AMENDMENT TO RULES COMMITTEE PRINT 118–36 OFFERED BY Ms. SHERRILL OF NEW JERSEY

At the end of subtitle C of title VII, insert the following:

1	SEC. 7 I ILOI I ROGRAM ON REMOTE BLOOD I RESSURE
2	MONITORING FOR CERTAIN PREGNANT AND
3	POST-PARTUM TRICARE BENEFICIARIES.
4	(a) Establishment .—
5	(1) In General.—Not later than 180 days
6	after the date of enactment of this act, the Secretary
7	of Defense, acting through the Defense Health
8	Agency, shall establish a pilot program on blood
9	pressure monitoring for at-risk pregnant and
10	postpartum TRICARE beneficiaries in order to in-
11	crease the rate of early detection of hypertensive dis-
12	order related to pregnancy and postpartum.
13	(2) Model.—The Secretary may model the
14	pilot program on a pilot program for blood pressure
15	self-monitoring of the Healthy Start Program oper-
16	ated by the Health Resources and Services Adminis-
17	tration of the Department of Health and Human
18	Services.

1	(b) Sites.—The Secretary shall select sites for the
2	pilot program in accordance with the following:
3	(1) The pilot program shall operate at not
4	fewer than two military medical treatment facilities
5	of each of the Army, Navy, Marine Corps, Air
6	Force, and Space Force.
7	(2) Sites shall be geographically diverse, includ-
8	ing locations in rural and urban areas.
9	(3) The Secretary shall give priority to a mili-
10	tary medical treatment facility that has a large num-
11	ber of obstetric patients or a history of maternal
12	health programs.
13	(c) Participants.—
14	(1) Eligibility.—An eligible participant for
15	the pilot program, is an individual—
16	(A) who is enrolled in TRICARE;
17	(B) who is pregnant or postpartum;
18	(C) who receives health care through a
19	military medical treatment facility selected
20	under subsection (b); and
21	(D) whom the Secretary determines is at
22	risk (based on evidence and current medical
23	standards and recommendations) of a hyper-
24	tensive disorder of pregnancy or negative health

1	outcomes as a result of a hypertensive disorder
2	of pregnancy.
3	(2) Voluntary.—Participation in the pilot
4	program shall be voluntary.
5	(d) EQUIPMENT.—A participant in the pilot program
6	shall receive—
7	(1) a device approved by the Food and Drug
8	Administration for the digital monitoring of blood
9	pressure, capable of remote monitoring and data
10	transmission; and
11	(2) educational materials and instructions on
12	the use of such device from a health care provider
13	of the Department of Defense.
14	(e) Providers.—In carrying out the pilot program,
15	the Secretary shall use primary care and obstetric care
16	providers of eligible participants, to the extent practicable.
17	(f) Materials.—The Secretary shall develop sup-
18	porting materials for health care providers who facilitate
19	the pilot program, including the following:
20	(1) Guidance on how to identify eligible partici-
21	pants for the pilot program;
22	(2) Evidence-based—
23	(A) educational materials regarding mater-
24	nal health best practices for eligible partici-
25	pants; and

1	(B) guidance for sharing information be-
2	tween members of a participant's care team.
3	(g) TERM.—The pilot program shall terminate five
4	years after the date on which the Secretary establishes
5	such pilot program.
6	(h) Report.—Not later than 180 days after the ter-
7	mination of the pilot program, the Secretary shall submit
8	to the Committees on Armed Services of the House of
9	Representatives and Senate a report on the pilot program,
10	and publish such report on the website of the Department
11	of Defense. The report shall include the following ele-
12	ments, disaggregated by the Armed Force, sex, age, race,
13	and ethnicity of participants:
14	(1) The number of participants in the pilot pro-
15	gram.
16	(2) The percentage of such participants who
17	used the monitors as prescribed.
18	(3) A summary of how participants under-used
19	the monitors.
20	(4) The percentage of participants who had
21	blood pressure readings of concern.
22	(5) The percentage of participants described in
23	paragraph (4) who received medical attention based
24	on such readings.

1	(6) A summary of provider and participant
2	feedback, including percentages of—
3	(A) providers that found the program in-
4	fluenced patient care; and
5	(B) participants who found the program
6	was helpful in managing their own care.
7	(7) Recommendations of the Secretary whether
8	the pilot program should be altered, expanded, or
9	made permanent.

