View in browser

NoBoCap

About Community Contacts

NOBOCAP TRAINING

Unlock the Path to MDR/IVDR Success with NoBoCap Training

Navigating the Medical Devices Regulation (MDR) and In Vitro Diagnostic Regulation (IVDR) can be challenging - but NoBoCap is here to guide you.

Through our training sessions, we empower innovators, startups, and notified bodies to confidently address regulatory requirements and accelerate market access for medical technologies.

2025 marks another year of comprehensive courses designed to equip you with the tools and insights needed for success.

Stay close to us to find out when registrations open for our upcoming sessions!

Your journey continues with NoBoCap - let’s build a stronger future together!

Register Now

Short-Term Courses and Interventions Update

Empowering Leaders and Innovators in MedTech with NoBoCap Short Courses!

NoBoCap continues to support the MedTech industry with specialized short courses tailored to address the most pressing challenges in the regulatory landscape.

- **NoBoCap Short Course for C-level Management**: Designed to equip executives with essential regulatory knowledge and strategic tools to navigate MDR/IVDR requirements.
- **NoBoCap Short Course on AI-supported Medical Devices**: The first cycle is now closed and will take place on 10-12 February 2025. Pre-registrations are already open for the next cycle - don’t miss your chance to gain important insights into evaluating AI-driven medical technologies.

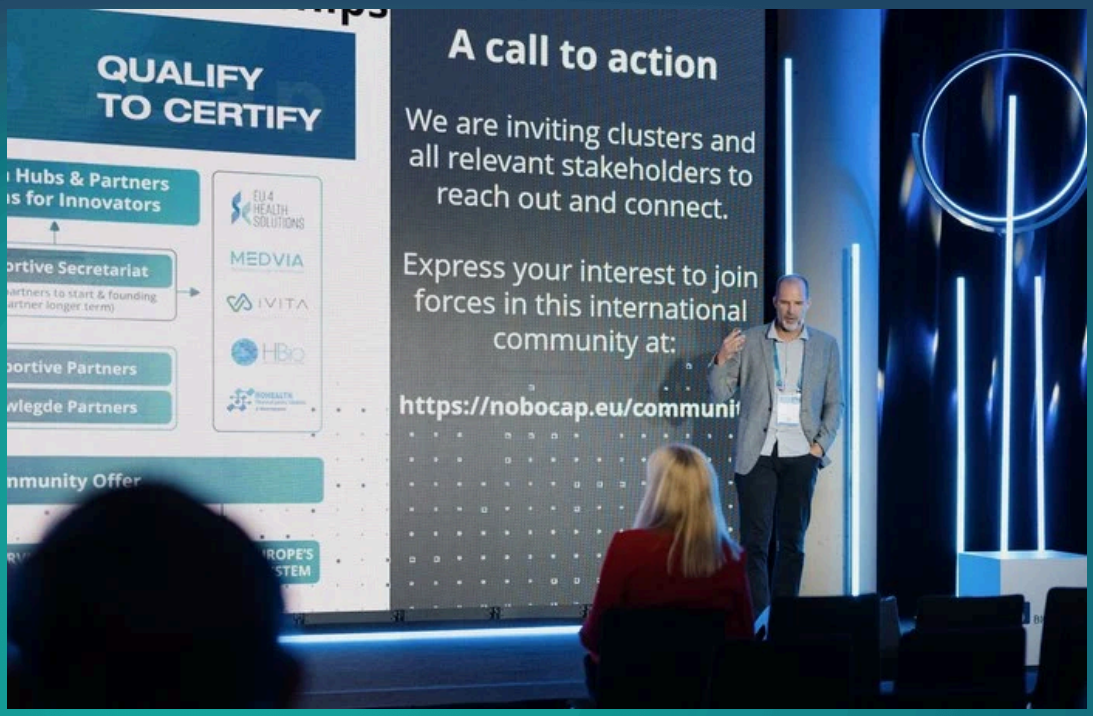
Keep an eye on our updates to find out more about the upcoming sessions and how to register.

Visit and Pre-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

Certification of Medical Devices in Lithuania: Challenges and Opportunities

In a rapidly evolving regulatory landscape, understanding the challenges and opportunities within each country's certification process is key to navigating the European MedTech ecosystem. A recent article on NoBoCap’s website delves into the complexities of medical device certification in Lithuania, shedding light on the unique hurdles faced by innovators and manufacturers.

Lithuania, as a growing hub for MedTech, presents both opportunities and challenges in ensuring regulatory compliance with the European Medical Device Regulation (MDR) and In-vitro Diagnostic Regulation (IVDR). This insightful piece explores the local regulatory environment, highlights the collaboration between notified bodies and innovators, and discusses how Lithuania is

working to streamline its processes while maintaining high standards of safety and efficacy for medical devices.

READ THE FULL ARTICLE HERE to gain a deeper understanding of how Lithuania is positioning itself in the European MedTech landscape and the ways in which companies can navigate these evolving regulations effectively.

Stay informed, stay compliant, and learn more about how NoBoCap is supporting the MedTech community in adapting to these dynamic changes.

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

Happy Winter Holidays!

With joy of life, health and laughs of child!

May the coming year be filled with success, growth, and exciting new opportunities.

Thank you for being a part of the NoBoCap journey!

Merry Christmas! Happy New Year 2025!

NoBoCap

www.nobocap.eu

You received this email because you signed up on our website

Unsubscribe

Co-funded by the European Union

Funded by the European Union. Views and opinions expressed are however those of the author(s) only and do not necessarily reflect those of the European Union or the European Health and Digital Executive Agency (HADEA). Neither the European Union nor the granting authority can be held responsible for them