NoBoCap

NOBOCAP TRAINING

Exciting Developments in Long-Term Educational Programs!

There’s still time to register for Module 3: Generating Data for Technical Documentation (IVDR)

Prepare yourself for the specialized world of In Vitro Diagnostic Regulation (IVDR) in Module 3 of our Postgraduate Certificate series. This module delves into the construction of Technical Documentation and Performance Evaluation Reports under the IVDR framework, building upon the foundational knowledge from Module 1.

Applications for Module 3 are already open until **August 30, 2024**.

The module offers a 10-week program delivered entirely online, with a combination of virtual classroom sessions and online assessments. Upon completion, you’ll receive a Postgraduate CPD certificate worth 5 credits, accredited by Technological University Dublin (TU Dublin).

Don’t miss this opportunity to excel in IVDR knowledge! Follow us to stay up-to-date with the latest updates.

There’s still time to **register for Module 3** on the following link:

Register Now

Short-Term Courses and Interventions Update

More Short-Courses Coming Soon!

We're thrilled to announce that we will be offering even more short-courses in the coming months! These courses are designed to provide you with the latest insights and practical strategies to excel in your field.

Stay tuned to learn more about:

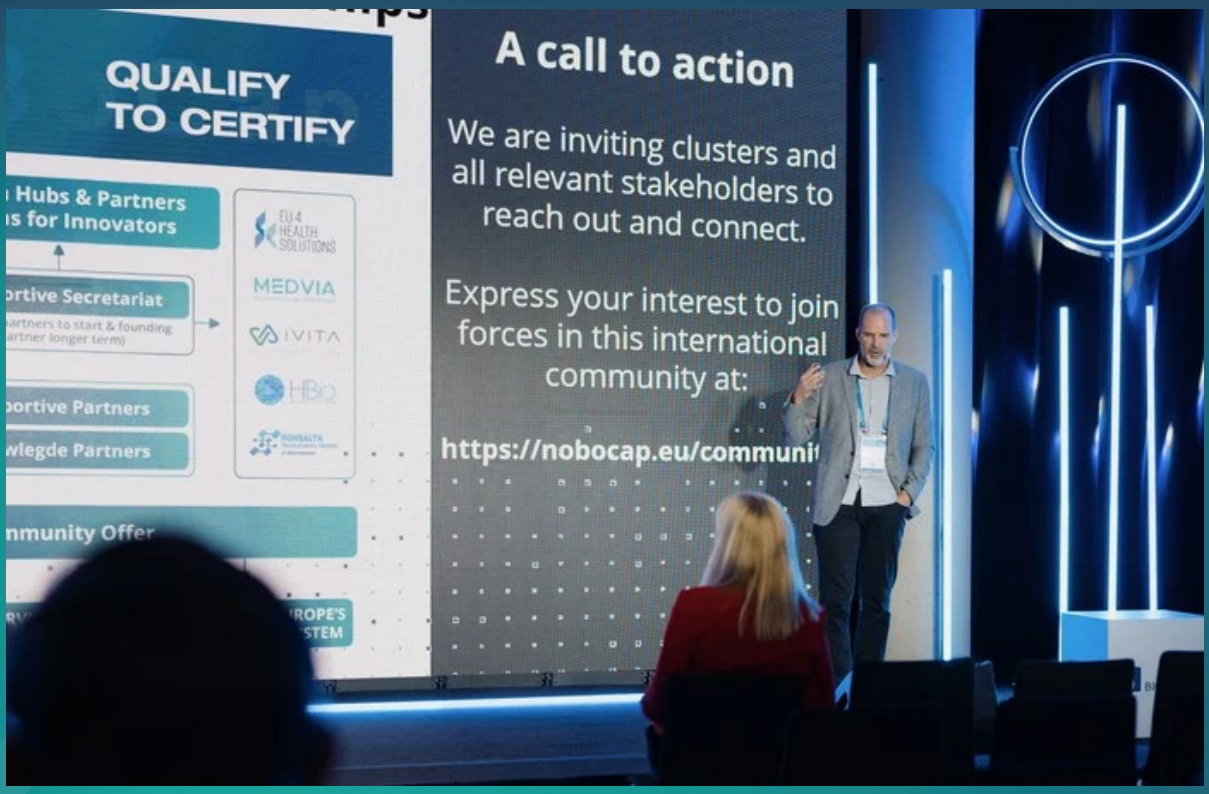
- New and innovative topics tailored for professionals at all levels
- Interactive sessions with industry experts
- Practical tools and resources for immediate application

Whether you're looking to enhance your skills or stay updated with industry trends, our short-courses are the perfect opportunity. Keep an eye on our updates to find out more about the upcoming sessions and how to register.

NOBOCAP COMMUNITY

Calling All Innovators!

Join NoBoCap Community



EU regulatory challenges, innovative solutions and initiatives toward strengthening the MedTech industry by enhancing the efficiency of Notified Bodies and Market Operators is more important than ever before.

We are calling Innovators to join the NoBoCap Community and become a part of our network driving change to Qualify to Certify! This could be the beginning of an unbelievable journey.

At NoBoCap Community, our pioneering and agile spirit keeps us interconnected with EU Regulatory challenges trying to increase knowledge and preparedness of Market Operators (MOs) in the application of MDR and IVDR.

NoBoCap Community allows you to be part of an innovation ecosystem designed to Unlock the EU regulation. Be one of those striving to bring innovation into healthcare and join the NoBoCap Community today!

Be innovative, bring unexpected ideas, push the boundaries, and contribute to solving complex challenges!

Visit our page

If interest to join the community, please visit the What to Offer page and complete the application form.

Application form

NEWS

Summit by NoBoCap Community Unlocking the MDR/IVDR Regulation for Innovators in Europe

Mark your calendars for **September 24-25, 2024**, and join us in **Brussels** for two days of high-level dialogues and sessions on MDR/IVDR regulations and innovation in MedTech.

Hear the speakers discuss building your knowledge about MDR\IVDR requirements, gain insights from EC Representatives on the evolving EU regulatory environment, and participate in workshops and panel discussions.

Connect with innovators, SMEs, and Notified Bodies through engaging sessions and networking opportunities.

Join NoBoCap Community Summit to stay ahead in the evolving regulatory landscape and ensure your innovations comply with MDR/IVDR requirements.

Register here

Stay informed! European Commission’s website

In the realm of Medical Devices and In Vitro Diagnostics, staying informed of regulatory updates is essential for manufacturers, healthcare professionals, and patients. The European Commission’s Directorate-General for Health and Food Safety (DG SANTE) release regularly updates and reports that provides valuable insights into the status of certifications and applications under the MDR and IVDR Regulations. Visit the European Union’s Health Website because it serves as an important resource for accessing the latest information and insights regarding the Medical Device Regulations (MDR) and the In Vitro Diagnostic Regulations (IVDR).

Visit website

Online Market Operators Survey

We have designed an anonymous survey aimed at understanding the training needs of medical device market operators across the European Union. Your participation will provide important insights that can shape future training programs and improve industry standards.

Your cooperation in completing this survey is highly appreciated. Your input will help us better serve the needs of online market operators in the medical device industry.

Thank you in advance for your participation and support! Let’s work together to enhance the effectiveness and efficiency of medical device markets in the EU.

Start the survey

NoBoCap

www.nobocap.eu

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