HOUSE BILL No. 1439

DIGEST OF INTRODUCED BILL

Citations Affected: IC 12-7-2; IC 12-15; IC 16-38-6-2.

Synopsis: FSSA matters. Allows a Medicaid recipient who is incarcerated to have the recipient's Medicaid suspended for up to two years instead of one year before terminating the recipient's Medicaid eligibility. Defines "comprehensive risk contract" and "managed care organization" for purposes of Medicaid. Specifies that if a provision of Indiana insurance law is inconsistent with a law applying to a managed care organization with respect to the managed care organization's Medicaid responsibilities, the law applying to the managed care organization with respect to the Medicaid responsibilities is controlling. Changes language in the Medicaid law to reflect the existence of more than one risk based managed care program. Removes obsolete references to "primary care case management". Removes references to "insurer", "insurance", and "health maintenance organization" in the law concerning the healthy Indiana plan to reflect the sole use of managed care organizations to provide coverage under the plan. Makes conforming amendments. Makes a technical correction to a federal Code citation.

Effective: July 1, 2017.

Kirchhofer

January 17, 2017, read first time and referred to Committee on Public Health.



First Regular Session of the 120th General Assembly (2017)

PRINTING CODE. Amendments: Whenever an existing statute (or a section of the Indiana Constitution) is being amended, the text of the existing provision will appear in this style type, additions will appear in this style type, and deletions will appear in this style type.

Additions: Whenever a new statutory provision is being enacted (or a new constitutional provision adopted), the text of the new provision will appear in **this style type**. Also, the word **NEW** will appear in that style type in the introductory clause of each SECTION that adds a new provision to the Indiana Code or the Indiana Constitution.

Conflict reconciliation: Text in a statute in *this style type* or *this style type* reconciles conflicts between statutes enacted by the 2016 Regular Session of the General Assembly.

HOUSE BILL No. 1439

A BILL FOR AN ACT to amend the Indiana Code concerning human services.

Be it enacted by the General Assembly of the State of Indiana:

1	SECTION 1. IC 12-7-2-40.4 IS ADDED TO THE INDIANA CODE
2	AS A NEW SECTION TO READ AS FOLLOWS [EFFECTIVE JULY
3	1,2017]: Sec. 40.4. "Comprehensive risk contract" has the meaning
4	set forth in 42 CFR 438.2.
5	SECTION 2. IC 12-7-2-126.9 IS ADDED TO THE INDIANA
6	CODE AS A NEW SECTION TO READ AS FOLLOWS
7	[EFFECTIVE JULY 1, 2017]: Sec. 126.9. "Managed care
8	organization" means a person that has a comprehensive risk
9	contract with the office of Medicaid policy and planning under
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0	IC 12-15.
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0	IC 12-15.
0	IC 12-15. SECTION 3. IC 12-15-1-20.4, AS AMENDED BY P.L.185-2015,
0 1 2	IC 12-15. SECTION 3. IC 12-15-1-20.4, AS AMENDED BY P.L.185-2015, SECTION 9, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
0 1 2 3	IC 12-15. SECTION 3. IC 12-15-1-20.4, AS AMENDED BY P.L.185-2015, SECTION 9, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 20.4. (a) If a Medicaid recipient is:
0 1 2 3 4	IC 12-15. SECTION 3. IC 12-15-1-20.4, AS AMENDED BY P.L.185-2015, SECTION 9, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 20.4. (a) If a Medicaid recipient is: (1) adjudicated to be a delinquent child and placed in:
0 1 2 3 4 5	IC 12-15. SECTION 3. IC 12-15-1-20.4, AS AMENDED BY P.L.185-2015, SECTION 9, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 20.4. (a) If a Medicaid recipient is: (1) adjudicated to be a delinquent child and placed in: (A) a community based correctional facility for children;



caring institution under IC 31-27; or (2) incarcerated in a prison or jail; and ineligible to participate in the Medicaid program during the placement described in subdivision (1) or (2) because of federal Medicaid law, the division of family resources, upon notice that a child has been adjudicated to be a delinquent child and placed in a facility described in subdivision (1) or upon notice that a person is incarcerated in a prison or jail and placed in a facility described in subdivision (2), shall suspend the person's participation in the Medicaid program for up to one (1) year two (2) years before terminating the person's eligibility. (b) If the division of family resources receives: (1) a dispositional decree under IC 31-37-19-28; or (2) a modified disposition order under IC 31-37-22-9; and the department of correction gives the division at least forty (40) days notice that a person will be released from a facility described in subsection (a)(1)(C) or (a)(2), the division of family resources shall take action necessary to ensure that a person described in subsection (a) is eligible to participate in the Medicaid program upon the person's release, if the person is eligible to participate. SECTION 4. IC 12-15-2-14, AS AMENDED BY THE TECHNICAL CORRECTIONS BILL OF THE 2017 GENERAL ASSEMBLY, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 14. (a) An individual: (1) who is less than nineteen (19) years of age; (2) who is not described in 42 U.S.C. 1396a(a)(10)(A)(f); and (3) whose family income does not exceed the income level established in subsection (b); is eligible to receive Medicaid. (b) An individual described in this section is eligible to receive Medicaid, subject to 42 U.S.C. 1396a et seq., if the individual's family income does not exceed one hundred fifty percent (150%) of the federal income poverty level for the same size family. (c) The office may apply a resource standard in determining the eligibility of an individual described in this section. This subsection expires Decembe		
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25 (2) who is not described in 42 U.S.C. 1396a(a)(10)(A)(I); 42 26 U.S.C. 1396a(a)(10)(A)(i)(I); and 27 (3) whose family income does not exceed the income level 28 established in subsection (b); 29 is eligible to receive Medicaid. 30 (b) An individual described in this section is eligible to receive 31 Medicaid, subject to 42 U.S.C. 1396a et seq., if the individual's family 32 income does not exceed one hundred fifty percent (150%) of the 33 federal income poverty level for the same size family. 34 (c) The office may apply a resource standard in determining the 35 eligibility of an individual described in this section. This subsection 36 expires December 31, 2013. 37 SECTION 5. IC 12-15-5-5, AS AMENDED BY P.L.101-2005, 38 SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE 39 JULY 1, 2017]: Sec. 5. (a) The office may provide a prescription drug 40 benefit to a Medicaid recipient in the a Medicaid risk based managed 41 care program.	24	
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41 care program.		- ''
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	42	(b) If the office provides a prescription drug benefit to a Medicaid



1	recipient in the a Medicaid risk based managed care program:
2	(1) the office shall develop a procedure and provide the recipient's
3	risk based managed care provider with information concerning
4	the recipient's prescription drug utilization for the risk based
5	managed care provider's case management program; and
6	(2) the provisions of IC 12-15-35.5 apply.
7	(c) If the office does not provide a prescription drug benefit to a
8	Medicaid recipient in the a Medicaid risk based managed care
9	program, a Medicaid managed care organization shall provide coverage
10	and reimbursement for outpatient single source legend drugs subject to
11	IC 12-15-35-46, IC 12-15-35-47, and IC 12-15-35.5.
12	SECTION 6. IC 12-15-5-13, AS AMENDED BY P.L.8-2016
13	SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
14	JULY 1, 2017]: Sec. 13. (a) The office shall provide coverage for
15	treatment of opioid or alcohol dependence that includes the following:
16	(1) Counseling services that address the psychological and
17	behavioral aspects of addiction.
18	(2) When medically indicated, drug treatment involving agents
19	approved by the federal Food and Drug Administration for the:
20	(A) treatment of opioid or alcohol dependence; or
21	(B) prevention of relapse to opioids or alcohol after
22	detoxification.
23	(3) Inpatient detoxification:
24	(A) in accordance with:
25	(i) the most current edition of the American Society of
26	Addiction Medicine Patient Placement Criteria; or
27	(ii) other clinical criteria that are determined by the office
28	and are evidence based and peer reviewed; and
29	(B) when determined by the treatment plan to be medically
30	necessary.
31	(b) The office shall:
32	(1) develop quality measures to ensure; and
33	(2) require a Medicaid managed care organization to report;
34	compliance with the coverage required under subsection (a).
35	(c) The office may implement quality capitation withholding of
36	reimbursement to ensure that a Medicaid managed care organization
37	has provided the coverage required under subsection (a).
38	(d) The office shall report the clinical use of the medications
39	covered under this section to the mental health Medicaid quality
40	advisory committee established by IC 12-15-35-51. The mental health
41	Medicaid quality advisory committee may make recommendations to
42	the office concerning this section.
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SECTION 7. IC 12-15-11.5-0.5 IS AMENDED TO READ AS
FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 0.5. This chapter does
not apply to a managed care contractor organization that, on or before
July 1, 2000, did not directly contract with a hospital (as defined in
section 1 of this chapter) for the provision of services under the office's
managed care program.
SECTION 8. IC 12-15-11.5-2 IS AMENDED TO READ AS
FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 2. The office's managed
care contractor organization shall regard a hospital as a contracted
provider in the office's managed care services program, which provides
a capitated prepayment managed care system, for the provision of
medical services to each individual who:
(1) is eligible to receive services under IC 12-15 and has enrolled

- in the office's managed care services program;
- (2) resides in the same city in which the hospital is located; and
- (3) has selected a primary care provider who:
 - (A) is a contracted provider with the office's managed care contractor; organization; and
 - (B) has medical staff privileges at the hospital.

SECTION 9. IC 12-15-11.5-6 IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 6. A claim for reimbursement for services shall be treated as a disputed claim under this chapter if:

- (1) it is submitted within one hundred twenty (120) days after the date that services are rendered;
- (2) it is denied by the managed care contractor; organization;
- (3) the hospital submits a written notice of dispute for the claim to the managed care contractor organization not more than sixty (60) days after the receipt of the denial notice;
- (4) it is appealed in accordance with the managed care
- contractor's organization's internal appeals process; and
- (5) payment for the claim is denied by the managed care contractor organization following its internal appeals process.

SECTION 10. IC 12-15-11.5-7 IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 7. The office's managed care contractor organization must conclude an appeal under section 6(4) of this chapter and notify the hospital of its decision not more than thirty-five (35) days after the managed care contractor organization receives a notice from the hospital disputing the managed care contractor's organization's denial of a claim.

SECTION 11. IC 12-15-11.5-8 IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 8. (a) A contract



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entered into by a hospital with the office's managed care contractor organization for the provision of services under the office's managed care services program must include a dispute resolution procedure for all disputed claims. Unless agreed to in writing by the hospital and the office's managed care contractor; organization, the dispute resolution procedure must include the following requirements: (1) That submission of disputed claims must be made to an independent arbitrator selected under subsection (b). (2) Each claim must set forth with specificity the issues to be arbitrated, the amount involved, and the relief sought. (3) That the hospital and the office's managed care contractor organization shall attempt in good faith to resolve all disputed claims. (4) The hospital shall submit to the arbitrator any claims that remain in dispute sixty (60) calendar days after the hospital receives written notice as provided under section 7 of this chapter. (5) That resolution of disputes by the arbitrator must occur not later than ninety (90) calendar days after submission of disputed claims to the arbitrator, unless the parties mutually agree otherwise. (6) That determinations of the arbitrator are final and binding and not subject to any appeal or review procedure. (7) That the arbitrator does not have the authority to award any punitive or exemplary damages or to vary or ignore the terms of any contract between the parties and shall be bound by controlling law. (8) That judgment upon the award rendered by the arbitrator may be entered and enforced in and is subject to the jurisdiction of a court with jurisdiction in Indiana. (9) That the cost of the arbitrator must be shared equally by the parties, and each party must bear its own attorney and witness fees. (b) The parties to a contract described in subsection (a) shall mutually agree on an independent arbitrator, or, if the parties are unable to reach agreement on an independent arbitrator, the following procedure must be followed: (1) Each party sha		
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33 (b) The parties to a contract described in subsection (a) shall mutually agree on an independent arbitrator, or, if the parties are unable to reach agreement on an independent arbitrator, the following procedure must be followed: 37 (1) Each party shall select an independent representative, and the independent representatives shall select a panel of three (3) independent arbitrators who have experience in institutional and professional health care delivery practices and procedures and		
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37 (1) Each party shall select an independent representative, and the 38 independent representatives shall select a panel of three (3) 39 independent arbitrators who have experience in institutional and 40 professional health care delivery practices and procedures and		
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independent arbitrators who have experience in institutional and professional health care delivery practices and procedures and		
40 professional health care delivery practices and procedures and		



arbitrator.

1	(2) The parties will each strike one (1) arbitrator from the panel
2	selected under subdivision (1), and the remaining arbitrator serves
3	as the arbitrator of the disputed claims under subsection (a).
4	(3) The procedures for selecting an arbitrator under this section
5	must be completed not later than twenty (20) calendar days after
6	the hospital provides written notice of at least one (1) disputed
7	claim.
8	SECTION 12. IC 12-15-11.5-9 IS AMENDED TO READ AS
9	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 9. The arbitration
10	process described in section 8 of this chapter shall also be followed for
11	resolution of disputed claims between a hospital and the office's
12	managed care contractor, organization, if the hospital is not a
13	contracted provider in the office's managed health care services
14	program.
15	SECTION 13. IC 12-15-11.5-10, AS ADDED BY P.L.220-2011,
16	SECTION 265, IS AMENDED TO READ AS FOLLOWS
17	[EFFECTIVE JULY 1, 2017]: Sec. 10. A hospital and the managed
18	care contractor organization of the office shall use the arbitration
19	procedure in section 8 of this chapter for the resolution of all disputed
20	claims that have accrued as of March 17, 2000.
21	SECTION 14. IC 12-15-12-0.9 IS ADDED TO THE INDIANA
22	CODE AS A NEW SECTION READ AS FOLLOWS [EFFECTIVE
23	JULY 1, 2017]: Sec. 0.9. (a) This section applies only with respect
24	to the responsibilities of a managed care organization under:
25	(1) this article;
26	(2) IC 12-17.6;
27	(3) 42 CFR 438; or
28	(4) a rule adopted under a law described in subdivision (1) or
29	(2).
30	(b) If a provision of, or rule adopted under, IC 27 is inconsistent
31	with a law described in subsection (a), the law described in
32	subsection (a) is controlling.
33	SECTION 15. IC 12-15-12-15 IS AMENDED TO READ AS
34	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 15. The office, for
35	purposes of the primary care case management program, and A
36	managed care eontractor, for purposes of the risk-based managed care
37	program, organization shall:
38	(1) cover and pay for all medically necessary screening services
39	provided to an individual who presents to an emergency
40	department with an emergency medical condition; and
41	(2) beginning July 1, 2001, not neither deny or nor fail to process
42	a claim for reimbursement for emergency services on the basis



1	that the enrollee's primary care provider's authorization code for
2	the services was not obtained before or after the services were
3	rendered.
4	SECTION 16. IC 12-15-12-17 IS AMENDED TO READ AS
5	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 17. (a) This section
6	applies to post-stabilization care services provided to an individual
7	enrolled in
8	(1) the a Medicaid risk-based risk based managed care program.
9	Of "
10	(2) the Medicaid primary care case management program.
11	(b) The office, if the individual is enrolled in the primary care case
12	management program, or the managed care organization if the through
13	which an individual is enrolled in the a risk-based risk based managed
14	care program, is financially responsible for the following services
15	provided to an the enrollee:
16	(1) Post-stabilization care services that are pre-approved
17	preapproved by a representative of the office or the managed
18	care organization. as applicable.
19	(2) Post-stabilization care services that are not pre-approved
20	preapproved by a representative of the office or the managed
21	care organization, as applicable, but that are administered to
22	maintain the enrollee's stabilized condition within one (1) hour of
23	a request to the office or the managed care organization for
24	pre-approval preapproval of further post-stabilization care
25	services.
26	(3) Post-stabilization care services provided after an enrollee is
27	stabilized that are not pre-approved preapproved by a
28	representative of the office or the managed care organization, as
29	applicable, but that are administered to maintain, improve, or
30	resolve the enrollee's stabilized condition if the office or the
31	managed care organization:
32	(A) does not respond to a request for preapproval within one
33	(1) hour;
34	(B) cannot be contacted; or
35	(C) cannot reach an agreement with the enrollee's treating
36	physician concerning the enrollee's care, and a physician
37	representing the office or the managed care organization as
38	applicable, is not available for consultation.
39	(c) If the conditions described in subsection (b)(3)(C) exist, the
40	office or the managed care organization as applicable, shall give the
41	enrollee's treating physician an opportunity to consult with a physician
42	representing the office or the managed care organization. The enrollee's



1	treating physician may continue with care of the enrollee until a
2	physician representing the office or the managed care organization as
3	applicable, is reached or until one (1) of the following criteria is met:
4	(1) A physician:
5	(A) representing the office or the managed care organization;
6	as applicable; and
7	(B) who has privileges at the treating hospital;
8	assumes responsibility for the enrollee's care.
9	(2) A physician representing the office or the managed care
10	organization as applicable, assumes responsibility for the
11	enrollee's care through transfer.
12	(3) A representative of the office or the managed care
13	organization as applicable, and the treating physician reach an
14	agreement concerning the enrollee's care.
15	(4) The enrollee is discharged from the treating hospital.
16	(d) This subsection applies to post-stabilization care services
17	provided under subsection (b)(1), (b)(2), and (b)(3) to an individual
18	enrolled in the a Medicaid risk-based risk based managed care
19	program by a provider who has not contracted with a Medicaid
20	risk-based the individual's managed care organization to provide
21	post-stabilization care services under subsection (b)(1), (b)(2), and
22	(b)(3) to the individual. Payment for post-stabilization care services
23	provided under subsection (b)(1), (b)(2), and (b)(3) must be in an
24	amount equal to one hundred percent (100%) of the current Medicaid
25	fee for service reimbursement rates for such services.
26	(e) This section does not prohibit a managed care organization from
27	entering into a subcontract with another Medicaid risk-based managed
28	care organization providing for the latter managed care organization
29	to assume financial responsibility for making the payments required
30	under this section.
31	(f) This section does not limit the ability of the office or the
32	managed care organization to:
33	(1) review; and
34	(2) make a determination of;
35	the medical necessity of the post-stabilization care services provided
36	to an enrollee for purposes of determining coverage for such services.
37	SECTION 17. IC 12-15-12-18 IS AMENDED TO READ AS
38	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 18. (a) Except as
39	provided in subsection (b), this section applies to:
40	(1) emergency services provided to an individual enrolled in the
41	a Medicaid risk-based risk based managed care program; and
42	(2) medically necessary screening services provided to an



1	individual enrolled in the a Medicaid risk-based risk based
2	managed care program;
3	who presents to an emergency department with an emergency medical
4	condition.
5	(b) This section does not apply to emergency services or screening
6	services provided to an individual enrolled in the a Medicaid
7	risk-based risk based managed care program by a provider who has
8	contracted with a Medicaid risk-based the individual's managed care
9	organization to provide emergency services to the individual.
10	(c) Payment for emergency services and medically necessary
11	screening services in the emergency department of a hospital licensed
12	under IC 16-21 must be in an amount equal to one hundred percent
13	(100%) of the current Medicaid fee for service reimbursement rates for
14	such services.
15	(d) Payment under subsection (c) is the responsibility of the
16	enrollee's risk-based managed care organization. This subsection does
17	not prohibit the risk-based managed care organization from entering
18	into a subcontract with another Medicaid risk-based managed care
19	organization providing for the latter managed care organization to
20	assume financial responsibility for making the payments required under
21	this section.
22	(e) This section does not limit the ability of the managed care
23	organization to:
24	(1) review; and
25	(2) make a determination of;
26	the medical necessity of the services provided in a hospital's emergency
27	department for purposes of determining coverage for such services.
28	SECTION 18. IC 12-15-12-19 IS REPEALED [EFFECTIVE JULY
29	1, 2017]. Sec. 19. (a) This section applies to an individual who is a
30	Medicaid recipient.
31	(b) Subject to subsection (c), the office shall develop the following
32	programs regarding individuals described in subsection (a):
33	(1) A disease management program for recipients with any of the
34	following chronic diseases:
35	(A) Asthma.
36	(B) Diabetes.
37	(C) Congestive heart failure or coronary heart disease.
38	(D) Hypertension.
39	(E) Kidney disease.
40	(2) A case management program for recipients described in
41	subsection (a) who are at high risk of chronic disease, that is
42	based on a combination of cost measures, elinical measures, and



1	health outcomes identified and developed by the office with input
2	and guidance from the state department of health and other
3	experts in health care case management or disease management
4	programs.
5	(c) The office shall implement:
6	(1) a pilot program for at least two (2) of the diseases listed in
7	subsection (b) not later than July 1, 2003; and
8	(2) a statewide chronic disease program as soon as practicable
9	after the office has done the following:
10	(A) Evaluated a pilot program described in subdivision (1).
11	(B) Made any necessary changes in the program based on the
12	evaluation performed under clause (A).
13	(d) The office shall develop and implement a program required
14	under this section in cooperation with the state department of health
15	and shall use the following persons to the extent possible:
16	(1) Community health centers.
17	(2) Federally qualified health centers (as defined in 42 U.S.C.
18	1396d(1)(2)(B)).
19	(3) Rural health clinics (as defined in 42 U.S.C. 1396d(1)(1)).
20	(4) Local health departments.
21	(5) Hospitals.
22	(6) Public and private third party payers.
23	(e) The office may contract with an outside vendor or vendors to
24	assist in the development and implementation of the programs required
25	under this section.
26	(f) The office and the state department of health shall provide the
27	interim study committee on public health, behavioral health, and
28	human services established by IC 2-5-1.3-4 in an electronic format
29	under IC 5-14-6 with an evaluation and recommendations on the costs,
30	benefits, and health outcomes of the pilot programs required under this
31	section. The evaluations required under this subsection must be
32	provided not more than twelve (12) months after the implementation
33	date of the pilot programs.
34	(g) The office and the state department of health shall report to the
35	interim study committee on public health, behavioral health, and
36	human services established by IC 2-5-1.3-4 in an electronic format
37	under IC 5-14-6 not later than November 1 of each year regarding the
38	programs developed under this section.
39	(h) The disease management program services for a recipient
40	diagnosed with diabetes or hypertension must include education for the
41	recipient on kidney disease and the benefits of having evaluations and

treatment for chronic kidney disease according to accepted practice



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1	guidelines.
2	SECTION 19. IC 12-15-12-20, AS ADDED BY P.L.135-2005,
3	SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
4	JULY 1, 2017]: Sec. 20. The office shall develop the following:
5	(1) A measure to evaluate the performance of a Medicaid
6	managed care organization in screening a child who is less than
7	six (6) years of age for lead poisoning.
8	(2) A system to maintain the results of an evaluation under
9	subdivision (1) in written form.
10	(3) A performance incentive program for Medicaid managed care
11	organizations evaluated under subdivision (1).
12	SECTION 20. IC 12-15-12-21, AS ADDED BY P.L.113-2008,
13	SECTION 6, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
14	JULY 1, 2017]: Sec. 21. (a) Not later than January 1, 2011, the
15	following must be accredited by the National Committee for Quality
16	Assurance or its successor:
17	(1) A managed care organization that has contracted with the
18	office before July 1, 2008, to provide Medicaid services under the
19	a risk based managed care program.
20	(2) A behavioral health managed care organization that has
21	contracted before July 1, 2008, with a managed care organization
22	described in subdivision (1).
23	(b) A:
24	(1) managed care organization that has contracted with the office
25	after June 30, 2008, to provide Medicaid services under the a risk
26	based managed care program; or
27	(2) behavioral health managed care organization that has
28	contracted after June 30, 2008, with a managed care organization
29	described in subdivision (1);
30	must begin the accreditation process and obtain accreditation by the
31	National Committee for Quality Assurance or its successor at the
32	earliest time that the National Committee for Quality Assurance allows
33	a managed care organization to be accredited.
34	SECTION 21. IC 12-15-12-22, AS ADDED BY P.L.113-2008,
35	SECTION 7, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
36	JULY 1, 2017]: Sec. 22. A:
37	(1) managed care organization that has a contract with the office
38	to provide Medicaid services under the a risk based managed care
39	program; or
40	(2) behavioral health managed care organization that has
41	contracted with a managed care organization described in



subdivision (1);

shall accept, receive, and process claims for payment that are filed electronically by a Medicaid provider.

SECTION 22. IC 12-15-13-6, AS AMENDED BY P.L.153-2011, SECTION 17, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 6. (a) Except as provided by IC 12-15-35-50, a notice or bulletin that is issued by:

(1) the office;

- (2) a contractor of the office; or
- (3) a managed care plan under the office; organization; concerning a change to the Medicaid program, including a change to prior authorization, claims processing, payment rates, and medical policies, that does not require use of the rulemaking process under IC 4-22-2 may not become effective until thirty (30) days after the date the notice or bulletin is communicated to the parties affected by the notice or bulletin.
- (b) The office must provide a written notice or bulletin described in subsection (a) within five (5) business days after the date on the notice or bulletin.
- (c) If the office, a contractor of the office, or a managed care plan under the office **organization** does not comply with the requirements in subsections (a) and (b):
 - (1) the notice or bulletin is void;
 - (2) a claim may not be denied because the claim does not comply with the void notice or bulletin; and
 - (3) the office, a contractor of the office, or a managed care plan under the office organization may not reissue the bulletin or notice for thirty (30) days unless the change is required by the federal government to be implemented earlier.
- SECTION 23. IC 12-15-15-2.5 IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 2.5. (a) Payment for physician services provided in the emergency department of a hospital licensed under IC 16-21 must be at a rate of one hundred percent (100%) of rates payable under the Medicaid fee structure.
- (b) The payment under subsection (a) must be calculated using the same methodology used for all other physicians participating in the Medicaid program.
- (c) For services rendered and documented in an individual's medical record, physicians must be reimbursed for federally required medical screening exams that are necessary to determine the presence of an emergency using the appropriate Current Procedural Terminology (CPT) codes 99281, 99282, or 99283 described in the Current Procedural Terminology Manual published annually by the American



1	Medical Association, without authorization by the enrollee's primary
2	medical provider.
3	(d) Payment for all other physician services provided in an
4	emergency department of a hospital to enrollees in the Medicaid
5	primary care case management program must be at a rate of one
6	hundred percent (100%) of the Medicaid fee structure rates, provided
7	the service is authorized, prospectively or retrospectively, by the
8	enrollee's primary medical provider.
9	(e) (d) This section does not apply to a person enrolled in the a
10	Medicaid risk-based risk based managed care program.
11	SECTION 24. IC 12-15-30-3 IS AMENDED TO READ AS
12	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 3. The office shall
13	select an approach to finance and administer Medicaid claims
14	consisting of one (1) of the following:
15	(1) A direct provider payment plan administered by the office.
16	(2) A direct provider payment plan administered by a fiscal agent.
17	(3) A Medicaid insurance plan administered by a health insurer.
18	managed care organization.
19	(4) Any combination of the plans described in this section.
20	SECTION 25. IC 12-15-35-18.7 IS AMENDED TO READ AS
21	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 18.7. A formulary
22	established by a Medicaid managed care organization is subject to
23	sections 46 and 47 of this chapter.
24	SECTION 26. IC 12-15-35-20 IS AMENDED TO READ AS
25	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 20. The board is
26	composed of the following:
27	(1) Four (4) individuals licensed and actively engaged in the
28	practice of medicine or osteopathic medicine in Indiana under
29	IC 25-22.5.
30	(2) Four (4) individuals licensed under IC 25-26 and actively
31	engaged in the practice of pharmacy in Indiana.
32	(3) One (1) individual with expertise in the rapeutic pharmacology
33	who is neither a physician or a pharmacist.
34	(4) A representative of the office who shall serve as an ex-officio
35	nonvoting member of the board.
36	(5) One (1) individual who:
37	(A) is employed by a health maintenance organization that has
38	a pharmacy benefit; and
39	(B) has expertise in formulary development and pharmacy
40	benefit administration.
41	The individual appointed under this subdivision may not be
42	employed by a health maintenance organization that is under



1 2	contract or subcontract with the state to provide services to Medicaid recipients under this article. a managed care
3	organization.
4	(6) One (1) individual who is a health economist.
5	SECTION 27. IC 12-15-35-28, AS AMENDED BY P.L.210-2015,
6	SECTION 50, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
7	JULY 1, 2017]: Sec. 28. (a) The board has the following duties:
8	(1) The implementation of a Medicaid retrospective and
9	prospective DUR program as outlined in this chapter, including
10	the approval of software programs to be used by the pharmacist
11	for prospective DUR and recommendations concerning the
12	provisions of the contractual agreement between the state and any
13	other entity that will be processing and reviewing Medicaid drug
14	claims and profiles for the DUR program under this chapter.
15	(2) The development and application of the predetermined criteria
16	and standards for appropriate prescribing to be used in
17	retrospective and prospective DUR to ensure that such criteria
18	and standards for appropriate prescribing are based on the
19	compendia and developed with professional input with provisions
20	for timely revisions and assessments as necessary.
21	(3) The development, selection, application, and assessment of
22	interventions for physicians, pharmacists, and patients that are
23	educational and not punitive in nature.
24	(4) The publication of an annual report that must be subject to
25	public comment before issuance to the federal Department of
26	Health and Human Services and to the Indiana legislative council
27	by December 1 of each year. The report issued to the legislative
28	council must be in an electronic format under IC 5-14-6.
29	(5) The development of a working agreement for the board to
30	clarify the areas of responsibility with related boards or agencies,
31	including the following:
32	(A) The Indiana board of pharmacy.
33	(B) The medical licensing board of Indiana.
34	(C) The SURS staff.
35	(6) The establishment of a grievance and appeals process for
36	physicians or pharmacists under this chapter.
37	(7) The publication and dissemination of educational information
38	to physicians and pharmacists regarding the board and the DUR
39	program, including information on the following:
40	(A) Identifying and reducing the frequency of patterns of
41	fraud, abuse, gross overuse, or inappropriate or medically
42	unnecessary care among physicians, pharmacists, and



1	recipients.
2	(B) Potential or actual severe or adverse reactions to drugs.
3	(C) Therapeutic appropriateness.
4	(D) Overutilization or underutilization.
5	(E) Appropriate use of generic drugs.
6	(F) Therapeutic duplication.
7	(G) Drug-disease contraindications.
8	(H) Drug-drug interactions.
9	(I) Incorrect drug dosage and duration of drug treatment.
10	(J) Drug allergy interactions.
11	(K) Clinical abuse and misuse.
12	(8) The adoption and implementation of procedures designed to
13	ensure the confidentiality of any information collected, stored,
14	retrieved, assessed, or analyzed by the board, staff to the board, or
15	contractors to the DUR program that identifies individual
16	physicians, pharmacists, or recipients.
17	(9) The implementation of additional drug utilization review with
18	respect to drugs dispensed to residents of nursing facilities shall
19	not be required if the nursing facility is in compliance with the
20	drug regimen procedures under 410 IAC 16.2-3.1 and 42 CFR
21	483.60.
22 23 24	(10) The research, development, and approval of a preferred drug
23	list for:
	(A) Medicaid's fee for service program;
25	(B) Medicaid's primary care case management program;
26	(C) Medicaid's (B) a risk based managed care program, if the
27	office provides a prescription drug benefit and subject to
28	IC 12-15-5; and
29	(D) (C) the children's health insurance program under
30	IC 12-17.6;
31	in consultation with the therapeutics committee.
32	(11) The approval of the review and maintenance of the preferred
33	drug list at least two (2) times per year.
34	(12) The preparation and submission of a report concerning the
35	preferred drug list at least one (1) time per year to the interim
36	study committee on public health, behavioral health, and human
37	services established by IC 2-5-1.3-4 in an electronic format under
38	IC 5-14-6.
39	(13) The collection of data reflecting prescribing patterns related
40	to treatment of children diagnosed with attention deficit disorder
41	or attention deficit hyperactivity disorder.
12	(14) Advising the Indiana comprehensive health insurance



1	association established by IC 27-8-10-2.1 concerning
2	implementation of chronic disease management and
3	pharmaceutical management programs under IC 27-8-10-3.5.
4	(b) The board shall use the clinical expertise of the therapeutics
5	committee in developing a preferred drug list. The board shall also
6	consider expert testimony in the development of a preferred drug list.
7	(c) In researching and developing a preferred drug list under
8	subsection (a)(10), the board shall do the following:
9	(1) Use literature abstracting technology.
10	(2) Use commonly accepted guidance principles of disease
11	management.
12	(3) Develop therapeutic classifications for the preferred drug list.
13	(4) Give primary consideration to the clinical efficacy or
14	appropriateness of a particular drug in treating a specific medical
15	condition.
16	(5) Include in any cost effectiveness considerations the cost
17	implications of other components of the state's Medicaid program
18	and other state funded programs.
19	(d) Prior authorization is required for coverage under a program
20	described in subsection (a)(10) of a drug that is not included on the
21	preferred drug list.
22	(e) The board shall determine whether to include a single source
23	covered outpatient drug that is newly approved by the federal Food and
24	Drug Administration on the preferred drug list not later than sixty (60)
25	days after the date on which the manufacturer notifies the board in
26	writing of the drug's approval. However, if the board determines that
27	there is inadequate information about the drug available to the board
28	to make a determination, the board may have an additional sixty (60)
29	days to make a determination from the date that the board receives
30	adequate information to perform the board's review. Prior authorization
31	may not be automatically required for a single source drug that is newly
32	approved by the federal Food and Drug Administration, and that is:
33	(1) in a therapeutic classification:
34	(A) that has not been reviewed by the board; and
35	(B) for which prior authorization is not required; or
36	(2) the sole drug in a new therapeutic classification that has not
37	been reviewed by the board.
38	(f) The board may not exclude a drug from the preferred drug list
39	based solely on price.
40	(g) The following requirements apply to a preferred drug list
41	developed under subsection (a)(10):
42	(1) Except as provided by IC 12-15-35.5-3(b) and



1	IC 12-15-35.5-3(c), the office or the board may require prior
2	authorization for a drug that is included on the preferred drug list
3	under the following circumstances:
4	(A) To override a prospective drug utilization review alert.
5	(B) To permit reimbursement for a medically necessary brand
6	name drug that is subject to generic substitution under
7	IC 16-42-22-10.
8	(C) To prevent fraud, abuse, waste, overutilization, or
9	inappropriate utilization.
10	(D) To permit implementation of a disease management
11	program.
12	(E) To implement other initiatives permitted by state or federal
13	law.
14	(2) All drugs described in IC 12-15-35.5-3(b) must be included on
15	the preferred drug list.
16	(3) The office may add a drug that has been approved by the
17	federal Food and Drug Administration to the preferred drug list
18	without prior approval from the board.
19	(4) The board may add a drug that has been approved by the
20	federal Food and Drug Administration to the preferred drug list.
21	(h) At least one (1) time each year, the board shall provide a report
22	to the interim study committee on public health, behavioral health, and
23	human services established by IC 2-5-1.3-4 in an electronic format
24 25	under IC 5-14-6. The report must contain the following information:
25	(1) The cost of administering the preferred drug list.
26	(2) Any increase in Medicaid physician, laboratory, or hospital
27	costs or in other state funded programs as a result of the preferred
28	drug list.
29	(3) The impact of the preferred drug list on the ability of a
30	Medicaid recipient to obtain prescription drugs.
31	(4) The number of times prior authorization was requested, and
32	the number of times prior authorization was:
33	(A) approved; and
34	(B) disapproved.
35	(i) The board shall provide the first report required under subsection
36	(h) not later than six (6) months after the board submits an initial
37	preferred drug list to the office.
38	SECTION 28. IC 12-15-35-45, AS AMENDED BY P.L.101-2005,
39	SECTION 4, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
40	JULY 1, 2017]: Sec. 45. (a) The chairman of the board, subject to the
41	approval of the board members, may appoint an advisory committee to
42	make recommendations to the board on the development of a Medicaid



outpatient drug formulary.

2	(b) If the office decides to establish a Medicaid outpatient drug
3	formulary, the formulary shall be developed by the board.
4	(c) A formulary, preferred drug list, or prescription drug benefit
5	used by a Medicaid managed care organization is subject to
6	IC 12-15-5-5, IC 12-15-35.5, and sections 46 and 47 of this chapter.
7	SECTION 29. IC 12-15-35-46 IS AMENDED TO READ AS
8	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 46. (a) This section
9	applies to a managed care organization that enters into an initial
10	contract with the office to be a Medicaid managed care organization
11	after May 13, 1999.
12	(b) Before a Medicaid managed care organization described in
13	subsection (a) implements a formulary, the managed care organization
14	shall submit the formulary to the office at least thirty-five (35) days
15	before the date that the managed care organization implements the
16	formulary for Medicaid recipients.
17	(c) The office shall forward the formulary to the board for the
18	board's review and recommendation.
19	(d) The office shall provide at least thirty (30) days notification to
20	the public that the board will review a Medicaid managed care
21	organization's proposed formulary at a particular board meeting. The
22	notification shall contain the following information:
23	(1) A statement of the date, time, and place at which the board
24	meeting will be convened.
25	(2) A general description of the subject matter of the board
26	meeting.
27	(3) An explanation of how a copy of the formulary to be discussed
28	may be obtained.
29	The board shall meet to review the formulary at least thirty (30) days
30	but not more than sixty (60) days after the notification.
31	(e) In reviewing the formulary, the board shall do the following:
32	(1) Make a determination, after considering evidence and credible
33	information provided to the board by the office and the public,
34	that the use of the formulary will not:
35	(A) impede the quality of patient care in the Medicaid
36	program; or
37	(B) increase costs in other parts of the Medicaid program,
38	including hospital costs and physician costs.
39	(2) Make a determination that:
40	(A) there is access to at least two (2) alternative drugs within
41	each therapeutic classification, if available, on the formulary;
42	(B) a process is in place through which a Medicaid member



1	has access to medically necessary drugs; and
2	(C) the managed care organization otherwise meets the
3	requirements of IC 27-13-38.
4	(f) The board shall consider:
5	(1) health economic data;
6	(2) cost data; and
7	(3) the use of formularies in the non-Medicaid markets;
8	in developing its recommendation to the office.
9	(g) Within thirty (30) days after the board meeting, the board shall
0	make a recommendation to the office regarding whether the proposed
1	formulary should be approved, disapproved, or modified.
2	(h) The office shall rely significantly on the clinical expertise of the
3	board. If the office does not agree with the recommendations of the
4	board, the office shall, at a public meeting, discuss the disagreement
5	with the board and present any additional information to the board for
6	the board's consideration. The board's consideration of additional
7	information must be conducted at a public meeting.
8	(i) Based on the final recommendations of the board, the office shall
9	approve, disapprove, or require modifications to the Medicaid managed
0.	care organization's proposed formulary. The office shall notify the
21	managed care organization of the office's decision within fifteen (15)
22	days of receiving the board's final recommendation.
23 24	(j) The managed care organization must comply with the office's
	decision within sixty (60) days after receiving notice of the office's
25 26	decision.
	(k) Notwithstanding the other provisions of this section, the office
27	may temporarily approve a Medicaid managed care organization's
28	proposed formulary pending a final recommendation from the board.
.9	SECTION 30. IC 12-15-35-47 IS AMENDED TO READ AS
0	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 47. (a) This section
1	applies to the following changes to a formulary used by a Medicaid
2	managed care organization for Medicaid recipients:
3	(1) Removing one (1) or more drugs from the formulary.
4	(2) Placing new restrictions on one (1) or more drugs on the
5	formulary.
6	(b) Before a Medicaid managed care organization makes a change
7	described in subsection (a), the managed care organization shall submit
8	the proposed change to the office.
9	(c) The office shall forward the proposed change to the board for the
-0	board's review and recommendation

(d) The office shall provide at least thirty (30) days notification to

the public that the board will:



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1	(1) review the proposed change; and
2	(2) consider evidence and credible information provided to the
3	board;
4	at the board's regular board meeting before making a recommendation
5	to the office regarding whether the proposed change should be
6	approved or disapproved.
7	(e) Based on the final recommendation of the board, the office may
8	approve or disapprove the proposed change. If a proposed change is not
9	disapproved within ninety (90) days after the date the managed care
0	organization submits the proposed change to the office, the managed
1	care organization may implement the change to the formulary.
2	(f) A Medicaid managed care organization:
3	(1) may add a drug to the managed care organization's formulary
4	without the approval of the office; and
5	(2) shall notify the office of any addition to the managed care
6	organization's formulary within thirty (30) days after making the
7	addition.
8	SECTION 31. IC 12-15-35-48, AS AMENDED BY P.L.53-2014,
9	SECTION 106, IS AMENDED TO READ AS FOLLOWS
20	[EFFECTIVE JULY 1, 2017]: Sec. 48. (a) The board shall review the
21	prescription drug program of a managed care organization that
22	participates in the state's risk-based a risk based managed care
23	program at least one (1) time per year. The board's review of a
22 23 24 25 26	prescription drug program must include the following:
2.5	(1) An analysis of the single source drugs requiring prior
26	authorization, including the number of drugs requiring prior
27	authorization in comparison to other managed care organizations'
28	prescription drug programs that participate in the state's Medicaid
29	program.
0	(2) A determination and analysis of the number and the type of
1	drugs subject to a restriction.
52	(3) A review of the rationale for:
3	(A) the prior authorization of a drug described in subdivision
4	(1); and
5	(B) a restriction on a drug.
6	(4) A review of the number of requests a managed care
7	organization received for prior authorization, including the
8	number of times prior authorization was approved and the number
9	of times prior authorization was disapproved.
$\cdot 0$	(5) A review of:
-1	(A) patient and provider satisfaction survey reports; and
-2	(B) pharmacy-related grievance data for a twelve (12) month



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1	period.
2	(b) A managed care organization described in subsection (a) shall
3	provide the board with the information necessary for the board to
4	conduct its review under subsection (a).
5	(c) The board shall report to the interim study committee on public
6	health, behavioral health, and human services established by
7	IC 2-5-1.3-4 in an electronic format under IC 5-14-6 at least one (1)
8	time per year on the board's review under subsection (a).
9	SECTION 32. IC 12-15-44.5-3, AS AMENDED BY P.L.30-2016,
0	SECTION 27, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
1	JULY 1, 2017]: Sec. 3. (a) The healthy Indiana plan is established.
2	(b) The office shall administer the plan.
3	(c) The following individuals are eligible for the plan:
4	(1) The adult group described in 42 CFR 435.119.
5	(2) Parents and caretaker relatives eligible under 42 CFR 435.110.
6	(3) Low income individuals who are:
7	(A) at least nineteen (19) years of age; and
8	(B) less than twenty-one (21) years of age;
9	and eligible under 42 CFR 435.222.
20	(4) Individuals, for purposes of receiving transitional medical
21	assistance.
22 23 24 25	An individual must meet the Medicaid residency requirements under
23	IC 12-15-4-4 and this article to be eligible for the plan.
.4	(d) The following individuals are not eligible for the plan:
25	(1) An individual who participates in the federal Medicare
26	program (42 U.S.C. 1395 et seq.).
27	(2) An individual who is otherwise eligible and enrolled for
28	medical assistance.
9	(e) The department of insurance and the office of the secretary shall
0	provide oversight of the marketing practices of the plan.
1	(f) The office shall promote the plan and provide information to
2	potential eligible individuals who live in medically underserved rural
3	areas of Indiana.
4	(g) The office shall, to the extent possible, ensure that enrollment in
5	the plan is distributed throughout Indiana in proportion to the number
6	of individuals throughout Indiana who are eligible for participation in
7	the plan. (b) The office shall establish standards for consumer materials.
8	(h) The office shall establish standards for consumer protection,
9	including the following: (1) Quality of core standards
-0 -1	(1) Quality of care standards.
2	(2) A uniform process for participant grievances and appeals.(3) Standardized reporting concerning provider performance.
· _	1.31 Standardized reporting concerning provider performance.



1	consumer experience, and cost.
2	(i) A health care provider that provides care to an individual who
3	receives health insurance coverage under the plan shall also participate
4	in the Medicaid program under this article.
5	(j) The following do not apply to the plan:
6	(1) IC 12-15-6.
7	(1) IC 12-13-0. (2) IC 12-15-12.
8	(3) IC 12-15-13.
9	(4) IC 12-15-14.
10	(4) IC 12-13-14. (5) IC 12-15-15.
11	(6) IC 12-15-13. (6) IC 12-15-21.
12	(6) IC 12-13-21. (7) IC 12-15-26.
13	(8) IC 12-15-31.1.
13	(8) IC 12-13-31.1. (9) IC 12-15-34.
15	(10) IC 12-15-34. (10) IC 12-15-35.
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17	(11) IC 16-42-22-10.
18	SECTION 33. IC 12-15-44.5-4.7, AS ADDED BY P.L.30-2016,
	SECTION 31, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
19	JULY 1, 2017]: Sec. 4.7. (a) To participate in the plan, an individual
20	must apply for the plan on a form prescribed by the office. The office
21	may develop and allow a joint application for a household.
22	(b) A pregnant woman is not subject to the cost sharing provisions
23	of the plan. Subsections (c) through (g) do not apply to a pregnant
24	woman participating in the plan.
25	(c) An applicant who is approved to participate in the plan does not
26	begin benefits under the plan until a payment of at least:
27	(1) one-twelfth $(1/12)$ of the two percent (2%) of annual income
28	contribution amount; or
29	(2) ten dollars (\$10);
30	is made to the individual's health care account established under
31	section 4.5 of this chapter for the individual's participation in the plan.
32	To continue to participate in the plan, an individual must contribute to
33	the individual's health care account at least two percent (2%) of the
34	individual's annual household income per year but not less than one
35	dollar (\$1) per month.
36	(d) If an applicant who is approved to participate in the plan fails to
37	make the initial payment into the individual's health care account, at
38	least the following must occur:
39	(1) If the individual has an annual income that is at or below one
40	hundred percent (100%) of the federal poverty income level, the
41	individual's benefits are reduced as specified in subsection (e)(1).

(2) If the individual has an annual income of more than one



1	hundred percent (100%) of the federal poverty income level, the
2	individual is not enrolled in the plan.
3 4	(e) If an enrolled individual's required monthly payment to the plan is not made within sixty (60) days after the required payment date, the
5	following, at a minimum, occur:
6	(1) For an individual who has an annual income that is at or below
7	one hundred percent (100%) of the federal income poverty level,
8	the individual is:
9	(A) transferred to a plan that has a material reduction in
10	benefits, including the elimination of benefits for vision and
11	dental services; and
12	(B) required to make copayments for the provision of services
13	that may not be paid from the individual's health care account.
14	(2) For an individual who has an annual income of more than one
15	hundred percent (100%) of the federal poverty income level, the
16	individual shall be terminated from the plan and may not reenroll
17	in the plan for at least six (6) months.
18	(f) The state shall contribute to the individual's health care account
19	the difference between the individual's payment required under this
20	section and the plan deductible set forth in section 4.5(c) of this
21	chapter.
22	(g) A member shall remain enrolled with the same health plan
23	managed care organization during the member's benefit period. A
24	member may change health plans managed care organizations as
25	follows:
26	(1) Without cause:
27	(A) before making a contribution or before finalizing
28	enrollment in accordance with subsection (d)(1); or
29	(B) during the annual plan renewal process.
30	(2) For cause, as determined by the office.
31	SECTION 34. IC 12-15-44.5-5, AS ADDED BY P.L.213-2015,
32	SECTION 136, IS AMENDED TO READ AS FOLLOWS
33	[EFFECTIVE JULY 1, 2017]: Sec. 5. (a) An insurer or health
34	maintenance A managed care organization that contracts with the
35	office to provide health insurance coverage, dental coverage, or vision
36	coverage to an individual who participates in the plan:
37	(1) is responsible for the claim processing for the coverage;
38	(2) shall reimburse providers at a rate that is not less than the rate
39	established by the secretary. The rate set by the secretary must be
40	based on a reimbursement formula that is:
41	(A) comparable to the federal Medicare reimbursement rate
42	for the service provided by the provider; or
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1	(B) one hundred thirty percent (130%) of the Medicaid
2	reimbursement rate for a service that does not have a Medicare
3	reimbursement rate; and
4	(3) may not deny coverage to an eligible individual who has been
5	approved by the office to participate in the plan.
6	(b) An insurer or health maintenance A managed care organization
7	that contracts with the office to provide health insurance coverage
8	under the plan must incorporate cultural competency standards
9	established by the office. The standards must include standards for
0	non-English speaking, minority, and disabled populations.
1	SECTION 35. IC 12-15-44.5-8, AS ADDED BY P.L.213-2015,
2	SECTION 136, IS AMENDED TO READ AS FOLLOWS
3	[EFFECTIVE JULY 1, 2017]: Sec. 8. The following requirements
4	apply to funds appropriated by the general assembly to the plan and the
5	incremental fee used for purposes of IC 16-21-10-13.3:
6	(1) At least eighty-seven percent (87%) of the funds must be used
7	to fund payment for health care services.
8	(2) An amount determined by the office of the secretary to fund:
9	(A) administrative costs of; and
0.	(B) any profit made by;
1	an insurer or a health maintenance managed care organization
22	under a contract with the office to provide health insurance
23 24	coverage under the plan. The amount determined under this
4	subdivision may not exceed thirteen percent (13%) of the funds.
25	SECTION 36. IC 16-38-6-2 IS AMENDED TO READ AS
26	FOLLOWS [EFFECTIVE JULY 1, 2017]: Sec. 2. The state
27	department, with the cooperation of the office of Medicaid policy and
28	planning, shall establish a chronic disease registry for the purpose of
9	(1) recording chronic disease cases that are diagnosed or treated
0	in Indiana; and
1	(2) compiling necessary and appropriate information determined
2	by the state department concerning cases described in subdivision
3	(1) in order to do the following:
4	(A) Conduct epidemiologic and environmental surveys of
5	chronic disease and use appropriate preventive and control
6	measures.
7	(B) Inform citizens regarding programs designed to manage
8	chronic disease.
9	(C) Provide guidance to the office of Medicaid policy and
0	planning to identify and develop cost and clinical measures for
-1	use in a program required by IC 12-15-12-19.

