NoBoCap

ABOUT NOBOCAP

Notified Body Increased Capacity

NoBoCap project – a groundbreaking initiative under the wings of the EU4Health Programme. We are committed to fostering a conducive environment for swift and efficient advancement in the Medical Technology (MedTech) sector.

> Funding Source: European Union

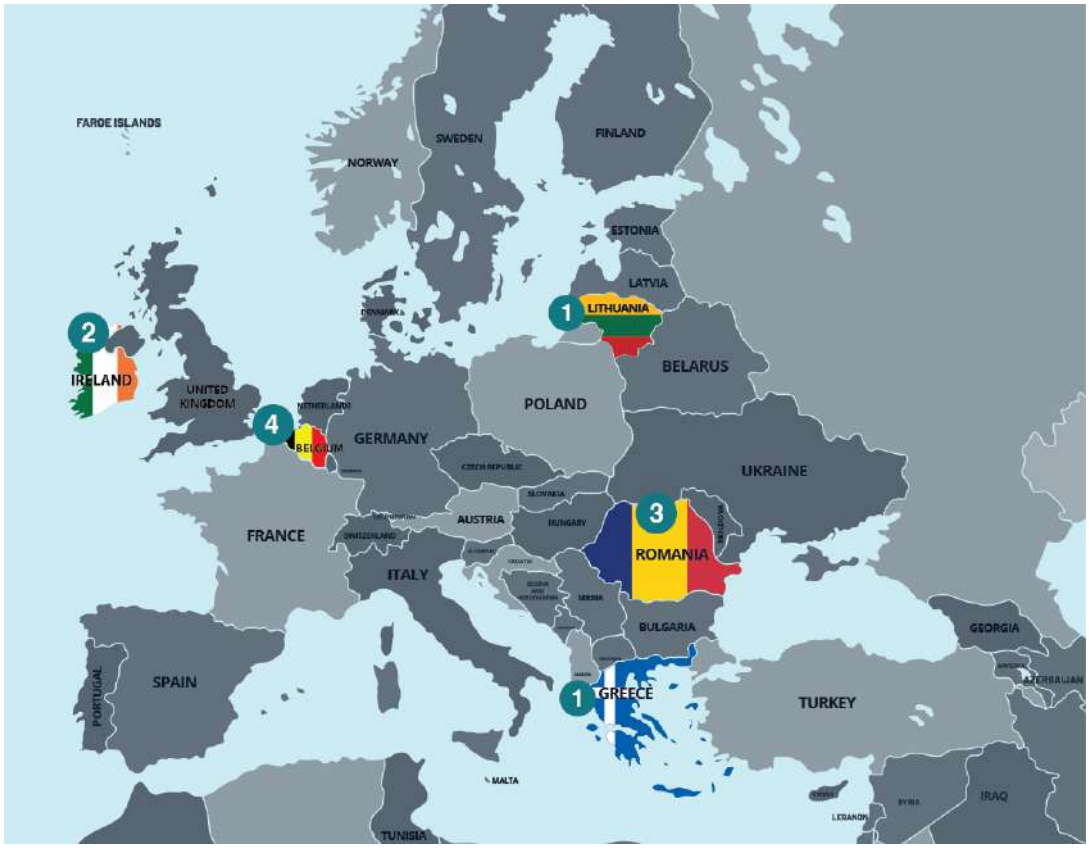
> Programme: EU4Health

> Reference: 1 0 1 1 0 1 2 6 9

Visit our page

Europe working together to qualify to certify!

NoBoCap is a unique consortium of 11 members comprising: 3 academia, 4 health clusters, 2 SMEs, and 2 global leader NBs.



ROHEALTH The Health and Bioeconomy Cluster · DUBLIN · iVITA HEALTH TECHNOLOGY CLUSTER · ALTFACTOR EDUCATIONALLY YOURS · MEDVIA

SGS · GHENT UNIVERSITY · HBIO · EU 4 HEALTH SOLUTIONS

NOBOCAP TRAINING

1st training module/results

The NoBoCap project, recently completed the first cohort of its ambitious MDR (Medical Device Regulation) master's Level Module 1 "Implementing Regulatory Requirements for Medical Devices" led by SGS (NB 1639) and TU Dublin.

Read more

Exciting Developments in Long-Term Educational Programs!

We are thrilled to bring you the latest updates on our range of long-term courses designed to enhance your skills and knowledge in MDR and iVDR. Let's dive into the details:

Module 1 Implementing regulatory requirements (MDR and IVDR) for medical devices - Cohort 3 - Anticipation is building for the start of Module 1 Cohort 3, expected in April 2024 mirroring last year's schedule.

Target Market: Quality, regulatory, and other relevant personnel at manufacturers

Aim: Provide full overview for required implementation of QMS and MDR requirements.

Module: 2. Generating data for technical documentation (MDR)

Mark your calendars! The first cohort for Module 2 is set to commence at the beginning of March 2024

Target Market: Quality, regulatory and other relevant personnel at manufacturers

Aim: Provide full overview for required implementation of QMS and MDR requirements.

Module: 3 – Generating data for technical documentation (IVDR) – it is set to begin in September 2024

Target Market: Quality, regulatory and other relevant personnel at manufacturers.

Aim: Provide full overview for required implementation of QMS and iVDR requirements.

Short-Term Courses and Interventions Update

In addition to our long-term courses, we're also excited to share updates on our short-term courses and interventions:

AI Training Course - Slated to begin between February and March 2024

Stay tuned for more updates as we continue to shape these exciting educational opportunities!

Survey for Market Operators in MedTech/HealtTech sectors

We Need Your Insights for shaping the training curriculum developed under the NoBoCap Project!

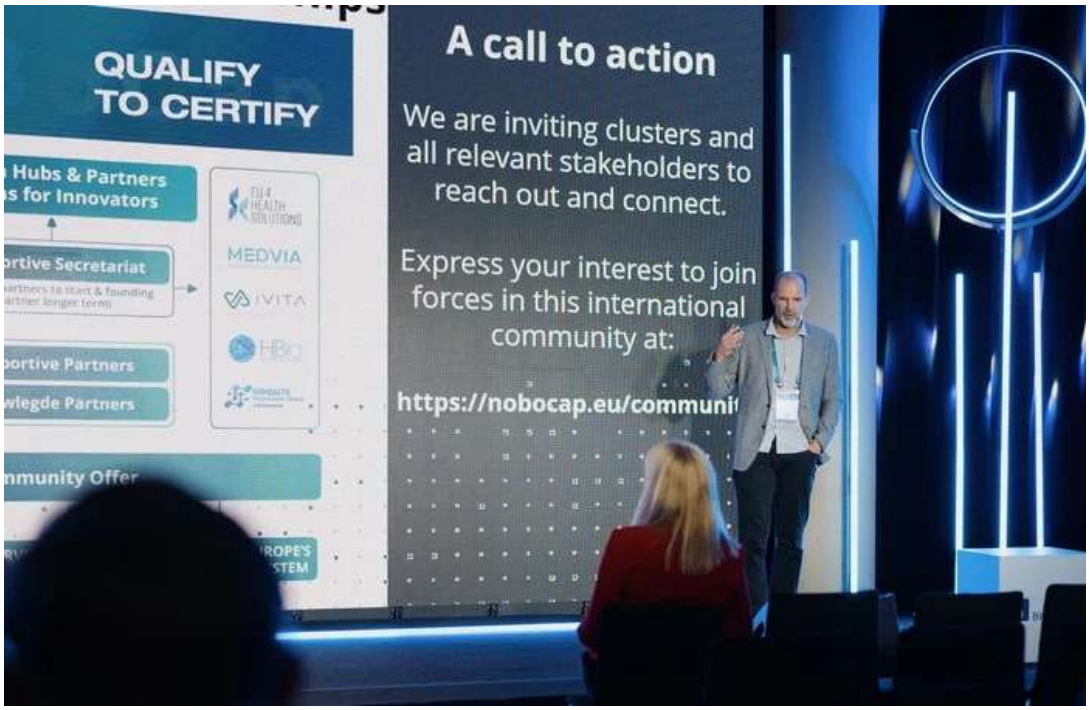
Participate in Our Brief Online Survey!

Your voice is crucial. By taking a few minutes to fill out our anonymous survey, you provide us with essential insights that will directly shape the future of our industry.

Start a survey

NOBOCAP COMMUNITY

The Community of Clusters, Innovation Hubs & Partners in Europe



We are establishing a vibrant NoBoCap community, uniting clusters, innovation hubs, supportive entities, and knowledge partners. This collaborative network will collectively contribute valuable insights and expertise to shape the MDR and IVDR legislation, facilitate training, promote matchmaking, and cultivate a nurturing ecosystem for innovation within the Medtech, Life Science, and HealthTech sectors.

Visit our page

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

Application form

NEWS

Start-ups left hanging in EU medical device regulation snarl-up

Delays in getting the CE mark for medical devices are stretching the resources of European start-ups and forcing some to turn their attention to the US.

Read more

The certification process and increased costs slow the growth of the medical devices market

The European Commission has taken measures to solve these problems.

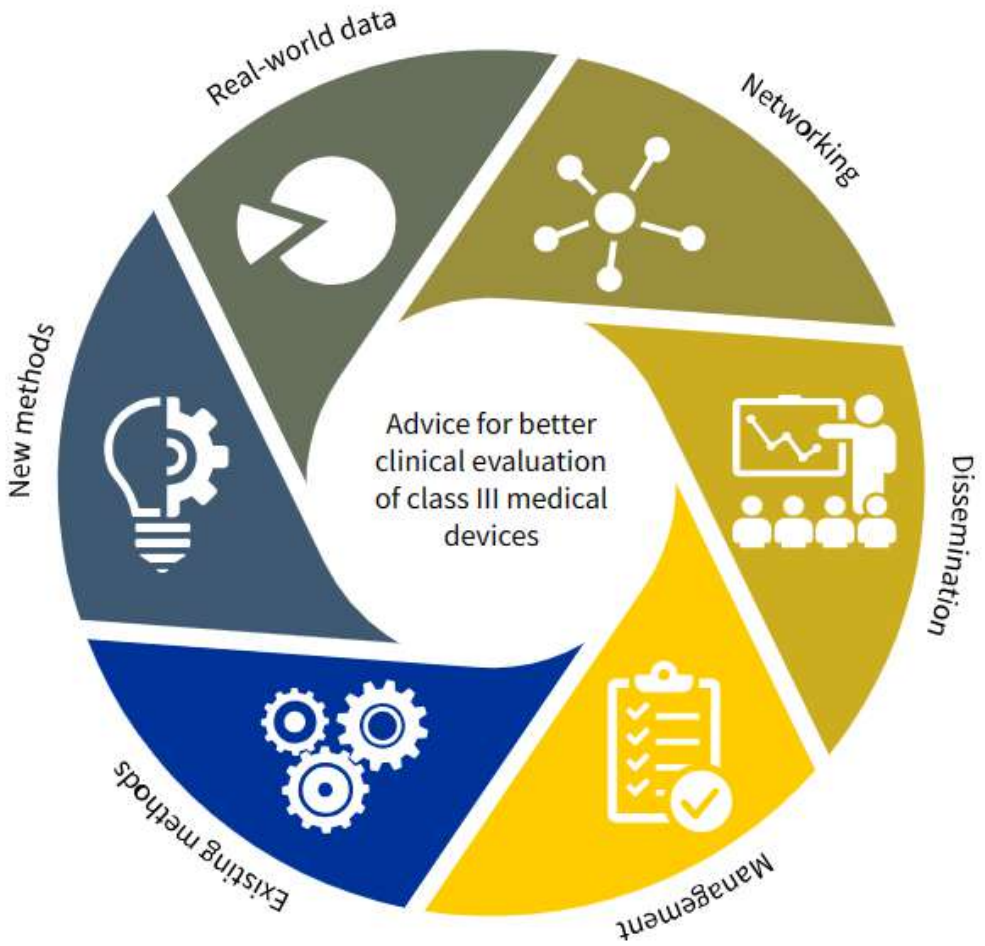
Webinar: Regulation 2023/607 MDD Extension and its Impact on Manufacturers

Join a free of charge webinar to learn about the impact on conformity assessment and marketing authorization of this regulation.

Read more

Read more

INITIATIVES



The Coordinating Research and Evidence for Medical Devices

The NoBoCap project cooperates with CORE-MD, contributing European added value to actions and innovations.

By fostering cooperation with existing related initiatives, we aim to

create a collaborative ecosystem to address challenges in medical devices.

Read more

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 2024!

NoBoCap

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