smooth muscle of the muscularis is inhibited by direct administration of progesterone, then the frequency of polyspermic fertilization rises (Day and Polge, 1968; Hunter and Leglise, 1971; Hunter, 1973). Thus, a functional isthmic sperm reservoir may also serve to prevent polyspermy by restricting the rate of sperm release into the ampulla to only a few at a time.

Once sperm have undergone the processes of capacitation and hyperactivation in preparation for fertilization, they appear to lose their binding affinity. Capacitation modifies the proteins on the surface of the sperm cell membrane, which could alter or reduce binding sites. Uncapacitated boar and bull sperm binds more avidly to tubal epithelium than after capacitation (Lefebvre and Suarez, 1996; Fazeli et al., 1999). Hyperactivation can supply the force necessary for the physical disruption of sperm–endosalpingeal attachment and may assist in the detachment of human sperm (Pacey et al., 1995b). Thus, maturational changes within the sperm themselves appear to regulate their binding and subsequent release from the tubal epithelium. Once released, ciliary motion may assist in transport to the site of fertilization within the ampulla, as co-culture of sperm with tubal epithelial cells has been shown to increase CBF (Morales et al., 1996).

Regulation of ciliary beat

Ciliary reactivity is affected by a variety of hormonal and neuronal stimuli. CBF is a calcium-dependent process which requires hydrolysis of ATP. In the absence of calcium in the culture medium, ciliary motility in vitro ceases (Verdugo, 1980). ATP increases CBF in a dose-dependent manner in vitro. It has been suggested that ATP is released by the secretory cells in vivo and subsequently acts upon the ciliated cells in a paracrine manner (Villalon and Cardina-Danovaro, 1994). Beta-adrenergic stimulation increases ciliary activity, an effect which can be blocked by the β-adrenergic receptor blocker, propranolol, confirming the receptor specificity of this response (Verdugo et al., 1980b).

Angiotensin II in nanomolar concentrations has been shown to stimulate CBF in vitro. This effect is inhibited by the specific type 1 angiotensin II receptor antagonist, losartan (Saridogan et al., 1996). The presence of a renin–angiotensin system has been described in the human Fallopian tube, and angiotensin II receptors are present in the tubal mucosa, with immunostaining being most prominent in the proliferative phase. The role of angiotensin II in tubal function has not yet been elucidated.

It has been hypothesized that the increase in CBF in vitro is because of rising progesterone levels in an estrogen-rich environment, which may increase the release of ATP from apically situated mitochondria within the ciliated cells, and thus increase CBF (Jansen, 1984). However, high-dose progesterone has been shown to inhibit CBF in vitro by up to 63% (Mahmood et al., 1998; Paltieli et al., 2000). This inhibition can be reversed by the progesterone receptor antagonist, mifepristone (Mahmood et al., 1998). The ovarian hormones are present in the Fallopian tube mucosa in much higher concentrations than in the general circulation, because of a countercurrent exchange mechanism between the ovarian artery and the venous plexuses along the mesosalpinx (Bendz et al., 1982). Also, the Fallopian tube cilia are directly exposed to high levels of ovarian steroids at midcycle with the influx of follicular fluid. During the secretory phase,
estrogen and progesterone levels remain high in the peritoneal fluid, which is in direct communication with the tubal lumen, thus prolonging the ciliary exposure to raised ovarian hormone levels. Because high concentrations of progesterone decrease CBF, these findings suggest that alternative factors may be responsible for the increase in CBF detected post-ovulation.

PGE$_2$ and F$_2$α both stimulate fimbrial CBF in the rabbit oviduct in vitro, an effect believed to be mediated by PG-induced release of calcium ions (Verdugo, 1980; Verdugo et al., 1980a). It has been hypothesized that PGs released locally in the tubal mucosa or synthesized by the cumulus complex surrounding the transported oocyte stimulate ciliary activity, acting through the release of calcium ions from intracellular storage sites or the extracellular space (Jansen, 1984).

Follicular fluid exerts a significant stimulatory effect on CBF of human Fallopian tube explants in vitro (Lyons et al., 2006). The follicular fluid of human pre-ovulatory ovarian follicles contains high concentrations of E$_2$, progesterone and PGs (Edwards et al., 1972; McNatty et al., 1979; Seibel et al., 1984). At ovulation, the fimbrial end of the Fallopian tube is in close apposition to the dominant ovarian follicle (Doyle, 1956). Once the follicle ruptures, the oovum is transported into the Fallopian tube by the flow of follicular fluid (Harper, 1982). Follicular fluid thus becomes the major constituent of tubal fluid immediately post-ovulation. It appears that PGs or other factors in follicular fluid may provide the stimulus for the increase in CBF observed in the secretory phase and that this may aid ovum pick-up and transport.

Pathological effectors of CBF

Ectopic pregnancy

Fallopian tubes containing an ectopic pregnancy demonstrate a marked reduction in the number of ciliated cells in comparison with those of women with an intrauterine gestation. Marked deciliation is also sometimes seen subsequent to an ectopic pregnancy and in biopsies from women undergoing tubal surgery who later develop a tubal pregnancy (Vasquez et al., 1983). This effect suggests that pathological processes affecting the tubal cilia may predispose to ectopic gestation.

Smoking

Women who smoke have a markedly increased risk of ectopic pregnancy and increased incidence of tubal infertility. Compared with women who have never smoked, women who smoke more than 20 cigarettes a day have an almost four-fold risk of ectopic gestation, similar to the increase in risk associated with a past history of pelvic inflammatory disease (Bouyer et al., 2003). Even prenatal exposure to tobacco smoke has been postulated to increase the prevalence of tubal disease, suggesting that tobacco smoke has a permanent detrimental effect on the developing Fallopian tubes (Matthews et al., 2002).

In animal models, nicotine alters tubal motility (Neri and Marcus, 1972) and decreases tubal blood flow (Mitchell and Hammer, 1985). Exposure of hamsters to doses of cigarette smoke within the range received by active or passive human smokers causes a small but significant increase in the secretory-to-ciliated cell ratio within the infundibulum (Magers et al., 1995). Acute in vitro exposure of the hamster infundibulum to smoke solutions causes a rapid reduction in CBF which is reversible upon washout of the smoke solution (Knoll et al., 1995).

Oocyte cumulus pick-up rate by the hamster oviduct is inhibited in a dose-dependent manner by smoke solutions, and this effect is not easily reversed by washout of the solution, demonstrating that the effect of smoking on ovum pick-up is separate to the effect on CBF. The likely explanation for this is that smoke solutions disrupt the adhesion between the negatively charged tips of the cilia and the oocyte cumulus complex, probably by the binding of an as-yet-undetermined smoke component (Knoll and Talbot, 1998).

Animal data demonstrating reduced efficacy of ovum pick-up and delayed transport along the tube because of decreased CBF may explain the higher rates of infertility and ectopic gestation seen in women who smoke. It is of concern that even fetal exposure to tobacco appears to affect tubal function, suggesting that the consequent tubal damage may be long-term, if not permanent.

Endometriosis

The association between endometriosis and infertility is well recognized, although whether or not endometriosis causes infertility remains controversial. It is easy to understand how severe endometriosis with the formation of ovarian endometriomas, adhesions and gross distortion of the pelvic anatomy may be associated with infertility. However, the link between mild endometriosis and infertility is as yet unexplained.

The peritoneal microenvironment in women with endometriosis differs from normal fertile controls. There is substantial evidence that the levels of activated macrophages in the peritoneal cavity are raised in women with endometriosis (Halme et al., 1983, 1987; Zeller et al., 1987). There appears to be an inverse relationship between stage of endometriosis and the presence of inflammatory mediators, with a tendency towards higher peritoneal macrophage levels in women with minimal-to-mild disease (Olive et al., 1985; Haney et al., 1991). Cytokines secreted from endometriotic deposits or activated macrophages have been detected at higher concentrations in the peritoneal fluid of women with endometriosis and have been shown to have a deleterious effect on fertility parameters (Fakhri et al., 1987; Sueldo et al., 1990).

There is a marked inhibitory effect of peritoneal fluid from women with mild-and-moderate endometriosis on CBF in vitro (Lyons et al., 2002b). Additionally, a macromolecular ovum capture inhibitor has also been detected in the peritoneal fluid of women with endometriosis (Suginami and Yano, 1988). This forms a membrane over the fimbrial cilia, causing a complete loss of ovum capture activity. It is possible that a similar mechanism of deposition of filamentous material may inhibit the activity of cilia, and thus tubal transport, in women with endometriosis. Furthermore, one or more of the constituents of the proinflammatory peritoneal fluid found in women with endometriosis may affect ciliary beat directly. The factor(s) in endometriosis peritoneal fluid responsible for this reduction in CBF are unknown. Possible mediators include macrophages or their various secretion products.

Interestingly, human sperm–endosalpingeal interaction in vitro appears to be altered in explants of tubal epithelium from women with a diagnosis of endometriosis, with more sperm bound in tissue from affected women (Reeve et al., 2005). This suggests that a reduction in free and motile sperm within the tubal lumen may...
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reduce fertilization rates. Although little is known about sperm–endosalpingeal interaction in humans, tubal integrins have been implicated in the process (Reeve et al., 2003), and integrin expression is known to be aberrant within the endometrium and endometriotic deposits of sufferers (Lessey et al., 1994; Puy et al., 2002). The impairment of tubal transport of gametes and embryos may thus represent one mechanism by which endometriosis exerts its detrimental effect on fertility.

Effects of micro-organisms on the Fallopian tube

*Neisseria gonorrhoeae*

Exposure of the human Fallopian tube epithelium *in vitro* to fresh isolates of *N. gonorrhoeae* and to gonococcal endotoxin results in reduction and subsequent cessation of ciliary activity. This ciliostatic effect occurs before any ultrastructural changes are apparent on scanning electron microscopy (Mardh et al., 1979). Gonococci invade the non-ciliated cells of the tubal mucosa, but destroy predominantly the ciliated cells, primarily by causing sloughing (McGee et al., 1981).

The two compounds likely to mediate this damage are gonococcal lipopolysaccharide (LPS), released from the outer membrane of the organism, and monomers of peptidoglycan, which are generated by the bacterium (Melly et al., 1984; Woods and McGee, 1986; Stephens et al., 1987). Immediately after addition to human Fallopian tube organ cultures, gonococcal LPS aggregates on the tips of the cilia. By 2 h after exposure, LPS is evident in the cytoplasm of both ciliated and non-ciliated cells in vesicle-like structures, and, by 12 h, sloughing of the ciliated cells has occurred (Cooper et al., 1986). Neutralization of the lipid A moiety of LPS abolishes this effect, confirming that the destructive action of *N. gonorrhoeae* is mediated indirectly by the release of toxins (Stephens et al., 1987).

Gonococcal infection of the human Fallopian tube mucosa results in production of tumor necrosis factor (TNF-α) by the mucosa. The extent of loss of ciliated cells from the tubal epithelium correlates with the mucosal tissue concentration of TNF-α (McGee et al., 1999), and blocking gonococcal production of TNF-α limits epithelial damage (McGee et al., 1992). Direct treatment of tubal epithelial cultures with recombinant TNF-α also causes sloughing of ciliated cells in the manner typical of gonococcal (McGee et al., 1992).

This suggests that induction of this proinflammatory cytokine by gonococcal infection may be another mechanism by which tubal damage occurs. As cytokines, including TNF-α, have already been implicated in the reduction in CBF produced by endometriotic peritoneal fluid, this is an area worthy of further investigation.

*Chlamydia*

Fallopian tube damage and tubal factor infertility are common sequelae of upper genital tract infection with *Chlamydia trachomatis*. This pathogen causes a direct cytotoxic effect on the mucosa of the human Fallopian tube, which results in loss of microvilli and disruption of cell junctions, associated with rupture of the epithelial cells (Cooper et al., 1990). However, it is believed to be the consequent immune response which results in permanent tissue scarring, with persistent or repeated infection exacerbating this destruction. Although the precise mechanism by which prolonged exposure to chlamydial antigens causes tubal damage is not fully elucidated, the 60 kDa chlamydial heat shock protein (hsp60) is believed to be the major antigen triggering this pathogenic immune response. Several studies have demonstrated a correlation between the level of immune response to chlamydial hsp60 and the extent of tubal damage (Brunham and Peeling, 1994; Eckert et al., 1997; Peeling et al., 1997). The detection of antibodies to hsp60 is strongly correlated to the presence of tubal factor infertility (Toye et al., 1993; Ault et al., 1998).

Additional chlamydial antigens have been implicated in the immunopathological mechanism of tubal damage. Immune response to hsp10 has been associated with tubal infertility in a population of women exposed to chlamydia (LaVerda et al., 2000). Recently, evidence of genetic protection against chlamydia-mediated tubal damage has emerged, with the description of two class II alleles detected less commonly in chlamydia-seropositive women with tubal infertility (Cohen et al., 2003).

Chlamydial hsp60 is capable of provoking both a humoral and cell-mediated immune response, with intense macrophage activation (LaVerda et al., 1999). Activated macrophages secrete proinflammatory cytokines, and chlamydial infection induces significantly higher secretion of TNF-α, interleukin-1β and interferon (IFN)-γ (Toth et al., 1992; Ojcius et al., 1998; Kinnunen et al., 2003). Apoptosis of both infected epithelial cells and macrophages occurs, releasing more proinflammatory cytokines and exacerbating the host inflammatory response (Ault et al., 1996; Ojcius et al., 1998).

Under different conditions, *C. trachomatis* appears to be capable of either promoting or inhibiting apoptosis. Chlamydial proteins in infected cells possess diverse antiapoptotic properties, including inhibition of caspases, the enzyme system primarily responsible for triggering apoptosis (Fan et al., 1998). Macrophage-released IFN-γ has also been shown to inhibit apoptosis of infected cells (Dean and Powers, 2001; Perfettini et al., 2002). This inhibition of apoptosis of infected cells is likely to contribute to the development of persistent infection (Witkin, 2002).

The pathways leading to permanent tubal damage have not yet been fully elucidated. The production of endothelial cell adhesion molecules within the Fallopian tube can be induced directly by chlamydial LPS or by the cytokines of the inflammatory response (Kelly et al., 2001). These cell adhesion molecules mediate the extravasation of lymphocytes from the circulation into the site of infection and thus promote the inflammatory response (Butcher et al., 1999). Enhanced production of PGs and collagen, with increased expression of integrin and transforming growth factor-β, contributes to fibrosis and scarring (Perfettini et al., 2000). Hsp60 also causes the induction of matrix metalloproteinases, which may produce tubal scarring as a result of increased turnover and repair of the extracellular matrix (LaVerda et al., 1999; Ault et al., 2002).

Salpingitis is associated with distal occlusion of the Fallopian tube and deciliation, which can be extensive (Patton et al., 1983, 1987; Westrom and Wolner-Hanssen, 1993). This deciliation appears to be permanent (Donnez et al., 1984). It is possible that the function of the remaining cilia may not be affected (Cooper et al., 1990). In women with tubal infertility, serological evidence of chlamydial infection is not associated with changes in CBF. However, there is preliminary evidence that particular chlamydial serotypes, such as serotypes C and E, may be associated with zero
or reduced levels of CBF respectively, although the numbers of subjects sampled are small, and these results must be interpreted with caution (Leng et al., 1998). CBF is significantly lower in the surviving cilia of Fallopian tubes showing evidence of oedema, erythema or distal obstruction (Patton et al., 1989; Leng et al., 1998). Therefore, it appears that chlamydia causes deciliation and the associated chronic inflammation reduces CBF, although it is possible that certain serotypes may affect CBF directly and may presage a particularly poor prognosis for subsequent fertility.

Other pathogens

Inoculation of Escherichia coli into the Fallopian tubes of rabbits results in a dose-dependent deciliation, with the remaining cilia being swollen, shortened and adherent throughout their entire length. The secretory cells demonstrate loss of microvilli. However, regeneration does occur, commencing 2 weeks after the initial injury, and being complete by eight weeks (Laufer et al., 1984).

Bacterial vaginosis (BV) is characterized by an overgrowth of vaginal anaerobic and facultative bacteria. It is associated with the development of endometritis and pelvic inflammatory disease, especially in the presence of other sexually transmitted diseases (Soper et al., 1994; Hillier et al., 1996; Wiesenfeld et al., 2002). BV is strongly associated with tubal factor infertility (Gaudoin et al., 1999). Some of the micro-organisms involved in the condition, including Mycoplasma hominis, Mobiluncus and Bacteroides ureolyticus, have been shown to cause direct tubal damage or altered ciliary activity.

Infection with M. hominis results in ciliostasis and swelling of the cilia (Mardh et al., 1976). Infected human tubal mucosa cultures demonstrate either zero or reduced ciliary activity (Baldetorp et al., 1983). Species of Mobiluncus (Mobiluncus curtisi and Mobiluncus mulieris) produce cytotoxins that reduce ciliary activity within 60 h in cultures of bovine oviduct. This is associated with loss of cilia and bloating and detachment of ciliated cells (Taylor-Robinson et al., 1993). LPS endotoxins released from B. ureolyticus (Prevotella) also damage the mucosa of the human Fallopian tube, causing sloughing of cells and loss of ciliary activity (Fontaine et al., 1988).

These various micro-organisms ascend into the upper genital tract where they frequently cause subclinical pelvic inflammation. There is an associated destruction of the ciliated cells combined with a general ciliostatic effect because of the release of endotoxins. These could be mechanisms by which BV contributes to the pathogenesis of tubal disease and infertility.

Conclusion

The Fallopian tube plays an essential role in tubal transport of both gametes and embryos and in early embryogenesis. The tube undergoes cyclical changes in morphology and ciliary activity in response to ovarian hormones. Whilst the varying contributions to tubal transport of ciliary activity, muscle contractions and secretary activity remain undetermined, there is emerging evidence that muscle contractions may play a role in mixing of secretions rather than in propulsion of gametes and embryos.

Ciliary activity is more vigorous in the secretory phase of the menstrual cycle. Many pathological conditions associated with infertility and ectopic pregnancy have been shown either to destroy cilia or to reduce ciliary motion or both. Gonococcal infection produces both destruction of the ciliated cells and reduced ciliary activity, whereas chlamydia also destroys the tubal mucosa. Although the micro-organism itself does not appear to alter ciliary beat, the inflammation and oedema associated with chlamydial salpingitis has been shown to reduce CBF. Peritoneal fluid from women with mild-and-moderate endometriosis reduces CBF significantly in vitro. An ‘ovum capture inhibitor’ has been described in the peritoneal fluid of women with endometriosis, which covers the fimbrial cilia resulting in a complete but reversible loss of ovum capture ability. The ‘immotile cilia syndrome’ is known to be associated with subfertility. Deciliation is found in Fallopian tubes of women with a past history of ectopic gestation. These women are at increased risk of future tubal pregnancies. This evidence suggests an important role for the tubal cilia in the mechanism of gamete and embryo transport.

Further research needs to be undertaken to investigate the functioning of the cilia in vivo. Only one study has measured physiological CBF in vivo (Paltieli et al., 1995), and this needs to be extended to the effect of pathological states on CBF. Direct examination of the effect of conditions such as endometriosis or pelvic inflammatory disease on ovum transport may be possible in animal models using laparoscopy to investigate ovum pick-up and fallopescoppy to study ovum transit along the tube. It is only as we begin to understand more about the complex interactions of the effectors of tubal transport that we approach the possibility of being able to improve tubal transport in women afflicted with tubal infertility.

References


Role of cilia in human Fallopian tube


