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**SENATE BILL NO. 1262**

Offered January 14, 2015

Prefiled January 14, 2015

*A BILL to amend and reenact §§ 2.2-2818, 32.1-325, and 32.1-330.2 of the Code of Virginia and to amend the Code of Virginia by adding a section numbered 38.2-3407.15:2, relating to health benefits plans and programs; carrier business practices; contract preauthorization provisions.*

Patrons—Newman and Watkins

Referred to Committee on Commerce and Labor

**Be it enacted by the General Assembly of Virginia:**

**1. That §§ 2.2-2818, 32.1-325, and 32.1-330.2 of the Code of Virginia are amended and reenacted and that the Code of Virginia is amended by adding a section numbered 38.2-3407.15:2 as follows:**

**§ 2.2-2818. Health and related insurance for state employees.**

A. The Department of Human Resource Management shall establish a plan, subject to the approval of the Governor, for providing health insurance coverage, including chiropractic treatment, hospitalization, medical, surgical and major medical coverage, for state employees and retired state employees with the Commonwealth paying the cost thereof to the extent of the coverage included in such plan. The same plan shall be offered to all part-time state employees, but the total cost shall be paid by such part-time employees. The Department of Human Resource Management shall administer this section. The plan chosen shall provide means whereby coverage for the families or dependents of state employees may be purchased. Except for part-time employees, the Commonwealth may pay all or a portion of the cost thereof, and for such portion as the Commonwealth does not pay, the employee, including a part-time employee, may purchase the coverage by paying the additional cost over the cost of coverage for an employee.

Such contribution shall be financed through appropriations provided by law.

B. The plan shall:

1. Include coverage for low-dose screening mammograms for determining the presence of occult breast cancer. Such coverage shall make available one screening mammogram to persons age 35 through 39, one such mammogram biennially to persons age 40 through 49, and one such mammogram annually to persons age 50 and over and may be limited to a benefit of \$50 per mammogram subject to such dollar limits, deductibles, and coinsurance factors as are no less favorable than for physical illness generally.

The term "mammogram" shall mean an X-ray examination of the breast using equipment dedicated specifically for mammography, including but not limited to the X-ray tube, filter, compression device, screens, film, and cassettes, with an average radiation exposure of less than one rad mid-breast, two views of each breast.

In order to be considered a screening mammogram for which coverage shall be made available under this section:

a. The mammogram shall be (i) ordered by a health care practitioner acting within the scope of his licensure and, in the case of an enrollee of a health maintenance organization, by the health maintenance organization provider; (ii) performed by a registered technologist; (iii) interpreted by a qualified radiologist; and (iv) performed under the direction of a person licensed to practice medicine and surgery and certified by the American Board of Radiology or an equivalent examining body. A copy of the mammogram report shall be sent or delivered to the health care practitioner who ordered it;

b. The equipment used to perform the mammogram shall meet the standards set forth by the Virginia Department of Health in its radiation protection regulations; and

c. The mammography film shall be retained by the radiologic facility performing the examination in accordance with the American College of Radiology guidelines or state law.

2. Include coverage for postpartum services providing inpatient care and a home visit or visits that shall be in accordance with the medical criteria, outlined in the most current version of or an official update to the "Guidelines for Perinatal Care" prepared by the American Academy of Pediatrics and the American College of Obstetricians and Gynecologists or the "Standards for Obstetric-Gynecologic Services" prepared by the American College of Obstetricians and Gynecologists. Such coverage shall be provided incorporating any changes in such Guidelines or Standards within six months of the publication of such Guidelines or Standards or any official amendment thereto.

3. Include an appeals process for resolution of complaints that shall provide reasonable procedures for the resolution of such complaints and shall be published and disseminated to all covered state

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SB1262

59 employees. The appeals process shall be compliant with federal rules and regulations governing  
60 nonfederal, self-insured governmental health plans. The appeals process shall include a separate  
61 expedited emergency appeals procedure that shall provide resolution within time frames established by  
62 federal law. For appeals involving adverse decisions as defined in § 32.1-137.7, the Department shall  
63 contract with one or more independent review organizations to review such decisions. Independent  
64 review organizations are entities that conduct independent external review of adverse benefit  
65 determinations. The Department shall adopt regulations to assure that the independent review  
66 organization conducting the reviews has adequate standards, credentials and experience for such review.  
67 The independent review organization shall examine the final denial of claims to determine whether the  
68 decision is objective, clinically valid, and compatible with established principles of health care. The  
69 decision of the independent review organization shall (i) be in writing, (ii) contain findings of fact as to  
70 the material issues in the case and the basis for those findings, and (iii) be final and binding if  
71 consistent with law and policy.

72 Prior to assigning an appeal to an independent review organization, the Department shall verify that  
73 the independent review organization conducting the review of a denial of claims has no relationship or  
74 association with (i) the covered person or the covered person's authorized representative; (ii) the treating  
75 health care provider, or any of its employees or affiliates; (iii) the medical care facility at which the  
76 covered service would be provided, or any of its employees or affiliates; or (iv) the development or  
77 manufacture of the drug, device, procedure or other therapy that is the subject of the final denial of a  
78 claim. The independent review organization shall not be a subsidiary of, nor owned or controlled by, a  
79 health plan, a trade association of health plans, or a professional association of health care providers.  
80 There shall be no liability on the part of and no cause of action shall arise against any officer or  
81 employee of an independent review organization for any actions taken or not taken or statements made  
82 by such officer or employee in good faith in the performance of his powers and duties.

83 4. Include coverage for early intervention services. For purposes of this section, "early intervention  
84 services" means medically necessary speech and language therapy, occupational therapy, physical therapy  
85 and assistive technology services and devices for dependents from birth to age three who are certified by  
86 the Department of Behavioral Health and Developmental Services as eligible for services under Part H  
87 of the Individuals with Disabilities Education Act (20 U.S.C. § 1471 et seq.). Medically necessary early  
88 intervention services for the population certified by the Department of Behavioral Health and  
89 Developmental Services shall mean those services designed to help an individual attain or retain the  
90 capability to function age-appropriately within his environment, and shall include services that enhance  
91 functional ability without effecting a cure.

92 For persons previously covered under the plan, there shall be no denial of coverage due to the  
93 existence of a preexisting condition. The cost of early intervention services shall not be applied to any  
94 contractual provision limiting the total amount of coverage paid by the insurer to or on behalf of the  
95 insured during the insured's lifetime.

96 5. Include coverage for prescription drugs and devices approved by the United States Food and Drug  
97 Administration for use as contraceptives.

98 6. Not deny coverage for any drug approved by the United States Food and Drug Administration for  
99 use in the treatment of cancer on the basis that the drug has not been approved by the United States  
100 Food and Drug Administration for the treatment of the specific type of cancer for which the drug has  
101 been prescribed, if the drug has been recognized as safe and effective for treatment of that specific type  
102 of cancer in one of the standard reference compendia.

103 7. Not deny coverage for any drug prescribed to treat a covered indication so long as the drug has  
104 been approved by the United States Food and Drug Administration for at least one indication and the  
105 drug is recognized for treatment of the covered indication in one of the standard reference compendia or  
106 in substantially accepted peer-reviewed medical literature.

107 8. Include coverage for equipment, supplies and outpatient self-management training and education,  
108 including medical nutrition therapy, for the treatment of insulin-dependent diabetes, insulin-using  
109 diabetes, gestational diabetes and noninsulin-using diabetes if prescribed by a health care professional  
110 legally authorized to prescribe such items under law. To qualify for coverage under this subdivision,  
111 diabetes outpatient self-management training and education shall be provided by a certified, registered or  
112 licensed health care professional.

113 9. Include coverage for reconstructive breast surgery. For purposes of this section, "reconstructive  
114 breast surgery" means surgery performed on and after July 1, 1998, (i) coincident with a mastectomy  
115 performed for breast cancer or (ii) following a mastectomy performed for breast cancer to reestablish  
116 symmetry between the two breasts. For persons previously covered under the plan, there shall be no  
117 denial of coverage due to preexisting conditions.

118 10. Include coverage for annual pap smears, including coverage, on and after July 1, 1999, for  
119 annual testing performed by any FDA-approved gynecologic cytology screening technologies.

120 11. Include coverage providing a minimum stay in the hospital of not less than 48 hours for a patient

following a radical or modified radical mastectomy and 24 hours of inpatient care following a total mastectomy or a partial mastectomy with lymph node dissection for treatment of breast cancer. Nothing in this subdivision shall be construed as requiring the provision of inpatient coverage where the attending physician in consultation with the patient determines that a shorter period of hospital stay is appropriate.

12. Include coverage (i) to persons age 50 and over and (ii) to persons age 40 and over who are at high risk for prostate cancer, according to the most recent published guidelines of the American Cancer Society, for one PSA test in a 12-month period and digital rectal examinations, all in accordance with American Cancer Society guidelines. For the purpose of this subdivision, "PSA testing" means the analysis of a blood sample to determine the level of prostate specific antigen.

13. Permit any individual covered under the plan direct access to the health care services of a participating specialist (i) authorized to provide services under the plan and (ii) selected by the covered individual. The plan shall have a procedure by which an individual who has an ongoing special condition may, after consultation with the primary care physician, receive a referral to a specialist for such condition who shall be responsible for and capable of providing and coordinating the individual's primary and specialty care related to the initial specialty care referral. If such an individual's care would most appropriately be coordinated by such a specialist, the plan shall refer the individual to a specialist. For the purposes of this subdivision, "special condition" means a condition or disease that is (i) life-threatening, degenerative, or disabling and (ii) requires specialized medical care over a prolonged period of time. Within the treatment period authorized by the referral, such specialist shall be permitted to treat the individual without a further referral from the individual's primary care provider and may authorize such referrals, procedures, tests, and other medical services related to the initial referral as the individual's primary care provider would otherwise be permitted to provide or authorize. The plan shall have a procedure by which an individual who has an ongoing special condition that requires ongoing care from a specialist may receive a standing referral to such specialist for the treatment of the special condition. If the primary care provider, in consultation with the plan and the specialist, if any, determines that such a standing referral is appropriate, the plan or issuer shall make such a referral to a specialist. Nothing contained herein shall prohibit the plan from requiring a participating specialist to provide written notification to the covered individual's primary care physician of any visit to such specialist. Such notification may include a description of the health care services rendered at the time of the visit.

14. Include provisions allowing employees to continue receiving health care services for a period of up to 90 days from the date of the primary care physician's notice of termination from any of the plan's provider panels. The plan shall notify any provider at least 90 days prior to the date of termination of the provider, except when the provider is terminated for cause.

For a period of at least 90 days from the date of the notice of a provider's termination from any of the plan's provider panels, except when a provider is terminated for cause, a provider shall be permitted by the plan to render health care services to any of the covered employees who (i) were in an active course of treatment from the provider prior to the notice of termination and (ii) request to continue receiving health care services from the provider.

Notwithstanding the provisions of this subdivision, any provider shall be permitted by the plan to continue rendering health services to any covered employee who has entered the second trimester of pregnancy at the time of the provider's termination of participation, except when a provider is terminated for cause. Such treatment shall, at the covered employee's option, continue through the provision of postpartum care directly related to the delivery.

Notwithstanding the provisions of this subdivision, any provider shall be permitted to continue rendering health services to any covered employee who is determined to be terminally ill (as defined under § 1861(dd)(3)(A) of the Social Security Act) at the time of a provider's termination of participation, except when a provider is terminated for cause. Such treatment shall, at the covered employee's option, continue for the remainder of the employee's life for care directly related to the treatment of the terminal illness.

A provider who continues to render health care services pursuant to this subdivision shall be reimbursed in accordance with the carrier's agreement with such provider existing immediately before the provider's termination of participation.

15. Include coverage for patient costs incurred during participation in clinical trials for treatment studies on cancer, including ovarian cancer trials.

The reimbursement for patient costs incurred during participation in clinical trials for treatment studies on cancer shall be determined in the same manner as reimbursement is determined for other medical and surgical procedures. Such coverage shall have durational limits, dollar limits, deductibles, copayments and coinsurance factors that are no less favorable than for physical illness generally.

For purposes of this subdivision:

182 "Cooperative group" means a formal network of facilities that collaborate on research projects and  
183 have an established NIH-approved peer review program operating within the group. "Cooperative group"  
184 includes (i) the National Cancer Institute Clinical Cooperative Group and (ii) the National Cancer  
185 Institute Community Clinical Oncology Program.

186 "FDA" means the Federal Food and Drug Administration.

187 "Multiple project assurance contract" means a contract between an institution and the federal  
188 Department of Health and Human Services that defines the relationship of the institution to the federal  
189 Department of Health and Human Services and sets out the responsibilities of the institution and the  
190 procedures that will be used by the institution to protect human subjects.

191 "NCI" means the National Cancer Institute.

192 "NIH" means the National Institutes of Health.

193 "Patient" means a person covered under the plan established pursuant to this section.

194 "Patient cost" means the cost of a medically necessary health care service that is incurred as a result  
195 of the treatment being provided to a patient for purposes of a clinical trial. "Patient cost" does not  
196 include (i) the cost of nonhealth care services that a patient may be required to receive as a result of the  
197 treatment being provided for purposes of a clinical trial, (ii) costs associated with managing the research  
198 associated with the clinical trial, or (iii) the cost of the investigational drug or device.

199 Coverage for patient costs incurred during clinical trials for treatment studies on cancer shall be  
200 provided if the treatment is being conducted in a Phase II, Phase III, or Phase IV clinical trial. Such  
201 treatment may, however, be provided on a case-by-case basis if the treatment is being provided in a  
202 Phase I clinical trial.

203 The treatment described in the previous paragraph shall be provided by a clinical trial approved by:

- 204 a. The National Cancer Institute;
- 205 b. An NCI cooperative group or an NCI center;
- 206 c. The FDA in the form of an investigational new drug application;
- 207 d. The federal Department of Veterans Affairs; or
- 208 e. An institutional review board of an institution in the Commonwealth that has a multiple project  
209 assurance contract approved by the Office of Protection from Research Risks of the NCI.

210 The facility and personnel providing the treatment shall be capable of doing so by virtue of their  
211 experience, training, and expertise.

212 Coverage under this subdivision shall apply only if:

- 213 (1) There is no clearly superior, noninvestigational treatment alternative;
- 214 (2) The available clinical or preclinical data provide a reasonable expectation that the treatment will  
215 be at least as effective as the noninvestigational alternative; and
- 216 (3) The patient and the physician or health care provider who provides services to the patient under  
217 the plan conclude that the patient's participation in the clinical trial would be appropriate, pursuant to  
218 procedures established by the plan.

219 16. Include coverage providing a minimum stay in the hospital of not less than 23 hours for a  
220 covered employee following a laparoscopy-assisted vaginal hysterectomy and 48 hours for a covered  
221 employee following a vaginal hysterectomy, as outlined in Milliman & Robertson's nationally recognized  
222 guidelines. Nothing in this subdivision shall be construed as requiring the provision of the total hours  
223 referenced when the attending physician, in consultation with the covered employee, determines that a  
224 shorter hospital stay is appropriate.

225 17. Include coverage for biologically based mental illness.

226 For purposes of this subdivision, a "biologically based mental illness" is any mental or nervous  
227 condition caused by a biological disorder of the brain that results in a clinically significant syndrome  
228 that substantially limits the person's functioning; specifically, the following diagnoses are defined as  
229 biologically based mental illness as they apply to adults and children: schizophrenia, schizoaffective  
230 disorder, bipolar disorder, major depressive disorder, panic disorder, obsessive-compulsive disorder,  
231 attention deficit hyperactivity disorder, autism, and drug and alcoholism addiction.

232 Coverage for biologically based mental illnesses shall neither be different nor separate from coverage  
233 for any other illness, condition or disorder for purposes of determining deductibles, benefit year or  
234 lifetime durational limits, benefit year or lifetime dollar limits, lifetime episodes or treatment limits,  
235 copayment and coinsurance factors, and benefit year maximum for deductibles and copayment and  
236 coinsurance factors.

237 Nothing shall preclude the undertaking of usual and customary procedures to determine the  
238 appropriateness of, and medical necessity for, treatment of biologically based mental illnesses under this  
239 option, provided that all such appropriateness and medical necessity determinations are made in the same  
240 manner as those determinations made for the treatment of any other illness, condition or disorder  
241 covered by such policy or contract.

242 18. Offer and make available coverage for the treatment of morbid obesity through gastric bypass  
243 surgery or such other methods as may be recognized by the National Institutes of Health as effective for

the long-term reversal of morbid obesity. Such coverage shall have durational limits, dollar limits, deductibles, copayments and coinsurance factors that are no less favorable than for physical illness generally. Access to surgery for morbid obesity shall not be restricted based upon dietary or any other criteria not approved by the National Institutes of Health. For purposes of this subdivision, "morbid obesity" means (i) a weight that is at least 100 pounds over or twice the ideal weight for frame, age, height, and gender as specified in the 1983 Metropolitan Life Insurance tables, (ii) a body mass index (BMI) equal to or greater than 35 kilograms per meter squared with comorbidity or coexisting medical conditions such as hypertension, cardiopulmonary conditions, sleep apnea, or diabetes, or (iii) a BMI of 40 kilograms per meter squared without such comorbidity. As used herein, "BMI" equals weight in kilograms divided by height in meters squared.

19. Include coverage for colorectal cancer screening, specifically screening with an annual fecal occult blood test, flexible sigmoidoscopy or colonoscopy, or in appropriate circumstances radiologic imaging, in accordance with the most recently published recommendations established by the American College of Gastroenterology, in consultation with the American Cancer Society, for the ages, family histories, and frequencies referenced in such recommendations. The coverage for colorectal cancer screening shall not be more restrictive than or separate from coverage provided for any other illness, condition or disorder for purposes of determining deductibles, benefit year or lifetime durational limits, benefit year or lifetime dollar limits, lifetime episodes or treatment limits, copayment and coinsurance factors, and benefit year maximum for deductibles and copayments and coinsurance factors.

20. On and after July 1, 2002, require that a prescription benefit card, health insurance benefit card, or other technology that complies with the requirements set forth in § 38.2-3407.4:2 be issued to each employee provided coverage pursuant to this section, and shall upon any changes in the required data elements set forth in subsection A of § 38.2-3407.4:2, either reissue the card or provide employees covered under the plan such corrective information as may be required to electronically process a prescription claim.

21. Include coverage for infant hearing screenings and all necessary audiological examinations provided pursuant to § 32.1-64.1 using any technology approved by the United States Food and Drug Administration, and as recommended by the national Joint Committee on Infant Hearing in its most current position statement addressing early hearing detection and intervention programs. Such coverage shall include follow-up audiological examinations as recommended by a physician, physician assistant, nurse practitioner or audiologist and performed by a licensed audiologist to confirm the existence or absence of hearing loss.

22. Notwithstanding any provision of this section to the contrary, every plan established in accordance with this section shall comply with the provisions of § 2.2-2818.2.

23. *On and after January 1, 2016, any contract between the Department of Human Resource Management and a third party that provides for the administration of a plan established under this section, under which a party has the right or obligation to require preauthorization for a drug benefit, shall contain provisions that are substantively identical to the provisions required to be contained in a contract requiring preauthorization as set out in § 38.2-3407.15:2, mutatis mutandis.*

C. Claims incurred during a fiscal year but not reported during that fiscal year shall be paid from such funds as shall be appropriated by law. Appropriations, premiums and other payments shall be deposited in the employee health insurance fund, from which payments for claims, premiums, cost containment programs and administrative expenses shall be withdrawn from time to time. The funds of the health insurance fund shall be deemed separate and independent trust funds, shall be segregated from all other funds of the Commonwealth, and shall be invested and administered solely in the interests of the employees and their beneficiaries. Neither the General Assembly nor any public officer, employee, or agency shall use or authorize the use of such trust funds for any purpose other than as provided in law for benefits, refunds, and administrative expenses, including but not limited to legislative oversight of the health insurance fund.

D. For the purposes of this section:

"Peer-reviewed medical literature" means a scientific study published only after having been critically reviewed for scientific accuracy, validity, and reliability by unbiased independent experts in a journal that has been determined by the International Committee of Medical Journal Editors to have met the Uniform Requirements for Manuscripts submitted to biomedical journals. Peer-reviewed medical literature does not include publications or supplements to publications that are sponsored to a significant extent by a pharmaceutical manufacturing company or health carrier.

"Standard reference compendia" means:

1. American Hospital Formulary Service - Drug Information;
2. National Comprehensive Cancer Network's Drugs & Biologics Compendium; or
3. Elsevier Gold Standard's Clinical Pharmacology.

"State employee" means state employee as defined in § 51.1-124.3; employee as defined in

§ 51.1-201; the Governor, Lieutenant Governor and Attorney General; judge as defined in § 51.1-301 and judges, clerks and deputy clerks of regional juvenile and domestic relations, county juvenile and domestic relations, and district courts of the Commonwealth; and interns and residents employed by the School of Medicine and Hospital of the University of Virginia, and interns, residents, and employees of the Virginia Commonwealth University Health System Authority as provided in § 23-50.16:24.

E. Provisions shall be made for retired employees to obtain coverage under the above plan, including, as an option, coverage for vision and dental care. The Commonwealth may, but shall not be obligated to, pay all or any portion of the cost thereof.

F. Any self-insured group health insurance plan established by the Department of Human Resource Management that utilizes a network of preferred providers shall not exclude any physician solely on the basis of a reprimand or censure from the Board of Medicine, so long as the physician otherwise meets the plan criteria established by the Department.

G. The plan shall include, in each planning district, at least two health coverage options, each sponsored by unrelated entities. No later than July 1, 2006, one of the health coverage options to be available in each planning district shall be a high deductible health plan that would qualify for a health savings account pursuant to § 223 of the Internal Revenue Code of 1986, as amended.

In each planning district that does not have an available health coverage alternative, the Department shall voluntarily enter into negotiations at any time with any health coverage provider who seeks to provide coverage under the plan.

This subsection shall not apply to any state agency authorized by the Department to establish and administer its own health insurance coverage plan separate from the plan established by the Department.

H. Any self-insured group health insurance plan established by the Department of Human Resource Management that includes coverage for prescription drugs on an outpatient basis may apply a formulary to the prescription drug benefits provided by the plan if the formulary is developed, reviewed at least annually, and updated as necessary in consultation with and with the approval of a pharmacy and therapeutics committee, a majority of whose members are actively practicing licensed (i) pharmacists, (ii) physicians, and (iii) other health care providers.

If the plan maintains one or more drug formularies, the plan shall establish a process to allow a person to obtain, without additional cost-sharing beyond that provided for formulary prescription drugs in the plan, a specific, medically necessary nonformulary prescription drug if, after reasonable investigation and consultation with the prescriber, the formulary drug is determined to be an inappropriate therapy for the medical condition of the person. The plan shall act on such requests within one business day of receipt of the request.

Any plan established in accordance with this section shall be authorized to provide for the selection of a single mail order pharmacy provider as the exclusive provider of pharmacy services that are delivered to the covered person's address by mail, common carrier, or delivery service. As used in this subsection, "mail order pharmacy provider" means a pharmacy permitted to conduct business in the Commonwealth whose primary business is to dispense a prescription drug or device under a prescriptive drug order and to deliver the drug or device to a patient primarily by mail, common carrier, or delivery service.

I. Any plan established in accordance with this section requiring preauthorization prior to rendering medical treatment shall have personnel available to provide authorization at all times when such preauthorization is required.

J. Any plan established in accordance with this section shall provide to all covered employees written notice of any benefit reductions during the contract period at least 30 days before such reductions become effective.

K. No contract between a provider and any plan established in accordance with this section shall include provisions that require a health care provider or health care provider group to deny covered services that such provider or group knows to be medically necessary and appropriate that are provided with respect to a covered employee with similar medical conditions.

L. The Department of Human Resource Management shall appoint an Ombudsman to promote and protect the interests of covered employees under any state employee's health plan.

The Ombudsman shall:

1. Assist covered employees in understanding their rights and the processes available to them according to their state health plan.

2. Answer inquiries from covered employees by telephone and electronic mail.

3. Provide to covered employees information concerning the state health plans.

4. Develop information on the types of health plans available, including benefits and complaint procedures and appeals.

5. Make available, either separately or through an existing Internet web site utilized by the Department of Human Resource Management, information as set forth in subdivision 4 and such additional information as he deems appropriate.

6. Maintain data on inquiries received, the types of assistance requested, any actions taken and the disposition of each such matter.

7. Upon request, assist covered employees in using the procedures and processes available to them from their health plan, including all appeal procedures. Such assistance may require the review of health care records of a covered employee, which shall be done only in accordance with the federal Health Insurance Portability and Accountability Act privacy rules. The confidentiality of any such medical records shall be maintained in accordance with the confidentiality and disclosure laws of the Commonwealth.

8. Ensure that covered employees have access to the services provided by the Ombudsman and that the covered employees receive timely responses from the Ombudsman or his representatives to the inquiries.

9. Report annually on his activities to the standing committees of the General Assembly having jurisdiction over insurance and over health and the Joint Commission on Health Care by December 1 of each year.

M. The plan established in accordance with this section shall not refuse to accept or make reimbursement pursuant to an assignment of benefits made to a dentist or oral surgeon by a covered employee.

For purposes of this subsection, "assignment of benefits" means the transfer of dental care coverage reimbursement benefits or other rights under the plan. The assignment of benefits shall not be effective until the covered employee notifies the plan in writing of the assignment.

N. Beginning July 1, 2006, any plan established pursuant to this section shall provide for an identification number, which shall be assigned to the covered employee and shall not be the same as the employee's social security number.

O. Any group health insurance plan established by the Department of Human Resource Management that contains a coordination of benefits provision shall provide written notification to any eligible employee as a prominent part of its enrollment materials that if such eligible employee is covered under another group accident and sickness insurance policy, group accident and sickness subscription contract, or group health care plan for health care services, that insurance policy, subscription contract or health care plan may have primary responsibility for the covered expenses of other family members enrolled with the eligible employee. Such written notification shall describe generally the conditions upon which the other coverage would be primary for dependent children enrolled under the eligible employee's coverage and the method by which the eligible enrollee may verify from the plan that coverage would have primary responsibility for the covered expenses of each family member.

P. Any plan established by the Department of Human Resource Management pursuant to this section shall provide that coverage under such plan for family members enrolled under a participating state employee's coverage shall continue for a period of at least 30 days following the death of such state employee.

Q. The plan established in accordance with this section that follows a policy of sending its payment to the covered employee or covered family member for a claim for services received from a nonparticipating physician or osteopath shall (i) include language in the member handbook that notifies the covered employee of the responsibility to apply the plan payment to the claim from such nonparticipating provider, (ii) include this language with any such payment sent to the covered employee or covered family member, and (iii) include the name and any last known address of the nonparticipating provider on the explanation of benefits statement.

R. The Department of Human Resource Management shall report annually, by November 30 of each year, on cost and utilization information for each of the mandated benefits set forth in subsection B, including any mandated benefit made applicable, pursuant to subdivision B 22, to any plan established pursuant to this section. The report shall be in the same detail and form as required of reports submitted pursuant to § 38.2-3419.1, with such additional information as is required to determine the financial impact, including the costs and benefits, of the particular mandated benefit.

**§ 32.1-325. Board to submit plan for medical assistance services to U.S. Secretary of Health and Human Services pursuant to federal law; administration of plan; contracts with health care providers.**

A. The Board, subject to the approval of the Governor, is authorized to prepare, amend from time to time, and submit to the U.S. Secretary of Health and Human Services a state plan for medical assistance services pursuant to Title XIX of the United States Social Security Act and any amendments thereto. The Board shall include in such plan:

1. A provision for payment of medical assistance on behalf of individuals, up to the age of 21, placed in foster homes or private institutions by private, nonprofit agencies licensed as child-placing agencies by the Department of Social Services or placed through state and local subsidized adoptions to the extent permitted under federal statute;

428 2. A provision for determining eligibility for benefits for medically needy individuals which  
429 disregards from countable resources an amount not in excess of \$3,500 for the individual and an amount  
430 not in excess of \$3,500 for his spouse when such resources have been set aside to meet the burial  
431 expenses of the individual or his spouse. The amount disregarded shall be reduced by (i) the face value  
432 of life insurance on the life of an individual owned by the individual or his spouse if the cash surrender  
433 value of such policies has been excluded from countable resources and (ii) the amount of any other  
434 revocable or irrevocable trust, contract, or other arrangement specifically designated for the purpose of  
435 meeting the individual's or his spouse's burial expenses;

436 3. A requirement that, in determining eligibility, a home shall be disregarded. For those medically  
437 needy persons whose eligibility for medical assistance is required by federal law to be dependent on the  
438 budget methodology for Aid to Families with Dependent Children, a home means the house and lot used  
439 as the principal residence and all contiguous property. For all other persons, a home shall mean the  
440 house and lot used as the principal residence, as well as all contiguous property, as long as the value of  
441 the land, exclusive of the lot occupied by the house, does not exceed \$5,000. In any case in which the  
442 definition of home as provided here is more restrictive than that provided in the state plan for medical  
443 assistance services in Virginia as it was in effect on January 1, 1972, then a home means the house and  
444 lot used as the principal residence and all contiguous property essential to the operation of the home  
445 regardless of value;

446 4. A provision for payment of medical assistance on behalf of individuals up to the age of 21, who  
447 are Medicaid eligible, for medically necessary stays in acute care facilities in excess of 21 days per  
448 admission;

449 5. A provision for deducting from an institutionalized recipient's income an amount for the  
450 maintenance of the individual's spouse at home;

451 6. A provision for payment of medical assistance on behalf of pregnant women which provides for  
452 payment for inpatient postpartum treatment in accordance with the medical criteria outlined in the most  
453 current version of or an official update to the "Guidelines for Perinatal Care" prepared by the American  
454 Academy of Pediatrics and the American College of Obstetricians and Gynecologists or the "Standards  
455 for Obstetric-Gynecologic Services" prepared by the American College of Obstetricians and  
456 Gynecologists. Payment shall be made for any postpartum home visit or visits for the mothers and the  
457 children which are within the time periods recommended by the attending physicians in accordance with  
458 and as indicated by such Guidelines or Standards. For the purposes of this subdivision, such Guidelines  
459 or Standards shall include any changes thereto within six months of the publication of such Guidelines  
460 or Standards or any official amendment thereto;

461 7. A provision for the payment for family planning services on behalf of women who were  
462 Medicaid-eligible for prenatal care and delivery as provided in this section at the time of delivery. Such  
463 family planning services shall begin with delivery and continue for a period of 24 months, if the woman  
464 continues to meet the financial eligibility requirements for a pregnant woman under Medicaid. For the  
465 purposes of this section, family planning services shall not cover payment for abortion services and no  
466 funds shall be used to perform, assist, encourage or make direct referrals for abortions;

467 8. A provision for payment of medical assistance for high-dose chemotherapy and bone marrow  
468 transplants on behalf of individuals over the age of 21 who have been diagnosed with lymphoma, breast  
469 cancer, myeloma, or leukemia and have been determined by the treating health care provider to have a  
470 performance status sufficient to proceed with such high-dose chemotherapy and bone marrow transplant.  
471 Appeals of these cases shall be handled in accordance with the Department's expedited appeals process;

472 9. A provision identifying entities approved by the Board to receive applications and to determine  
473 eligibility for medical assistance, which shall include a requirement that such entities obtain accurate  
474 contact information, including the best available address and telephone number, from each applicant for  
475 medical assistance, to the extent required by federal law and regulations;

476 10. A provision for breast reconstructive surgery following the medically necessary removal of a  
477 breast for any medical reason. Breast reductions shall be covered, if prior authorization has been  
478 obtained, for all medically necessary indications. Such procedures shall be considered noncosmetic;

479 11. A provision for payment of medical assistance for annual pap smears;

480 12. A provision for payment of medical assistance services for prostheses following the medically  
481 necessary complete or partial removal of a breast for any medical reason;

482 13. A provision for payment of medical assistance which provides for payment for 48 hours of  
483 inpatient treatment for a patient following a radical or modified radical mastectomy and 24 hours of  
484 inpatient care following a total mastectomy or a partial mastectomy with lymph node dissection for  
485 treatment of disease or trauma of the breast. Nothing in this subdivision shall be construed as requiring  
486 the provision of inpatient coverage where the attending physician in consultation with the patient  
487 determines that a shorter period of hospital stay is appropriate;

488 14. A requirement that certificates of medical necessity for durable medical equipment and any  
489 supporting verifiable documentation shall be signed, dated, and returned by the physician, physician



assistant, or nurse practitioner and in the durable medical equipment provider's possession within 60 days from the time the ordered durable medical equipment and supplies are first furnished by the durable medical equipment provider;

15. A provision for payment of medical assistance to (i) persons age 50 and over and (ii) persons age 40 and over who are at high risk for prostate cancer, according to the most recent published guidelines of the American Cancer Society, for one PSA test in a 12-month period and digital rectal examinations, all in accordance with American Cancer Society guidelines. For the purpose of this subdivision, "PSA testing" means the analysis of a blood sample to determine the level of prostate specific antigen;

16. A provision for payment of medical assistance for low-dose screening mammograms for determining the presence of occult breast cancer. Such coverage shall make available one screening mammogram to persons age 35 through 39, one such mammogram biennially to persons age 40 through 49, and one such mammogram annually to persons age 50 and over. The term "mammogram" means an X-ray examination of the breast using equipment dedicated specifically for mammography, including but not limited to the X-ray tube, filter, compression device, screens, film and cassettes, with an average radiation exposure of less than one rad mid-breast, two views of each breast;

17. A provision, when in compliance with federal law and regulation and approved by the Centers for Medicare & Medicaid Services (CMS), for payment of medical assistance services delivered to Medicaid-eligible students when such services qualify for reimbursement by the Virginia Medicaid program and may be provided by school divisions;

18. A provision for payment of medical assistance services for liver, heart and lung transplantation procedures for individuals over the age of 21 years when (i) there is no effective alternative medical or surgical therapy available with outcomes that are at least comparable; (ii) the transplant procedure and application of the procedure in treatment of the specific condition have been clearly demonstrated to be medically effective and not experimental or investigational; (iii) prior authorization by the Department of Medical Assistance Services has been obtained; (iv) the patient selection criteria of the specific transplant center where the surgery is proposed to be performed have been used by the transplant team or program to determine the appropriateness of the patient for the procedure; (v) current medical therapy has failed and the patient has failed to respond to appropriate therapeutic management; (vi) the patient is not in an irreversible terminal state; and (vii) the transplant is likely to prolong the patient's life and restore a range of physical and social functioning in the activities of daily living;

19. A provision for payment of medical assistance for colorectal cancer screening, specifically screening with an annual fecal occult blood test, flexible sigmoidoscopy or colonoscopy, or in appropriate circumstances radiologic imaging, in accordance with the most recently published recommendations established by the American College of Gastroenterology, in consultation with the American Cancer Society, for the ages, family histories, and frequencies referenced in such recommendations;

20. A provision for payment of medical assistance for custom ocular prostheses;

21. A provision for payment for medical assistance for infant hearing screenings and all necessary audiological examinations provided pursuant to § 32.1-64.1 using any technology approved by the United States Food and Drug Administration, and as recommended by the national Joint Committee on Infant Hearing in its most current position statement addressing early hearing detection and intervention programs. Such provision shall include payment for medical assistance for follow-up audiological examinations as recommended by a physician, physician assistant, nurse practitioner, or audiologist and performed by a licensed audiologist to confirm the existence or absence of hearing loss;

22. A provision for payment of medical assistance, pursuant to the Breast and Cervical Cancer Prevention and Treatment Act of 2000 (P.L. 106-354), for certain women with breast or cervical cancer when such women (i) have been screened for breast or cervical cancer under the Centers for Disease Control and Prevention (CDC) Breast and Cervical Cancer Early Detection Program established under Title XV of the Public Health Service Act; (ii) need treatment for breast or cervical cancer, including treatment for a precancerous condition of the breast or cervix; (iii) are not otherwise covered under creditable coverage, as defined in § 2701 (c) of the Public Health Service Act; (iv) are not otherwise eligible for medical assistance services under any mandatory categorically needy eligibility group; and (v) have not attained age 65. This provision shall include an expedited eligibility determination for such women;

23. A provision for the coordinated administration, including outreach, enrollment, re-enrollment and services delivery, of medical assistance services provided to medically indigent children pursuant to this chapter, which shall be called Family Access to Medical Insurance Security (FAMIS) Plus and the FAMIS Plan program in § 32.1-351. A single application form shall be used to determine eligibility for both programs;

24. A provision, when authorized by and in compliance with federal law, to establish a public-private

long-term care partnership program between the Commonwealth of Virginia and private insurance companies that shall be established through the filing of an amendment to the state plan for medical assistance services by the Department of Medical Assistance Services. The purpose of the program shall be to reduce Medicaid costs for long-term care by delaying or eliminating dependence on Medicaid for such services through encouraging the purchase of private long-term care insurance policies that have been designated as qualified state long-term care insurance partnerships and may be used as the first source of benefits for the participant's long-term care. Components of the program, including the treatment of assets for Medicaid eligibility and estate recovery, shall be structured in accordance with federal law and applicable federal guidelines; and

25. A provision for the payment of medical assistance for otherwise eligible pregnant women during the first five years of lawful residence in the United States, pursuant to § 214 of the Children's Health Insurance Program Reauthorization Act of 2009 (P.L. 111-3); and

26. *A provision requiring any contract under which providers are reimbursed for medical assistance services on a fee-for-service basis to contain provisions that are substantively identical to the provisions required to be contained in a contract requiring preauthorization as set out in § 38.2-3407.15:2, mutatis mutandis.*

B. In preparing the plan, the Board shall:

1. Work cooperatively with the State Board of Health to ensure that quality patient care is provided and that the health, safety, security, rights and welfare of patients are ensured.

2. Initiate such cost containment or other measures as are set forth in the appropriation act.

3. Make, adopt, promulgate and enforce such regulations as may be necessary to carry out the provisions of this chapter.

4. Examine, before acting on a regulation to be published in the Virginia Register of Regulations pursuant to § 2.2-4007.05, the potential fiscal impact of such regulation on local boards of social services. For regulations with potential fiscal impact, the Board shall share copies of the fiscal impact analysis with local boards of social services prior to submission to the Registrar. The fiscal impact analysis shall include the projected costs/savings to the local boards of social services to implement or comply with such regulation and, where applicable, sources of potential funds to implement or comply with such regulation.

5. Incorporate sanctions and remedies for certified nursing facilities established by state law, in accordance with 42 C.F.R. § 488.400 et seq. "Enforcement of Compliance for Long-Term Care Facilities With Deficiencies."

6. On and after July 1, 2002, require that a prescription benefit card, health insurance benefit card, or other technology that complies with the requirements set forth in § 38.2-3407.4:2 be issued to each recipient of medical assistance services, and shall upon any changes in the required data elements set forth in subsection A of § 38.2-3407.4:2, either reissue the card or provide recipients such corrective information as may be required to electronically process a prescription claim.

C. In order to enable the Commonwealth to continue to receive federal grants or reimbursement for medical assistance or related services, the Board, subject to the approval of the Governor, may adopt, regardless of any other provision of this chapter, such amendments to the state plan for medical assistance services as may be necessary to conform such plan with amendments to the United States Social Security Act or other relevant federal law and their implementing regulations or constructions of these laws and regulations by courts of competent jurisdiction or the United States Secretary of Health and Human Services.

In the event conforming amendments to the state plan for medical assistance services are adopted, the Board shall not be required to comply with the requirements of Article 2 (§ 2.2-4006 et seq.) of Chapter 40 of Title 2.2. However, the Board shall, pursuant to the requirements of § 2.2-4002, (i) notify the Registrar of Regulations that such amendment is necessary to meet the requirements of federal law or regulations or because of the order of any state or federal court, or (ii) certify to the Governor that the regulations are necessitated by an emergency situation. Any such amendments that are in conflict with the Code of Virginia shall only remain in effect until July 1 following adjournment of the next regular session of the General Assembly unless enacted into law.

D. The Director of Medical Assistance Services is authorized to:

1. Administer such state plan and receive and expend federal funds therefor in accordance with applicable federal and state laws and regulations; and enter into all contracts necessary or incidental to the performance of the Department's duties and the execution of its powers as provided by law.

2. Enter into agreements and contracts with medical care facilities, physicians, dentists and other health care providers where necessary to carry out the provisions of such state plan. Any such agreement or contract shall terminate upon conviction of the provider of a felony. In the event such conviction is reversed upon appeal, the provider may apply to the Director of Medical Assistance Services for a new agreement or contract. Such provider may also apply to the Director for reconsideration of the agreement or contract termination if the conviction is not appealed, or if it is not reversed upon appeal.

3. Refuse to enter into or renew an agreement or contract, or elect to terminate an existing agreement or contract, with any provider who has been convicted of or otherwise pled guilty to a felony, or pursuant to Subparts A, B, and C of 42 C.F.R. Part 1002, and upon notice of such action to the provider as required by 42 C.F.R. § 1002.212.

4. Refuse to enter into or renew an agreement or contract, or elect to terminate an existing agreement or contract, with a provider who is or has been a principal in a professional or other corporation when such corporation has been convicted of or otherwise pled guilty to any violation of § 32.1-314, 32.1-315, 32.1-316, or 32.1-317, or any other felony or has been excluded from participation in any federal program pursuant to 42 C.F.R. Part 1002.

5. Terminate or suspend a provider agreement with a home care organization pursuant to subsection E of § 32.1-162.13.

6. (Effective January 1, 2015; Expires January 1, 2020) Provide payments or transfers pursuant to § 457 of the Internal Revenue Code to the deferred compensation plan described in § 51.1-602 on behalf of an individual who is a dentist or an oral and maxillofacial surgeon providing services as an independent contractor pursuant to a Medicaid agreement or contract under this section. Notwithstanding the provisions of § 51.1-600, an "employee" for purposes of Chapter 6 (§ 51.1-600 et seq.) of Title 51.1 shall include an independent contractor as described in this subdivision.

For the purposes of this subsection, "provider" may refer to an individual or an entity.

E. In any case in which a Medicaid agreement or contract is terminated or denied to a provider pursuant to subsection D, the provider shall be entitled to appeal the decision pursuant to 42 C.F.R. § 1002.213 and to a post-determination or post-denial hearing in accordance with the Administrative Process Act (§ 2.2-4000 et seq.). All such requests shall be in writing and be received within 15 days of the date of receipt of the notice.

The Director may consider aggravating and mitigating factors including the nature and extent of any adverse impact the agreement or contract denial or termination may have on the medical care provided to Virginia Medicaid recipients. In cases in which an agreement or contract is terminated pursuant to subsection D, the Director may determine the period of exclusion and may consider aggravating and mitigating factors to lengthen or shorten the period of exclusion, and may reinstate the provider pursuant to 42 C.F.R. § 1002.215.

F. When the services provided for by such plan are services which a marriage and family therapist, clinical psychologist, clinical social worker, professional counselor, or clinical nurse specialist is licensed to render in Virginia, the Director shall contract with any duly licensed marriage and family therapist, duly licensed clinical psychologist, licensed clinical social worker, licensed professional counselor or licensed clinical nurse specialist who makes application to be a provider of such services, and thereafter shall pay for covered services as provided in the state plan. The Board shall promulgate regulations which reimburse licensed marriage and family therapists, licensed clinical psychologists, licensed clinical social workers, licensed professional counselors and licensed clinical nurse specialists at rates based upon reasonable criteria, including the professional credentials required for licensure.

G. The Board shall prepare and submit to the Secretary of the United States Department of Health and Human Services such amendments to the state plan for medical assistance services as may be permitted by federal law to establish a program of family assistance whereby children over the age of 18 years shall make reasonable contributions, as determined by regulations of the Board, toward the cost of providing medical assistance under the plan to their parents.

H. The Department of Medical Assistance Services shall:

1. Include in its provider networks and all of its health maintenance organization contracts a provision for the payment of medical assistance on behalf of individuals up to the age of 21 who have special needs and who are Medicaid eligible, including individuals who have been victims of child abuse and neglect, for medically necessary assessment and treatment services, when such services are delivered by a provider which specializes solely in the diagnosis and treatment of child abuse and neglect, or a provider with comparable expertise, as determined by the Director.

2. Amend the Medallion II waiver and its implementing regulations to develop and implement an exception, with procedural requirements, to mandatory enrollment for certain children between birth and age three certified by the Department of Behavioral Health and Developmental Services as eligible for services pursuant to Part C of the Individuals with Disabilities Education Act (20 U.S.C. § 1471 et seq.).

3. Utilize, to the extent practicable, electronic funds transfer technology for reimbursement to contractors and enrolled providers for the provision of health care services under Medicaid and the Family Access to Medical Insurance Security Plan established under § 32.1-351.

I. The Director is authorized to negotiate and enter into agreements for services rendered to eligible recipients with special needs. The Board shall promulgate regulations regarding these special needs patients, to include persons with AIDS, ventilator-dependent patients, and other recipients with special needs as defined by the Board.

674 J. Except as provided in subdivision A 1 of § 2.2-4345, the provisions of the Virginia Public  
675 Procurement Act (§ 2.2-4300 et seq.) shall not apply to the activities of the Director authorized by  
676 subsection I of this section. Agreements made pursuant to this subsection shall comply with federal law  
677 and regulation.

678 **§ 32.1-330.2. Medicaid managed care programs; program information documents; plain**  
679 **language required.**

680 A. Whenever medical assistance services pursuant to this chapter are furnished through managed care  
681 programs, the Board of Medical Assistance Services shall require that all program information  
682 documents furnished recipients covered thereunder shall be written in nontechnical, readily  
683 understandable language, using words of common, everyday usage.

684 B. Each sponsor or administrator of any such managed care program shall test the readability of its  
685 program information documents by use of the Flesch Readability Formula, as set forth in Rudolf Flesch,  
686 The Art of Readable Writing (1949, as revised 1974), and no program information document shall be  
687 used unless it achieves a Flesch total readability score of forty or more. The requirements of this  
688 subsection shall not apply to language which is mandated by federal or state laws, regulations or  
689 agencies.

690 C. All program information documents within the scope of this section, and all amendments thereto,  
691 shall be filed with the Department of Medical Assistance Services in advance of their use and  
692 distribution, accompanied by certificates setting forth the Flesch scores and certifying compliance with  
693 the requirements of this section. Any program information document which is exempt from the  
694 requirements of subsection B shall be accompanied by a documentation of the federal or state law,  
695 regulation or agency mandate that authorizes the exemption.

696 D. For the purpose of this section, the term "program information documents" means all forms,  
697 brochures, handbooks or other documentation (i) provided recipients covered under Medicaid managed  
698 care programs, and (ii) describing the programs' medical care coverages and the rights and  
699 responsibilities of recipients covered thereunder. Further, the term "recipient" shall include potential  
700 recipients and recipients.

701 E. *Whenever medical assistance services pursuant to this chapter are furnished through managed*  
702 *care programs, the Board of Medical Assistance Services shall require that the managed care contract*  
703 *contains provisions that are substantively identical to the provisions required to be contained in a*  
704 *contract requiring preauthorization as set out in § 38.2-3407.15:2, mutatis mutandis.*

705 **§ 38.2-3407.15:2. Carrier contracts; required provisions regarding preauthorization.**

706 A. As used in this section, unless the context requires a different meaning:

707 "Carrier" has the same meaning ascribed thereto in subsection A of § 38.2-3407.15.

708 "Chronic disease management drug" means any drug used to treat a permanent or recurring medical  
709 condition.

710 "Common preauthorization form" means the form developed by the Commission pursuant to  
711 subsection E for use in preauthorization.

712 "Contract requiring preauthorization" means any (i) contract between a carrier and its intermediary  
713 pursuant to which the intermediary has the right or obligation to require preauthorization for a drug  
714 benefit, (ii) provider contract between a carrier and a participating health care provider or its  
715 contracting agent pursuant to which the carrier has the right or obligation to require preauthorization  
716 for a drug benefit, or (iii) contract between a carrier and an individual or group health benefit plan  
717 policyholder pursuant to which the carrier has the right or obligation to require preauthorization for a  
718 drug benefit.

719 "Mental health drug" means any drug used to treat an individual's psychological or emotional  
720 health.

721 "Preauthorization" means the mechanism for approval by a carrier before certain drug benefits may  
722 be provided.

723 "Provider contract" has the same meaning ascribed thereto in subsection A of § 38.2-3407.15.

724 "Step therapy restriction" means a covered benefit restriction placed on certain drug benefits by a  
725 carrier or its intermediary requiring the use of alternative therapies prior to approval of a drug benefit  
726 subject to preauthorization.

727 "Supplementation" means an electronic request communicated by the carrier or its intermediary to  
728 the provider for additional information necessary to approve or deny a preauthorization or waiver  
729 request.

730 B. Any contract requiring preauthorization shall contain specific provisions:

731 1. Providing for preauthorization by means of a common preauthorization form, which  
732 preauthorization shall be valid for a period of not less than one year;

733 2. Providing for the electronic submission of preauthorization in a manner that is compatible with  
734 e-prescribing systems, electronic health records, and health information exchange platforms;

735 3. Waiving preauthorization requirements for chronic disease management drug benefits upon written

certification by the prescriber or his agent that a patient (i) is medically stable on the prescribed drug, (ii) has changed coverage by or between carriers and is medically stable on the prescribed drug, or (iii) has completed prior step therapy requirements for the same or similar drug;

4. Waiving preauthorization requirements for mental health drug benefits if a prescribed medication has become subject to step therapy restrictions since the patient's last refill in all instances, including when the patient has changed coverage by or between carriers;

5. Providing that (i) unless an initial request for supplementation of a preauthorization request or a waiver request is communicated electronically to the provider within 48 hours of the requested preauthorization or waiver, the requested preauthorization or waiver is deemed approved and (ii) unless a subsequent request for supplementation of a preauthorization request or a waiver request, made after an initial request for supplementation pursuant to clause (i), is communicated electronically to the provider within 24 hours of receipt of initial request for supplementation, the requested preauthorization or waiver is deemed approved;

6. Requiring that, in the absence of a request for supplementation within 48 hours of requested preauthorization or waiver that is communicated as described in subdivision 5, preauthorization requests be either approved or denied within 48 hours of receipt of either the common preauthorization form or the carrier's or its agent's proprietary form;

7. Requiring that if a preauthorization or waiver request is denied as set forth in subdivision 6, the reasons for the denial shall be communicated electronically to the prescriber within the 48-hour period set forth in subdivision 6;

8. Prohibiting preauthorization restrictions for generic drug benefits; and

9. Requiring that information identifying all drug benefits subject to preauthorization and all procedures required for compliance with the preauthorization process be (i) posted on the carrier's website and (ii) kept current.

C. The Commission shall have no jurisdiction to adjudicate individual controversies arising out of this section.

D. This section shall apply with respect to any contract requiring preauthorization that is entered into, amended, extended, or renewed on or after January 1, 2016.

E. The Commission shall develop, in consultation with appropriate health care provider and carrier stakeholders, a common preauthorization form. The form shall (i) contain a check box for the provider to enter patient specific information and (ii) enable the provider to submit a renewal request by marking the form to indicate that there has been no change in the patient's condition since the last request. The Commission shall make the form available not later than January 1, 2016.

F. The provisions of subsection B shall apply to (i) any state plan for medical assistance services that provides for reimbursement to providers on a fee-for-service basis as set forth in subdivision A 26 of § 32.1-325, (ii) any Medicaid managed care health program as set forth in subsection E of § 32.1-330.2, and (iii) any health insurance plan created for the Commonwealth's employees as set forth in subdivision B 23 of § 2.2-2818.