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Part B: Information about the Nominee

(to be completed by the Nominee)

NOTE>>>The applicants for Group and Region Focused Training Program are required to fill in "Every Item". As for the applications for Country Focused Training Program including Counterpart Training Program and some specified International Dialogue Programs, it is required to fill in the designated "required" items as is shown below.

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2) Nationality (as shown in the passport)				Ge	Georgia				1000000	5) Date of Birth (pleas month in English as in					
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4) Re	ligion				Ort	Orthodox Christianity				* . * .	7	Ma	y	1977	41
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7) Ty	pe of	Orga	nizati	on											
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9) Contact Information

11.7.799	Address: Tsereteli ave. 144, Tbilisi, 0119						
Office	TEL:	Mobile (Cell Phone): +995 577 591591					
	FAX:	E-mail: tjikia@moh.gov.ge					
	Address: Beliashvili 12, flat 10, Tbilisi						
Home	TEL:	Mobile (Cell Phone): +995 599992662					
	FAX:	E-mail: teajikia77@yahoo.com					
	Name: Merab Urushadze						
Contact person	Relationship to you: Husband						
in emergency	Address: Beliashvili 12, flat 10, Tbilisi						
	TEL:	Mobile (Cell Phone): +995 599473332					
	FAX:	E-mail: merabiur@yahoo.com					

10) Others (if necessary)			

4. Career Record

1) Job Record (After graduation)

_	City/	Pe	eriod		
Organization	Country	From Month/Year	To Month/Year	Position or Title	Brief Job Description
Pharmaceutical Activities Department at State Regulation Agency for Medical Activities Division of Marketing	Tbilisi	11.2011	12.2018	Deputy Head	In charge of coordination and overall supervision of specialized units of department
Authorization, Pharmaceutical Activities Department at State Regulation Agency for Medical Activities	Tbilisi	01.2009	11.2011	Head Specialist	Peer reviewer, Assessment of scientific technical dossier for MA
Division of Marketing Authorization, Pharmaceutical Activities Department at State Regulation Agency for Medical Activities	Tbilisi	03.2008	12.2008	Senior specialist	Assessment of scientific technical dossier for MA
Department of Inspections. Drug Agency of Georgia	Tbilisi	01.2007	02.2008	Senior specialist	Inspections of retailers and wholesalers
Department of GMP Inspections. Drug Agency of Georgia		06.2006	12.2006	Specialist	Inspections of pharmaceutical manufacturers

2) Educational Record (Higher Education)(required)

=) Educational Reco	a (mgne	Education	n)(required)	
	City/	Pe	riod	2002	
Institution	Country	From Month/Year	To Month/Year	Degree obtained	Major
Georgian Academy of Sciences, lovel Kutateladze Institute of Pharmacochemistry and Tbilisi State Medical University	Tbilisi	09.2001	12.2006	Candidate of Pharmaceu- tical Sciences (Scientific title awarded by the Georgian Academy of Sciences, equal to PhD)	Pharmaceutical technology, development of solid inhalation dosage forms



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Tbilisi State Medical University, Faculty of Pharmacy.	Tbilisi	09.1996	07.2001	Degree in a Pharmacy equated to Master degree	Basic and advanced pharmaceutical disciplines

3) Training or Study in Foreign Countries; please write your past visits to Japan specifically as much as possible, if any

much as possible, if ar	ıy.		2	
			eriod	
Institution	City/ Country	From Month/Ye ar	To Month/Year	Field of Study / Program Title
WHO PQT	Denmark / Copenha gen	01.2012	01.2012	Training for quality assessors (theoretical training with written exam)
WHO PQT	Denmark / Copenha gen	05.2011	05.2011	Training for quality assessors (practical)
WHO PQT	Denmark / Copenha gen	01.2011	01.2011	Training for quality assessors (theoretical training with written exam)
Centre for Drug Policy and Pharmacy Practice Development, "Pharmakon"	Denmark /Hillerod	08.2006	09.2006	The first training module for pharmaceutical inspectors.
Mediterranean University, Faculty of Pharmacy, Laboratory of Galenic Preparations, Industrial Pharmacy and Cosmetology	Marseille / France	10.2003	07.2004	Pharmaceutical technology, development of solid inhalation dosage forms – dry powder inhalation formulation
"Technologie Servier", Drug Delivery Research Department	Orleans/ France	09.2004	04.2005	Pharmaceutical technology, development of solid inhalation dosage forms – dry powder inhalation formulation
Laboratory of Microbiology, Virology and Hygiene. Timone Hospital,	Marseille / France	05.2005	08.2005	Pharmaceutical technology, development of solid inhalation dosage forms – dry powder inhalation formulation, microbiological part

5. Language Proficiency (required)

or manager remoterion (req	(all ca)				
1) Language to be used in the progr	am (as in GI)				
Listening	(×) Excellent	() Good	() Fair	() Poor	
Speaking	(×) Excellent	() Good	() Fair	() Poor	
Reading	(x) Excellent	() Good	() Fair	() Poor	
Writing	(×) Excellent	() Good	() Fair	() Poor	
Certificate (Examples: TOEFL, TOEIC)	· ,		•		
2) Mother Tongue	Georgian	ě		и	
3)Other languages (Russian)	(×) Excellent	() Good	() Fair	() Poor	



				Application of the second
4) French	() Excellent	() Good	() Fair	(×) Poor
¹ Excellent: Refined fluency skills and to deal with various essay types, including	pic-controlled discus narrative, compariso	sions, debates & pre n, cause-effect & arç	sentations. Formula jumentative essays.	tes strategies to

Good: Conversational accuracy & fluency in a wide range of situations: discussions, short presentations & interviews.
 Compound complex sentences. Extended essay formation.
 Fair: Broader range of language related to expressing opinions, giving advice, making suggestions. Limited compound and complex sentences & expanded paragraph formation.
 Poor: Simple conversation level, such as self-introduction, brief question & answer using the present and past tenses.





6. Expectation on the applied training and dialogue program

1) Personal Goal: Describe what you intend to achieve in the applied training and dialogue program in relation to the organizational purpose described in Part A-2.

My personal goal is to gain more experience, more knowledge, develop new skills, to get acquainted with stringent pharmaceutical regulatory system established in Japan. Training course is an excellent opportunity to meet colleagues, share experience, to discuss challenges in countries with well-established regulatory systems and in countries where pharmaceutical regulatory system is insufficient. As all of us share the main goal — to establish functional pharmaceutical regulatory system in the world, all of us should be properly armed with the knowledge, experience and appropriate information in order to build functional regulatory system in our own countries and ensure access to high-quality medicines. Georgia is facing many challenges as we are in the initial stage of the regulatory system reform and international cooperation is very important for successful implementation of the goals.

2) Relevant Experience: Describe your previous vocational experiences which are highly relevant in the themes of the applied training and dialogue program. (required)

Almost all topics of the course are directly related to the main functions of Agency where I'm working since 2006. I've started to work at Agency as Inspector. Later, approximately during 4 years, I was performing quality part assessment of technical dossier for marketing authorization of medicines, including biologicals. Since 2011, coordination and supervision of main units of the regulatory body was my daily work.

3) Area of Interest: Describe your subject of particular interest with reference to the contents of the applied training and dialogue program. (required)

Pharmaceutical Administration in Japan, Japan's Medical Insurance system and Drug price, generic medicines market, Health Technology Assessment (HTA) system in Japan, Regulation of traditional herbal medicines, International cooperation.

*7. Declaration (to be signed by the Nominee) (required)

I certify that the statements I have made in this form are true and correct to the best of my knowledge. If accepted for the program, I agree:

- (a) not to bring or invite any member of my family (except for a program whose period is one year or more),
- (b) to carry out such instructions and abide by such conditions as may be stipulated by both the nominating government and the Japanese Government regarding the program,
- (c) to follow the program, and abide by the rules of the institution or establishment that implements said program,
- (d) to refrain from engaging in political activity or any form of employment for profit or gain,
- (e) to return to my home country at the end of the activities in Japan on the designated flight schedule arranged by JICA,
- (f) to discontinue the program if JICA and the applying organization agree on any reason for such discontinuation and not to claim any cost or damage due to the said discontinuation.



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- (g) to consent to waive any copyright holder's rights for documents or products produced during the project, against duplication and/or translation by JICA, as long as they are used for the purposes of the program.
- (h) to approve the privacy policy and the copyright policy mentioned in the Guidelines of Application. JICA's Information Security Policy in relation to Personal Information Protection
 - JICA will properly and safely manage personal information collected through this application form in accordance with JICA's privacy policy and the relevant laws of Japan concerning protection of personal information and take protection measures to prevent divulgation, loss or damages of such personal information.
 - Unless otherwise obtained approval from an applicant itself or there are valid reasons such as disclosure under laws and ordinances, etc.; and except for the following 1.-3., JICA will neither provide nor disclose personal information to any third party. JICA will use personal information provided only for the purposes in the following 1.-3 and will not use for any purpose other than the following 1.-3 without prior approval of an applicant itself.
 - 1. To provide technical training to technical training participants from developing countries.
 - 2. To provide technical training to technical training trainees from developing countries under the Citizens' Cooperation Activities..
 - 3. In addition to 1. and 2. above, if the government of Japan or JICA determines necessary in the course of technical cooperation.
- (i) to observe Japanese laws and ordinances during my stay, if I violate Japanese laws and ordinances, I will return the total amount or a part of the expenditure required for the training depending on the extent of the violation.
- (j) to understand that JICA does not assure issuance of Japan entry visa even after JICA decide to accept me. I understand the Embassy of Japan will decide it according to necessary formalities upon the submission of visa application from each participant.

Date: 29.03.2019	Signature: T. TRA	
	Print Name: Tea Jikia	



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MEDICAL HISTORY

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3. Other M	edical Probler	ne)		
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condition m	nay not be fina	ancially compens	sated by JICA	and may	result in	termination	of the
program.							
Data: 00.00.0	2010			11			
Date: 29.03.2	2019	Signature:	T. T	TRI	a		
		Print Name: Tea					