PUBLIC CONSULTATION FEEDBACK FORM

**Global Competency Framework for Regulators of Medical Products**

(WHO/GCF/DRAFT v1.6.4 July 2022)

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| **Contact Information**Name (First Last):Affiliation:Country:Email: |
| **About You or Your Organization**\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ |

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| **Feedback Questions** |
| 1. Does the Global Competency Framework for Regulators of Medical Products (GCF) provide appropriate level of details to facilitate national regulatory agencies in establishing competency-based recruitment, performance assessment, training and development?
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| 1. Do proficiency levels (Foundational, intermediate, advanced) for each competency domain or activity domain follow a logical progression?
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| **General Comments** |
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| **Sections/page and line No.** | **Original Text** | **Comment** | **Suggested Amendment** | **Internal Use Only****[blank]** |
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| **Glossary** |
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| **1 Introduction** |
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| **2 Overview of the Competency Model for Regulators** |
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| **3 Implementation** |
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|  |  |  |  |  |
| **4 Organizational level competencies** |
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| **5 Role-specific competencies** |
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|  |  |  |  |  |
|  **5.1 Reviewers** |
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|  **5.2 Inspectors** |
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|  **5.3 Laboratory Analysts** |
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|  **5.4 Vigilance Professionals** |
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| **Other Comments:** |
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