



## 899 NORTH CAPITOL ST. NE $-2^{ND}$ FLR. WASHINGTON, DC 20002

October 3, 2019

9:30 AM – 10:49 AM

## **OPEN SESSION MEETING MINUTES**

## **Board of Pharmacy Mission Statement:**

"To protect and improve the public health through the efficient and effective regulation of the practice of Pharmacy and Pharmaceutical Detailing; through the licensure of Pharmacists, Pharmaceutical Detailers, Pharmacy Interns, and Pharmacy Technicians."

## CALL TO ORDER:

## PRESIDING: MR. JAMES APPLEBY, R.PH VICE CHAIR

## BOARD MEMBERSHIP/ATTENDANCE:

BOARD MEMBERS:		
	DR .TAMARA MCCANTS, PHARM.D. R.PH, CHAIRPERSON	Absent
	MR. JAMES APPLEBY, R.PH VICE CHAIR	PRESENT
	DR. BENJAMIN MILES, PHARM.D. R.PH	Present
	MR. ALAN FRIEDMAN, R.PH	Present
	DR. ASHLEE BOW, PHARM.D. R.PH	Present
	CHIKITA SANDERS, CONSUMER BOARD MEMBER	ABSENT
STAFF:	SHAUNA WHITE, EXECUTIVE DIRECTOR	Present
	KARIN BARRON, HEALTH LICENSING SPECIALIST	Present
	LUANNE GREENAWAY, HEALTH LICENSING SPECIALIST	ABSENT UNTIL 10:05
LEGAL STAFF:	CARLA WILLIAMS, ASSISTANT GENERAL COUNSEL	Present
VISITORS:	TIFFANIE TAYLOR, WALGREENS	
	DENISE FELLUCA, WALGREENS DON ZOWADA, DC PUBLIC	
	RYAN MOUTON, GWMFA	
	MONET STANFORD, KAISER PERMANENTE	
	CHARLENE FAIRFAX, DC DHCF	
	CHARLENET AIRPAA, DC DITCI	

# Open Session Agenda

## Quorum:

Introduction:	None
1003-0-01	Approval of the Open Session Meeting Minutes         September 5, 2019         Motion: Board Member Dr. Benjamin Miles moves the Board to approves         the August 1, 2019 open session minutes.         Seconded by: Dr. Ashlee Bow.         Abstentions: None.         Motion Carried.
	August 1, 2019 Motion: Board Member Dr. Benjamin Miles moves the Board to approves the August 1, 2019 open session minutes Seconded by: Dr. Ashlee Bow. Abstentions: Mr. Alan Friedman Motion Carried.
Consent Agenda	None
Executive Director Report	Licensing Report         Statistical Report on pharmacy professionals in the District of Columbia         > Pharmacists: 2,096         > Pharmacists with Vaccination and Immunization Authority: 662         > Pharmacy Interns: 524         > Pharmacy Technicians: 902         > Pharmacy Technician Trainees: 79         > Pharmacy Technician Training Programs: 11         > Pharmacy Technician Training Programs: 11         > Pharmaceutical Detailers: 862         Prescription Monitoring Program Update         > Licensed pharmacists in the District of Columbia are encouraged to observe the mandatory registration for the Prescription Drug Monitoring Program.         > Executive Director, Dr. Shauna Whites requests the public's assistance in encouraging practicing pharmacists to register for the program.         > The next step of the program is to encourage pharmacists, working in a clinical or dispensing capacity, to use the system when appropriate to practice.
	Opioid Strategic Plan <u>https://livelong.dc.gov/page/about-live-long-dc</u> , which is the designated website regarding the District of Columbia's opioid

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· · ·	I. For current information on the District's	
5 5	sis, the program, the strategic plan, and	
community partners, go	) to the website.	
NARRUndatas		
NABP Updates	ree (3) CEUs at the District 1 and 2 meeting	
this year:	ree (3) CLOS at the District 1 and 2 meeting	
	State Regulations.	
	s of Pharmacists in Changing the Health Care	
	fically value based care models.	
iii. Compounding.		
<ul> <li>For further information,</li> </ul>	, please go to <u>https://nabp.pharmacy/</u>	
DC Board of Pharmacy CE Audit	t Update	
<b>3 3 -</b>	meeting, the District of Columbia Board of	
	ate practitioner audits of licensed	
-	utical Detailers and Pharmacy Technicians. ess, Executive Director Dr. White will invite	
	ne audit upon receipt of CEU transcripts.	
	le dout opointeceipt of CEO transcripts.	
DEA Take Back Day		
	DEA's Take Back Day scheduled for October	
	ttps://takebackday.dea.gov/	
DC Health's NARCAN Pilot		
	le at pharmacies throughout the District of	
	ion does not require a prescription and is	
dispensed to the custon		
The pharmacies particip	pating in the pilot program are:	
Morgan Pharmacy	3001 P Street NW	
intergant narmacy	Washington, DC 20007	
Grubbs Pharmacy	326 East Capitol Street NE	
,	Washington, DC 20003	
Grubbs Pharmacy	1800 Martin Luther King, Jr.	
	Avenue SE	
	Washington, DC 20020	
Grubbs Pharmacy	1517 17 <sup>th</sup> Street NW	
	Washington, DC 20036	
Kalorama Pharmacy	1841 Columbia Road NW	
	Washington, DC 20009	
Good Care Pharmacy	2910 Martin Luther King Jr. Avenue	
	SE Washington DC appag	
Excel Pharmany	Washington, DC 20032	
Excel Pharmacy	Washington, DC 20032 3923 South Capitol Street SW	
	Washington, DC 20032 3923 South Capitol Street SW Washington, DC 20032	
Excel Pharmacy CVS Pharmacy, #22	Washington, DC 20032 3923 South Capitol Street SW Washington, DC 20032 320 40 <sup>th</sup> Street NE	
CVS Pharmacy, #22	Washington, DC 200323923 South Capitol Street SWWashington, DC 20032320 40th Street NEWashington, DC 20019	
	Washington, DC 20032 3923 South Capitol Street SW Washington, DC 20032 320 40 <sup>th</sup> Street NE	

	CVS Pharmacy, #1360	2834 Alabama Avenue SE	
		Washington, DC 20020	
	CVS Pharmacy, #1364	6514 Georgia Avenue NW	
		Washington, DC 20012	
	CVS Pharmacy, #1354	2601 Connecticut Avenue NW	
		Washington, DC 20008	
	CVS Pharmacy, #2834	303114 <sup>th</sup> Street NW	
		Washington, DC 20009	
	Safeway Pharmacy, #1445	2845 Alabama Avenue SE	
		Washington, DC 20020	
	Walgreens Pharmacy, #15360	8017 <sup>th</sup> Street NW	
		Washington, DC 20001	
	Walgreens Pharmacy, #16049	Howard University Location	
		2041 Georgia Avenue NW	
		Washington, DC 20060	
	Giant Pharmacy, #384	1535 Alabama Avenue SE	
		Washington, DC 20032	
	The pilot program will be evaluated as a second	valuated after a year.	
<u>Assistant General</u> <u>Counsel Report</u>			Carla Williams
1003-0-02	Clinical Lab Practitioner Advisory C	ommittee Updates	
	<ul> <li>Clinical Lab Practitioner Ad amending DC Law 20-272: Amendment Act of 2014, as clinical [laboratory] practice</li> <li>A draft of the amended DC Practitioners Amendment A 2019 meeting, with sugges amendments to the law, ar Consequently, regulations regulate clinical laboratory</li> <li>The proposed amendments i. Add in the qualifica Director.</li> <li>add in language for various clinical laboratory piv. Add in language the Advisory Committee concerning matters</li> </ul>	Law 20-272 the Clinical Laboratory ct of 2014 was presented at the October tions and recommendation for ad by extension, legislation. will authorize the District to license and practitioners. s are as follows: tions to serve as the Clinical Laboratory r accuracy and definition regarding the pratory practices. or clarification of regulations regarding	
	concerning matters practitioners."	s pertaining to clinical laboratory	

	Motion: Board Member Mr. Alan Friedman moves that The Board approves amendments to the District of Columbia's <i>Health Occupations Revision Act</i> of 1985 relative to DC Law 20-272: the Clinical Laboratory Practitioners Amendment Act of 2014. Seconded by: Dr. Benjamin Miles. Abstentions: None. Motion Carried.	
<u>Subcommittee</u> <u>Report</u>		
<u>Report</u>	Legislative and Regulatory	
1003-O-03	<ul> <li>Defending Access to Women's Health Care Services Amendment Act of 2017</li> <li>The Board of Medicine approved the materials and regulations to reflect the Board of Pharmacy's approvals of the Act, which was reported at the August 2019 meeting.</li> <li>Executive Director, Dr. Shauna White, will present the updated Act during the Open Session of the December 2019 Board of Pharmacy Meeting.</li> </ul>	Dr. Shauna White and Mr. Alan Friedman
<u>Addendum</u>		
	<ul> <li>Collaborative Practice Agreement Regulations</li> <li>Section 10001.7 of the <i>Requirements for Participation in a Collaborative</i> <i>Practice Agreement</i> of the <i>District of Columbia Municipal Regulations</i> states the following educational requirement:</li> <li>A Pharmacist participating in a collaborative practice agreement is required to have obtained: <ul> <li>(1) A minimum of three (3) years of relevant clinical experience, if the pharmacist holds an academic degree of Doctor of Pharmacy; <u>or</u></li> <li>(2) A minimum of five (5) years of relevant clinical experience, if the pharmacist holds an academic degree of Bachelor of Science in Pharmacy; <u>and</u></li> <li>(A) A residency accredited by the American Society of Health Systems Pharmacists, a body approved by the Board of Pharmacy or offered by a body accredited by the Accreditation Council for Pharmacy Education; <u>or</u></li> <li>(B) A certificate program approved by the Board of Pharmacy.</li> </ul> </li> <li>The Legislative and Regulatory Subcommittee proposes a change in this requirement to reflect the following:</li> </ul>	

(1) A minimum of three (3) years of relevant clinical experience, if the pharmacist holds an academic degree of Doctor of Pharmacy; <u>or</u>
(2) A minimum of five (5) years of relevant clinical experience, if the pharmacist holds an academic degree of Bachelor of Science in Pharmacy; <b>or</b>
(A) A residency accredited by the American Society of Health Systems Pharmacists, a body approved by the Board of Pharmacy or offered by a body accredited by the Accreditation Council for Pharmacy Education; <u>or</u>
(B) A certificate program approved by the Board of Pharmacy.
The Subcommittee proposes the change because it recognizes that the current regulation prohibits professionals who have completed residencies; to participate in collaborative practice agreements. The Subcommittee dissents that the regulation limits professionals in the community setting, who may be qualified to engage in a collaborative practice agreement. The proposed change would allow the professionals who have not completed a residency but have had adequate work experience to engage in a collaborative practice agreement in a hospital setting as well as a community setting.
Motion: Board Member Dr. Benjamin Miles moves that The Board approves a change to the training requirements set forth in Section 10001.7 of the <i>Requirements for Participation in a Collaborative Practice Agreement</i> of the <i>District of Columbia Municipal Regulations</i> , from <u>and</u> to <u>or</u> in subsections 10009.9(a)(2)(B) and 10001.7 (a)(2)(B) thereby allowing certification <u>or</u> residency <u>or</u> experience [as acceptable qualification to engage in collaborative practice agreements]. Seconded by: Mr. Alan Friedman. Abstentions: None. Motion Carried.
Collaborative Practice Agreement Regulations
During the August 2019 Board of Pharmacy meeting, the following was discussed:
The Board of Medicine reviewed the findings from the Collaborative Practice Subcommittee at its June 2019 meeting, and voted to approve the following categories as not needing approval by the Boards:
<ul> <li>a. Parenteral Nutrition</li> <li>b. Vancomycin</li> <li>c. Aminoglycosides</li> <li>d. Phenytoin</li> </ul>
e. Anticoagulation/Warfarin f. Renal Dose Adjustments

For the other categories identified below, the Board of Medicine voted to continue discussion and research on the topics.

- a. Pain Management (e.g., management of medication dosing)
- b. HIV
- c. Diabetes
- d. Hypertension
- e. Hyperlipidemia
- f. Depression

Additional suggestion [was disclosed] for consideration is Smoking Cessation.

Mr. Alan Friedman questions the recommendations proposed by the Board of Medicine disease states where the Board of Medicine agrees that there is no requirement to satisfy in establishing a collaborative practice agreement. Mr. Friedman also notes that for other disease states, the Board of Medicine considers approval or no approval to establish a collaborative practice agreement. However, collaborative practice regulations were written and approved expressly stating that there are no requirements for either Board to approve collaborative practice agreements that are deemed "standard of care." The "standard of care" is also defined in the regulations. If the intent of the Board of Medicine is to further consider disease states for approval or non-approval, then a regulatory change is necessary.

Assistant General Counsel suggests that Executive Director, Dr. Shauna White, must discuss this matter with the Board of Medicine to determine the intent of that Board.

### No motion entered.

### Collaborative Practice Agreements.

The Legislative and Regulatory Subcommittee would like to explore review of regulations of pharmacies and pharmacists as the Board of Pharmacy has oversight of the [pharmacists, pharmacy technicians and pharmaceutical detailers]. The Subcommittee suggests updating the regulations, to include a methodical process and propose updates for the public's consideration.

Motion: Board Member Mr. Alan Friedman moves that The Board tasks the Legislative and Regulatory Subcommittee with reviewing current chapters of regulation that provide broad oversight of pharmacists, pharmacy technicians, pharmaceutical detailers and collaborative practice agreements; and that the committee will make recommendations to The Board for possible amendments. Seconded by: Dr. Ashlee Bow. Abstentions: None. Motion Carried.

1003-0-04	Communications Subcommittee Report	
NABP E- Newsletter	<u>September 11, 2019</u>	Dr. Shauna White
	NABP Launches New Consumer Website	
	Missouri Board of Pharmacy Accepting Applications for CS Drug Take Back Program	
	CDC Awards New Funds to Reduce Opioid Overdose Deaths	
	Plastikon Healthcare Recalls Milk of Magnesia Oral Suspension Due to Microbial Contamination	
	June 19, 2019	
	Hurricane Dorian Prompts State of Emergency in Five States (Florida, Georgia , North Carolina, South Carolina, and Virginia)	
	NABP Model Act Updated to Address Suspicious Orders, PMP Reporting for Veterinarians, and More	
	DEA Announces Steps to Improve Access to Marijuana Research	
	Forty-One People Face Charges for Diversion of Prescription Medication Press Release: <u>https://www.justice.gov</u>	
	Note to the Public: To receive weekly updates from NABP, please sign up by using the following link: <u>https://nabp.pharmacy/newsroom/news/</u>	
<u>Matters for Board</u> <u>Consideration</u>		
1003-0-05	Access to Biosimilars Amendment Act of 2019	
	This legislation was developed by the Medical Society of the District of Columbia. The bill will amend the District of Columbia Prescription Drug Price Information Act to authorize licensed pharmacists to dispense interchangeable biological products, and to require notifications to physicians when such interchangeable biological products are dispensed.	
	This information is brought before The Board for review and feedback.	
	<ul> <li>Mr. Alan Friedman suggests that the language, including the notification requirements and in particular, the language regarding interoperable systems, is standard across many states of the nation.</li> <li>Mr. Friedman poses the following questions to The Board for consideration:         <ol> <li>With regard to the definition of interchangeable biological product: Why the reference to the FDA's orange book versus its</li> </ol> </li> </ul>	
	product: Why the reference to the FDA's orange book versus its purple book, where the purple book is related to biologicals and biosimilars?	

1003-0-06	product: the Board of Pharmacy does not currently publish a list and pharmacists in practice consult references wherever needed. No motion entered. License Examination Testing Accommodation Policy The DC Board of Pharmacy is interested in establishing guidelines for candidates that are requesting for exam testing accommodations. The Board thought it would be beneficial to contact the Virginia Board of Pharmacy for guidelines in establishing a policy regarding this matter. Virginia's Regulation regarding testing accommodations: 18VAC110-20-60 F. If an applicant requests a testing accommodation for either examination
	<ol> <li>While the narrative on interoperability is acceptable, electronic communication and more documentation between pharmacy systems and medical records will cause a problem where no system exists. Patient notifications to the physicians therefore, would demand a manual process. Pharmacies that are systematically connected to providers would not need to notify a physician. However, for pharmacies outside of the network, how would they remember to notify a provider? The [inconsistency] then, may create a gap and would therefore, bring into question the compliance of the pharmacies that are not systematically connected to providers.</li> <li>Assistant General Counsel, Ms. Carla Williams informs The Board that this bill was introduced for legislation on September 17, 2019.</li> <li>Mr. Friedman suggests the following actions by The Board:         <ol> <li>Research the technicalities between [the FDA's] purple book versus its orange book.</li> <li>Determine the feasibility of patient notification compliance.</li> </ol> </li> <li>Executive Director, Dr. Shauna White requests that board members submit all suggestions regarding this bill by Monday, October 7, 2019.</li> <li>With regard to maintaining a link on the websites of the Boards of medicine and pharmacy, which discloses the current list of biological products determined by the FDA to be interchangeable with a specific biological products determined by the FDA to be interchangeable with a specific biological products determined by the FDA to be interchangeable with a specific biological products determined by the FDA to be interchangeable with a specific biological products determined by the FDA to be interchangeable with a specific biological products determined by the FDA to be interchangeable with a specific biological products determined by the FDA to be interchangeable with a specific biological products determined by the FDA to be intherchangeable with a specific biological products determined by</li></ol>
	2. With regard to the patient notification requirements: if the idea behind interchangeabilities of biosimilars is no different from those of brand generic, conceptually all is the same. Why [then], would we have to notify the physician, when we currently do not to the same for brand generic?

more major life activities, subject to the Americans with Disabilities Act, the board may approve a reasonable accommodation that does not compromise the security or integrity of the examination.	
1. Supporting documentation shall be provided by the applicant to include the following to be considered for review: a. A letter of request from the candidate that specifies the testing accommodation requested;	
b. A written report of an evaluation (educational, psychological, or physical) within the preceding two years from a qualified professional that states a diagnosis of the disability, describes the disability, recommends specific accommodations, and provides justification that the accommodation is appropriate and necessary for the diagnosed disability. If the comprehensive evaluation was done more than two years ago and the condition is one that is not subject to change, the original evaluation report may be submitted along with a current letter from the qualified professional stating that there has been no change in the condition since the time of the evaluation; and	
c. A written statement from the appropriate person at the applicant's school of pharmacy that describes any testing accommodations made while the student was enrolled, if applicable.	
3. The applicant will be notified in writing of the decision. If the request for accommodation is granted, the approval information will be forwarded to the examination contractor and the form of the accommodation will be coordinated with the contractor.	
In recognition of the Americans with Disabilities Act (ADA), Executive Director, Dr. Shauna White states that a drafted policy will be presented at the December 2019 Board of Pharmacy monthly meeting.	
No motion entered.	
<u>Dr. Tiffany Taylor, Pharmacy Resident, Walgreens (Howard University</u> <u>Location)</u>	
Dr. Taylor informs The Board of the success of the NARCAN Pilot Program as follows:	
<ul> <li>As a participant in the NARCAN Pilot Program, the Walgreens Pharmacy at Howard University is receiving great feedback on the program.</li> </ul>	
<ul> <li>on the program as well as the dispensing of medication.</li> <li>The Walgreens Pharmacy at Howard University has already dispensed eight (8) NARCAN kits for the month of October.</li> </ul>	
Dr. Taylor will submit a more detailed data report on the NARCAN Pilot Program specific to the Walgreens Pharmacy at Howard University during the Open Session of the December 2019 Board of Pharmacy meeting.	
	<ul> <li>board may approve a reasonable accommodation that does not compromise the security or integrity of the examination.</li> <li>1. Supporting documentation shall be provided by the applicant to include the following to be considered for review: a. A letter of request from the candidate that specifies the testing accommodation requested;</li> <li>b. A written report of an evaluation (educational, psychological, or physical) within the preceding two years from a qualified professional that states a diagnosis of the disability, describes the disability, recommends specific accommodations, and provides justification that the accommodation is appropriate and necessary for the diagnosed disability. If the comprehensive evaluation was done more than two years ago and the condition is one that is not subject to change, the original evaluation report may be submitted along with a current letter from the qualified professional stating that there has been no change in the condition since the time of the evaluation, and</li> <li>c. A written statement from the appropriate person at the applicant's school of pharmacy that describes any testing accommodations made while the student was enrolled, if applicable.</li> <li>3. The applicant will be notified in writing of the decision. If the request for accommodation is granted, the approval information will be forwarded to the examination contractor and the form of the accommodation will be coordinated with the contractor.</li> <li>In recognition of the Americans with Disabilities Act (ADA), Executive Director, Dr. Shauna White states that a drafted policy will be presented at the December 2019 Board of Pharmacy monthly meeting.</li> <li>No motion entered.</li> <li>Dr. Taylor informs The Board of the success of the NARCAN Pilot Program as follows:</li> <li>As a participant in the NARCAN Pilot Program, the Walgreens Pharmacy at Howard University is receiving great feedback on the program.</li> <li>Overall, patients are welcoming and appreciative of the information on the pr</li></ul>

the Open Session	"Mr. Chairperson, I move that the Board close the Open Public session portion of the meeting and move into the Closed Executive Session portion of the meeting pursuant to D.C. Official Code § 2-575(b) for the following purposes: to discuss disciplinary matters pursuant to § 2-575(b)(9); to seek the advice of counsel to the board, to preserve the attorney-client privilege, or to approve settlement agreements pursuant to § 2-575(b)(4); and to plan, discuss, or hear reports concerning ongoing or planned investigations
	pursuant to § 2-575(b)(14)." (Roll Call Vote)
	Seconded by: Board Member, Mr. Alan Friedman.
	Roll Call Vote: Dr. Benjamin Miles: Votes in favor of the motion. Mr. Alan Friedman: Votes in favor of the motion. Dr. Ashlee Bow: Votes in favor of the motion. Mr. James Appleby: Votes in favor of the motion. Abstentions: None Motion Carried

This concludes the Public Open Session of the meeting. The Board will now move into the Closed Executive Session portion of the meeting pursuant to D.C. Official Code § 2-575(b) for the reasons set forth in the motion.

## Open Session Meeting Adjourned at 10:49 AM

This meeting is governed by the Open Meetings Act. Please address any questions or complaints arising under this meeting to the Office of Open Government at opengovoffice@dc.gov.