

077607440

HOUSE BILL NO. 2155

Offered January 10, 2007

Prefiled January 9, 2007

A BILL to amend and reenact §§ 32.1-102.1, 32.1-102.1:1, 32.1-102.2, 32.1-102.3, 32.1-102.3:2, 32.1-102.4, 32.1-102.6, 32.1-102.10, 32.1-102.11, 32.1-102.12, and 32.1-122.05 of the Code of Virginia, relating to certificates of public need.

Patron—O'Bannon

Referred to Committee on Health, Welfare and Institutions

Be it enacted by the General Assembly of Virginia:

1. That §§ 32.1-102.1, 32.1-102.1:1, 32.1-102.2, 32.1-102.3, 32.1-102.3:2, 32.1-102.4, 32.1-102.6, 32.1-102.10, 32.1-102.11, 32.1-102.12 and 32.1-122.05 of the Code of Virginia are amended and reenacted as follows:

§ 32.1-102.1. Definitions.

As used in this article, unless the context indicates otherwise:

"Certificate" means a certificate of public need for a project required by this article.

"Clinical health service" means a single diagnostic, therapeutic, rehabilitative, preventive or palliative procedure or a series of such procedures that may be separately identified for billing and accounting purposes.

"Health planning region" means a contiguous geographical area of the Commonwealth with a population base of at least 500,000 persons which is characterized by the availability of multiple levels of medical care services, reasonable travel time for tertiary care, and congruence with planning districts.

"Medical care facility," as used in this title, means any institution, place, building or agency, whether or not licensed or required to be licensed by the Board or the State Mental Health, Mental Retardation and Substance Abuse Services Board, whether operated for profit or nonprofit and whether privately owned or privately operated or owned or operated by a local governmental unit, (i) by or in which health services are furnished, conducted, operated or offered for the prevention, diagnosis or treatment of human disease, pain, injury, deformity or physical condition, whether medical or surgical, of two or more nonrelated mentally or physically sick or injured persons, or for the care of two or more nonrelated persons requiring or receiving medical, surgical or nursing attention or services as acute, chronic, convalescent, aged, physically disabled or crippled or (ii) which is the recipient of reimbursements from third-party health insurance programs or prepaid medical service plans. For purposes of this article, only the following medical care facilities shall be subject to review:

1. General hospitals.

2. Sanitariums.

3. Nursing homes.

4. Intermediate care facilities, except those intermediate care facilities established for the mentally retarded that have no more than 12 beds and are in an area identified as in need of residential services for people with mental retardation in any plan of the Department of Mental Health, Mental Retardation and Substance Abuse Services.

5. Extended care facilities.

6. Mental hospitals.

7. Mental retardation facilities.

8. Psychiatric hospitals and intermediate care facilities established primarily for the medical, psychiatric or psychological treatment and rehabilitation of alcoholics or drug addicts.

9. Specialized centers or clinics or that portion of a physician's office developed for the provision of outpatient or ambulatory surgery, cardiac catheterization, computed tomographic (CT) scanning, ~~gamma knife surgery, lithotripsy, stereotactic radiosurgery,~~ magnetic resonance imaging (MRI), magnetic source imaging (MSI), positron emission tomographic (PET) scanning, radiation therapy, ~~nuclear medicine imaging,~~ except for the purpose of nuclear cardiac imaging, or such other specialty services as may be designated by the Board by regulation.

10. Rehabilitation hospitals.

11. Any facility licensed as a hospital.

The term "medical care facility" shall not include any facility of (i) the Department of Mental Health, Mental Retardation and Substance Abuse Services; (ii) any nonhospital substance abuse residential treatment program operated by or contracted primarily for the use of a community services board under the Department of Mental Health, Mental Retardation and Substance Abuse Services' Comprehensive

INTRODUCED

HB2155

59 Plan; (iii) an intermediate care facility for the mentally retarded that has no more than 12 beds and is in
60 an area identified as in need of residential services for people with mental retardation in any plan of the
61 Department of Mental Health, Mental Retardation and Substance Abuse Services; (iv) a physician's
62 office, except that portion of a physician's office described above in subdivision 9 of the definition of
63 "medical care facility"; or (v) the Woodrow Wilson Rehabilitation Center of the Department of
64 Rehabilitative Services. "Medical care facility" shall also not include that portion of a physician's office
65 dedicated to providing nuclear cardiac imaging.

66 "Project" means:

67 1. Establishment of a medical care facility;

68 2. An increase in the total number of beds or operating rooms in an existing medical care facility;

69 3. Relocation at the same site of 10 beds or 10 percent of the beds, whichever is less, from one
70 existing physical facility to another in any two-year period; however, a hospital shall not be required to
71 obtain a certificate for the use of 10 percent of its beds as nursing home beds as provided in § 32.1-132;

72 43. Introduction into an existing medical care facility of any new nursing home service, such as
73 intermediate care facility services, extended care facility services, or skilled nursing facility services,
74 regardless of the type of medical care facility in which those services are provided;

75 54. Introduction into an existing medical care facility of any new cardiac catheterization, computed
76 tomographic (CT) scanning, gamma knife surgery, lithotripsy, stereotactic radiosurgery, magnetic
77 resonance imaging (MRI), magnetic source imaging (MSI), medical rehabilitation, neonatal special care,
78 obstetrical, open heart surgery, positron emission tomographic (PET) scanning, psychiatric, organ or
79 tissue transplant service, radiation therapy, nuclear medicine imaging, except for the purpose of nuclear
80 cardiac imaging, substance abuse treatment, or such other specialty clinical services as may be
81 designated by the Board by regulation, which the facility has never provided or has not provided in the
82 previous 12 months;

83 65. Conversion of beds in an existing medical care facility to medical rehabilitation beds or
84 psychiatric beds;

85 76. The addition by an existing medical care facility of any medical equipment for the provision of
86 cardiac catheterization, computed tomographic (CT) scanning, gamma knife surgery, lithotripsy,
87 stereotactic radiosurgery, magnetic resonance imaging (MRI), magnetic source imaging (MSI), open
88 heart surgery, positron emission tomographic (PET) scanning, radiation therapy, or other specialized
89 service designated by the Board by regulation valued at \$500,000 or more. The addition by an existing
90 medical care facility of any medical equipment for the provision of cardiac catheterization, computed
91 tomographic (CT) scanning, stereotactic radiosurgery, magnetic resonance imaging (MRI), magnetic
92 source imaging (MSI), open heart surgery, positron emission tomographic (PET) scanning, radiation
93 therapy, or other specialized service designated by the Board by regulation valued at less than
94 \$500,000, or Replacement replacement of existing equipment shall not constitute a project for the
95 purposes of this article and shall not require a certificate of public need; or

96 87. Any capital expenditure of \$5 \$15 million or more, not defined as reviewable in subdivisions 1
97 through 76 of this definition, by or in behalf of a medical care facility. However, capital expenditures
98 between \$1\$5 and \$5 \$15 million not defined as reviewable in subdivisions 1 through 6 of this
99 definition, by or on behalf of a medical facility, shall be registered with the Commissioner pursuant to
100 regulations developed by the Board.

101 "Regional health planning agency" means the regional agency, including the regional health planning
102 board, its staff and any component thereof, designated by the Virginia Health Planning Board to perform
103 the health planning activities set forth in this chapter within a health planning region.

104 "State Medical Facilities Plan" means the planning document adopted by the Board of Health which
105 shall include, but not be limited to, (i) methodologies for projecting need for medical care facility beds
106 and services; (ii) statistical information on the availability of medical care facilities and services; and
107 (iii) procedures, criteria and standards for review of applications for projects for medical care facilities
108 and services.

109 "Virginia Health Planning Board" means the statewide health planning body established pursuant to
110 § 32.1-122.02 which serves as the analytical and technical resource to the Secretary of Health and
111 Human Resources in matters requiring health analysis and planning.

112 § 32.1-102.1:1. Equipment registration required.

113 Within thirty calendar days of becoming contractually obligated to acquire any medical equipment for
114 the provision of cardiac catheterization, computed tomographic (CT) scanning, gamma knife surgery,
115 lithotripsy, stereotactic radiosurgery, magnetic resonance imaging (MRI), magnetic source imaging
116 (MSI), open heart surgery, positron emission tomographic (PET) scanning, radiation therapy, or other
117 specialized service designated by the Board by regulation regardless of the value of such equipment, any
118 person shall register such purchase with the Commissioner and the appropriate health planning agency.

119 § 32.1-102.2. Regulations.

120 A. The Board shall promulgate regulations which are consistent with this article and:

1. Shall establish concise procedures for the prompt review of applications for certificates consistent with the provisions of this article which may include a structured batching process which incorporates, but is not limited to, authorization for the Commissioner to request proposals for certain projects. In any structured batching process established by the Board, applications, combined or separate, for computed tomographic (CT) scanning, magnetic resonance imaging (MRI), positron emission tomographic (PET) scanning, radiation therapy ~~or~~ and nuclear imaging shall be considered in the radiation therapy batch. A single application may be filed for a combination of (i) radiation therapy and (ii) any or all of the computed tomographic (CT) scanning, magnetic resonance imaging (MRI), positron emission tomographic (PET) scanning, and nuclear medicine imaging;

2. May classify projects and may eliminate one or more or all of the procedures prescribed in § 32.1-102.6 for different classifications;

3. May provide for exempting from the requirement of a certificate projects determined by the Commissioner, upon application for exemption, to be subject to the economic forces of a competitive market or to have no discernible impact on the cost or quality of health services;

4. Shall establish specific criteria for determining need in rural areas, giving due consideration to distinct and unique geographic, socioeconomic, cultural, transportation, and other barriers to access to care in such areas and providing for weighted calculations of need based on the barriers to health care access in such rural areas in lieu of the determinations of need used for the particular proposed project within the relevant health systems area as a whole; and

5. May establish, on or after July 1, 1999, a schedule of fees for applications for certificates to be applied to expenses for the administration and operation of the certificate of public need program. Such fees shall not be less than \$1,000 nor exceed the lesser of one percent of the proposed expenditure for the project or \$20,000. Until such time as the Board shall establish a schedule of fees, such fees shall be one percent of the proposed expenditure for the project; however, such fees shall not be less than \$1,000 or more than \$20,000.

B. The Board shall promulgate regulations providing for time limitations for schedules for completion and limitations on the exceeding of the maximum capital expenditure amount for all reviewable projects. The Commissioner shall not approve any such extension or excess unless it complies with the Board's regulations.

C. The Board shall also promulgate regulations authorizing the Commissioner to condition approval of a certificate on the agreement of the applicant to provide a level of care at a reduced rate to indigents or accept patients requiring specialized care. In addition, the Board's licensure regulations shall direct the Commissioner to condition the issuing or renewing of any license for any applicant whose certificate was approved upon such condition on whether such applicant has complied with any agreement to provide a level of care at a reduced rate to indigents or accept patients requiring specialized care.

§ 32.1-102.3. Certificate required; criteria for determining need.

A. No person shall commence any project without first obtaining a certificate issued by the Commissioner. No certificate may be issued unless the Commissioner has determined that a public need for the project has been demonstrated. If it is determined that a public need exists for only a portion of a project, a certificate may be issued for that portion and any appeal may be limited to the part of the decision with which the appellant disagrees without affecting the remainder of the decision. Any decision to issue or approve the issuance of a certificate shall be consistent with the most recent applicable provisions of the State Medical Facilities Plan; however, if the Commissioner finds, upon presentation of appropriate evidence, that the provisions of such plan are not relevant to a rural locality's needs, inaccurate, outdated, inadequate or otherwise inapplicable, the Commissioner, consistent with such finding, may issue or approve the issuance of a certificate and shall initiate procedures to make appropriate amendments to such plan.

B. In determining whether a public need for a project has been demonstrated, the Commissioner shall consider:

1. ~~The recommendation and the reasons therefor of the appropriate health planning agency.~~

21. The relationship of the project to the applicable health plans of the Board ~~and the health planning agency.~~

32. The relationship of the project to the long-range development plan, if any, of the person applying for a certificate.

43. The need that the population served or to be served by the project has for the project, including, but not limited to, the needs of rural populations in areas having distinct and unique geographic, socioeconomic, cultural, transportation, and other barriers to access to care.

54. The extent to which the project will be accessible to all residents of the area proposed to be served and the effects on accessibility of any proposed relocation of an existing service or facility.

65. The area, population, topography, highway facilities and availability of the services to be provided by the project in the particular part of the health service area in which the project is proposed,

182 in particular, the distinct and unique geographic, socioeconomic, cultural, transportation, and other
183 barriers to access to care.

184 76. Less costly or more effective alternate methods of reasonably meeting identified health service
185 needs.

186 87. The immediate and long-term financial feasibility of the project.

187 98. The relationship of the project to the existing health care system of the area in which the project
188 is proposed; however, for projects proposed in rural areas, the relationship of the project to the existing
189 health care services in the specific rural locality shall be considered.

190 109. The availability of resources for the project.

191 110. The organizational relationship of the project to necessary ancillary and support services.

192 121. The relationship of the project to the clinical needs of health professional training programs in
193 the area in which the project is proposed.

194 132. The special needs and circumstances of an applicant for a certificate, such as a medical school,
195 hospital, multidisciplinary clinic, specialty center or regional health service provider, if a substantial
196 portion of the applicant's services or resources or both is provided to individuals not residing in the
197 health service area in which the project is to be located.

198 143. The special needs and circumstances of health maintenance organizations. When considering
199 the special needs and circumstances of health maintenance organizations, the Commissioner may grant a
200 certificate for a project if the Commissioner finds that the project is needed by the enrolled or
201 reasonably anticipated new members of the health maintenance organization or the beds or services to be
202 provided are not available from providers which are not health maintenance organizations or from other
203 health maintenance organizations in a reasonable and cost-effective manner.

204 154. The special needs and circumstances for biomedical and behavioral research projects which are
205 designed to meet a national need and for which local conditions offer special advantages.

206 165. In the case of a construction project, the costs and benefits of the proposed construction.

207 176. The probable impact of the project on the costs of and charges for providing health services by
208 the applicant for a certificate and on the costs and charges to the public for providing health services by
209 other persons in the area.

210 187. Improvements or innovations in the financing and delivery of health services which foster
211 competition and serve to promote quality assurance and cost effectiveness.

212 198. In the case of health services or facilities proposed to be provided, the efficiency and
213 appropriateness of the use of existing services and facilities in the area similar to those proposed,
214 including, in the case of rural localities, any distinct and unique geographic, socioeconomic, cultural,
215 transportation, and other barriers to access to care.

216 209. The need and the availability in the health service area for osteopathic and allopathic services
217 and facilities and the impact on existing and proposed institutional training programs for doctors of
218 osteopathy and medicine at the student, internship, and residency training levels.

219 § 32.1-102.3:2. Certificates of public need; applications to be filed in response to Requests For
220 Applications (RFAs).

221 A. Except for applications for continuing care retirement community nursing home bed projects filed
222 by continuing care providers registered with the State Corporation Commission pursuant to Chapter 49
223 (§ 38.2-4900 et seq.) of Title 38.2 which comply with the requirements established in this section, the
224 Commissioner of Health shall only approve, authorize or accept applications for the issuance of any
225 certificate of public need pursuant to this article for any project which would ~~result in an increase in the~~
226 ~~number of beds in a planning district in which nursing facility beds, establish new or extended care~~
227 ~~services, establish new radiation therapy services or stereotactic radiosurgery services or increase the~~
228 ~~number of radiation therapy services or stereotactic radiosurgery services at an existing medical care~~
229 ~~facility, establish new neonatal special care services, establish new obstetrical services, establish new~~
230 ~~medical rehabilitation services, establish new psychiatric services or increase the number of psychiatric~~
231 ~~beds at an existing psychiatric facility, or establish a new long-term care or acute care hospital in a~~
232 ~~planning district are provided~~ when such applications are filed in response to Requests For Applications
233 (RFAs).

234 B. The Board of Health shall adopt regulations establishing standards for the approval and issuance
235 of Requests for Applications by the Commissioner of Health. The standards shall include, but shall not
236 be limited to, a requirement that determinations of need take into account any limitations on access to
237 existing nursing home beds, *extended care services, radiation therapy services, stereotactic radiosurgery*
238 *services, neonatal special care services, obstetrical services, medical rehabilitation services, psychiatric*
239 *services, or long-term care or acute care hospitals* in the planning districts. The RFAs, which shall be
240 published at least annually, shall be jointly developed by the Department of Health and the Department
241 of Medical Assistance Services and based on analyses of the need, or lack thereof, for increases in the
242 nursing home ~~bed~~ *beds, extended care services, radiation therapy services, stereotactic radiosurgery*
243 *services, neonatal special care services, obstetrical services, medical rehabilitation services, psychiatric*

services, or long-term care or acute care hospital supply in each of the Commonwealth's planning districts in accordance with standards adopted by the Board of Health by regulation. The Commissioner shall only accept for review applications in response to such RFAs which conform with the geographic and bed need determinations of the specific RFA.

C. Sixty days prior to the Commissioner's approval and issuance of any Request For Applications, the Board of Health shall publish the proposed RFA in the Virginia Register for public comment together with an explanation of (i) the regulatory basis for the planning district bed needs set forth in the RFA and (ii) the rationale for the RFA's planning district designations. Any person objecting to the contents of the proposed RFA may notify, within fourteen days of the publication, the Board and the Commissioner of his objection and the objection's regulatory basis. The Commissioner shall prepare, and deliver by registered mail, a written response to each such objection within two weeks of the date of receiving the objection. The objector may file a rebuttal to the Commissioner's response in writing within five days of receiving the Commissioner's response. If objections are received, the Board may, after considering the provisions of the RFA, any objections, the Commissioner's responses, and if filed, any written rebuttals of the Commissioner's responses, hold a public hearing to receive comments on the specific RFA. Prior to making a decision on the Request for Applications, the Commissioner shall consider any recommendations made by the Board.

D. Except for a continuing care retirement community applying for a certificate of public need pursuant to provisions of subsections A, B, and C above, applications for continuing care retirement community nursing home bed projects shall be accepted by the Commissioner of Health only if the following criteria are met: (i) the facility is registered with the State Corporation Commission as a continuing care provider pursuant to Chapter 49 (§ 38.2-4900 et seq.) of Title 38.2, (ii) the number of new nursing home beds requested in the initial application does not exceed the lesser of twenty percent of the continuing care retirement community's total number of beds that are not nursing home beds or sixty beds, (iii) the number of new nursing home beds requested in any subsequent application does not cause the continuing care retirement community's total number of nursing home beds to exceed twenty percent of its total number of beds that are not nursing home beds, and (iv) the continuing care retirement community has established a qualified resident assistance policy.

E. The Commissioner of Health may approve an initial certificate of public need for nursing home beds in a continuing care retirement community not to exceed the lesser of sixty beds or twenty percent of the total number of beds that are not nursing home beds which authorizes an initial one-time, three-year open admission period during which the continuing care retirement community may accept direct admissions into its nursing home beds. The Commissioner of Health may approve a certificate of public need for nursing home beds in a continuing care retirement community in addition to those nursing home beds requested for the initial one-time, three-year open admission period if (i) the number of new nursing home beds requested in any subsequent application does not cause the continuing care retirement community's total number of nursing home beds to exceed twenty percent of its total number of beds that are not nursing beds, (ii) the number of licensed nursing home beds within the continuing care retirement community does not and will not exceed twenty percent of the number of occupied beds that are not nursing beds, and (iii) no open-admission period is allowed for these nursing home beds. Upon the expiration of any initial one-time, three-year open admission period, a continuing care retirement community which has obtained a certificate of public need for a nursing facility project pursuant to subsection D may admit into its nursing home beds (i) a standard contract holder who has been a bona fide resident of the non-nursing home portion of the continuing care retirement community for at least thirty days, or (ii) a person who is a standard contract holder who has lived in the non-nursing home portion of the continuing care retirement community for less than thirty days but who requires nursing home care due to change in health status since admission to the continuing care retirement community, or (iii) a person who is a family member of a standard contract holder residing in a non-nursing home portion of the continuing care retirement community.

F. Any continuing care retirement community applicant for a certificate of public need to increase the number of nursing home beds shall authorize the State Corporation Commission to disclose such information to the Commissioner as may be in the State Corporation Commission's possession concerning such continuing care retirement community in order to allow the Commissioner of Health to enforce the provisions of this section. The State Corporation Commission shall provide the Commissioner with the requested information when so authorized.

G. For the purposes of this section:

"Family member" means spouse, mother, father, son, daughter, brother, sister, aunt, uncle or cousin by blood, marriage or adoption.

"One-time, three-year open admission period" means the three years after the initial licensure of nursing home beds during which the continuing care retirement community may take admissions directly into its nursing home beds without the signing of a standard contract. The facility or a related facility on

305 the same campus shall not be granted any open admissions period for any subsequent application or
306 authorization for nursing home beds.

307 "Qualified resident assistance policy" means a procedure, consistently followed by a facility, pursuant
308 to which the facility endeavors to avoid requiring a resident to leave the facility because of inability to
309 pay regular charges and which complies with the requirements of the Internal Revenue Service for
310 maintenance of status as a tax exempt charitable organization under § 501 (c) (3) of the Internal
311 Revenue Code. This policy shall be (i) generally made known to residents through the resident contract
312 and (ii) supported by reasonable and consistent efforts to promote the availability of funds, either
313 through a special fund, separate foundation or access to other available funds, to assist residents who are
314 unable to pay regular charges in whole or in part.

315 This policy may (i) take into account the sound financial management of the facility, including
316 existing reserves, and the reasonable requirements of lenders and (ii) include requirements that residents
317 seeking such assistance provide all requested financial information and abide by reasonable conditions,
318 including seeking to qualify for other assistance and restrictions on the transfer of assets to third parties.

319 A qualified resident assistance policy shall not constitute the business of insurance as defined in
320 Chapter 1 (§ 38.2-100 et seq.) of Title 38.2.

321 "Standard contract" means a contract requiring the same entrance fee, terms, and conditions as
322 contracts executed with residents of the non-nursing home portion of the facility, if the entrance fee is
323 no less than the amount defined in § 38.2-4900.

324 H. This section shall not be construed to prohibit or prevent a continuing care retirement community
325 from discharging a resident (i) for breach of nonfinancial contract provisions, (ii) if medically
326 appropriate care can no longer be provided to the resident, or (iii) if the resident is a danger to himself
327 or others while in the facility.

328 I. The provisions of subsections D, E, and H of this section shall not affect any certificate of public
329 need issued prior to July 1, 1998; however, any certificate of public need application for additional
330 nursing home beds shall be subject to the provisions of this act.

331 § 32.1-102.4. Conditions of certificates; monitoring; revocation of certificates; reporting.

332 A. A certificate shall be issued with a schedule for the completion of the project and a maximum
333 capital expenditure amount for the project. The schedule may not be extended and the maximum capital
334 expenditure may not be exceeded without the approval of the Commissioner in accordance with the
335 regulations of the Board.

336 B. The Commissioner shall monitor each project for which a certificate is issued to determine its
337 progress and compliance with the schedule and with the maximum capital expenditure. The
338 Commissioner shall also monitor all continuing care retirement communities for which a certificate is
339 issued authorizing the establishment of a nursing home facility or an increase in the number of nursing
340 home beds pursuant to § 32.1-102.3:2 and shall enforce compliance with the conditions for such
341 applications which are required by § 32.1-102.3:2. Any willful violation of a provision of § 32.1-102.3:2
342 or conditions of a certificate of public need granted under the provisions of § 32.1-102.3:2 shall be
343 subject to a civil penalty of up to \$100 per violation per day until the date the Commissioner determines
344 that such facility is in compliance.

345 C. A certificate may be revoked when:

346 1. Substantial and continuing progress towards completion of the project in accordance with the
347 schedule has not been made;

348 2. The maximum capital expenditure amount set for the project is exceeded;

349 3. The applicant has willfully or recklessly misrepresented intentions or facts in obtaining a
350 certificate; or

351 4. A continuing care retirement community applicant has failed to honor the conditions of a
352 certificate allowing the establishment of a nursing home facility or granting an increase in the number of
353 nursing home beds in an existing facility which was approved in accordance with the requirements of
354 § 32.1-102.3:2.

355 D. Further, the Commissioner shall not approve an extension for a schedule for completion of any
356 project or the exceeding of the maximum capital expenditure of any project unless such extension or
357 excess complies with the limitations provided in the regulations promulgated by the Board pursuant to
358 § 32.1-102.2.

359 E. Any person willfully violating the Board's regulations establishing limitations for schedules for
360 completion of any project or limitations on the exceeding of the maximum capital expenditure of any
361 project shall be subject to a civil penalty of up to \$100 per violation per day until the date of
362 completion of the project.

363 F. The Commissioner may condition, pursuant to the regulations of the Board, the approval of a
364 certificate (i) upon the agreement of the applicant to provide a level of care at a reduced rate to
365 indigents or accept patients requiring specialized care or (ii) upon the agreement of the applicant to
366 facilitate the development and operation of primary medical care services in designated medically

underserved areas of the applicant's service area.

Any person willfully refusing, failing, or neglecting to honor such agreement shall be subject to a civil penalty of up to \$100 per violation per day until the date of compliance. *Failure to comply with any condition of a certificate shall be grounds for denial of future applications until such time as the applicant is able to demonstrate full compliance.*

G. Any facility seeking or holding a certificate shall report (i) patient volumes, (ii) gross patient revenues, (iii) net patient revenues, (iv) charity care volume and (v) gross charity care expenditures to the Virginia Department of Health. Failure to report shall (a) render the facility, the facility's parent corporation, the facility's owners, and the parent corporation's owners ineligible to apply for additional certificates until all reporting is made current from the later of the start of the service or January 1, 2007, and (b) cause capacity at nonreporting facilities to be excluded from calculations to determine need.

GH. For the purposes of this section, "completion" means conclusion of construction activities necessary for the substantial performance of the contract.

§ 32.1-102.6. Administrative procedures.

A. At least 30 calendar days before any person is contractually obligated to acquire an existing medical care facility, the cost of which is \$600,000 or more, that person shall notify the Commissioner of the intent, the services to be offered in the facility, the bed capacity in the facility, and the projected impact that the cost of the acquisition will have upon the charges for services to be provided. If clinical services or beds are proposed to be added as a result of the acquisition, the Commissioner may require the proposed new owner to obtain a certificate prior to the acquisition.

B. To obtain a certificate for a project, the applicant shall file a completed application for a certificate with the Department and the appropriate health planning agency. In order to verify the date of the Department's and the appropriate health planning agency's receipt of the application, the applicant shall transmit the document electronically, by certified mail or a delivery service, return receipt requested, or shall deliver the document by hand, with signed receipt to be provided.

Within 10 calendar days of the date on which the document is received, the Department and the appropriate health planning agency shall determine whether the application is complete or not and complete. If the application is complete, the Department shall notify the applicant that the application has been accepted for review; if the application is not complete, the Department shall notify the applicant (i) of the information needed to complete the application, (ii) that the application will not be accepted for the current review cycle, and (iii) of the dates of the next available review cycle. If the application is not determined to be complete within 10 calendar days of the date of submission, the application shall be refiled in the next batch of like projects.

At least 30 calendar days before any person is contractually obligated to acquire an existing medical care facility, the cost of which is \$600,000 or more, that person shall notify the Commissioner and the appropriate health planning agency of the intent, the services to be offered in the facility, the bed capacity in the facility and the projected impact that the cost of the acquisition will have upon the charges for services to be provided. If clinical services or beds are proposed to be added as a result of the acquisition, the Commissioner may require the proposed new owner to obtain a certificate prior to the acquisition.

B. The appropriate health planning agency shall review each completed application for a certificate within 60 calendar days of the day which begins the appropriate batch review cycle as established by the Board by regulation pursuant to subdivision A 1 of § 32.1-102.2, such cycle not to exceed 190 days in duration. The health planning agency shall hold one public hearing on each application in a location in the county or city in which the project is proposed or a contiguous county or city. The health planning agency shall cause notice of the public hearing to be published in a newspaper of general circulation in the county or city where a project is proposed to be located at least nine calendar days prior to the public hearing. Prior to the public hearing, the health planning agency shall notify the local governing bodies in the planning district. The health planning agency shall consider the comments of such governing bodies and all other public comments in making its decision. Such comments shall be part of the record provided to the Department. In no case shall a health planning agency hold more than two meetings on any application, one of which shall be the public hearing conducted by the board of the health planning agency or a subcommittee of the board. The applicant shall be given the opportunity, prior to the vote by the board of the health planning agency or a committee of the agency, if acting for the board, on its recommendation, to respond to any comments made about the project by the health planning agency staff, any information in a staff report, or comments by those voting; however, such opportunity shall not increase the 60-calendar-day period designated herein for the health planning agency's review unless the applicant or applicants request a specific extension of the health planning agency's review period.

The health planning agency shall submit its recommendations on each application and its reasons

therefor to the Department within 10 calendar days after the completion of its 60-calendar-day review or such other period in accordance with the applicant's request for extension.

If the health planning agency has not completed its review within the specified 60 calendar days or such other period in accordance with the applicant's request for extension and submitted its recommendations on the application and the reasons therefor within 10 calendar days after the completion of its review, the Department shall, on the eleventh calendar day after the expiration of the health planning agency's review period, proceed as though the health planning agency has recommended project approval without conditions or revision.

C. After commencement of any public hearing and before a decision is made there shall be no ex parte contacts concerning the subject certificate or its application between (i) any person acting on behalf of the applicant or holder of a certificate or any person opposed to the issuance or in favor of revocation of a certificate of public need and (ii) any person in the Department who has authority to make a determination respecting the issuance or revocation of a certificate of public need, unless the Department has provided advance notice to all parties referred to in (i) of the time and place of such proposed contact.

DC. The Department shall commence the review of each completed application upon the day which begins the appropriate batch review cycle and simultaneously with the review conducted by the health planning agency as established by regulations promulgated pursuant to § 32.1-102.2. Such review shall not exceed 190 calendar days.

A determination whether a public need exists for a project shall be made by the Commissioner within 190 calendar days of the day which begins the appropriate batch cycle.

The 190-calendar-day review period shall begin on the date upon which the application is determined to be complete within the batching process specified in subdivision A 1 of § 32.1-102.2.

If the application is not determined to be complete within 40 calendar days from submission, the application shall be refiled in the next batch for like projects.

D. The Department shall hold one public hearing on each application in a location in the county or city in which the project is proposed or a contiguous county or city. The Department shall cause notice of the public hearing to be published in a newspaper of general circulation in the county or city where a project is proposed to be located at least nine calendar days prior to the public hearing. Prior to the public hearing, the Department shall notify the local governing bodies in the planning district. The Department shall consider the comments of such governing bodies and all other public comments in making its decision. Such comments shall be part of the record provided to the Department. In no case shall the Department hold more than two meetings on any application.

E. The Commissioner shall make determinations in accordance with the provisions of Article 3 (§ 2.2-4018 et seq.) of Chapter 40, of Title 2.2 (§ 2.2-4000 et seq.) except for those parts of the determination process for which timelines and specifications are delineated in subsection EG of this section. Further, if an informal fact-finding conference is determined to be necessary by the Department or is requested by a person seeking good cause standing, the parties to the case shall include only the applicant, any person showing good cause, and any third-party payor providing health care insurance or prepaid coverage to five percent or more of the patients in the applicant's service area, and the relevant health planning agency.

EF. After commencement of any public hearing and before a decision is made there shall be no ex parte contact concerning the subject certificate or its application between (i) any person acting on behalf of the applicant or holder of a certificate or any person opposed to the issuance or in favor of revocation of a certificate of public need and (ii) any person in the Department who has authority to make a determination respecting the issuance or revocation of a certificate of public need, unless the Department has provided advance notice to all parties referred to in clause (i) of the time and place of the proposed contact.

G. Upon entry of each completed application or applications into the appropriate batch review cycle:

1. The Department shall establish, for every application, a date between the eightieth and ninetieth calendar days within the 190-calendar-day review period for holding an informal fact-finding conference, if such conference is necessary. Once scheduled by the Department, the informal fact-finding conference shall be held on the scheduled date unless rescheduled by the presiding adjudication officer upon a showing of special, unavoidable circumstances by the party seeking to reschedule, with the concurrence of all parties.

2. The Department shall review every application at or before the seventy-fifth calendar day within the 190-calendar-day review period to determine whether an informal fact-finding conference is necessary.

3. Any person seeking to be made a party to the case for good cause at an informal fact-finding conference shall notify the Department of his request and the basis therefor on or before the eightieth calendar day following the day which begins the appropriate batch review cycle.

4. In any case in which an informal fact-finding conference is held, a date shall be established for

the closing of the record which shall not be more than 30 calendar days after the date for holding the informal fact-finding conference. deliberations shall be based solely on (i) the material in the record on the sixtieth day of the review cycle, (ii) material submitted prior to the informal fact-finding conference by a party with good cause, and (iii) any informal fact-finding conference testimony made regarding the material in the record. Following the sixtieth day of the review cycle, only (a) the analysis and recommendation of the Division of Certificate of Public Need, (b) the transcript of the informal fact-finding conference, (c) the analysis and recommendation of the adjudication officer, and (d) the Commissioner's decision may be added to the record. The date for the closing of the record shall be not more than 45 calendar days after the date on which the informal fact-finding conference is concluded.

5. In any case in which an informal fact-finding conference is not held, the record shall be closed on the earlier of (i) the date established for holding the informal fact-finding conference or (ii) the date that the Department determines an informal fact-finding conference is not necessary. Only the Commissioner's decision may be added to the record after the date on which the record is closed.

6. The provisions of subsection D of § 2.2-4019 notwithstanding, if a determination whether a public need exists for a project is not made by the Commissioner within 45 calendar days of the closing of the record, the Commissioner shall notify the applicant or applicants and any persons seeking to show good cause, in writing, that the application or the application of each shall be deemed approved 25 calendar days after expiration of such 45-calendar-day period, unless the receipt of recommendations from the person performing the hearing officer functions permits the Commissioner to issue his case decision within that 25-calendar-day period. The validity or timeliness of the aforementioned notice shall not, in any event, prevent, delay or otherwise impact the effectiveness of this subdivision 6 of § 32.1-102.6.

7. In any case when a determination whether a public need exists for a project is not made by the Commissioner within 70 calendar days after the closing of the record, the application shall be deemed to be approved and the certificate shall be granted.

8. If a determination whether a public need exists for a project is not made by the Commissioner within 45 calendar days of the closing of the record, any applicant who is competing in the relevant batch or who has filed an application in response to the relevant Request For Applications issued pursuant to § 32.1-102.3:2 may, prior to the application being deemed approved, petition for immediate injunctive relief pursuant to § 2.2-4030, naming as respondents the Commissioner and all parties to the case. During the pendency of the proceeding, no applications shall be deemed to be approved. In such a proceeding, the provisions of § 2.2-4030 shall apply.

FH. Deemed approvals shall be construed as the Commissioner's case decision on the application pursuant to the Administrative Process Act (§ 2.2-4000 et seq.) and shall be subject to judicial review on appeal as the Commissioner's case decision in accordance with such act.

Any person who has sought to participate in the Department's review of such deemed-to-be-approved application as a person showing good cause who has not received a final determination from the Commissioner concerning such attempt to show good cause shall be deemed to be a person showing good cause for purposes of appeal of the deemed approval of the certificate.

In any appeal of the Commissioner's case decision granting a certificate of public need pursuant to a Request for Applications issued pursuant to § 32.1-102.3:2, the court may require the appellant to file a bond pursuant to § 8.01-676.1, in such sum as shall be fixed by the court for protection of all parties interested in the case decision, conditioned on the payment of all damages and costs incurred in consequence of such appeal.

GI. For purposes of this section, "good cause" shall mean that (i) there is significant relevant information not previously presented at and not available at the time of the public hearing, (ii) there have been significant changes in factors or circumstances relating to the application subsequent to the public hearing, or (iii) there is a substantial material mistake of fact or law in the Department staff's report on the application or in the report submitted by the health planning agency.

HJ. The project review procedures shall provide for separation of the project review manager functions from the hearing officer functions. No person serving in the role of project review manager shall serve as a hearing officer.

IK. The applicants, and only the applicants, shall have the authority to extend any of the time periods specified in this section. If all applicants consent to extending any time period in this section, the Commissioner, with the concurrence of the applicants, shall establish a new schedule for the remaining time periods.

§ 32.1-102.10. Commencing project without certificate grounds for refusing to issue license.

Commencing any project without a certificate required by this article shall constitute grounds for refusing to issue a certificate or license for such project.

§ 32.1-102.11. Application of article.

A. On and after July 1, 1992, every project of an existing or proposed medical care facility, as

defined in § 32.1-102.1, shall be subject to all provisions of this article unless, with respect to such project, the owner or operator of an existing medical care facility or the developer of a proposed medical care facility (i) has, by February 1, 1992, purchased or leased equipment subject to registration pursuant to former § 32.1-102.3:4, (ii) has, by February 1, 1992, initiated construction requiring a capital expenditure exceeding one million dollars, or (iii) has made or contracted to make or otherwise legally obligated to make, during the three years ending February 1, 1992, preliminary expenditures of \$350,000 or more for a formal plan of construction of the specific project, including expenditures for site acquisition, designs, preliminary or working drawings, construction documents, or other items essential to the construction of the specific project.

Any project exempted pursuant to subdivisions (ii) and (iii) of this subsection shall be limited to such construction, services, and equipment as specifically identified in the formal plan of construction which shall have existed and been formally committed to by February 1, 1992. Further, the equipment to be exempted pursuant to subdivisions (ii) and (iii) shall be limited to the number of units and any types of medical equipment, in the case of medical equipment intended to provide any services included in subdivision 65 of the definition of project in § 32.1-102.1, as are specifically identified in such plan and, in the case of all other equipment, such equipment as is appropriate for the construction and services included in such plan.

None of the exemptions provided in this subsection shall be applicable to projects which required a certificate of public need pursuant to this article on January 1, 1992.

B. Any medical care facility or entity claiming to meet one of the conditions set forth in subsection A of this section shall file a completed application for an exemption from the provisions of this article with the Commissioner by August 1, 1992. Forms for such application shall be made available by the Commissioner no later than April 1, 1992. The Commissioner may deny an exemption if the application is not complete on August 1, 1992, and the medical care facility or entity has not filed a completed application within forty-five days after notice of deficiency in the filing of the completed application. After receiving a completed application, the Commissioner shall determine whether the project has met one of the criteria for an exemption and is, therefore, exempt or has not met any of the criteria for an exemption and is, therefore, subject to all provisions of this article and shall notify the medical care facility or entity of his determination within sixty days of the date of filing of the completed application. If it is determined that an exemption exists for only a portion of a project, the Commissioner may approve an exemption for that portion and any appeal may be limited to the part of the decision with which the appellant disagrees without affecting the remainder of the decision. The Commissioner's determination shall be made in accordance with the provisions of the Administrative Process Act (§ 2.2-4000 et seq.), except that parties to the case shall include only those parties specified in § 32.1-102.6.

C. For the purposes of this section:

"Formal plan of construction" means documentary evidence indicating that the facility, the owner or operator of the facility, or the developer of a proposed facility was formally committed to the project by February 1, 1992, and describing the specific project in sufficient detail to reasonably define and confirm the scope of the project including estimated cost, intended location, any clinical health services to be involved and any types of equipment to be purchased. Such documentary evidence shall include designs, preliminary or working drawings, construction documents or other documents which have been used to explicitly define and confirm the scope of the project for the purposes of seeking architectural or construction plans or capital to the extent that such capital was committed or agreed to be provided for such project prior to February 1, 1992.

"Initiated construction" means an owner or operator of an existing facility or the developer of a proposed facility can present evidence for a specific project that (i) a construction contract has been executed; (ii) if applicable, short-term financing has been completed; (iii) if applicable, a commitment for long-term financing has been obtained; and (iv) if the project is for construction of a new facility or expansion of an existing facility, predevelopment site work and building foundations have been completed.

"Leased" means that the owner or operator of an existing medical care facility or the developer of a proposed facility has a legally binding commitment to lease the equipment pursuant to an agreement providing for fixed, periodic payments commencing no later than June 30, 1992, including a lease-purchase agreement in which the owner or operator of the facility or developer has an option to purchase the equipment for less than fair market value upon conclusion of the lease or an installment sale agreement with fixed periodic payments commencing no later than June 30, 1992.

"Purchased" means that the equipment has been acquired by the owner or operator of an existing medical care facility or the developer of a proposed medical care facility, or the owner or operator of the facility or the developer can present evidence of a legal obligation to acquire the equipment in the form of an executed contract or appropriately signed order or requisition and payment has been made in full by June 30, 1992.

§ 32.1-102.12. Report required.

The Commissioner shall annually report to the Governor and the General Assembly on the status of Virginia's certificate of public need program. The report shall be issued by October 1 of each year and shall include, but need not be limited to:

1. A summary of the Commissioner's actions during the previous fiscal year pursuant to this article;
2. A five-year schedule for analysis of all project categories which provides for analysis of at least three project categories per year;
3. An analysis of the appropriateness of continuing the certificate of public need program for at least three project categories in accordance with the five-year schedule for analysis of all project categories;
4. An analysis of the effectiveness of the application review procedures used by the ~~health systems agencies and the Department required by~~ *pursuant to* § 32.1-102.6, which details the review time required during the past year for various project categories, the number of contested or opposed applications and the project categories of these contested or opposed projects, ~~the number of applications upon which the health systems agencies have failed to act in accordance with the timelines of § 32.1-102.6 B, and the number of deemed approvals from the Department because of their failure to comply with the timelines required by~~ § 32.1-102.6 EG, and any other data determined by the Commissioner to be relevant to the efficient operation of the program;
5. An analysis of health care market reform in the Commonwealth and the extent, if any, to which such reform obviates the need for the certificate of public need program;
6. An analysis of the accessibility by the indigent to care provided by the medical care facilities regulated pursuant to this article and the relevance of this article to such access;
7. An analysis of the relevance of this article to the quality of care provided by medical care facilities regulated pursuant to this article; and
8. An analysis of equipment registrations required pursuant to § 32.1-102.1:1, including the type of equipment, whether an addition or replacement, and the equipment costs.

§ 32.1-122.05. Regional health planning agencies; boards; duties and responsibilities.

A. For the purpose of representing the interests of health planning regions and performing health planning activities at the regional level, there are hereby created such regional health planning agencies as may be designated by the Board of Health.

B. Each regional health planning agency shall be governed by a regional health planning board to be composed of not more than thirty residents of the region. The membership of the regional health planning boards shall include, but not be limited to, consumers, providers, a director of a local health department, a director of a local department of social services or welfare, a director of a community services board, a director of an area agency on aging and representatives of health care insurers, local governments, the business community and the academic community. The majority of the members of each regional health planning board shall be consumers. Consumer members shall be appointed in a manner that ensures the equitable geographic and demographic representation of the region. Provider members shall be solicited from professional organizations, service and educational institutions and associations of service providers and health care insurers in a manner that assures equitable representation of provider interest.

The members of the regional health planning boards shall be appointed for no more than two consecutive terms of four years or, when appointed to fill an unexpired term of less than four years, for three consecutive terms consisting of one term of less than four years and two terms of four years. The boards shall not be self-perpetuating. The Board of Health shall establish procedures requiring staggered terms. The composition and the method of appointment of the regional health planning boards shall be established in the regulations of the Board of Health. In addition, the Board of Health shall require, pursuant to regulations, each regional health planning board to report and maintain a record of its membership, including, but not limited to, the names, addresses, dates of appointment, years served, number of consecutive and nonconsecutive terms, and the group represented by each member. These membership reports and records shall be public information and shall be published in accordance with the regulations of the Board.

C. An agreement shall be executed between the Commissioner, in consultation with the Board of Health, and each regional health planning board to delineate the work plan and products to be developed with state funds. Funding for the regional health planning agencies shall be contingent upon meeting these obligations and complying with the Board's regulations.

D. Each regional health planning agency shall assist the Board of Health by: (i) conducting data collection, research and analyses as required by the Board; (ii) preparing reports and studies in consultation and cooperation with the Board; (iii) reviewing and commenting on the components of the State Health Plan; (iv) conducting needs assessments as appropriate and serving as a technical resource to the Board; (v) identifying gaps in services, inappropriate use of services or resources and assessing accessibility of critical services; ~~and (vi) reviewing applications for certificates of public need and~~

674 making recommendations to the Department thereon as provided in § ~~32.1-102.6~~; and (vii) conducting
675 such other functions as directed by the regional health planning board. All regional health planning
676 agencies shall demonstrate and document accountability for state funds through annual budget
677 projections and quarterly expenditure and activity reports that shall be submitted to the Commissioner. A
678 regional health planning agency may designate membership and activities at subarea levels as deemed
679 appropriate by its regional health planning board. Each regional health planning board shall adopt
680 bylaws for its operation and for the election of its chairman and shall maintain and publish a record of
681 its membership and any subarea levels as required by this section and the regulations of the Board of
682 Health.