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HOUSE BILL NO. 1497

Offered January 24, 2000

A BILL to amend and reenact § 2.1-20.1 of the Code of Virginia and to amend the Code of Virginia by adding a section numbered 38.2-3407.13:1, relating to accident and sickness insurance policies; coordination of benefits.

Patron—Devolites

Referred to Committee on Corporations, Insurance and Banking

Be it enacted by the General Assembly of Virginia:

1. That § 2.1-20.1 of the Code of Virginia is amended and reenacted, and that the Code of Virginia is amended by adding a section numbered 38.2-3407.13:1 as follows:

§ 2.1-20.1. Health and related insurance for state employees.

A. 1. The Governor shall establish a plan 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 Department of Personnel and Training shall administer this section. The plan chosen shall provide means whereby coverage for the families or dependents of state employees may be purchased. The Commonwealth may pay all or a portion of the cost thereof, and for such portion as the Commonwealth does not pay, the employee may purchase the coverage by paying the additional cost over the cost of coverage for an employee.

2. Such contribution shall be financed through appropriations provided by law.

B. The plan shall:

1. a. 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 thirty-five through thirty-nine, one such mammogram biennially to persons age forty through forty-nine, and one such mammogram annually to persons age fifty and over and may be limited to a benefit of fifty dollars 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.

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

(1) The mammogram must 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 physician, (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 must be sent or delivered to the health care practitioner who ordered it;

(2) 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

(3) 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 the treatment of breast cancer by dose-intensive chemotherapy with autologous bone marrow transplants or stem cell support when performed at a clinical program authorized to provide such therapies as a part of clinical trials sponsored by the National Cancer Institute. For persons previously covered under the plan, there shall be no denial of coverage due to the existence of a preexisting condition.

3. Include coverage for postpartum services providing inpatient care and a home visit or visits which 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.

4. a. Include an appeals process for resolution of written complaints concerning denials or partial denials of claims that shall provide reasonable procedures for resolution of such written complaints and

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60 shall be published and disseminated to all covered state employees. Such appeals process shall include a
61 separate expedited emergency appeals procedure which shall provide resolution within one business day
62 of receipt of a complaint concerning situations requiring immediate medical care. For appeals involving
63 adverse decisions as defined in § 32.1-137.7, the Department shall contract with one or more impartial
64 health entities to review such decisions. Impartial health entities may include medical peer review
65 organizations and independent utilization review companies. The Department shall adopt regulations to
66 assure that the impartial health entity conducting the reviews has adequate standards, credentials and
67 experience for such review. The impartial health entity shall examine the final denial of claims to
68 determine whether the decision is objective, clinically valid, and compatible with established principles
69 of health care. The decision of the impartial health entity shall (i) be in writing, (ii) contain findings of
70 fact as to 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 b. Prior to assigning an appeal to an impartial health entity, the Department shall verify that the
73 impartial health entity conducting the review of a denial of claims has no relationship or association
74 with (i) the covered employee, (ii) the treating health care provider, or any of its employees or affiliates,
75 (iii) the medical care facility at which the covered service would be provided, or any of its employees or
76 affiliates, or (iv) the development or manufacture of the drug, device, procedure or other therapy which
77 is the subject of the final denial of a claim. The impartial health entity shall not be a subsidiary of, nor
78 owned or controlled by, a health plan, a trade association of health plans, or a professional association
79 of health care providers. There shall be no liability on the part of and no cause of action shall arise
80 against any officer or employee of an impartial health entity for any actions taken or not taken or
81 statements made by such officer or employee in good faith in the performance of his powers and duties.

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

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

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

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

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

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

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

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

119 12. Include coverage providing a minimum stay in the hospital of not less than forty-eight hours for
120 a patient following a radical or modified radical mastectomy and twenty-four hours of inpatient care
121 following a total mastectomy or a partial mastectomy with lymph node dissection for treatment of breast

122 cancer. Nothing in this subdivision shall be construed as requiring the provision of inpatient coverage
123 where the attending physician in consultation with the patient determines that a shorter period of
124 hospital stay is appropriate.

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

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

151 15. a. Include provisions allowing employees to continue receiving health care services for a period
152 of up to ninety days from the date of the primary care physician's notice of termination from any of the
153 plan's provider panels.

154 b. The plan shall notify any provider at least ninety days prior to the date of termination of the
155 provider, except when the provider is terminated for cause.

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

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

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

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

175 16. a. Include coverage for patient costs incurred during participation in clinical trials for treatment
176 studies on cancer, including ovarian cancer trials.

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

181 c. 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 d. 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 e. The treatment described in clause d shall be provided by a clinical trial approved by:

204 (1) The National Cancer Institute;

205 (2) An NCI cooperative group or an NCI center;

206 (3) The FDA in the form of an investigational new drug application;

207 (4) The federal Department of Veterans Affairs; or

208 (5) 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 f. The facility and personnel providing the treatment shall be capable of doing so by virtue of their
211 experience, training, and expertise.

212 g. Coverage under this section 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 17. Include coverage providing a minimum stay in the hospital of not less than twenty-three hours
220 for a covered employee following a laparoscopy-assisted vaginal hysterectomy and forty-eight hours for
221 a covered employee following a vaginal hysterectomy, as outlined in Milliman & Robertson's nationally
222 recognized guidelines. Nothing in this subdivision shall be construed as requiring the provision of the
223 total hours referenced when the attending physician, in consultation with the covered employee,
224 determines that a shorter hospital stay is appropriate.

225 18. (Effective until July 1, 2004) a. Include coverage for biologically based mental illness.

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

237 d. 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 e. In no case, however, shall coverage for mental disorders provided pursuant to this section be
243 diminished or reduced below the coverage in effect for such disorders on January 1, 1999.

244 C. Claims incurred during a fiscal year but not reported during that fiscal year shall be paid from

245 such funds as shall be appropriated by law. Appropriations, premiums and other payments shall be
 246 deposited in the employee health insurance fund, from which payments for claims, premiums, cost
 247 containment programs and administrative expenses shall be withdrawn from time to time. The funds of
 248 the health insurance fund shall be deemed separate and independent trust funds, shall be segregated from
 249 all other funds of the Commonwealth, and shall be invested and administered solely in the interests of
 250 the employees and beneficiaries thereof. Neither the General Assembly nor any public officer, employee,
 251 or agency shall use or authorize the use of such trust funds for any purpose other than as provided in
 252 law for benefits, refunds, and administrative expenses, including but not limited to legislative oversight
 253 of the health insurance fund.

254 D. For the purposes of this section:

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

261 "Standard reference compendia" means the American Medical Association Drug Evaluations, the
 262 American Hospital Formulary Service Drug Information, or the United States Pharmacopoeia Dispensing
 263 Information.

264 "State employee" means state employee as defined in § 51.1-124.3, employee as defined in
 265 § 51.1-201, the Governor, Lieutenant Governor and Attorney General, judge as defined in § 51.1-301
 266 and judges, clerks and deputy clerks of regional juvenile and domestic relations, county juvenile and
 267 domestic relations, and district courts of the Commonwealth, interns and residents employed by the
 268 School of Medicine and Hospital of the University of Virginia, and interns, residents, and employees of
 269 the Medical College of Virginia Hospitals Authority as provided in § 23-50.16:24.

270 E. Provisions shall be made for retired employees to obtain coverage under the above plan. The
 271 Commonwealth may, but shall not be obligated to, pay all or any portion of the cost thereof.

272 F. Any self-insured group health insurance plan established by the Department of Personnel and
 273 Training which utilizes a network of preferred providers shall not exclude any physician solely on the
 274 basis of a reprimand or censure from the Board of Medicine, so long as the physician otherwise meets
 275 the plan criteria established by the Department.

276 G. The plan established by the Department shall include, in each planning district, at least two health
 277 coverage options, each sponsored by unrelated entities. In each planning district that does not have an
 278 available health coverage alternative, the Department shall voluntarily enter into negotiations at any time
 279 with any health coverage provider who seeks to provide coverage under the plan. This section shall not
 280 apply to any state agency authorized by the Department to establish and administer its own health
 281 insurance coverage plan separate from the plan established by the Department.

282 H. 1. Any self-insured group health insurance plan established by the Department of Personnel that
 283 includes coverage for prescription drugs on an outpatient basis may apply a formulary to the prescription
 284 drug benefits provided by the plan if the formulary is developed, reviewed at least annually, and updated
 285 as necessary in consultation with and with the approval of a pharmacy and therapeutics committee, a
 286 majority of whose members are actively practicing licensed (i) pharmacists, (ii) physicians, and (iii)
 287 other health care providers.

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

294 I. Any plan established by the Department of Personnel and Training requiring preauthorization prior
 295 to rendering medical treatment shall have personnel available to provide authorization at all times when
 296 such preauthorization is required.

297 J. Any plan established by the Department of Personnel and Training shall provide to all covered
 298 employees written notice of any benefit reductions during the contract period at least thirty days before
 299 such reductions become effective.

300 K. No contract between a provider and any plan established by the Department of Personnel and
 301 Training shall include provisions which require a health care provider or health care provider group to
 302 deny covered services that such provider or group knows to be medically necessary and appropriate that
 303 are provided with respect to a covered employee with similar medical conditions.

304 L. 1. The Department of Personnel and Training shall appoint an Ombudsman to promote and protect
 305 the interests of covered employees under any state employee's health plan.

306 2. The Ombudsman shall:
307 a. Assist covered employees in understanding their rights and the processes available to them
308 according to their state health plan.
309 b. Answer inquiries from covered employees by telephone and electronic mail.
310 c. Provide to covered employees information concerning the state health plans.
311 d. Develop information on the types of health plans available, including benefits and complaint
312 procedures and appeals.
313 e. Make available, either separately or through an existing Internet web site utilized by the
314 Department of Personnel and Training, information as set forth in clause d and such additional
315 information as he deems appropriate.
316 f. Maintain data on inquiries received, the types of assistance requested, any actions taken and the
317 disposition of each such matter.
318 g. Upon request, assist covered employees in using the procedures and processes available to them
319 from their health plan, including all appeal procedures. Such assistance may require the review of health
320 care records of a covered employee, which shall be done only with that employee's express written
321 consent. The confidentiality of any such medical records shall be maintained in accordance with the
322 confidentiality and disclosure laws of the Commonwealth.
323 h. Ensure that covered employees have access to the services provided by the Ombudsman and that
324 the covered employees receive timely responses from the Ombudsman or his representatives to the
325 inquiries.
326 i. Report annually on his activities to the standing committees of the General Assembly having
327 jurisdiction over insurance and over health and the Joint Commission on Health Care by December 1 of
328 each year.
329 M. 1. The plan established by the Department of Personnel and Training shall not refuse to accept or
330 make reimbursement pursuant to an assignment of benefits made to a dentist or oral surgeon by a
331 covered employee.
332 2. For purposes of this subsection, "assignment of benefits" means the transfer of dental care
333 coverage reimbursement benefits or other rights under the plan. The assignment of benefits shall not be
334 effective until the covered employee notifies the plan in writing of the assignment.
335 *N. The administrator of any self-insured group health insurance plan established by the Department*
336 *of Personnel and Training that contains a coordination of benefits provision shall provide written*
337 *notification to any covered employee within thirty days after obtaining information indicating that such*
338 *covered employee is covered under another accident and sickness insurance policy, accident and*
339 *sickness subscription contract, or health care plan for health care services. Such written notification*
340 *shall inform the covered employee (i) that multiple coverage exists and (ii) which coverage has primary*
341 *responsibility for the covered expenses of each family member.*
342 *§ 38.2-3407.13:1. Coordination of benefits; notice of priority of coverage.*
343 *Each (i) insurer issuing individual or group accident and sickness insurance policies providing*
344 *hospital, medical and surgical or major medical coverage on an expense-incurred basis, (ii) corporation*
345 *providing individual or group accident and sickness subscription contracts, and (iii) health maintenance*
346 *organization providing a health care plan for health care services, whose policy, contract or plan,*
347 *including any certificate or evidence of coverage issued in connection with any such policy, contract or*
348 *plan, contains a coordination of benefits provision shall provide written notification to the insured,*
349 *subscriber or member within thirty days after obtaining information indicating that such insured,*
350 *subscriber or member is covered under another policy, contract or plan. Such written notification shall*
351 *inform the insured, subscriber or member (i) that multiple coverage exists and (ii) which coverage has*
352 *primary responsibility for the covered expenses of each family member. The provisions of this section*
353 *shall not be construed to abrogate any coordination of benefits provision authorized pursuant to*
354 *subsection B of § 38.2-3405.*