Regulatory Analysis Form

(Completed by Promulgating Agency)

(All Comments submitted on this regulation will appear on IRRC's website)

INDEPENDENT REGULATORY REVIEW COMMISSION

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(1) Agency: **Department of State, Bureau of Professional and Occupational Affairs, State Board of Pharmacy**

(2) Agency Number: **16A**

Identification Number: **5424**

IRRC Number: 2963

(3) PA Code Cite: **49 Pa. Code § 27.26**

(4) Short Title: **Pharmacy Internship**

(5) Agency Contacts (List Telephone Number and Email Address):

Primary Contact: **Kerry E. Maloney, Board Counsel, State Board of Pharmacy, phone: (717) 783-7200; fax: (717) 787-0251; email: kmaloney@pa.gov**
Secondary Contact: **Cynthia K. Montgomery, Regulatory Counsel, Department of State, phone: (717) 783-7200; fax (717) 787-0251; email: cymontgome@pa.gov**

(6) Type of Rulemaking (check applicable box):

X PROPOSED REGULATION
☐ Final Regulation
☐ Final Omitted Regulation
☐ Emergency Certification Regulation;
☐ Certification by the Governor
☐ Certification by the Attorney General

(7) Briefly explain the regulation in clear and nontechnical language. (100 words or less)

The proposed rulemaking would amend the Board's regulations at § 27.26 (relating to pharmacy internship). Its main objectives are: (1) to permit pharmacy interns to complete more of the required internship through an internship through a school of pharmacy while a student rather than in a pharmacy, (2) to permit (with Board approval) a non-traditional internship (other than in a pharmacy or through a school), and (3) to provide that a pharmacy internship registration will automatically terminate if the intern permanently leaves pharmacy school prior to graduation. Additionally, the rulemaking would revise the regulation to use current terms of art and to reflect Board practice.

(8) State the statutory authority for the regulation. Include specific statutory citation.

This rulemaking is authorized by sections 3(c) and 6(k)(9) of the Pharmacy Act (act) (63 P.S. §§ 390-3(c) and 390-6(k)(9)).

(9) Is the regulation mandated by any federal or state law or court order, or federal regulation? Are there any relevant state or federal court decisions? If yes, cite the specific law, case or regulation as well as, any deadlines for action.

The proposed rulemaking is not mandated by any Federal or State law or court order or Federal regulation.

(10) State why the regulation is needed. Explain the compelling public interest that justifies the regulation. Describe who will benefit from the regulation. Quantify the benefits as completely as possible and approximate the number of people who will benefit.

In addition to graduation from an approved school of pharmacy and successful completion of the licensure examination, completion of an internship prepares the applicant to function competently and effectively upon licensure as a pharmacist. Traditionally, the entire internship would be completed after graduation. However, the historical trend in pharmacy education is for greater participation by students in an academic internship as part of the educational process. Often, the pharmacy student will rotate through internships in a variety of practice settings. Rather than duplicating the educational internship, this rulemaking would permit a pharmacy intern to complete more of the internship requirements through academic internships or non-traditional internships.

Additionally, a pharmacy intern who permanently leaves a school of pharmacy without graduating might be able to continue working as a pharmacy intern. The position of pharmacy intern is intended to prepare the person to become a licensed pharmacist and not simply to provide subordinate pharmacy services, such as a pharmacy technician or unlicensed person. This rulemaking would better assure that only those aspiring pharmacists who continue the path to licensure will be permitted to continue working as pharmacy interns.

(11) If data is the basis for this regulation, please provide a description of the data, explain in detail how the data was obtained, and how it meets the acceptability standard for empirical, replicable and testable data that is supported by documentation, statistics, reports, studies or research. Please submit data or supporting materials with the regulatory package. If the material exceeds 50 pages, please provide it in a searchable electronic format or provide a list of citations and internet links that, where possible, can be accessed in a searchable format in lieu of the actual material. If other data was considered but not used, please explain why that data was determined not to be acceptable.

The proposed rulemaking is not based upon any scientific data, studies, or references.

(12) Describe who and how many people will be adversely affected by the regulation. How are they affected?

The Board does not foresee any groups being adversely affected by the proposed rulemaking.

(13) List the persons, groups or entities that will be required to comply with the regulation. Approximate the number of people who will be required to comply.

All pharmacy interns, including those pharmacy students participating in an internship, would be required to comply with the provisions of this rulemaking. There are approximately 4,200 pharmacy interns currently registered with the Board.

(14) Provide a specific estimate of the costs and/or savings to the **regulated community** associated with compliance, including any legal, accounting or consulting procedures which may be required. Explain how the dollar estimates were derived.

There are no costs or savings to the regulated community associated with compliance with the proposed rulemaking.

(15) Provide a specific estimate of the costs and/or savings to **local governments** associated with compliance, including any legal, accounting or consulting procedures which may be required. Explain how the dollar estimates were derived.

There are no costs or savings to local governments associated with compliance with the proposed rulemaking.

(16) Provide a specific estimate of the costs and/or savings to **state government** associated with the implementation of the regulation, including any legal, accounting, or consulting procedures which may be required. Explain how the dollar estimates were derived.

There are no costs or savings to state government associated with compliance with the proposed rulemaking.

(17) In the table below, provide an estimate of the fiscal savings and costs associated with implementation and compliance for the regulated community, local government, and state government for the current year and five subsequent years.

	Current FY Year	FY +1 Year	FY +2 Year	FY +3 Year	FY +4 Year	FY +5 Year
SAVINGS:	$	$	$	$	$	$
Regulated Community						
Local Government						
State Government						
Total Savings	N/A	N/A	N/A	N/A	N/A	N/A
COSTS:						
Regulated Community						
Local Government						
State Government						
Total Costs	N/A	N/A	N/A	N/A	N/A	N/A
REVENUE LOSSES:						
Regulated Community						
Local Government						
State Government						
Total Revenue Losses	N/A	N/A	N/A	N/A	N/A	N/A

(17a) Provide the past three year expenditure history for programs affected by the regulation.

Program	FY –3 (FY 07-08)	FY –2 (FY 08-09)	FY –1 (FY 09-10)	Current FY (FY 10-11)
Pa. State Board of Pharmacy	**actual $1,742,656**	**actual $1,695,150**	**projected $1,748,926**	**budgeted $2,226,000**

(18) Explain how the benefits of the regulation outweigh any cost and adverse effects.

No adverse effects or costs have been associated with compliance with the proposed rulemaking. Therefore, the above-identified benefits would outweigh any costs.

(19) Describe the communications with and input from the public and any advisory council/group in the development and drafting of the regulation. List the specific persons and/or groups who were involved.

The Board solicited and received input from the Commonwealth's schools of pharmacy and the proposed rulemaking was discussed at public meetings of the Board, which are routinely attended by members of the regulated community and their professional associations.

(20) Include a description of any alternative regulatory provisions which have been considered and rejected and a statement that the least burdensome acceptable alternative has been selected.

No alternative regulatory schemes were considered.

(21) Are there any provisions that are more stringent than federal standards? If yes, identify the specific provisions and the compelling Pennsylvania interest that demands stronger regulations.

The proposed rulemaking is not more stringent and does not overlap or conflict with any federal requirements.

(22) How does this regulation compare with those of other states? How will this affect Pennsylvania's ability to compete with other states?

In order to become licensed as a pharmacist in Delaware, an applicant must also complete a board-approved practical experience under the supervision of a licensed pharmacist of at least 1,500 hours, including at least 1,000 hours in community or hospital settings. The remaining hours may be earned working as an industrial pharmacist, drug information pharmacist, military pharmacist, mail order pharmacist, HMO pharmacist, consultant pharmacist (such as nursing home, infusion, Medicaid DUR), home health care pharmacist (such as durable medical equipment), nuclear pharmacist, compliance pharmacist, government pharmacist, clinical pharmacist, or contracted pharmacy services. Hours spent in a pharmacy school experience program may be counted as well.

In order to become licensed as a pharmacist in Maryland, an applicant must also complete either: (1) a school-supervised professional experience program of at least 1,000 hours in an accredited school of pharmacy, or (2) at least 1,560 hours of full-time training in a pharmacy under the direct supervision of a licensed pharmacist. A school program of less than 1,000 hours may be used to satisfy a portion of the hours required in full-time training.

In order to become licensed as a pharmacist in New Jersey, an applicant must also complete either: (1) subsequent to graduation from pharmacy school an internship of at least 1,000 hours over at least 24 weeks working between 20 and 45 hours per week, or (2) a structured externship and clinical pharmacy clerkship through a pharmacy school during the third professional year or later of at least 1,000 hours over no more than 45 hours per week during which the extern has direct involvement with consumers or patients or healthcare providers at least 75% of the time.

In order to become licensed as a pharmacist in New York, an applicant who has graduated from an approved school of pharmacy must also have at least 6 months' experience as a pharmacy intern in an internship program devoted to the preparing, compounding, preserving and dispensing of drugs, medicines and therapeutic devices and to the performance of the functions related thereto, such as the counseling of patients and the monitoring of drug regimens, under the supervision of a registered pharmacist. The experience may be obtained via full-time work at 40 hours per week and may be accumulated during the summer and winter semester breaks as well as after graduation and prior to licensure in another state. Intern hours accumulated concurrent with the school year are also not acceptable.

In order to become licensed as a pharmacist in Ohio, an applicant must also complete at least 1,500 hours of supervised practical experience, and may satisfy up to 500 hours of that requirement by a board-approved supervised experience outside a pharmacy (such as manufacturing, research, consulting, drug information, and drug utilization review). Practical experience obtained in a structured program for which academic credit is awarded, such as in an externship or clerkship, may be applied to satisfy this requirement. The intern must be actively working towards licensure as a pharmacist.

In order to become licensed as a pharmacist in West Virginia, an applicant must also complete an internship of at least 1,500 hours in a pharmacy under the supervision of a licensed pharmacist.

The proposed rulemaking would not put Pennsylvania at a competitive disadvantage.

(23) Will the regulation affect any other regulations of the promulgating agency or other state agencies? If yes, explain and provide specific citations.

This proposed regulation would not affect other regulations of the Board or other state agencies.

(24) Submit a statement of legal, accounting or consulting procedures and additional reporting, recordkeeping or other paperwork, including copies of forms or reports, which will be required for implementation of the regulation and an explanation of measures which have been taken to minimize these requirements.

This proposed rulemaking would not require any additional legal, accounting or consulting procedures or any additional reporting, recordkeeping or other paperwork.

(25) Please list any special provisions which have been developed to meet the particular needs of affected groups or persons including, but not limited to, minorities, elderly, small businesses, and farmers.

The Board has determined that there are no special needs of any subset of its applicants or licensees for whom special accommodations should be made.

(26) Include a schedule for review of the regulation including:

A. The date by which the agency must receive public comments: **Within 30 days of publication of the proposed rulemaking.**

B. The date or dates on which public meetings or hearings will be held: **The Board will review all comments on the proposed rulemaking at regularly scheduled board meetings. The Board generally meets on the third Tuesday of each month.**

C. The expected date of promulgation of the proposed regulation as a final-form regulation: **Spring 2012.**

D. The expected effective date of the final-form regulation: **Upon publication as final.**

E. The date by which compliance with the final-form regulation will be required: **Upon the effective date.**

F. The date by which required permits, licenses or other approvals must be obtained: **N/A**

(27) Provide the schedule for continual review of the regulation.

The Board continually reviews the efficacy of its regulations, as part of its annual review process under Executive Order 1996-1. The Board reviews its regulatory proposals at regularly scheduled public meetings, generally the third Tuesday of each month. More information can be found on the Department's website (www.dos.state.pa.us).

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FACE SHEET
FOR FILING DOCUMENTS
WITH THE LEGISLATIVE REFERENCE BUREAU

(Pursuant to Commonwealth Documents Law)

DO NOT WRITE IN THIS SPACE

Copy below is hereby approved as to form and legality. Attorney General

BY: _____
(DEPUTY ATTORNEY GENERAL)

JUL 26 2012
DATE OF APPROVAL

Copy below is hereby certified to be a true and correct copy of a document issued, prescribed or promulgated by:

State Board of Pharmacy
(AGENCY)

DOCUMENT/FISCAL NOTE NO. 16A-5424

DATE OF ADOPTION:

BY: Richard R. Smiga
Richard R. Smiga, RPh

TITLE: Chairperson
(EXECUTIVE OFFICER, CHAIRMAN OR SECRETARY)

Copy below is approved as to form and legality. Executive or ~~Independent~~ Agencies.

BY: Shawn E. Smith

JUN 27 2012
DATE OF APPROVAL

(Deputy General Counsel
(~~Chief Counsel~~,
~~Independent Agency~~
(Strike inapplicable title)

[ ] Check if applicable
Copy not approved.
Objections attached.

[ ] Check if applicable.
No Attorney General approval
or objection within 30 day
after submission.

PROPOSED RULEMAKING

COMMONWEALTH OF PENNSYLVANIA
DEPARTMENT OF STATE
BUREAU OF PROFESSIONAL AND OCCUPATIONAL AFFAIRS
STATE BOARD OF PHARMACY

49 Pa. Code § 27.26
PHARMACY INTERNSHIP

The State Board of Pharmacy (Board) proposes to amend § 27.26 (relating to pharmacy internship), to read as set forth in Annex A.

Effective date

The amendments will be effective upon publication of the final-form rulemaking in the *Pennsylvania Bulletin.*

Statutory Authority

The amendments are authorized under sections 3(c) and 6(k)(9) of the Pharmacy Act (act) (63 P.S. §§ 390-3(c) and 390-6(k)(9)).

Background and Need for the Amendment

Section 3(c) of the act requires that, "to insure proficiency in the practical aspects of pharmacy, the board shall, by regulation, prescribe internship requirements which must be satisfactorily completed prior to issuance of a pharmacist license." Section 27.26 sets forth standards for the pharmacy internship. In addition to graduation from an ACPE-accredited pharmacy degree program and successful completion of the licensure examination, completion of an internship prepares the applicant to function competently and effectively upon licensure. The current trend in pharmacy education is for greater participation by students in an academic internship as part of the educational process. This rulemaking intends to permit a pharmacy intern to complete more of the internship requirements through academic internships or non-traditional internships. Additionally, this rulemaking intends to better assure that only those aspiring pharmacists who continue the path to licensure will be permitted to continue to work as pharmacy interns.

Description of the Proposed Amendments

The proposed rulemaking would first amend § 27.26(a) to provide that the purpose of the internship is to provide the intern with experience that would enable the intern to begin "functioning competently and effectively upon licensure" rather than simply functioning competently "under the act and this chapter."

In addition to education in an ACPE-accredited pharmacy degree program, a pharmacy internship is a tremendously important phase of pharmacist training. An intern registration is valid for 6 years. However, if a pharmacy student chooses to permanently cease enrollment in a pharmacy program, the student is no longer progressing towards ultimate licensure as a pharmacist and should not be permitted to continue working as a pharmacy intern. The proposed rulemaking would amend § 27.26(c) to provide that a pharmacy intern registration will automatically become invalid if the intern permanently ceases enrollment in an accredited pharmacy degree program and to require the former intern to immediately return to the Board the pharmacy intern registration and preceptor approval documents.

The proposed rulemaking also would amend § 27.26 (b)(2) to clarify not only that an applicant must be enrolled or accepted into an ACPE-accredited pharmacy degree program, but also to expand the pre-pharmacy educational path by which an applicant may become so enrolled or accepted. So long as an applicant is enrolled or accepted as a first professional year (P1) student in an ACPE-accredited pharmacy degree program, the applicant is not required to have completed the 2 years of pre-pharmacy education at an ACPE-accredited school or college of pharmacy.

Section 27.26(d)(1) requires that a pharmacy intern serve an internship of at least 1,500 hours. The proposed rulemaking would amend § 27.26(d)(3) to decrease the minimum portion of that time that must be served in a pharmacy from 750 to 500 hours. Correspondingly, the proposed rulemaking would amend § 27.26(d)(4) to increase the maximum portion of that time that may be served in an academic internship from 750 to 1,000 hours.

Section 27.26(e) limits internship credit to activities related to the practice of pharmacy and provides examples of those activities. Because a pharmacist may take and fill a prescription by means of an oral order, telephone or otherwise, the proposed rulemaking would add to those examples of activities related to the practice of pharmacy "taking oral orders for prescriptions by telephone or otherwise."

The proposed rulemaking would add a new subsection to allow for an internship not served in a pharmacy or sponsored or approved by an accredited pharmacy degree program. As examples of a non-traditional internship, the Board has previously been asked to consider an intern serving with a preceptor pharmacist without being in a licensed pharmacy and also at a pharmaceutical company participating in a research project. Such non-traditional internships would have to be approved by the Board in advance. As a non-traditional internship by definition is not served in a pharmacy, it may not be used to satisfy the pharmacy hours of § 27.26(d)(3). In order to permit the Board adequate time to review the proposed internship, the request must be submitted at least 90 days before beginning the internship.

Finally, because the Board does not specifically approve a pharmacy in which a pharmacy intern serves an internship, the proposed rulemaking would amend existing §§ 27.26(g) and 27.26(h)(3) to refer to the pharmacy that is "utilized" for intern training, rather than a pharmacy that is "approved" for intern training.

Fiscal Impact and Paperwork Requirements

The proposed rulemaking would have no adverse fiscal impact on the Commonwealth or its political subdivisions. The rulemaking would impose no additional paperwork requirements upon the Commonwealth, political subdivisions or the private sector.

Sunset Date

The Board continuously monitors the effectiveness of its regulations. Therefore, no sunset date has been assigned.

Regulatory Review

Under section 5(a) of the Regulatory Review Act (71 P.S. § 745.5(a)), on August 8, 2012, the Board submitted a copy of this proposed rulemaking and a copy of a Regulatory Analysis Form to the Independent Regulatory Review Commission (IRRC) and to the Chairpersons of the Senate Consumer Protection and Professional Licensure Committee and the House Professional Licensure Committee. A copy of this material is available to the public upon request.

Under section 5(g) of the Regulatory Review Act, IRRC may convey comments, recommendations or objections to the proposed rulemaking within 30 days of the close of the public comment period. The comments, recommendations or objections shall specify the regulatory review criteria which have not been met. The Regulatory Review Act specifies detailed procedures for review, prior to final publication of the rulemaking, by the Board, the General Assembly and the Governor of comments, recommendations or objections raised.

Public Comment

Interested persons are invited to submit written comments, suggestions or objections regarding this proposed rulemaking to Kerry E. Maloney, Board Counsel, State Board of Pharmacy, by mail at P.O. Box 2649, Harrisburg, PA 17105-2649, or by email at st-pharmacy@state.pa.us, within 30 days following publication of this proposed rulemaking in the *Pennsylvania Bulletin*. Please reference No. 16A-5424 (pharmacy internship), when submitting comments.

Annex A

TITLE 49. PROFESSIONAL AND VOCATIONAL STANDARDS

PART I. DEPARTMENT OF STATE

Subpart A. PROFESSIONAL AND OCCUPATIONAL AFFAIRS

CHAPTER 27. STATE BOARD OF PHARMACY

* * * * *

PHARMACISTS

* * * * *

§ 27.26. Pharmacy internship.

(a) Pharmacy internship means the supervised practical experience required for licensure as a registered pharmacist. The purpose of the pharmacy internship program is to provide a registered intern with the knowledge and practical experience necessary for functioning competently [under the act and this chapter] and effectively upon licensure.

(b) [A certificate of registration] Registration as a pharmacy intern will be available to an individual of good moral character who has completed at least 2 years of [pharmacy] college [or an accredited program leading to transfer into the third year of a college in which the individual is enrolled or accepted] and is enrolled or accepted as a student of pharmacy in an ACPE -accredited pharmacy degree program. A person desiring to register as a pharmacy intern shall do the following:

* * * * *

(2) Forward to the Board [a letter or transcript certifying] acceptable documentation verifying that the applicant has successfully completed at least 2 years of [pharmacy] college [or an accredited program leading to transfer into the third year of a

pharmacy college in which the applicant is enrolled or accepted] and is enrolled or accepted as a student of pharmacy in an ACPE-accredited pharmacy degree program. Acceptable documentation must include a document bearing the school's seal received by the Board directly from the dean or registrar of the ACPE-accredited pharmacy degree program which includes the pharmacy student's name, address, social security number, and a statement indicating that the student has successfully completed at least 2 years of college and is enrolled or accepted as a student of pharmacy in, or has graduated from, the ACPE-accredited pharmacy degree program.

(c) The Board will register an applicant after it receives a completed application and other items in subsection (b). A pharmacy intern [certificate] registration is valid for 6 years from the date of issue exclusive of time spent in the military. A pharmacy intern registration will automatically become invalid if the pharmacy intern permanently ceases enrollment in an ACPE-accredited pharmacy degree program prior to graduation. A pharmacy intern whose registration becomes invalid under this subsection shall immediately return to the Board the pharmacy intern registration and all preceptor approval documents.

(d) The following applies to internship credit:

* * * * *

(3) An intern shall serve at least [750] 500 of the 1,500 hours in a pharmacy.

(4) An intern may earn up to [750] 1,000 of the 1,500 hours in an internship program sponsored or approved by [the pharmacy college] an ACPE-accredited pharmacy degree program [subject to the following conditions:

(i) The Board will determine the maximum number of hours available for each internship program sponsored or approved by a pharmacy college.

(ii) The Board will grant internship credit to an individual in an internship program sponsored or approved by a pharmacy college only if the following applies:

(A) The internship program is full-time.

(B) There is no concurrent academic course load.

(C) The individual achieves a passing grade in the program.

(iii) A pharmacy college which desires to sponsor or approve an internship program shall request approval from the Board.

(iv) The Board will monitor internship programs which are sponsored or approved by a pharmacy college].

* * * * *

(6) The Board will not grant internship credit for hours that an individual served in a pharmacy if the supervising pharmacist was not registered as a preceptor. An exception to the requirement that the supervising pharmacist register as a preceptor will be made for internship hours acquired in an internship program sponsored or approved by [a pharmacy college] an ACPE-accredited pharmacy degree program.

(e) The Board will grant internship credit only for activities related to the practice of pharmacy. The following are examples of these activities: scrutinizing prescriptions or drug orders, taking oral orders for prescriptions by telephone or otherwise, compounding medication and filling prescriptions. The Board will not grant internship credit for activities which are not related to the practice of pharmacy. [The following are examples of these activities: retail sales unrelated to pharmacy items, shelving or clerical functions unrelated to pharmacy.]

(f) An intern who wishes to receive credit for internship experience that is not in a pharmacy or sponsored or approved by an APCE-accredited pharmacy degree program must apply to the Board for approval before commencing such internship experience. Upon receipt of the application, the Board will review and determine how much, if any, credit will be given. Requests for approval should be submitted at least 90 days before the experience begins. Any credit given for a non-traditional internship may not be used to satisfy the requirement of paragraph (d)(3) pertaining to the minimum amount of time the internship must be served in a pharmacy.

(g) A person may not be eligible to become a candidate for registration to practice pharmacy unless the person receives instruction in practical pharmacy and pharmaceutical technique from an instructor, professor, or faculty member who is a registered pharmacist or from a faculty member who is a registered pharmacist at [a pharmacy college] an ACPE-accredited pharmacy degree program.

[(g)] (h) The following requirements are applicable to a pharmacy [approved] utilized for intern training:

* * * * *

[(h)] (i) The requirements for registration as a pharmacist preceptor are as follows:

* * * * *

(3) An applicant shall be working on a full-time basis in a pharmacy [approved] utilized for intern training.

* * * * *

[(i)] (j) * * *

[(j)] (k) * * *

[(k)] (l) * * *

* * * * *

COMMONWEALTH OF PENNSYLVANIA
DEPARTMENT OF STATE
BUREAU OF PROFESSIONAL AND OCCUPATIONAL AFFAIRS
STATE BOARD OF PHARMACY
Post Office Box 2649
Harrisburg, Pennsylvania 17105-2649
(717) 783-7156

August 8, 2012

The Honorable Silvan B. Lutkewitte, III, Chairman
INDEPENDENT REGULATORY REVIEW COMMISSION
14th Floor, Harristown 2, 333 Market Street
Harrisburg, Pennsylvania 17101

Re: Proposed Regulation
State Board of Pharmacy
16A-5424: Pharmacy Internship

Dear Chairman Lutkewitte:

Enclosed is a copy of a proposed rulemaking package of the State Board of Pharmacy pertaining to Pharmacy Internship.

The Board will be pleased to provide whatever information the Commission may require during the course of its review of the rulemaking.

Sincerely,

Edward J. Bechtel, R. Ph., Chairperson
State Board of Pharmacy

CKM/KEM:jsg
Enclosure
cc: Katie True, Commissioner
Bureau of Professional and Occupational Affairs
Rebecca Oyler, Director of Policy, Department of State
Steven V. Turner, Chief Counsel
Department of State
Cynthia Montgomery, Regulatory Counsel
Department of State
Kerry E. Maloney, Counsel
State Board of Pharmacy
State Board of Pharmacy

TRANSMITTAL SHEET FOR REGULATIONS SUBJECT TO THE REGULATORY REVIEW ACT

I.D. NUMBER: 16A-5424

SUBJECT: PHARMACY INTERNSHIP

AGENCY: DEPARTMENT OF STATE (STATE BOARD OF PHARMACY)
BUREAU OF PROFESSIONAL AND OCCUPATIONAL AFFAIRS

TYPE OF REGULATION

X Proposed Regulation

Final Regulation

Final Regulation with Notice of Proposed Rulemaking Omitted

120-day Emergency Certification of the Attorney General

120-day Emergency Certification of the Governor

Delivery of Tolled Regulation
a. With Revisions b. Without Revisions

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FILING OF REGULATION

DATE	SIGNATURE	DESIGNATION
8/8/12	Angie Kelly	HOUSE COMMITTEE ON PROFESSIONAL LICENSURE
		MAJORITY CHAIR Julie Harhart
		SENATE COMMITTEE ON CONSUMER PROTECTION & PROFESSIONAL LICENSURE
8/8/12	Jennifer Donelly	MAJORITY CHAIR Robert M. Tomlinson
8/8/12	K Cooper	INDEPENDENT REGULATORY REVIEW COMMISSION
		ATTORNEY GENERAL (for Final Omitted only)
8/8/12	Samantha Hasen	LEGISLATIVE REFERENCE BUREAU (for Proposed only)