House _____ **Amendment NO.**____

Offered By

_____ _____

AMEND House Committee Substitute for Senate Substitute for Senate Bill No. 608, Page 1, Section A, Line 2, by inserting after all of said section and line the following:

"96.192. 1. The board of trustees of any hospital authorized under subsection 2 of this section, and established and organized under the provisions of sections 96.150 to 96.229, may invest up to twenty-five percent of the hospital's funds not required for immediate disbursement in obligations or for the operation of the hospital in any United States investment grade fixed income funds or any diversified stock funds, or both.

2. The provisions of this section shall only apply if the hospital:

(1) Receives less than one percent of its annual revenues from municipal, county, or state taxes; and

(2) Receives less than one percent of its annual revenue from appropriated funds from the municipality in which such hospital is located.

167.638. The department of health and senior services shall develop an informational brochure relating to meningococcal disease that states that **[**an immunization**]** immunizations against meningococcal disease **[**is**]** are available. The department shall make the brochure available on its website and shall notify every public institution of higher education in this state of the availability of the brochure. Each public institution of higher education shall provide a copy of the brochure to all students and if the student is under eighteen years of age, to the student's parent or guardian. Such information in the brochure shall include:

(1) The risk factors for and symptoms of meningococcal disease, how it may be diagnosed, and its possible consequences if untreated;

(2) How meningococcal disease is transmitted;

(3) The latest scientific information on meningococcal disease immunization and its effectiveness, including information on all meningococcal vaccines receiving a Category A or B recommendation from the Advisory Committee on Immunization Practices; **[**and**]**

(4) A statement that any questions or concerns regarding immunization against meningococcal disease may be answered by contacting the individuals's health care provider; and

(5) A recommendation that the current student or entering student receive meningococcal vaccines in accordance with current Advisory Committee on Immunization Practices of the Centers for Disease Control and Prevention guidelines.

174.335. 1. Beginning with the 2004-05 school year and for each school year thereafter, every public institution of higher education in this state shall require all students who reside in on-campus housing to have received the meningococcal vaccine not more than five years prior to enrollment and in accordance with the latest recommendations of the Advisory Committee on Immunization Practices of the Centers for Disease Control and Prevention, unless a signed

Standing Action Taken_____ Date _____

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statement of medical or religious exemption is on file with the institution's administration. A student shall be exempted from the immunization requirement of this section upon signed certification by a physician licensed under chapter 334 indicating that either the immunization would seriously endanger the student's health or life or the student has documentation of the disease or laboratory evidence of immunity to the disease. A student shall be exempted from the immunization requirement of this section if he or she objects in writing to the institution's administration that immunization violates his or her religious beliefs.

2. Each public university or college in this state shall maintain records on the meningococcal vaccination status of every student residing in on-campus housing at the university or college.

3. Nothing in this section shall be construed as requiring any institution of higher education to provide or pay for vaccinations against meningococcal disease.

4. For purposes of this section, the term "on-campus housing" shall include, but not be limited to, any fraternity or sorority residence, regardless of whether such residence is privately owned, on or near the campus of a public institution of higher education.""; and

Further amend said bill, Page 2, Section 197.170, Line 53, by inserting after all of said section and line the following:

"197.315. 1. Any person who proposes to develop or offer a new institutional health service within the state must obtain a certificate of need from the committee prior to the time such services are offered.

2. Only those new institutional health services which are found by the committee to be needed shall be granted a certificate of need. Only those new institutional health services which are granted certificates of need shall be offered or developed within the state. No expenditures for new institutional health services in excess of the applicable expenditure minimum shall be made by any person unless a certificate of need has been granted.

3. After October 1, 1980, no state agency charged by statute to license or certify health care facilities shall issue a license to or certify any such facility, or distinct part of such facility, that is developed without obtaining a certificate of need.

4. If any person proposes to develop any new institutional health care service without a certificate of need as required by sections 197.300 to 197.366, the committee shall notify the attorney general, and he shall apply for an injunction or other appropriate legal action in any court of this state against that person.

5. After October 1, 1980, no agency of state government may appropriate or grant funds to or make payment of any funds to any person or health care facility which has not first obtained every certificate of need required pursuant to sections 197.300 to 197.366.

6. A certificate of need shall be issued only for the premises and persons named in the application and is not transferable except by consent of the committee.

7. Project cost increases, due to changes in the project application as approved or due to project change orders, exceeding the initial estimate by more than ten percent shall not be incurred without consent of the committee.

8. Periodic reports to the committee shall be required of any applicant who has been granted a certificate of need until the project has been completed. The committee may order the forfeiture of the certificate of need upon failure of the applicant to file any such report.

9. A certificate of need shall be subject to forfeiture for failure to incur a capital expenditure on any approved project within six months after the date of the order. The applicant may request an extension from the committee of not more than six additional months based upon substantial

expenditure made.

10. Each application for a certificate of need must be accompanied by an application fee. The time of filing commences with the receipt of the application and the application fee. The application fee is one thousand dollars, or one-tenth of one percent of the total cost of the proposed project, whichever is greater. All application fees shall be deposited in the state treasury. Because of the loss of federal funds, the general assembly will appropriate funds to the Missouri health facilities review committee.

11. In determining whether a certificate of need should be granted, no consideration shall be given to the facilities or equipment of any other health care facility located more than a fifteen-mile radius from the applying facility.

12. When a nursing facility shifts from a skilled to an intermediate level of nursing care, it may return to the higher level of care if it meets the licensure requirements, without obtaining a certificate of need.

13. In no event shall a certificate of need be denied because the applicant refuses to provide abortion services or information.

14. A certificate of need shall not be required for the transfer of ownership of an existing and operational health facility in its entirety.

15. A certificate of need may be granted to a facility for an expansion, an addition of services, a new institutional service, or for a new hospital facility which provides for something less than that which was sought in the application.

16. The provisions of this section shall not apply to facilities operated by the state, and appropriation of funds to such facilities by the general assembly shall be deemed in compliance with this section, and such facilities shall be deemed to have received an appropriate certificate of need without payment of any fee or charge. The provisions of this subsection shall not apply to hospitals operated by the state and licensed under chapter 197, except for department of mental health state-operated psychiatric hospitals.

17. Notwithstanding other provisions of this section, a certificate of need may be issued after July 1, 1983, for an intermediate care facility operated exclusively for the intellectually disabled.

18. To assure the safe, appropriate, and cost-effective transfer of new medical technology throughout the state, a certificate of need shall not be required for the purchase and operation of:

(1) Research equipment that is to be used in a clinical trial that has received written approval from a duly constituted institutional review board of an accredited school of medicine or osteopathy located in Missouri to establish its safety and efficacy and does not increase the bed complement of the institution in which the equipment is to be located. After the clinical trial has been completed, a certificate of need must be obtained for continued use in such facility; or

(2) Equipment that is to be used by an academic health center operated by the state in furtherance of its research or teaching missions.

198.054. Each year between October first and March first, all long-term care facilities licensed under this chapter shall assist their health care workers, volunteers, and other employees who have direct contact with residents in obtaining the vaccination for the influenza virus by either offering the vaccination in the facility or providing information as to how they may independently obtain the vaccination, unless contraindicated, in accordance with the latest recommendations of the Centers for Disease Control and Prevention and subject to availability of the vaccine. Facilities are encouraged to document that each health care worker, volunteer, and employee has been offered assistance in receiving a vaccination against the influenza virus and has either accepted or declined."; and

Further amend said bill, Page 4, Section 208.800, Line 3, by inserting after all of said section and line the following:

"338.200. 1. In the event a pharmacist is unable to obtain refill authorization from the prescriber due to death, incapacity, or when the pharmacist is unable to obtain refill authorization from the prescriber, a pharmacist may dispense an emergency supply of medication if:

(1) In the pharmacist's professional judgment, interruption of therapy might reasonably produce undesirable health consequences;

(2) The pharmacy previously dispensed or refilled a prescription from the applicable prescriber for the same patient and medication;

(3) The medication dispensed is not a controlled substance;

(4) The pharmacist informs the patient or the patient's agent either verbally, electronically, or in writing at the time of dispensing that authorization of a prescriber is required for future refills; and

(5) The pharmacist documents the emergency dispensing in the patient's prescription record, as provided by the board by rule.

2. (1) If the pharmacist is unable to obtain refill authorization from the prescriber, the amount dispensed shall be limited to the amount determined by the pharmacist within his or her professional judgment as needed for the emergency period, provided the amount dispensed shall not exceed a seven-day supply.

(2) In the event of prescriber death or incapacity or inability of the prescriber to provide medical services, the amount dispensed shall not exceed a thirty-day supply.

3. Pharmacists or permit holders dispensing an emergency supply pursuant to this section shall promptly notify the prescriber or the prescriber's office of the emergency dispensing, as required by the board by rule.

4. An emergency supply may not be dispensed pursuant to this section if the pharmacist has knowledge that the prescriber has otherwise prohibited or restricted emergency dispensing for the applicable patient.

5. The determination to dispense an emergency supply of medication under this section shall only be made by a pharmacist licensed by the board.

6. The board shall promulgate rules to implement the provisions of this section. Any rule or portion of a rule, as that term is defined in section 536.010, that is created under the authority delegated in this section shall become effective only if it complies with and is subject to all of the provisions of chapter 536 and, if applicable, section 536.028. This section and chapter 536 are nonseverable and if any of the powers vested with the general assembly pursuant to chapter 536 to review, to delay the effective date, or to disapprove and annul a rule are subsequently held unconstitutional, then the grant of rulemaking authority and any rule proposed or adopted after August 28, 2013, shall be invalid and void.

338.202. 1. Notwithstanding any other provision of law to the contrary, unless the prescriber has specified on the prescription that dispensing a prescription for a maintenance medication in an initial amount followed by periodic refills is medically necessary, a pharmacist may exercise his or her professional judgment to dispense varying quantities of maintenance medication per fill up to the total number of dosage units as authorized by the prescriber on the original prescription, including any refills. Dispensing of the maintenance medication based on refills authorized by the prescriber on the prescription shall be limited to no more than a ninety-day supply of the medication, and the maintenance medication shall have been previously prescribed to the patient for at least a three-month period.

2. For the purposes of this section "maintenance medication" is a medication prescribed for

chronic, long-term conditions and is taken on a regular, recurring basis, except that it shall not include controlled substances as defined in section 195.010.

376.379. 1. A health carrier or managed care plan offering a health benefit plan in this state that provides prescription drug coverage shall offer, as part of the plan, medication synchronization services developed by the health carrier or managed care plan that allow for the alignment of refill dates for an enrollee's prescription drugs that are covered benefits.

2. Under its medication synchronization services, a health carrier or managed care plan shall:

(1) Not charge an amount in excess of the otherwise applicable co-payment amount under the health benefit plan for dispensing a prescription drug in a quantity that is less than the prescribed amount if:

(a) The pharmacy dispenses the prescription drug in accordance with the medication synchronization services offered under the health benefit plan; and

(b) A participating provider dispenses the prescription drug; and

(2) Provide a full dispensing fee to the pharmacy that dispenses the prescription drug to the covered person.

3. For purposes of this section, the terms "health carrier", "managed care plan", "health benefit plan", "enrollee", and "participating provider" shall have the same meanings given to such terms under section 376.1350.

376.388. 1. As used in this section, unless the context requires otherwise, the following terms shall mean:

(1) "Contracted pharmacy" or "pharmacy", a pharmacy located in Missouri participating in the network of a pharmacy benefits manager through a direct or indirect contract;

(2) "Health carrier", an entity subject to the insurance laws and regulations of this state that contracts or offers to contract to provide, deliver, arrange for, pay for, or reimburse any of the costs of health care services, including a sickness and accident insurance company, a health maintenance organization, a nonprofit hospital and health service corporation, or any other entity providing a plan of health insurance, health benefits, or health services, except that such plan shall not include any coverage pursuant to a liability insurance policy, workers' compensation insurance policy, or medical payments insurance issued as a supplement to a liability policy;

(3) "Maximum allowable cost", the per unit amount that a pharmacy benefits manager reimburses a pharmacist for a prescription drug, excluding a dispensing or professional fee;

(4) "Maximum allowable cost list" or "MAC list", a listing of drug products that meet the standard described in this section;

(5) "Pharmacy", as such term is defined in chapter 338;

(6) "Pharmacy benefits manager", an entity that contracts with pharmacies on behalf of health carriers or any health plan sponsored by the state or a political subdivision of the state.

2. Upon each contract execution or renewal between a pharmacy benefits manager and a pharmacy or between a pharmacy benefits manager and a pharmacy's contracting representative or agent, such as a pharmacy services administrative organization, a pharmacy benefits manager shall, with respect to such contract or renewal:

(1) Include in such contract or renewal the sources utilized to determine maximum allowable cost and update such pricing information at least every seven days; and

(2) Maintain a procedure to eliminate products from the maximum allowable cost list of drugs subject to such pricing or modify maximum allowable cost pricing at least every seven days, if such drugs do not meet the standards and requirements of this section, in order to remain consistent with pricing changes in the marketplace.

3. A pharmacy benefits manager shall reimburse pharmacies for drugs subject to maximum

allowable cost pricing that has been updated to reflect market pricing at least every seven days as set forth under subdivision (1) of subsection 2 of this section.

4. A pharmacy benefits manager shall not place a drug on a maximum allowable cost list unless there are at least two therapeutically equivalent multisource generic drugs, or at least one generic drug available from at least one manufacturer, generally available for purchase by network pharmacies from national or regional wholesalers.

5. All contracts between a pharmacy benefits manager and a contracted pharmacy or between a pharmacy benefits manager and a pharmacy's contracting representative or agent, such as a pharmacy services administrative organization, shall include a process to internally appeal, investigate, and resolve disputes regarding maximum allowable cost pricing. The process shall include the following:

(1) The right to appeal shall be limited to fourteen calendar days following the reimbursement of the initial claim; and

(2) A requirement that the pharmacy benefits manager shall respond to an appeal described in this subsection no later than fourteen calendar days after the date the appeal was received by such pharmacy benefits manager.

6. For appeals that are denied, the pharmacy benefits manager shall provide the reason for the denial and identify the national drug code of a drug product that may be purchased by contracted pharmacies at a price at or below the maximum allowable cost and, when applicable, may be substituted lawfully.

7. If the appeal is successful, the pharmacy benefits manager shall:

(1) Adjust the maximum allowable cost price that is the subject of the appeal effective on the day after the date the appeal is decided;

(2) Apply the adjusted maximum allowable cost price to all similarly situated pharmacies as determined by the pharmacy benefits manager; and

(3) Allow the pharmacy that succeeded in the appeal to reverse and rebill the pharmacy benefits claim giving rise to the appeal.

8. Appeals shall be upheld if:

(1) The pharmacy being reimbursed for the drug subject to the maximum allowable cost pricing in question was not reimbursed as required under subsection 3 of this section; or

(2) The drug subject to the maximum allowable cost pricing in question does not meet the requirements set forth under subsection 4 of this section.

376.1237. 1. Each health carrier or health benefit plan that offers or issues health benefit plans which are delivered, issued for delivery, continued, or renewed in this state on or after January 1, 2014, and that provides coverage for prescription eye drops shall provide coverage for the refilling of an eye drop prescription prior to the last day of the prescribed dosage period without regard to a coverage restriction for early refill of prescription renewals as long as the prescribing health care provider authorizes such early refill, and the health carrier or the health benefit plan is notified.

2. For the purposes of this section, health carrier and health benefit plan shall have the same meaning as defined in section 376.1350.

3. The coverage required by this section shall not be subject to any greater deductible or co-payment than other similar health care services provided by the health benefit plan.

4. The provisions of this section shall not apply to a supplemental insurance policy, including a life care contract, accident-only policy, specified disease policy, hospital policy providing a fixed daily benefit only, Medicare supplement policy, long-term care policy, short-term major medical policies of six months' or less duration, or any other supplemental policy as determined by the director of the department of insurance, financial institutions and professional

registration.

5. The provisions of this section shall terminate on January 1, [2017] 2020.

Section B. Because immediate action is necessary to preserve access to quality health care facilities for the citizens of Missouri, the repeal and reenactment of section 197.315 of section A of this act is deemed necessary for the immediate preservation of the public health, welfare, peace, and safety, and is hereby declared to be an emergency act within the meaning of the constitution, and the repeal and reenactment of section 197.315 of section A of this act shall be in full force and effect upon its passage and approval."; and

Further amend said bill by amending the title, enacting clause, and intersectional references accordingly.