SGS/LG

SENATE STATE OF MINNESOTA NINETY-SECOND SESSION

S.F. No. 1884

(SENATE AUTI	HORS: PAPP	AS)
DATE	D-PG	OFFICIAL STATUS
03/08/2021		Introduction and first reading
		Referred to Health and Human Services Finance and Policy

1.1	A bill for an act
1.2 1.3 1.4 1.5 1.6	relating to health insurance; establishing supply requirements for prescription contraceptives; requiring health plans to cover contraceptive methods, sterilization, and related medical services, patient education, and counseling; establishing accommodations for eligible organizations; amending Minnesota Statutes 2020, section 256B.0625, subdivision 13; proposing coding for new law in Minnesota
1.7	Statutes, chapter 62Q.
1.8	BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MINNESOTA:
1.9	Section 1. [62Q.521] COVERAGE OF CONTRACEPTIVE METHODS AND
1.10	SERVICES.
1.11	Subdivision 1. Definitions. (a) The definitions in this subdivision apply to this section.
1.12	(b) "Closely held for-profit entity" means an entity that:
1.13	(1) is not a nonprofit entity;
1.14	(2) has more than 50 percent of the value of its ownership interest owned directly or
1.15	indirectly by five or fewer individuals, or has an ownership structure that is substantially
1.16	similar; and
1.17	(3) has no publicly traded ownership interest, having any class of common equity
1.18	securities required to be registered under United States Code, title 15, section 781.
1.19	For purposes of this paragraph:
1.20	(i) ownership interests owned by a corporation, partnership, estate, or trust are considered
1.21	owned proportionately by that entity's shareholders, partners, or beneficiaries;

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2.1	(ii) owne	ership interests ow	ned by a nonprofit	entity are considered ov	vned by a single
2.2	owner;				
2.3	(iii) own	ership interests ov	vned by an individ	al are considered owned	d, directly or
2.4	indirectly, b	y or for the individ	lual's family. For p	urposes of this item, "fa	mily" means
2.5	brothers and	l sisters, including	half-brothers and h	alf-sisters, a spouse, and	cestors, and lineal
2.6	descendants	; and			
2.7	(iv) if an	individual or enti	ty holds an option	to purchase an ownershi	p interest, the
2.8	individual o	r entity is consider	red to be the owner	of those ownership inte	erests.
2.9	<u>(c) "Con</u>	traceptive method"	means a drug, dev	ice, or other product app	roved by the Food
2.10	and Drug A	dministration to pr	event unintended	oregnancy.	
2.11	<u>(d)</u> "Con	traceptive service'	' means consultation	on, examination, procedu	ures, and medical
2.12	services rela	ted to the preventi	on of unintended p	regnancy. This includes	but is not limited
2.13	to voluntary	sterilization proce	edures, patient edu	cation, counseling on co	ntraceptives, and
2.14	follow-up se	rvices related to co	ontraceptive metho	ds or services, manageme	ent of side effects,
2.15	counseling f	or continued adhe	rence, and device i	nsertion or removal.	
2.16	<u>(e) "Elig</u>	ible organization"	means an organiza	tion that opposes provid	ling coverage for
2.17	some or all o	contraceptive meth	nods or services on	account of religious obj	jections and that
2.18	is:				
2.19	<u>(1) organ</u>	nized as a nonprofi	t entity and holds	tself as a religious organ	nization; or
2.20	(2) organ	nized and operates	as a closely held f	or-profit entity, and the o	organization's
2.21	highest gove	rning body has add	opted, under the org	ganization's applicable ru	les of governance
2.22	and consiste	nt with state law, a	a resolution or sim	ilar action establishing t	hat it objects to
2.23	covering son	ne or all contracep	tive methods or se	rvices on account of the	owners' sincerely
2.24	held religiou	is beliefs.			
2.25	<u>(f)</u> "Med	ical necessity" inc	ludes but is not lin	nited to considerations s	uch as severity of
2.26	side effects,	difference in perma	anence and reversa	bility of a contraceptive r	method or service,
2.27	and ability t	o adhere to the app	propriate use of the	contraceptive method c	or service, as
2.28	determined	by the attending p	covider.		
2.29	<u>(g)</u> "Reli	gious organizatior	" means an organi	zation that is organized	and operates as a
2.30	nonprofit en	tity and meets the r	equirements of sec	tion 6033(a)(3)(A)(i) or ((iii) of the Internal
2.31	Revenue Co	de of 1986, as am	ended.		

3.1	(h) "Therapeutic equivalent version" means a drug, device, or product that can be expected
3.2	to have the same clinical effect and safety profile when administered to a patient under the
3.3	conditions specified in the labeling, and that:
3.4	(1) is approved as safe and effective;
3.5	(2) is a pharmaceutical equivalent, (i) containing identical amounts of the same active
3.6	drug ingredient in the same dosage form and route of administration, and (ii) meeting
3.7	compendial or other applicable standards of strength, quality, purity, and identity;
3.8	(3) is bioequivalent in that:
3.9	(i) the drug, device, or product does not present a known or potential bioequivalence
3.10	problem and meets an acceptable in vitro standard; or
3.11	(ii) if the drug, device, or product does present a known or potential bioequivalence
3.12	problem, it is shown to meet an appropriate bioequivalence standard;
3.13	(4) is adequately labeled; and
3.14	(5) is manufactured in compliance with current manufacturing practice regulations.
3.15	Subd. 2. Required coverage; cost sharing prohibited. (a) A health plan must provide
3.16	coverage for contraceptive methods and services.
3.17	(b) A health plan company must not impose cost-sharing requirements, including co-pays,
3.18	deductibles, or co-insurance, for contraceptive methods or services.
3.19	(c) Notwithstanding paragraph (b), a health plan that is a high-deductible health plan in
3.20	conjunction with a health savings account must include cost-sharing for contraceptive
3.21	methods and services at the minimum level necessary to preserve the enrollee's ability to
3.22	make tax exempt contributions and withdrawals from the health savings account, as provided
3.23	by section 223 of the Internal Revenue Code of 1986, as amended.
3.24	(d) A health plan company must not impose any referral requirements, restrictions, or
3.25	delays for contraceptive methods or services.
3.26	(e) A health plan must include at least one of each type of Food and Drug Administration
3.27	approved contraceptive method in its formulary. If more than one therapeutic equivalent
3.28	version of a contraceptive method is approved, a health plan must include at least one
3.29	therapeutic equivalent version in its formulary, but is not required to include all therapeutic
3.30	equivalent versions.
3.31	(f) For each health plan, a health plan company must list the contraceptive methods and
3.32	services that are covered without cost-sharing in a manner that is easily accessible to

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4.1	enrollees, health care providers, and representatives of health care providers. The list for
4.2	each health plan must be promptly updated to reflect changes to the coverage.
4.3	(g) If an enrollee's attending provider recommends a particular contraceptive method or
4.4	service based on a determination of medical necessity for that enrollee, the health plan must
4.5	cover that contraceptive method or service without cost-sharing. The health plan company
4.6	issuing the health plan must defer to the attending provider's determination that the particular
4.7	contraceptive method or service is medically necessary for the enrollee.
4.8	Subd. 3. Religious employers; exempt (a) A religious employer is not required to cover
4.9	contraceptive methods or services if the employer has religious objections to the coverage.
4.10	A religious employer that chooses to not provide coverage for contraceptive methods and
4.11	services must notify employees as part of the hiring process and to all employees at least
4.12	30 days before:
4.13	(1) an employee enrolls in the health plan; or
7.15	
4.14	(2) the effective date of the health plan, whichever occurs first.
4.15	(b) If the religious employer provides coverage for some contraceptive methods or
4.16	services, the notice must provide a list of the contraceptive methods or services the employer
4.17	refuses to cover.
4.18	Subd. 4. Accommodation for eligible organizations. (a) A health plan established or
4.19	maintained by an eligible organization complies with the requirements of subdivision 2 to
4.20	provide coverage of contraceptive methods and services if the eligible organization provides
4.21	notice to any health plan company the eligible organization contracts with that it is an eligible
4.22	organization and that the eligible organization has a religious objection to coverage for all
4.23	or a subset of contraceptive methods or services.
4.24	(b) The notice from an eligible organization to a health plan company under paragraph
4.25	(a) must include (1) the name of the eligible organization, (2) a statement that it objects to
4.26	coverage for some or all of contraceptive methods or services, including a list of the
4.27	contraceptive methods or services the eligible organization objects to, if applicable, and (3)
4.28	the health plan name. The notice must be executed by a person authorized to provide notice
4.29	on behalf of the eligible organization.
4.30	(c) An eligible organization must provide a copy of the notice under paragraph (b) to
4.31	prospective employees as part of the hiring process and to all employees at least 30 days
4.32	before:
4.33	(1) an employee enrolls in the health plan; or

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5.1	(2) the e	ffective date of the	health plan, which	never occurs first.	
5.2	(d) A he	alth plan company	that receives a cor	by of the notice under pa	ragraph (a) with
5.3	<u> </u>			l by an eligible organizat	
5.4	(1) expre	essly exclude cover	age for some or al	l contraceptive methods	or services from
5.5	the health pl		8	I	
5.6			nts for any contrac	eptive methods or servic	as required to be
5.7	· / 2	· · ·	*	g as the enrollee remains	<u> </u>
5.8	health plan.				
5.9	(e) The h	nealth plan compan	y must not impose	any cost-sharing require	ements, including
5.10	co-pays, dec	luctibles, or co-insu	arance, or directly	or indirectly impose any	premium, fee, or
5.11	other charge	for contraceptive s	ervices or method	s on the eligible organiza	tion, health plan,
5.12	or enrollee.				
5.13	(f) On Ja	nuary 1, 2023, and	every year thereaf	fter a health plan compan	y must notify the
5.14	commission	er, in a manner to b	be determined by t	he commissioner, regard	ing the number
5.15	of eligible o	rganizations grante	d an accommodat	ion under this subdivisio	on.
5.16	EFFEC	FIVE DATE. This	section is effective	e January 1, 2022, and ap	plies to coverage
5.17	offered, sold	l, issued, or renewe	ed on or after that	date.	
5.18				CRIPTION CONTRAC	<u>'EPTIVES;</u>
5.19	SUPPLY R	EQUIREMENTS	<u>.</u>		
5.20	Subdivis	tion 1. Scope of co	verage. Except as	otherwise provided in se	ection 62Q.521,
5.21	subdivision	3, all health plans	hat provide presen	ription coverage must co	mply with the
5.22	requirement	s of this section.			
5.23	<u>Subd. 2.</u>	Definition. For pu	rposes of this sect	ion, "prescription contra	ceptive" means
5.24	any drug or	device that require	s a prescription an	d is approved by the Foo	od and Drug
5.25	Administrat	ion to prevent preg	nancy. Prescriptio	n contraceptive does not	include an
5.26	emergency of	contraceptive drug	that prevents preg	nancy when administere	d after sexual
5.27	contact.				
5.28	<u>Subd. 3.</u>	Required coverag	ge. (a) Health plan	coverage for a prescript	ion contraceptive
5.29	must provid	e a 12-month supp	ly for any prescrip	tion contraceptive, regar	dless of whether
5.30	the enrollee	was covered by the	e health plan at the	e time of the first dispens	sing.
5.31	<u>(b)</u> The p	prescribing health c	eare provider must	determine the appropria	te number of
5.32	months to p	rescribe the prescri	ption contraceptiv	es for, up to 12 months.	

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6.1	EFFEC	FIVE DATE. This	section is effective	ve January 1, 2022, and a	pplies to coverage	
6.2	EFFECTIVE DATE. This section is effective January 1, 2022, and applies to coverage offered, sold, issued, or renewed on or after that date.					
		<u> </u>				
6.3	Sec. 3. Mi	nnesota Statutes 20	020, section 256B	.0625, subdivision 13, is	amended to read:	
6.4	Subd. 13	. Drugs. (a) Medie	cal assistance cov	ers drugs, except for fert	ility drugs when	
6.5	specifically	used to enhance fe	rtility, if prescribe	ed by a licensed practitio	ner and dispensed	
6.6	by a license	d pharmacist, by a	physician enrolle	d in the medical assistan	ce program as a	
6.7	dispensing p	bhysician, or by a p	ohysician, a physi	cian assistant, or an adva	inced practice	
6.8	registered m	urse employed by	or under contract	with a community health	board as defined	
6.9	in section 14	15A.02, subdivisio	n 5, for the purpo	ses of communicable dis	sease control.	
6.10	(b) The c	lispensed quantity	of a prescription	drug must not exceed a 3	34-day supply,	
6.11	unless autho	rized by the comm	nissioner or as pro	wided in paragraph (h).		
6.12	(c) For th	ne purpose of this :	subdivision and s	ubdivision 13d, an "activ	e pharmaceutical	
6.13	ingredient"	s defined as a sub-	stance that is repr	esented for use in a drug	and when used in	
6.14	the manufacturing, processing, or packaging of a drug becomes an active ingredient of the					
6.15	drug product. An "excipient" is defined as an inert substance used as a diluent or vehicle					
6.16	for a drug. The commissioner shall establish a list of active pharmaceutical ingredients and					
6.17	excipients which are included in the medical assistance formulary. Medical assistance covers					
6.18	selected active pharmaceutical ingredients and excipients used in compounded prescriptions					
6.19	when the compounded combination is specifically approved by the commissioner or when					
6.20	a commercia	ally available prod	uct:			
6.21	(1) is not	t a therapeutic opti	on for the patient	;		
6.22	(2) does	not exist in the sar	ne combination o	f active ingredients in th	e same strengths	
6.23	as the comp	ounded prescriptio	n; and			
6.24	(3) canno	ot be used in place	of the active pha	rmaceutical ingredient in	the compounded	
6.25	prescription					
6.26	(d) Medi	cal assistance cove	ers the following o	over-the-counter drugs w	hen prescribed by	
6.27	a licensed pr	ractitioner or by a	licensed pharmac	ist who meets standards	established by the	
6.28	commission	er, in consultation	with the board of p	bharmacy: antacids, aceta	minophen, family	
6.29	planning pro	oducts, aspirin, ins	ulin, products for	the treatment of lice, vit	amins for adults	
6.30	with docum	ented vitamin defic	ciencies, vitamins	for children under the a	ge of seven and	
6.31	pregnant or	nursing women, ar	nd any other over	the-counter drug identif	ied by the	
6.32	commission	er, in consultation	with the Formula	ry Committee, as necess	ary, appropriate,	
6.33	and cost-effe	ective for the treat	ment of certain sp	ecified chronic diseases,	conditions, or	

disorders, and this determination shall not be subject to the requirements of chapter 14. A
pharmacist may prescribe over-the-counter medications as provided under this paragraph
for purposes of receiving reimbursement under Medicaid. When prescribing over-the-counter
drugs under this paragraph, licensed pharmacists must consult with the recipient to determine
necessity, provide drug counseling, review drug therapy for potential adverse interactions,
and make referrals as needed to other health care professionals.

7.7 (e) Effective January 1, 2006, medical assistance shall not cover drugs that are coverable under Medicare Part D as defined in the Medicare Prescription Drug, Improvement, and 7.8 Modernization Act of 2003, Public Law 108-173, section 1860D-2(e), for individuals eligible 7.9 for drug coverage as defined in the Medicare Prescription Drug, Improvement, and 7.10 Modernization Act of 2003, Public Law 108-173, section 1860D-1(a)(3)(A). For these 7.11 individuals, medical assistance may cover drugs from the drug classes listed in United States 7.12 Code, title 42, section 1396r-8(d)(2), subject to this subdivision and subdivisions 13a to 7.13 13g, except that drugs listed in United States Code, title 42, section 1396r-8(d)(2)(E), shall 7.14 not be covered. 7.15

(f) Medical assistance covers drugs acquired through the federal 340B Drug Pricing
Program and dispensed by 340B covered entities and ambulatory pharmacies under common
ownership of the 340B covered entity. Medical assistance does not cover drugs acquired
through the federal 340B Drug Pricing Program and dispensed by 340B contract pharmacies.

(g) Notwithstanding paragraph (a), medical assistance covers self-administered hormonal
contraceptives prescribed and dispensed by a licensed pharmacist in accordance with section
151.37, subdivision 14; nicotine replacement medications prescribed and dispensed by a
licensed pharmacist in accordance with section 151.37, subdivision 15; and opiate antagonists
used for the treatment of an acute opiate overdose prescribed and dispensed by a licensed
pharmacist in accordance with section 151.37, subdivision 16.

(h) Medical assistance coverage for a prescription contraceptive must provide a 12-month
supply for any prescription contraceptive, regardless of whether the enrollee was covered
by medical assistance or the health plan at the time of the first dispensing. The prescribing
health care provider must determine the appropriate number of months to prescribe the
prescription contraceptives for, up to 12 months.

7.31 For purposes of this paragraph, "prescription contraceptive" means any drug or device that

7.32 requires a prescription and is approved by the Food and Drug Administration to prevent

7.33 pregnancy. Prescription contraceptive does not include an emergency contraceptive drug

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8.1	approved to prev	vent pregnancy who	en administered after s	exual contact. For pu	urposes of this
8.2	paragraph, "hea	lth plan" has the m	eaning provided in sec	ction 62Q.01, subdiv	vision 3.

- 8.3 **EFFECTIVE DATE.** This section applies to medical assistance and MinnesotaCare
- 8.4 <u>coverage effective January 1, 2022.</u>