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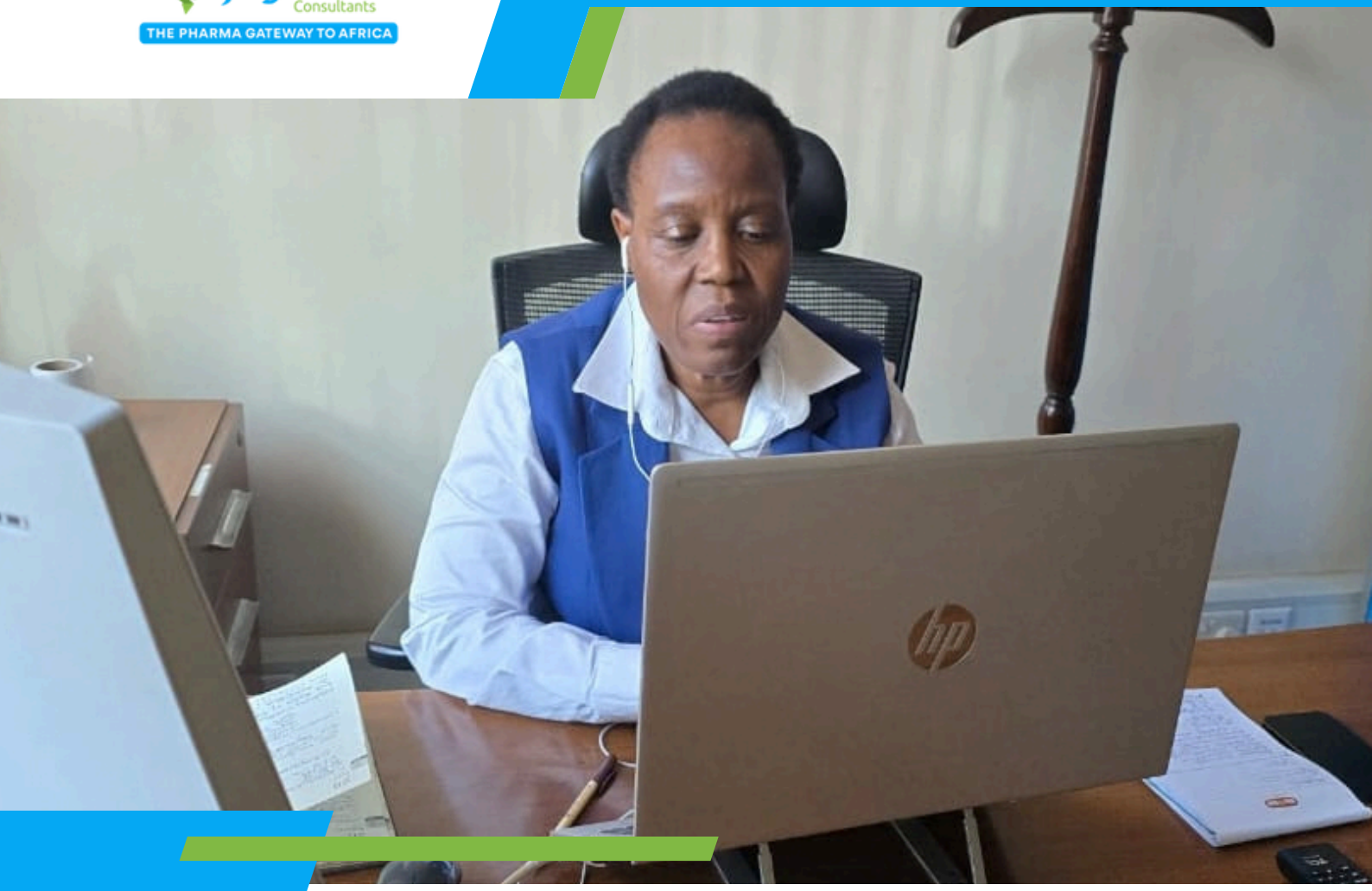
THE PHARMA GATEWAY TO AFRICA



Master Regulatory Excellence: Certificate in Regulatory Affairs

Join our intensive program to master regulatory frameworks that drive compliance, innovation, and market access in healthcare.

+254 798 725 500 jeyflex.com info@jeyflex.com

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In an evolving medical landscape, Regulatory Affairs is the heartbeat of compliance and innovation. This 30-hour intensive program is designed to transform health professionals into regulatory experts capable of navigating the complexities of Medical Products and Health Technologies (HPTs).

Who Should Enroll?

This course is perfect for:

- Pharmacovigilance Professionals seeking to deepen their expertise and stay compliant with global standards.
- Students eager to break into the pharmaceutical industry with a competitive edge.

Why Choose This Course?

01	**Industry-Aligned Curriculum:** Our content is mapped to current global and local regulatory standards for all Health Products and Technologies (HPTs).
02	**Practical Learning:** We move beyond theory with hands-on sessions for CTD dossier management and sample handling.
03	**Career Growth:** Equip yourself for high-demand roles in regulatory affairs, market access, and consultancy.
04	**Holistic Skill Development:** We integrate technical "hard" skills with essential "soft" skills like emotional intelligence and entrepreneurship.

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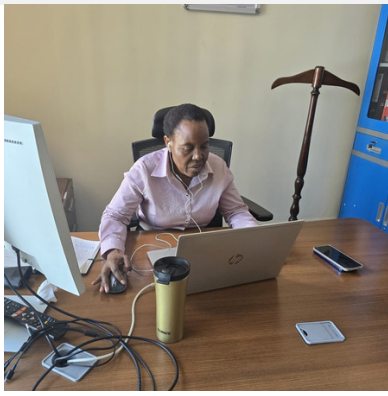
Course Module Highlights:

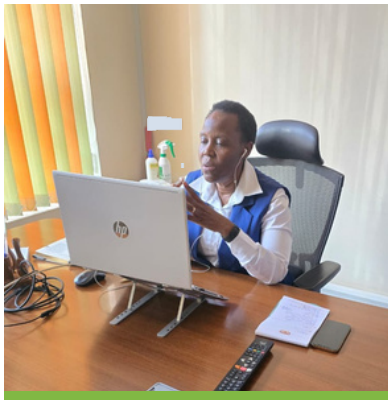


01 Foundations of Regulatory Affairs:

Understand the regulatory landscape, key frameworks, roles, and ethical responsibilities in the healthcare sector.

02 Cross-Functional Dynamics & Quality:

Learn how regulatory affairs interacts with pharmacovigilance, quality assurance, legal, and other departments.



03 Core Technical Processes (Core Focus):

Master HPT lifecycle management, CTD dossier preparation, and regulatory submissions.

Handle variations, renewals, and regulatory changes to maintain compliance.

04 Professional Development & Leadership

Build entrepreneurial thinking, communication skills, and emotional intelligence for career growth.

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Our Faculty

Learn from industry experts with extensive experience in pharmacovigilance and regulatory affairs:

Dr Jerusha Kamau

Lead Trainer | CEO, Jeyflex Consultants Ltd.

Dr. Jerusha Kamau is a pharmaceutical compliance expert and the CEO of Jeyflex Consultants Ltd, a Nairobi-based firm dedicated to advancing healthcare standards through specialized consulting. With over a decade of experience in pharmacovigilance, regulatory affairs, and professional capacity building, she equips professionals to navigate complex regulatory landscapes and enhance patient safety.

Beyond her role at Jeyflex, Dr. Kamau serves as Director for the Healthcare Sector at GS1 East Africa, leading innovative track-and-trace solutions for medical products. She is also the author of the upcoming Pharmaceutical Regulatory Handbook and a recognized thought leader committed to shaping the future of pharmaceutical compliance in the region.

Program Details:

- **Duration:** 7.5 weeks (30 contact hours)
- **Delivery Mode:** Virtual with interactive case studies and discussions.
- **Start date:** May 2026
- **Prerequisites:** Basic pharmacology/healthcare knowledge recommended (supplementary materials provided).
- **Assessment:** Quizzes, assignments, and a final evaluation (80% pass rate for certification).

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Investment:

- **Cost:** Ksh 30,000 (exclusive of VAT)
- **Pre-registration fee:** Ksh 5,000
- Balance payable within the first 3 weeks of training.

Payment Details

Pay via MPESA

Paybill Number: 4136453
Account Number: Student's Name

Join the Future of Regulatory Affairs!

Don't just watch the industry change, lead the transformation. Whether you are aiming to launch a new career or secure a seat at the leadership table, the JEYFLEX Certificate in Regulatory Affairs provides the technical depth and professional edge you need.

Secure your spot today and become the bridge between innovation and law!

ENROLL NOW