Hall of the House of Representatives

94th General Assembly - Regular Session, 2023

Amendment Form

Subtitle of House Bill No. 1121

CONCERNING COVERAGE FOR BIOMARKER TESTING FOR EARLY DETECTION AND MANAGEMENT FOR CANCER DIAGNOSES.

Amendment No. 1 to House Bill 1121

Amend House Bill No. 1121 as originally introduced:

Add Representatives K. Brown, Dalby, Evans, K. Ferguson, L. Johnson, Nicks, Pilkington, J. Richardson, Warren as cosponsors of the bill

AND

Add Senators D. Wallace, J. Boyd, Irvin, M. Johnson, R. Murdock as cosponsors of the bill

AND

Page 1, delete line 32, and substitute the following:
"intervention, including known gene-drug interactions for medications being considered for use or already being administered."

AND

Page 1, line 36, delete "or fluid biological specimen" and substitute "or other biospecimen"

AND

Page 2, delete lines 2 through 4, and substitute the following: "single-analyte tests, multiplex panel tests, protein expression, and whole exome, whole genome, and whole transcriptome sequencing;

(3) "Consensus statement" means a statement that:

(A) Is developed by an independent, multidisciplinary panel of experts that uses a transparent methodology and reporting structure that includes a conflict of interest policy;

(B) Is based on the best available evidence for the purpose of optimizing clinical care outcomes; and



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(4)(A) "Health benefit plan" means an individual, blanket, or"
AND
Page 2, delete lines 9 and 10, and substitute the following:
                  "(B) "Health benefit plan" includes:
                       (i) Indemnity and managed care plans; and
                        (ii) The Arkansas Medicaid Program."
AND
Page 2, line 26, delete "plan;" and substitute "plan; or"
AND
Page 2, line 28, delete "or"
AND
Page 2, delete lines 29 and 30
AND
Page 2, line 31, delete "(4)(A)" and substitute "(5)(A)"
AND
Page 3, line 5, delete "and"
AND
Page 3, line 6, delete "(5)" and substitute "(6)"
AND
Page 3, delete line 8, and substitute the following:
"health care in the ordinary course of the practice of his or her profession;
           (7) "Nationally recognized clinical practice guidelines" means
evidence-based clinical practice guidelines that:
                  (A) Are developed by independent organizations or medical
professional societies using a:
                       (i) Transparent methodology and reporting structure;
and
                        (ii) Conflict of interest policy; and
                  (B) Establish standards of care that are informed by:
                        (i) A systemic review of evidence; and
                        (ii) An assessment of the benefits and costs of
alternative care options that includes recommendations intended to optimize
patient care;
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(C) Is aimed at specific clinical circumstances;

- (8)(A) "Subscriber" means an individual eligible to receive coverage of healthcare services by a healthcare professional under a health benefit plan.
- (B) "Subscriber" includes a subscriber's legally authorized representative;
- (8) "Urgent healthcare service" means a healthcare service for a non-life-threatening condition that, in the opinion of a physician with knowledge of a subscriber's medical condition, requires prompt medical care in order to prevent:
 - (A) A serious threat to life, limb, or eyesight;
- (B) Worsening impairment of a bodily function that threatens the body's ability to regain maximum function;
- (C) Worsening dysfunction or damage of any bodily organ or part that threatens the body's ability to recover from the dysfunction or damage; or
- (D) Severe pain that cannot be managed without prompt medical care; and
- (10)(A) "Utilization review entity" means an individual or entity that performs prior authorization for at least one (1) of the following:
 - (i) A healthcare insurer;
- (ii) A preferred provider organization or health maintenance organization; or
- (iii) Any other individual or entity that provides, offers to provide, or administers hospital, outpatient, medical, or other health benefits to a person treated by a healthcare provider in this state under a policy, health benefit plan, or contract.
- (B) A healthcare insurer is a utilization review entity if the healthcare insurer performs prior authorization.
- (C) "Utilization review entity" does not include an insurer of automobile, homeowners, or casualty and commercial liability insurance or the insurer's employees, agents, or contractors."

AND

- Page 3, delete lines 13 through 36, and substitute the following: "state shall provide coverage for biomarker testing.
- (b) The evidence of coverage document provided with a health benefit plan under this subchapter shall include biomarker testing for the purpose of diagnosis, treatment, appropriate management, or ongoing monitoring of a subscriber's disease or condition to guide treatment decisions when the biomarker test is supported by medical and scientific evidence, including without limitation:
- (1) Labeled indications for tests that are approved or cleared by the United States Food and Drug Administration;
- (2) Indicated tests for a drug that is approved by the United States Food and Drug Administration;
- (3) Warnings and precautions on United States Food and Drug Administration-approved drug labels;

- (4) Centers for Medicare & Medicaid Services national coverage determinations or Medicare administrative contractor local coverage determinations; or
- (5) Nationally recognized clinical practice guidelines and consensus statements.
- (c) A health benefit plan shall ensure that coverage is provided in a manner that limits disruptions in care, including the need for multiple biopsies and biospecimen samples as determined by a healthcare professional.
- (d)(1) A subscriber and a subscriber's healthcare professional shall have access to a clear, readily available, and convenient process to request an exception to a health benefit plan under this subchapter.
- (2) The process under subdivision (d)(1) of this section shall be readily accessible on the health benefit plan's website.
- (3) This section shall not be construed to require a separate process if the health benefit plan's existing process complies with subdivision (d)(1) of this section.
- (e) A utilization review entity shall make a determination on a request for coverage of biomarker testing at the same scope, duration, and frequency as the health benefit plan otherwise provides to subscribers.
- (f) If prior authorization is required for biomarker testing, the utilization review entity shall approve or deny a prior authorization request and notify the subscriber, the subscriber's healthcare professional, and any entity requesting prior authorization of the healthcare service:
- (1) Within seventy-two (72) hours for request for nonurgent healthcare services; or
- (2) Within twenty-four (24) hours for requests for urgent healthcare services."

AND

Page 4, delete lines 1 through 9

The Amendment was read	
By: Representative F. Allen	
ANS/ANS - 02-14-2023 14:36:59	
ANS186	Chief Clerk