

SPERMATOGENESIS IN FISH UNDER THE INFLUENCE OF VARIOUS FACTORS WITH SPECIAL REFENCE TO INDIAN MAJOR CARPS

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Introduction:

Spermatogenesis is a dynamic process that follows a definite pattern in nature and ultimately produces haploid spermatozoa. The cellular basis of reproduction in males is the spermatozoon. They develop from testicular germinal epithelium [3] (spermatogonial stem cells), which first proliferate (spermatogonia), then differentiate into spermatocytes that undergo meiosis to become haploid spermatids that differentiate to become spermatozoa (spermiogenesis). The process as a whole is called spermatogenesis and requires a special environment. The gonadal function changes during annual reproductive cycles in male fish. During proliferation of diploid germ cells, the spermatogonia repopulate the testis lobules by mitotic cell division. Though spermatogenesis is a natural dynamic process, it has been modulated and manipulated by various factors to get maximum viable sperms for aquaculture practice. Environment and chemicals of natural and synthetic origin can interact with the endocrine system and alter the male cell generation germ and proliferation/inhibition in fish. In the present investigative review, an attempt has been made to ascertainvarious factors that influence the spermatogonial proliferation and maturation of fish with a special reference to Indian major carps (*Catla catla, Labeo rohita, Cirrhinus mrigala and Labeo calbasu*). Methods:

Various maturity stages of fish (Indian major carps) were sampled to estimate their testis somatic index (TSI) at different times of the year. The immature fish were subjected to several external factors (environment) and some chemical inducers such as hormones and other inducing agents to estimate the semen quantity and quality and assess the spermatogonial proliferation due to these and also the regulation of spermatogenesis annually. A comparative evaluation of spermatogenic cycles with other species has also been done.

Results and Discussion:

Fish species inhabiting diverse ecological niche offers an enormous challenge to generalize their reproductive characters that includes theirsexual dimorphism, spermatogenic cycle (*in vivo and in vitro*), spermatozoon structure and the external inducers that modulate the spermatogenesis in fish with particular reference to Indian major carps (IMCs).The testes in

Table 1. A comparative account of gonadal (spermatogenic cycle) in different fishes			
Species	Spermatogenesis	Inducement	Reference
Catfish,	Annual gonadal cycle	Natural cycle	[9]
H. fossilis			
Rainbow trout,	Twice in year (Autumn and summer)	Natural cycle	[1]
S. gairdneri			
Rainbow trout,	Advanced maturation and spawning by 6-12 weeks at 9°C	Photo period	[12]
S. gairdneri		manipulation	
Tilapia	Testicular activity restricted to autumn and winter, and testes	Natural	[4]
	are in resting phase from July-September in equatorial zone	photoperiod	
Indian major carps	Increasing gonadal recrudescence; males attain maturity earlier	Photo period	[10]
	than females	manipulation	
Indian major carps	Advancement of sexual maturation and off-season spawning of	Photothermal	[7]
	IMCs	manipulation	
Zebra fish	Combined duration of meiotic and spermiogenic phases is very	Natural cycle	[5]
Danio rerio	short in this species and lasts approximately 6 days		
C. fasciata	Spermatogenicphase 6 days	Natural cycle	[8]
Black molly,	Spermatogenic phase 21days	Natural cycle	[2]
P. sphenops			
IMCs &	Spermatogenic phase 15-21 days	GnRH based	Present study
L.calbasu		inducement	

Proceedings of 9th International Symposium on Reproductive Physiology of Fish, Cochin, India.



most teleost reported including that of the IMCs are bilobed, with spermatogonia dispersed throughout the gametogenic epithelium of the seminiferous tubules. The sexual dimorphism is clear in species during the reproductive season (perennial spawners in nature) however, some are very inconspicuous during nonspawning season (IMCs) [11, 6]. The spermatogenic cycle varies among the fish species (Table 1) and also the sperm density (40 to 53,000 millions/ml). With the advancements in reproductive biology of fishes, it is now possible that spermatogenesis can be modulated and manipulated by various factors to get maximum viable sperms for aquaculture practice.

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