Understanding sperm motility mechanisms and the implication of sperm surface molecules in promoting motility

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Abstract

Background: It is estimated that approximately 8–12% of couples globally face problems associated with infertility. A large number of men exhibit suboptimal sperm parameters. Sperm motility is one of the factors that is measured when analysing sperm parameters. The indication of several crucial sperm surface molecules, having the ability to modulate motility, has opened new avenues in understanding the complex processes involved in motility.

Main body of the abstract: There are various mechanisms that regulate and enhance sperm motility. Several surface molecules on sperm cells can also regulate motility, thus showing their possible application as a treatment for infertility caused by impaired motility. Sperm motility is regulated by intracellular and extracellular pH, along with calcium ions (Ca²⁺) and carbonate ion (HCO₃⁻) concentrations. Moreover, sperm cells have an array of surface proteins which play a critical role in their function and motility. The indication of surface molecules presented new opportunities for understanding sperm motility and the possibility of treating infertility caused by impaired sperm function. Infertility and problems associated with conception can cause underlying stress and mental trauma. Although there are several methods for treating infertility, most are complex, invasive, and expensive.

Conclusion: It is important to understand how surface molecules and proteins on the sperm cell regulate motility. This will enable us to treat anomalies associated with proper sperm function. This review highlights the general mechanisms that regulate sperm motility, and it stresses the importance and relevance of sperm surface molecules in regulating sperm motility.

Keywords: Sperm motility, Membrane proteins, Fertilization, Fertility regulation, Infertility, IVF

Introduction

Sperm cells are specialized haploid cells that act as the male gamete. During fertilization, sperm cells fuse with the ovum to form a zygote. It is estimated that approximately 8–12% of couples worldwide are affected by infertility and problems of impaired fecundity. Globally, 40–50% of the infertility cases are caused by male factor infertility, and as many as 2% of all men exhibit suboptimal sperm parameters. Semen analysis is routinely used to investigate male infertility. Sperm motility is a key factor in determining the quality of semen and a reliable predictor for fertilization success [1]. Male infertility is usually diagnosed on finding abnormal results on semen analysis. Descriptive diagnoses usually include “oligozoospermia” (reduced sperm count), “asthenozoospermia” (reduced sperm motility), and “teratozoospermia” (reduced percentage of sperm with normal morphology). One of the major causes of male infertility is idiopathic primary testicular dysfunction with abnormal spermatogenesis. Other conditions contributing to male infertility are testicular damage due to systemic chemotherapy for cancer, Klinefelter syndrome, genetic mutations, testicular cancer, pelvic irradiation or surgery, trauma,
cryptorchidism, infection, autoimmune destruction, and drugs [2].

Sperm cells are smaller than most cells in the body; they have a distinctive head, mid-piece, and tail region. Capacitation is an important process that leads to the destabilization of the acrosomal sperm head which allows the sperm to penetrate the ovum. During capacitation, several biochemical changes occur in the tail that enhances sperm motility, and there is a significant amount of efflux of cholesterol in the plasma membrane leading to an increase in membrane fluidity and permeability to bicarbonate and calcium ions, an increase in the polarization of the plasma membrane and changes in protein phosphorylation and protein kinase activity. There is also an increase in the intracellular concentrations of bicarbonate ions (HCO₃⁻), calcium ions (Ca²⁺), and cyclic adenosine monophosphate (cAMP) levels [3–5]. Studying sperm morphology and understanding each physiological process is crucial in eliminating various complications that may arise during conception. This narrative review highlights the factors that influence the motility of sperm cells, and it discusses the role of surface molecules present on the surface of the sperm cell that influences motility, and the role of these molecules as potential candidates for treating infertility caused by impaired sperm function.

Asthenozoospermia and its causes

Asthenozoospermia or asthenospermia is a condition that is characterized by reduced or no motility of sperm cells in the fresh ejaculate. Sperm cells undergo maturation in the epididymis and acquire motility during the epididymal transition [6]. Motility is a very important parameter for successful fertilization. Impaired motility can result in unsuccessful fertilization and can be caused by several reasons which are described in Table 1.

Sperm cells are transcriptionally and translationally inactive, but there are several specific metabolic pathways that are able to regulate their ability to fertilize an ovum. Sperm motility is also regulated by several signalling cascades and mechanisms. Most notable is the cAMP/protein kinase A and phosphoinositide 3-kinase signalling, which are mediated through Ca²⁺, HCO₃⁻, or both [12]. This review describes the relevance of each of the pathways in regulating sperm motility.

**Table 1** List of notable causes of male infertility and their relationship with impaired motility

| Notable causes of male infertility | Relationship with motility |
|-----------------------------------|---------------------------|
| Varicocele                         | Results in elevated levels of reactive oxygen species (ROS) which alters the testicular microenvironment, thereby causing reduced motility [7]. Damaged mitochondria result in abnormal expression of mitochondrial proteins, thereby decreasing ATP levels and altering calcium signalling cascade [8]. |
| Genetic abnormalities             | Defect in several genes coding for the proteins in the central axoneme apparatus, dynein proteins, as well as genes such as Spag 6, 16, 17 that are responsible for central axoneme function result in impaired motility [9–11]. |
| Lifestyle choices                 | Lifestyle factors such as smoking, stress, and alcoholism may affect sperm parameters. Tobacco smoke containing traces of tar, carbon monoxide, polycyclic aromatic hydrocarbons, and heavy metals are known to influence motility [12]. |
| Radiation                         | Causes genetic abnormalities and production of ROS. Catsper genes are known to be mutated due to exposure to radiation, thereby causing impaired calcium metabolism [13]. |
| Heat exposure                     | Heat exposure downregulates mitochondrial activity and decreases ATP levels. Heat exposure also decreases antioxidant levels, alters protein expression, and causes mitochondrial degeneration [14]. |
| Environmental Factors             | Environmental chemicals, such as pesticides, polychlorinated biphenyls, bisphenol A, glycol ethers, perfluorinated compounds, dioxins and dioxin-like compounds, phthalates, heavy metals, dichloro-diphenyl-trichloroethane, and plasticizers are known to affect sperm motility [12]. |
| Infections                        | Bacterial infections, leucocyte accumulation (leukocytospermia), antibody buildup, inflammation and oxidative stress are known to impair fertility. Infections reduce mitochondrial membrane potential and increases apoptosis [15]. |
| Psychological stress              | Hormones such as corticosterone suppress testosterone and inhibit, thereby altering the testicular microenvironment. Hormonal changes are known to affect motility [12]. |
using much less invasive methods. Therefore, it is important to elucidate the process of sperm maturation and maturational changes that spermatozoa undergo during epididymal transit [16].

The epididymis is grossly divided into three regions: the caput (head), corpus (body), and cauda (tail). Each region of the epididymis, as described in Fig. 1, performs distinctive functions. The caput contributes to the early maturation events, whereas the corpus participates in late maturation events. The cauda region serves as a reservoir for storing functionally mature cells. The primary cell type along the epididymal tubule remains the same from the proximal to the distal end; however, cells from each region exhibit different subsets of genes, thus contributing to the ever-changing luminal environment [17, 18]. A septum further subdivides the caput, corpus, and cauda epididymis into discrete intraregional segments and that region-specific gene expression may in fact be highly ordered and compartmentalized within these precise segments [18]. Several sperm-associated proteins such as ADAM2 (fertilin β), ADAM3 (cyritestin), ADAM24 (testase), and CE9 are proteolytically cleaved and activated by various proteases during the epididymal transition. Interestingly, several free radical scavenging enzymes such as γ glutamyl transpeptidase, glutathione peroxidases, and superoxide dismutase are produced in the epididymis to prevent oxidative damage of the sperm cell membrane in the oxygen-rich epididymal lumen. As the spermatozoa migrate from the proximal to the distal regions of the epididymis, they are exposed to segment-specific gene expression encoding signalling molecules, regulatory proteins, transporters, and receptors, thus contributing to the formation of a unique microenvironment in each segment [18–20].

**Altered structure of the flagellum impairs motility**

The four core elements of the sperm cell are described in Fig. 2. The vigorous beating of the flagella is crucial for penetrating through the corona radiata. It is clear that immotile sperm cannot pass through the cervical mucus. Moreover, the type of movement is also crucial. For example, sperm moving in tight circles cannot travel through the uterotubal tract. Only forward-moving sperm can successfully fertilize the ovum [21]. The flagellum propels the sperm through the cervical mucus in the female genital tract. It mainly consists of a structure known as the axoneme. This highly conserved microtubule-based structure is very similar to the internal cytoskeleton of motile cilia that are found at the surface of many cell types such as the epithelial cells from the airways, the fallopian tubes, or the brain ventricles.
The axoneme comprises nine outer doublet microtubules and central doublets (9 + 2) associated with radial spokes and dynein arms. The dynein arms within the axoneme provide the motor apparatus for the movement of the sperm tail [22]. Proper formation of the axoneme during spermatogenesis is crucial in sperm motility. The structure of the sperm tail axoneme resembles that of motile cilia. Therefore, male infertility caused by malformations of the axonemal structure is often associated with primary ciliary dyskinesia (PCD). However, male infertility is not systematically investigated and often not recorded in cases of PCD [23].

An association between gene mutation and sperm motility and male infertility has been considered in several reviews. Mutations in more than 30 genes have been identified in cases of PCD including dynein arm assembly genes. Defects in the axonemal outer dynein arms (ODA) genes, dynein axonemal heavy chain 5 (DNAH5), and dynein axonemal intermediate chain 1 (DNAI1) leads to reduced sperm motility, even though sperm axonemal structure appears intact [24, 25]. Mutations in inner dynein arms (IDA)-related coiled-coil domain-containing proteins 39 (CCDC39) and 40 (CCDC40) cause reduced sperm motility and absence of IDA [26]. Male infertility was also observed in cases showing defects in central pair-related genes. In humans, depletion of hydin causes PCD, and spermatozoa appear rigid and completely immotile [9]. It has been reported that several sperm-associated antigen (Spag) genes (Spag6, Spag16, and Spag17) in mice are important for central pair complex function. Infertility due to missing axonemal central pair and disorganized ODFs was caused by the total loss of Spag6 [10]. However, depletion of the SPAG16L isoform resulted only in sperm motility defects with intact axonemal structure [11]. A large number of people exhibit idiopathic asthenospermia. Understanding and screening for genetic anomalies that may hinder sperm motility and function are crucial in developing treatment strategies for such infertility cases.

**Role of calcium and bicarbonate ions on sperm motility**

Soluble adenylate cyclase (sACs) are activated by Ca$^{2+}$ and bicarbonate ions. They are the predominant ACs responsible for the generation of cAMP in spermatozoa and are involved in the increase in beat frequency in the spermatozoa. On the other hand, transmembrane ACs are not responsive to bicarbonate ions but are regulated by heterotrimeric G-proteins. Transmembrane ACs are involved in the basic mechanism for motility activation through cAMP-dependent protein phosphorylation and in progressive motility [27]. Calcium ions act as major secondary messengers and regulate the amount of cAMP in the cell [28]. The concentration of calcium is a crucial regulatory factor that affects capacitation, hyperactivation, and acrosome reaction. It has been observed that the flagellum beats asymmetrically when the intracellular Ca$^{2+}$ concentration is low. The waveform becomes more asymmetric, and sperm becomes...
hyperactivated with a gradual increase in Ca\(^{2+}\). However, an excessively high amount of intracellular concentration of calcium reduces motility [29]. This occurs due to a decrease in protein phosphorylation caused by the increased concentration of Ca\(^{2+}\), which prevents substrate-kinase interactions [30]. Calmodulin is a Ca\(^{2+}\) receptor that orchestrates Ca\(^{2+}\)-initiated signal transduction cascades leading to changes in cell function. It is a key axonemal Ca\(^{2+}\) sensor that mediates motility through direct interaction with protein kinases, phosphatases, and sAC [31]. The impact of the extracellular concentration of Ca\(^{2+}\) on motility has been well-debated. Several studies have reported that extracellular Ca\(^{2+}\) enhances sperm motility, whereas others have also reported that a high concentration of Ca\(^{2+}\) inhibits sperm motility. In conclusion, it is clear that the role of Ca\(^{2+}\) in sperm motility is paradoxical [28]. Bhoumik et al. have reported a biphasic role of extracellular calcium in the motility of caprine cauda epididymal spermatozoa. They observed that an optimum concentration of 10 μm of Ca\(^{2+}\) significantly enhanced motility, whereas concentrations above this hindered motility [28]. This observation is particularly important because seminal plasma usually contains about 11 mM of Ca\(^{2+}\) and about 0.24 mM of free Ca\(^{2+}\); the remainder is complexed with citrate. Such high levels of free calcium in ejaculated semen until capacitation reaction may actually hinder sperm motility. With this study, Bhoumik et al. demonstrated the importance of maintaining optimum levels of calcium for enhanced motility [23 28]. Furthermore, along with the optimum extracellular Ca\(^{2+}\), bicarbonate ions are important anions that are transported into sperm during capacitation; they are important in the influx of Ca\(^{2+}\) ions [32]. Wennemuth et al. have reported that in vitro treatment with bicarbonate ion induces an influx of Ca\(^{2+}\), subsequently increasing flagellar beat frequency but decreasing flagellar beat asymmetry [33]. Therefore, it is clear that calcium plays a critical role in regulating sperm motility. Moreover, maintaining optimum concentrations of calcium is particularly important in ART.

**Role of pH on sperm motility**

All biophysiological events are dependent on pH. Even a small alteration in the pH can either lead to a deviation in the function or even lead to the inhibition of a particular cell function [41]. Sperm cells encounter several changes in pH right from the maturation stage to the stage when they enter the uterotubal tract. Sperm cells encounter a pH of 7.2–7.4 in the mammalian seminiferous tubules, 6.5 in the caput epididymis, and 6.7–6.8 in the cauda epididymis. The pH of the semen during ejaculation becomes 7.2–7.4. On ejaculation into the vagina, the sperm encounters a pH of 4.5–7.5. Subsequently, the pH increases to 6.5–7.5 in the cervix and 7–7.8 in the uterus and fallopian tube [42]. Being the only cell that performs its activity outside the male body, sperm cells are largely affected by the surrounding environment. Critical processes such as motility, viability, capacitation, and acrosome reaction are vastly influenced by

**Reactive oxygen species and its role in sperm motility**

Reactive oxygen species (ROS) like superoxide anion, nitric oxide, and hydrogen peroxide play a crucial role in regulating sperm motility [34]. ROS are highly reactive and are known for their ability to initiate a cascade of chain reactions that lead to extensive cellular damage. Free radicals are usually by-products of different metabolic processes. An increase in ROS levels affects mitochondrial oxidative phosphorylation which further damages proteins and lipids. The plasma membrane in sperm contains lipids in the form of polyunsaturated fatty acids (PUFA). An increased amount of ROS results in lipid peroxidation thus affecting the integrity of the plasma membrane [7]. However, a small quantity of ROS is essential for normal sperm functioning. ROS can function as signalling molecules and are crucial in capacitation and acrosome reaction, as well as motility of mature sperm cells [35]. It is important that ROS is maintained at appropriate levels to ensure proper physiological function while preventing pathological damage to the sperm. Minute levels of superoxide ions trigger hyperactivation and capacitation [35, 36]. Generally, ROS originate in the ejaculate by the spontaneous production of nicotinamide adenine dinucleotide phosphate oxidase system at the sperm plasma membrane level [37] and by the mitochondria in sperm [15, 38] as well as from leukocytes that infiltrate into semen [39]. Leukocytospermia, chronic inflammation, genitourinary tract infections, and various conditions such as varicocele, orchitis, cryptorchidism, and ageing lead to an increased amount of ROS, thus causing a decrease in sperm motility [40, 41]. Antioxidants like glutathione reductase and peroxidase in the epididymis and testes protect the lipid components on the sperm, thus preserving viability and motility. Superoxide dismutase, catalase, and glutathione peroxidase are some of the important antioxidant enzyme systems present in the semen [7, 35]. Unhealthy lifestyles such as alcohol abuse, smoking, exposure to chemical pollutants, and electromagnetic radiation have increased instances of oxidative stress in the body. Sperm cells are highly susceptible to ROS-induced damage. ROS affects several critical processes like signal transductions in the sperm cell thereby contributing to impaired function. Therefore, it is critical to understand the mechanisms of how ROS and endogenous antioxidant systems affect sperm function.
Sperm motility is restored in alkaline conditions [44]. The sperm of Anodonta woodiana Pacifica Houde showed enhanced motility at pH 8.5, and there was a significant decline in the motility with the decrease in pH [45]. There are several mechanisms by which the pH in spermatozoa and its surrounding is regulated across different organisms. For instance, in fishes, the pituitary gland and gonadotropin stimulate the production of 17α-hydroxyprogesterone in testicular somatic cells resulting in a marked increase in the production of 7α, 2β-dihydroxy-4-pregnen-3-one in spermatozoa; this increases the pH in the sperm duct, as well as the intrasperm cAMP, which initiates motility [46]. The mechanism that regulates pH in mammalian spermatozoa is more complex. HCO$_3^-$ influx, voltage-gated proton channel (Hv1), and Na$^+$/H$^+$ exchanger (NHE) are the three mechanisms by which pH is regulated in spermatozoa [42]. The HCO$_3^-$ influx system involves the inward movement of HCO$_3^-$ ions. Activation of soluble ACs results in the production of cAMP, thus promoting alkalization and membrane hyperpolarization. Na$^+$ plays a crucial role in this system as removal of Na$^+$ ions prevents alkalization and hyperpolarization, indicating the presence of Na$^+$/HCO$_3^-$ co-transport mechanism in spermatozoa [47]. Carbonic anhydrase, an enzyme that catalyzes the reaction of CO2 and produces HCO$_3^-$, plays an important role in the entrance of HCO$_3^-$ [48]. Cystic fibrosis transmembrane is another regulator that plays a crucial role in the influx of HCO$_3^-$ during capacitation [49]. The NHE is another mechanism that regulates the pH in spermatozoa. The NHE is present in the flagellum; a study noted that male mice that lacked NHEs were sterile and showed diminished sperm motility [50]. Another important H$^+$ transporter that is present across the membrane of the spermatozoa is the Hv1 channel or the voltage-gated proton channel. This membrane transporter is abundantly localized in the flagellum of the spermatozoa. A characteristic feature of this voltage-gated proton channel is that it is activated at low intracellular pH. Moreover, it conducts protons much more rapidly and efficiently and conducts them unidirectionally to the extracellular space, thus regulating the internal pH [42]. Therefore, it is clear that the pH regulates sperm motility by various mechanisms. HCO$_3^-$/CO$_2$ inorganic ions, organic acids, sugars, lipids, steroids, amino acids, polyamines, nitrogenous bases, and proteins in the seminal plasma usually contribute to a buffering action. A disturbance in the pH of the seminal plasma contributes to impaired sperm motility and function. Therefore, understanding the effects of changes in seminal plasma pH may be useful in the treatment of impaired motility and infertility.

**Analysing sperm motility**

Sperm motility plays an important role in assisted reproductive technology. Semen analysis of infertile men is crucial as it reflects the overall functionality of sperm production by the testes and the patency of the genital tract, as well as the secretory activity of all accessory glands [51]. Ejaculate volume, sperm density, sperm motility, and sperm morphology are the typical parameters that are analysed during sperm analysis. The lower limits of normal as defined by the World Health Organization are as follows: semen volume 1.5 mL, total sperm number 39 million/ejaculate, sperm concentration 15 M/mL, total motility 40%, progressive motility 32%, and morphologically normal forms 4%. Hormonal evaluation is indicated in men with sperm concentration < 10 M/mL or with clinical evidence of an endocrinopathy. Genetic evaluation by karyotype and for Y chromosome microdeletions should be considered in men with sperm concentration < 10 M/mL, and cystic fibrosis genetic testing should be performed in men with congenital absence of the vas deferens. Sperm motility is predominantly assessed using microscopic methods. Different types of movements such as progressive motility, non-progressive motility, or no motility are observed when evaluating motility [52].

More recently, methods such as the light scattering method, laser beam method, and multiple exposure photographic method have come into use for measuring sperm motility. Measuring horizontal velocity is a major drawback when using the aforementioned methods. In the recent past, a highly automated instrument has been developed to measure sperm motility. Computer-assisted semen analysis (CASA) is a sophisticated system that utilizes complex software and hardware, a high-resolution camera, and a microscope to analyse sperm concentration, morphology, and motility. Although most methods measure horizontal velocity, measuring vertical velocity is also important as only a smaller percentage of the spermatozoa exhibit vertical velocity. The movement of sperm on the vertical plane is important because healthy and motile sperm cells are expected to be more active and functional, enabling them to travel the entire female reproductive tract to fertilize the egg cell. Saha et al. have developed a method of measuring vertical sperm velocity.
using a novel computer-assisted method. In this method, an electromechanical system comprising a modified cuvette holder and a stepper motor was used. With this novel method, vertically moving sperm cells are detected at different heights using newly developed software. Since no method describes a way of measuring vertical velocity, this method highlights the importance of the vertical movement of sperm [53].

Methods adopted in cases with inadequate sperm motility

Normal reproduction is an extremely complex process that involves an array of complicated steps. As mentioned earlier, the interaction between the sperm and the egg includes a complex set of reactions. The production of a sufficient number of sperm cells having adequate motility for it to travel through the vaginal canal into the fallopian tube and the ability to bring about fertilization is of utmost importance. These processes are highly regulated; any error at any stage drastically reduces the chances of conception. Assisted reproductive technologies (ARTs) are clinical methods that involve the in vitro handling of sperm, oocytes, and embryos for their use in reproduction [54]. ARTs came in as a ray of hope for individuals suffering from infertility, giving them an opportunity to conceive successfully.

The world’s first in vitro fertilization (IVF) baby, Louise Brown, was born at Oldham General hospital on 25th July 1978. In 2010, Robert G Edwards was awarded the Nobel Prize for Physiology and Medicine for the development of IVF [55]. Since the development of the IVF method, there has been a steady increase in the use of ARTs over the last decades. Initially, it was presumed that male factor infertility was a contraindication to IVF because abnormal sperm are less likely to fertilize oocytes than normal sperm. However, subsequent analyses performed over a decade ago revealed that fertilization and subsequent live births were possible despite impaired sperm quality [56].

As of 2018, as many as eight million babies have been conceived by ART [55]. ARTs are routinely used in veterinarian clinical settings. Artificial insemination is the most routinely used method in breeding programs because of its simplicity and effectiveness [57]. Despite the huge success and the ever-growing research on ARTs, its efficiency still has a lot of potential to improve. The American and European societies of reproduction and fertility have reported the efficiencies of ICSI or IVF to be 37% and 25% of pregnancies and deliveries, respectively, per embryo transfer [58]. There are several reasons that contribute to low efficiencies such as suboptimal in vitro conditions, injuries associated with gametes and embryo manipulation, subjacent male and female factors, etc. Although ART has been the solution to problems associated with conception, it is still an expensive and invasive method. Therefore, there is a need to develop and discover novel methods to tackle infertility caused by impaired motility.

The role of sperm surface molecules in sperm motility

As mentioned earlier, immature sperm undergo several biochemical changes as they pass through different parts of the epididymis during maturation. During the epididymal transit, there is a marked increase in the intrasperm level of cAMP and pH, thus suggesting that elevated intrasperm levels of cAMP and pH play an important role in the in vivo initiation of sperm forward progression. However, not much is known about the molecular basis of the initiation of flagellar motility while it is in the epididymis and its subsequent regulation. There are several cell surface molecules present on the sperm cell membrane. These molecules play a pivotal role in modulating cell–cell interactions, effector–receptor interactions, membrane permeability, membrane fluidity, transmembrane signalling, etc. Similarly, cell surface molecules present on sperm cells play a crucial role in processes like capacitation, acrosomal reaction, and fertilization [59–61]. A summary of these molecules is described in Fig. 3.

Lipids are important biomolecules that form the membrane bilayer permeability barrier of cells and organelles. Rana et al. have reported that there was a significant decrease in phospholipids (PL) and glycolipids (GL) of caprine sperm membrane during sperm maturation. Among the phospholipids, it was observed that the amounts of phosphatidylethanolamine showed the maximum decrease. However, an enhanced neutral lipid (NL) fraction was noted during the transit from caput to cauda. Sterol and steryl esters, which are the major constituents of NL, were enhanced during maturation while other membrane-bound neutral lipids decreased. Since cholesterol (CH) was the major component of the sterol fraction, the aforementioned changes led to appreciable enhancement in the cholesterol/phospholipid (CH/PL) ratio [62]. These findings suggest an important relationship between the lipid profile of the sperm plasma membrane and the maturation of sperm cells. However, the exact role of these changes in the lipid content on motility is still unknown. It is clear that cholesterol plays a crucial role in the capacitation reaction. During capacitation, a marked efflux of cholesterol results in an increase in membrane fluidity and permeability to bicarbonate and calcium ions which in turn affect motility.

Along with lipids, there are several regulatory protein complexes present on the surface of sperm cells. Majumdar et al. reported the presence of an ecto-cyclic AMP-dependent protein kinase on the external surface of rat
spermatozoa [63]. Subsequently, several publications reported the localization of cAMP-dependent protein kinases and cell surface phosphoproteins on the surface of rats and human spermatozoa [64, 65]. Extracellular cAMP binds to the cAMP-binding protein thereby disassociating the catalytic subunit. The catalytic subunit, now active, brings about the phosphorylation of exogenous proteins in the presence of ATP. The amount of ecto-cAMP-dependent protein kinase is low in immature sperm cells. However, it appears specifically during motility initiation in the mature cauda sperm, indicating that it is maturation specific. Forward-motile sperm cells are enriched in both kinases. However, the exact role of the ecto-cAMP-dependent protein kinases in the induction and regulation of sperm motility is largely unknown [65, 66].

Along with ecto-cAMP-dependent protein kinases, ecto-cAMP-independent protein kinases on the outer surfaces of goat epididymal spermatozoa are known to phosphorylate serine and threonine residues of multiple endogenous proteins localized on the sperm outer surface. The intact sperm-bound ectoenzyme is also capable of phosphorylating exogenous proteins such as casein, phosvitin, histone, and protamine [67]. It has been demonstrated that the incorporation of major physiological substrate (MPS) proteins of ecto-cAMP-independent kinases increases motility and forward motility. Moreover, at maximal MPS incorporation, the increments in motility and forward motility were also maximal [68]. Therefore, it can be conclusively said that MPS, the sperm membrane-bound phosphoprotein, serves as an activator of sperm forward motility.

Another interesting observation was noted by Roy et al., where they observed that the incorporation of Cu\(^{2+}\) enhanced forward motility of caprine spermatozoa. However, increasing the Cu\(^{2+}\) concentration beyond 5 μM reduced forward motility. This is a clear indication that Cu\(^{2+}\) exerts a biphasic regulation on sperm motility. Moreover, increasing the concentration of Cu\(^{2+}\) to beyond 100 μM led to sperm head-to-head agglutination [69]. Eventually, Roy et al. reported the presence of a novel copper-dependent sialic acid-specific lectin on the external surface of sperm cells that binds with its specific receptor of neighbouring cells thereby causing sperm–sperm agglutination. It was also noted that immature sperm cells do not undergo agglutination in the presence of copper, thus indicating that this unique copper-dependent lectin/receptor is acquired during epidydimal maturation. It has been proposed that the copper ion modulates cell surface lectin-sugar interactions. However, the mechanisms by which it affects motility need to be studied in detail [70].

Acott et al. demonstrated the induction of motility in immotile sperm cells after incubating the immotile cells

![Fig. 3](image_url)
with epididymal or seminal plasma [71]. This suggested the presence of factors in the epididymal or seminal plasma that have the ability to modulate sperm motility extracellularly.

Mandal et al. demonstrated for the first time the presence and the effect of a novel forward motility-promoting protein extracted from buffalo serum. This forward motility-stimulating factor (FMSF) is a 66-kDa heat-stable glycoprotein. It is also an Mg$^{2+}$-dependent monomeric protein, and both its protein and sugar parts of the protein are essential for its motility-promoting potential. Further studies revealed that this forward motility-stimulating protein through receptor/G-protein activation promotes the transmembrane AC activity in a dose-dependent manner to enhance intracellular cAMP and forward motility. Protein kinase A and tyrosine kinase are key players in regulating motility under the influence of forward-stimulating protein. The FMSF initiates a novel signalling cascade by stimulating tmAC activity which supplements intracellular cAMP and downstream cross-talk of phosphokinases, thus enhancing forward motility in mature spermatozoa. The FMSF binds to specific receptors that are present on the external cell surface, which leads to the activation of adenylate cyclase and consequently enhances intra-sperm cAMP level that triggers the flagellar movement through a series of cascade molecules including protein kinase A and tyrosine kinase [72, 73].

Saha et al. 2013 reported a novel motility stimulating protein (MSP) isolated and purified from caprine blood serum. This novel 66-kDa heat-stable protein-enhanced sperm horizontal forward motility as well as vertical velocity at a concentration of 0.9 μM. Furthermore, it was noted that these proteins were distributed on the surface of the sperm cells. It was also found that this motility stimulating protein showed cAMP-independent activity and that there must be alternative pathways through which motility-stimulating activity is mediated, which still needs to be identified [74].

A motility-initiating protein (MIP) and a motility-inhibiting factor (MIF) were then identified from caprine epididymal plasma [75, 76]. The addition of MIP induced forward motility, and there was a significant increase in the number of motile sperm cells. Furthermore, the addition of a rabbit polyclonal antibody raised against purified sperm motility-inhibiting factor (MIF-II) enhanced sperm motility by 75% compared with the control. A 40% increase in vertical velocity of MIF-II antibody-treated spermatozoa was observed as compared to the control serum. MIF-II antibody also enhanced the motility of immature caput spermatozoa under in vitro initiation media [77].

These novel motility-promoting proteins are promising candidates in enhancing motility in sperm samples that display poor motility traits. These proteins are physiological activators of sperm motility, and they can be used in biomedical applications in human infertility clinics and infertility management, as well as animal breeding and conservation centres. Currently, treatment of male infertility involves invasive procedures like surgery, hormonal treatment, ART, etc. which are not only expensive but also may have detrimental effects on the quality of life.

**Conclusion**

Human infertility is a social stigma in many cultures and causes mental stress and trauma, and it affects a large number of people globally. Studies show that about 40–50% of the infertility cases are attributed to male factor fertility, and as many as 2% of all men exhibit suboptimal sperm parameters. Currently, there are limited treatment strategies available for tackling infertility. There are several sophisticated ART methods (e.g., IVF and ICSI) that are routinely employed to tackle and treat individuals affected by infertility.

For several years, managing and treating male factor infertility was mostly based on “experience” and not “evidence”. Choosing the correct technique for tackling infertility is an important issue. This usually depends on the sperm parameters. However, most of these technologies have several disadvantages. They are invasive and expensive; moreover, their success rate is also very low. Most ARTs utilize hormonal treatments that have various side-effects on the individual and affect the quality of life. This review has described several molecular mechanisms that play a critical role in regulating sperm motility. Concentrations of calcium, hydrogen ions, pH, are some of the factors that affect sperm motility. An optimum concentration of extra- and intra-cellular Ca$^{2+}$, HCO$_3^{-}$, and ROS, along with the proper maintenance of pH is critical in maintaining proper sperm function. Moreover, along with the various biochemical mechanisms and the optimum microenvironments that are essential in regulating cellular functions, cell surface molecules also play a critical role in regulating various cellular processes such as transmembrane signalling, cell–cell interactions, and effector–receptor interactions. These molecular mechanisms can be exploited by using several key surface molecules to tackle impaired motility. Therefore, identifying sperm surface molecules and understanding their roles in sperm physiology are important as these will help in treating various anomalies associated with sperm motility and behaviour. These motility-promoting proteins have the potential for improving cattle and poultry breeding and conservation of endangered species. Understanding
sperm motility and the specific motility-promoting molecules may also be useful in human infertility clinics as a solution to problems associated with human infertility.

Utilizing motility-inducing proteins and other sperm surface proteins will negate hormonal treatments that may affect the quality of life of an individual. Understanding the role of spermatozoa and seminal plasma proteomes, along with surface proteins and molecules, will open up new avenues in understanding infertility and tackling it effectively.

Abbreviations

HCO3−: Bicarbonate ions; Ca2+: Calcium ions; cAMP: Cyclic adenosine monophosphate; ICSI: Intracytoplasmic sperm injection; PKA: Protein kinase A; PCD: Primary ciliary dyskinesia; ODA: Outer dynein arms; DNAH5: Dynein axonemal heavy chain 5; DNAI1: Dynein axonemal intermediate chain 1; IDA: Inner dynein arms; CDC: Coiled-coil domain-containing proteins; Spag: Sperm-associated antigen; ACs: Adenylate cyclase; ROS: Reactive oxygen species; PUFA: Polyunsaturated fatty acids; NHE: Na+/H+ exchanger; CASA: Computer-assisted semen analysis; ARTs: Assisted reproductive technologies; IVF: In vitro fertilization; PL: Phospholipids; GL: Glycolipids; NL: Neutral lipid; CH: Cholesterol; MPS: Major physiological substrate; FMSF: Forward motility-stimulating factor; WSP: Motility-stimulating protein; MIP: Motility-initiating protein; MIF: Motility-inhibiting factor.

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All authors contributed equally to this paper. Both SC and SS conceived and designed the theme of the review. SC wrote the paper under the guidance of SS. The authors read and approved the final manuscript.

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Availability of data and materials

Data sharing is not applicable to this article as no datasets were generated or analyzed during the current study.

Declarations

Ethics approval and consent to participate

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