I_135_0359-2

135th General Assembly Regular Session 2023-2024

Sub. H. B. No. 24

A BILL

То	enact sections 3902.64 and 5164.13 of the	1
	Revised Code to require health benefit plan and	2
	Medicaid program coverage of biomarker testing.	3

BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE OF OHIO:

Section 1. That sections 3902.64 and 5164.13 of the	4
Revised Code be enacted to read as follows:	5
Sec. 3902.64. (A) As used in this section, "biomarker,"	6
"biomarker testing," and "nationally recognized clinical	7
practice guidelines" have the same meanings as in section	8
5164.13 of the Revised Code.	9
(B) Notwithstanding section 3901.71 of the Revised Code, a	10
health benefit plan issued, renewed, or modified in this state	11
on or after the effective date of this section shall cover	12
biomarker testing for any of the following purposes:	
(1) Diagnosis;	14
(2) Treatment and appropriate management of a disease or	15
condition;	16



(3) Ongoing monitoring of a disease or condition.	17
(C) A health benefit plan shall cover biomarker testing	18
ordered and deemed medically necessary by the qualified treating	19
health care provider working within the provider's scope of	20
practice for the purposes included in division (B) of this	21
section when the test is supported by medical or scientific	22
evidence, as defined by section 3922.01 of the Revised Code,	23
including at least one of the following:	24
(1) Labeled indications for a United States food and drug	25
administration approved or cleared test;	26
(2) Indicated tests for a drug approved by the United	27
States food and drug administration;	28
(3) Warnings and precautions for United States food and	29
drug administration approved drug labels;	30
(4) National coverage determinations made by the United	31
States centers for medicare and medicaid services;	32
(5) Medicare administrative contractor local coverage	33
<pre>determinations;</pre>	34
(6) Nationally recognized clinical practice guidelines;	35
(7) Nationally recognized and peer reviewed studies	36
indicating that the test materially improves health outcomes.	37
(D) A health plan issuer shall ensure coverage as required	38
in division (B) of this section in a manner that limits	39
disruptions in care, including the need for multiple biopsies or	40
biospecimen samples.	41
(E) Any appeal of a biomarker testing coverage	42
determination shall be handled in accordance with the health	43

plan issuer's appeal policy and any other relevant provision of	4 4
law, including section 1751.82 or Chapter 3922. of the Revised	45
Code. The appeal process shall be made readily accessible to all	46
participating providers and recipients in writing and online.	47
(F) Nothing in this section shall be construed to require	48
coverage of biomarker testing for screening purposes.	49
Sec. 5164.13. (A) As used in this section:	50
(1) "Biomarker" means a characteristic that is objectively	51
measured and evaluated as an indicator of normal biological	52
processes, pathogenic processes, or pharmacologic responses to	53
specific therapeutic intervention, including known gene-drug	54
interactions for drugs being considered for use or already	55
available for use. Biomarkers include, but are not limited to,	56
gene mutations, characteristics of genes, or protein expression.	57
(2) "Biomarker testing" means the analysis of tissue,	58
blood, or another biospecimen for the presence of a biomarker,	59
and includes, but is not limited to, single-analyte tests,	60
multiplex panel tests, protein expression, and whole exome,	61
whole genome, and whole transcriptome sequencing.	62
(3) "Nationally recognized clinical practice guidelines"	63
are evidence-based clinical practice quidelines establishing	64
standards of care informed by a systematic review and assessment	65
of benefits and risks of alternative care options and include	66
recommendations intended to optimize patient care, developed by	67
independent organizations or medical professional societies	68
utilizing a transparent methodology and reporting structure and	69
with a conflict of interest policy.	70
(B) The medicaid program shall cover biomarker testing,	71
subject to division (C) of this section, for any of the	72

<pre>following purposes:</pre>	
(1) Diagnosis;	74
(2) Treatment and appropriate management of a disease or	75
<pre>condition;</pre>	
(3) Ongoing monitoring of a disease or condition.	77
(C) The medicaid program shall cover biomarker testing	78
ordered and deemed medically necessary by the qualified treating	79
health care provider working within the provider's scope of	
practice for the purposes included in division (B) of this	81
section when the test is supported by medical or scientific	82
evidence, as defined by section 3922.01 of the Revised Code,	83
including at least one of the following:	84
(1) Labeled indications for a United States food and drug	85
administration approved or cleared test;	86
(2) Indicated tests for a drug approved by the United	87
States food and drug administration;	88
(3) Warnings and precautions for United States food and	89
drug administration approved drug labels;	90
(4) National coverage determinations made by the United	91
States centers for medicare and medicaid services;	92
(5) Medicare administrative contractor local coverage	93
<pre>determinations;</pre>	94
(6) Nationally recognized clinical practice guidelines;	95
(7) Nationally recognized and peer reviewed studies	96
indicating that the test materially improves health outcomes.	97
(D) The Medicaid program shall ensure coverage as required	98
in division (B) of this section in a manner that limits	99