FDA-Approved Drugs that Can Negatively Affect Sperm

Drug category*	Generic name of drug	Adverse impact on human	Supported/refuted by PubMed publications**
Diag category	deficite fiame of all ag	spermatogenesis	bapported/related by rabbled publications
		Decrease in sperm motility and seminal	
		vesicle secretions, abnormal sperm	
Analgesic	Methadone hydrochloride	morphology	Supported in humans [12,13]
	Pregabalin	Epididymitis (rare)	Refuted in humans [14]
	Gabapentin	Epididymitis (rare)	Refuted in rats [15], no data for humans
Anti-arrhythmic agent	Amiodarone hydrochloride	Epididymitis (rare)	Supported in rats [16], no data for humans
Anti-bacterial agent	Lomefloxacin hydrochloride	Epididymitis, orchitis (<1% of patients)	No data published for animals or humans
		Spermatogenic arrest/decreased sperm	
	Nitrofurantoin	count (high doses)	Supported in humans [17]
	Dapsone	Orchitis, male infertility	Supported in rats [18], no data in humans
Anticonvulsant	Lamotrigine	Epididymitis (rare)	Supported in humans [19]
		Reversible reduction in ability to fertilize	
Antihypertensive agent	Nifedipine	ova	Supported in humans [23]
Anti-infective agent	Voriconazole	Epididymitis (<2% of all patients)	No data published for animals or humans
		Changes in the motility and number of	No specific data published for methylprednisone,
	Methylprednisolone/prednisone	spermatozoa	prednisone is supported in humans [30]
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	Sulfasalazine	Reversible oligozoospermia and infertility	Supported in humans [31]
		Changes in the motility and number of	
	Triamcinolone hexacetonide	spermatozoa	No data published for animals or humans
		Damage to spermatozoa and testicular	A
Anti Parkinson agent	Busulfan	tissue, azoospermia, testicular atrophy	Supported in humans [32]
	Chlorambucil	Azoospermia (prolonged or permanent)	Supported in humans [33]
		Interferes with spermatogenesis,	
		testicular atrophy, azoospermia,	
	Cyclophosphamide	oligozoospermia	Supported in humans [34]
	gyelepheephamae	Impaired spermatogenesis, decreased	Supported in Hamano [O 1]
	Dabrafenib mesylate	sperm count	No data published for animals or humans
	Degarelix	Testicular atrophy	No data published for animals or humans
	Degat en A	Damage to spermatozoa and testicular	The united published for unifficient
	Fludarabine phosphate	tissue	Supported in humans [35]
	Tradarabilic phosphate	tissue	Supported for rodents [38,39], conflicting data for
	Methotrexate sodium	Oligozoospermia (reversible)	humans ($[40]$ (-) vs. $[41]$ (+))
	Method cade Souldin	ongozoosperima (reversible)	Supported in mice [42], unclear for humans (all
	Procarbazine hydrochloride	Azoospermia	trials have used combination therapy $[43]$)
	Triptorelin pamoate	Testicular atrophy	Supported in humans [44]
	Triptorenni pannoate	resticular atrophy	Supported in rats [45] and in vitro for humans
	Vinblastine sulfate	Azoosparmia	Supported in rats $[\underline{45}]$ and m vitro for numans $[\underline{46}]$
	Vinorelbine tartrate	Azoospermia	No data published in animals or humans
		Damage to spermatozoa Orchitis	-
	Thalidomide		Supported in rabbits [47], no data for humans
Anti-Parkinson agent	Pramipexole dihydrochloride	Epididymitis, orchitis	No data published for animals or humans
Antipsychotic agent	Quetiapine fumarate	Orchitis (infrequent)	No data published for animals or humans

		Azoospermia or oligozoospermia (~1% of	
Anti-rejection drug	Everolimus	patients)	No data published for animals or humans
Antiviral agent	Delavirdine mesylate	Hematospermia, epididymitis	No data published for animals or humans
		Testicular hypotrophy, aspermatogenesis	
	Ganciclovir/ganciclovir sodium	(dose-dependent)	Supported in rats [48], no data in humans
			No specific data published, but is expected to
	Valganciclovir	Inhibition of spermatogenesis	result in the same effects as ganciclovir
Cardiovascular agent	Bosentan	Decreased sperm count	No data published for animals or humans
		Decreased spermatogenesis,	
Hormones, hormone substitutes and		abnormalities in semen volume, viscosity,	
hormone antagonists	Danazol	sperm count, and motility	Supported in humans [51]
		Decreased sperm count, semen volume,	
	Dutasteride	and sperm motility	Supported in humans [52]
		Decreased ejaculate volume and total	
	Finasteride	sperm per ejaculation (reversible)	Supported in humans [52]
		Interference with testosterone, decreased	Supported in mice [53], supported in combination
	Flutamide	sperm count	with other agents in humans [<u>54</u>]
			Supported in rats [55] and rhesus monkeys [56],
	Histrelin acetate	Testicular atrophy	no data for humans
			Supported in humans [57], also may be useful for
		Suppressed testicular steroidogenesis,	protecting/restoring fertility following toxic
	Leuprolide acetate	testicular atrophy	insults [<u>58,59</u>]
		Oligozoospermia, suppressed	Supported in dogs [60] and rats [61], no data for
	Methyltestosterone	spermatogenesis	humans
		Inhibition of testicular function, testicular	
		atrophy and oligozoospermia,	
	Nandrolone decanoate	epididymitis	Supported in humans [62]
	Nilutamide	Testicular atrophy	No data published for animals or humans
		Suppressed spermatogenesis, inhibition of	
		testicular function, testicular atrophy,	Supported in rats [63] and in a human case report
	Oxandrolone	oligozoospermia, epididymitis	[<u>64</u>]
		Inhibition of testicular function, testicular	
		atrophy, oligospermia, decreased seminal	Supported in mice [65] and rats [66], no data for
	Oxymetholone	volume, epididymitis	humans
		Suppressed	
	Testosterone/testosterone	spermatogenesis/oligozoospermia,	
	cypionate/testosterone	testosterone undecanoate may also cause	Supported in humans [<u>67</u> - <u>69</u>], but dose-
	enanthate/testosterone undecanoate	spermatocele formation	dependent and variable results
Immunosuppressant	Sirolimus	Azoospermia (reversible)	Supported in humans [70]
		Impairment of testicular	
Radioactive compound	Sodium iodide I 131	function/transient infertility	Supported in humans [73]

^{*}Some drugs can be classified into multiple categories. They have been classified here on the basis of their most common indication/target.

Source: FDA-approved drugs that have the potential to impair human spermatogenesis or cause more mutations in sperm Oncotarget. 2017 Feb 7; 8(6): 10714–10725.

^{**}Although the drugs noted to support the DailyMed labels were all found to affect some aspect of human spermatogenesis, the effect was not always the same as that listed in the drug label.