



EuroMedis



OŚRODEK BADAŃ KLINICZNYCH



PROGRESS

BREAKTHROUGH

INNOVATIONS

1. ABOUT US

2. RESOURCES

3. EXPERIENCE IN CLINICAL TRIALS

4. OPERATING RULES

1 ABOUT US



- **1997**
Prof A. Potemkowski establishes Individual Specialist Neurological Practice.
- **1999**
Prof A. Potemkowski signs a contract with NHS.
- **2000**
Prof A. Potemkowski conducts first SM clinical trials.
- **2004**
Dr M. Ratajczak and dr A. Ratajczak join Prof A. Potemkowski research team.
- **2005**
Dr. J. Kolenda joins the research team. Dr M. Ratajczak establishes the Individual Doctor's Practice.
- **2006**
The establishment of Memory Disorder Centre.
- **2009**
The establishment of EuroMedis. Transfer of the NHS contract to EuroMedis. Extension of the NHS contract. Expanding the scope of Clinical Trials. Adding Specialist Clinics to our offer.
- **2014**
Relocation to the new head office- further development of the Centre.
- **2021**
Opening a 24-hour diagnostic department.

LOCATION

- The nearest airport - Goleniów port serves domestic and international flights (UK, Scandinavia). The transfer to the airport takes about 40 minutes from Szczecin.
- Airport Berlin Schoenefeld - 2 hours by car from Szczecin.



OUR MISSION

We believe that by providing health services focused on the needs of the patient and his family, we improve the quality of life, and because of our commitment to the development of new ideas and technologies we contribute to overcoming the limitations of current medicine.



OUR VISION

We want our services to be widely available, comprehensive and improve the quality of life so that everyone can enjoy the health.





ORGANIZATION



EuroMedis



**Over 50 specialists,
2 600 m² of space in two locations.**

EuroMedis Medical Centre provides own diagnostics facilities.

Sampling for laboratory analysis, Ultrasonography, EEG, ECG with Holter monitoring and ergospirometry are all provided onsite.

Patients are also provided with a fully equipped ophthalmological laboratory where, for instance, perimetry is performed.

EEG**USG****Doppler
USG****ECG****RR
Holter****ECG
Holter****Perimetry****Ergospirometry**

DIAGNOSTICS

Within external contracts, the facility may carry out the following examinations: Computed Tomography, Magnetic Resonance Imaging, X-rays, PET, endoscopy, spirometry or other laboratory test.

The terms of the contracts determine the fast time to complete the procedures, which facilitates diagnosis and eventual AE checking at a rapid pace in patients from clinical trials.

CT**MRI****PET****X-RAY****LAB****OCT****Spirometry****Endoscopy**

2 RESOURCES



CLINICAL TRIAL TEAM



14

RESEARCHES

8

COORDINATORS

5

NURSES

6

RATERS

CONSTANT RESEARCHES TEAM



prof. dr n. med. Andrzej Potemkowski

neurologist

22 years of experience in
clinical trials/
67 clinical trials



dr n. med. Marcin Ratajczak

neurologist

16 years of experience in
clinical trials/
78 clinical trials



lek. Anna Ratajczak

pediatrician

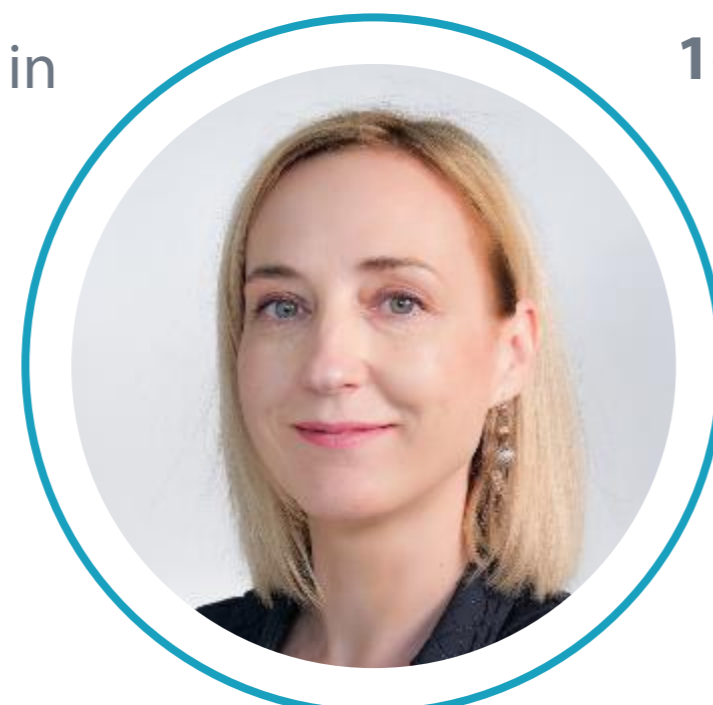
16 years of experience in
clinical trials/
47 clinical trials



lek. Patrycja Sendorek

neurologist

14 years of experience in
clinical trials/
34 clinical trials



dr n. med. Aneta Wódecka

neurologist

8 years of experience in
clinical trials/
20 clinical trials



lek. Joanna Adamiak

psychiatrist

4 years of experience in
clinical trials/
6 clinical trials



dr n. med. Lilianna Osowicz-Korolonek

endocrinologist

4 years of experience in
clinical trials/
3 clinical trials

COORDINATORS

**mgr inż. Nina Omelańska**

8 years of experience in
clinical trials /
55 clinical trials

**mgr inż. Karolina Sorbian-Gajewska**

8 years of experience in
clinical trials /
59 clinical trials

**mgr Beata Miąsek**

8 years of experience in
clinical trials /
41 clinical trials

**mgr Magdalena Draj**

6 years of experience in
clinical trials /
22 clinical trials

**mgr inż. Agata Piłatowska**

5 years of experience in
clinical trials /
36 clinical trials

**mgr Jakub Burdanowski**

4 years of experience in
clinical trials /
28 clinical trials

**mgr Maciej Kaczmarek**

4 years of experience in
clinical trials /
30 clinical trials

**dr n. med. Małgorzata Ryder-Woźniak**

2 years of experience in
clinical trials /
20 clinical trials

NURSES



Izabela Zamilska

12 years of experience in
clinical trials /
72 clinical trials



mgr Anna Kozińska

7 years of experience in
clinical trials /
54 clinical trials



mgr Jolanta Rudzińska

6 years of experience in
clinical trials /
38 clinical trials



mgr inż Natalia Laskowska

4 years of experience in
clinical trials /
28 clinical trials



mgr Kinga Bakalarczyk

2 years of experience in
clinical trials /
17 clinical trials



mgr Paulina Ceglaz

2 years of experience in
clinical trials /
3 clinical trials

RATERS**dr n. med. Jerzy Kolenda**

14 years of experience in
clinical trials /
52 clinical trials

**mgr Dąbrówka Miler**

7 years of experience in
clinical trials /
17 clinical trials

**mgr Agata Zapala**

7 years of experience in
clinical trials /
15 clinical trials

**mgr Emilia Bastrzyk**

5 years of experience in
clinical trials /
10 clinical trials

**mgr Malwina Prajwowska**

4 years of experience in
clinical trials /
11 clinical trials

**mgr Anna Iwińska**

2 years of experience in
clinical trials /
7 clinical trials

QUALITY

- **SOP (STANDARD OPERATING PROCEDURES)** – A SET OF DETAILED WRITTEN INSTRUCTIONS, STEP BY STEP, TO ACHIEVE THE UNITY OF A SPECIFIC FUNCTION
- **QUALITY CONTROL**



We have certified equipment with a technical passports and full documentation.

- **FREEZER UP TO (-80°C)**
- **FREEZERS UP TO (-25°C)**
- **LABORATORY CENTRIFUGES**
- **COOLED CETRIFUGE UP TO DO 2°C**
- **REFRIGERATORS FROM 2°C TO 8°C**
- **ECG**
- **DEFIBRILLATOR**
- **SPHYGMOMANOMETERS**
- **INFUSION PUMPS**
- **WEIGHT WITH STADIOMETER**
- **LAMINAR CHAMBER**



SECURITY

Our facility is also equipped with electronic system for temperature monitoring in medicines storehouse. The test product as well as biological samples are monitored.



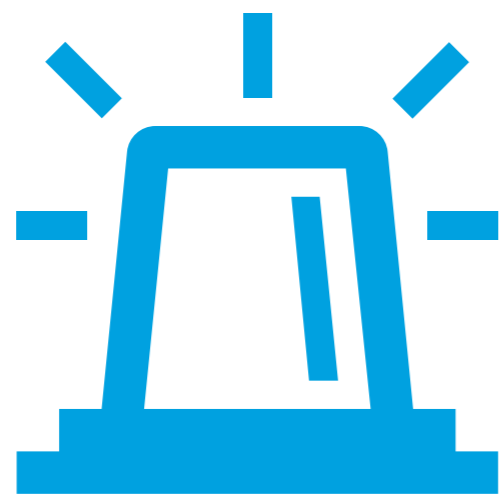
The measurement is performed for 24 hours and notification about exceed is sent via text message or e-mail. **The measurement is supervised by the rapid response team (up to 1 hour after the alarm occurred).** It is possible to print reports on temperature control of any period of time.





SECURITY

The facility has a power generator in the event of a power failure.



3 EXPERIENCE IN CLINICAL TRIALS



EM

AREAS OF CLINICAL TRIALS

NEUROLOGY

PSYCHIATRY

DERMATOLOGY

OPHTHALMOLOGY

NEPHROLOGY

GASTROENTEROLOGY

DIABETOLOGY

PRIMARY HEALTH CARE



EM

COOPERATION

WE ARE OPEN TO NEW AREAS
OF CLINICAL TRIALS!



EM

EXPERIENCE

91

CONDUCTED RESEARCH

9

TOP RECRUITER



EM

EXPERIENCE

1604

RECRUITED PATIENTS



EM

EXPERIENCE

1648

PATIENTS SUBJECTED
TO SCREENING



The logo consists of the letters 'EM' in a bold, white, sans-serif font, enclosed within a blue square.

EXPERIENCE

24

COOPERATING SPONSORS

19

COOPERATING CRO



EM

EXPERIENCE

15

AUDITS

1

FDA AUDIT

APPROVED

EM

EXPERIENCE

18

CURRENTLY CONDUCTED
CLINICAL TRIALS

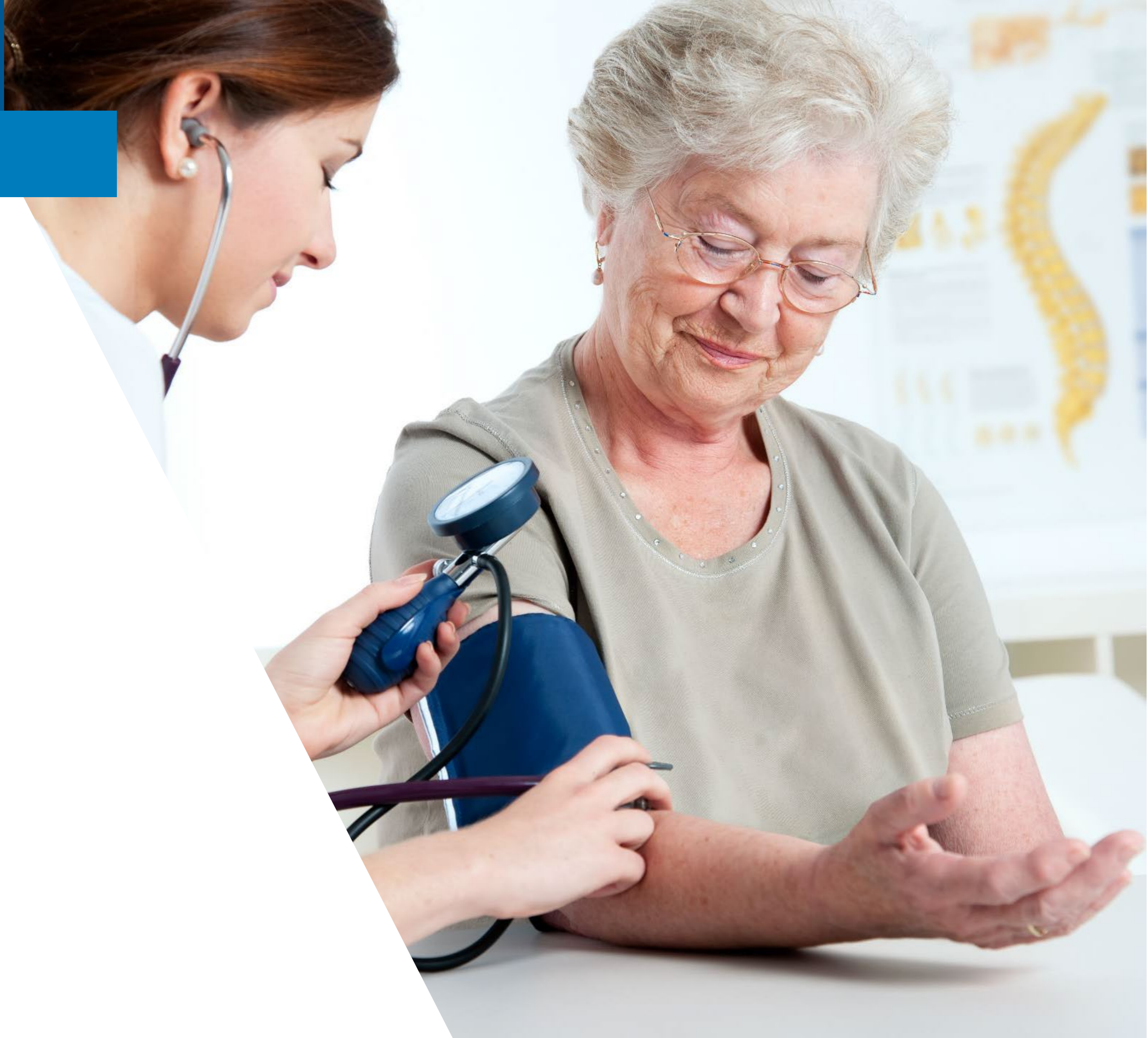


The logo consists of the letters 'EM' in a bold, white, sans-serif font, set against a dark blue square background.A horizontal blue banner with the word 'EXPERIENCE' in white, uppercase, sans-serif font.The number '157' in a large, bold, blue, sans-serif font.

PATIENTS IN CURRENTLY
CONDUCTED TRIALS

The number '250' in a large, bold, blue, sans-serif font.

MONTHLY VISITS



4 OPERATING RULES



REGULAR MEETINGS

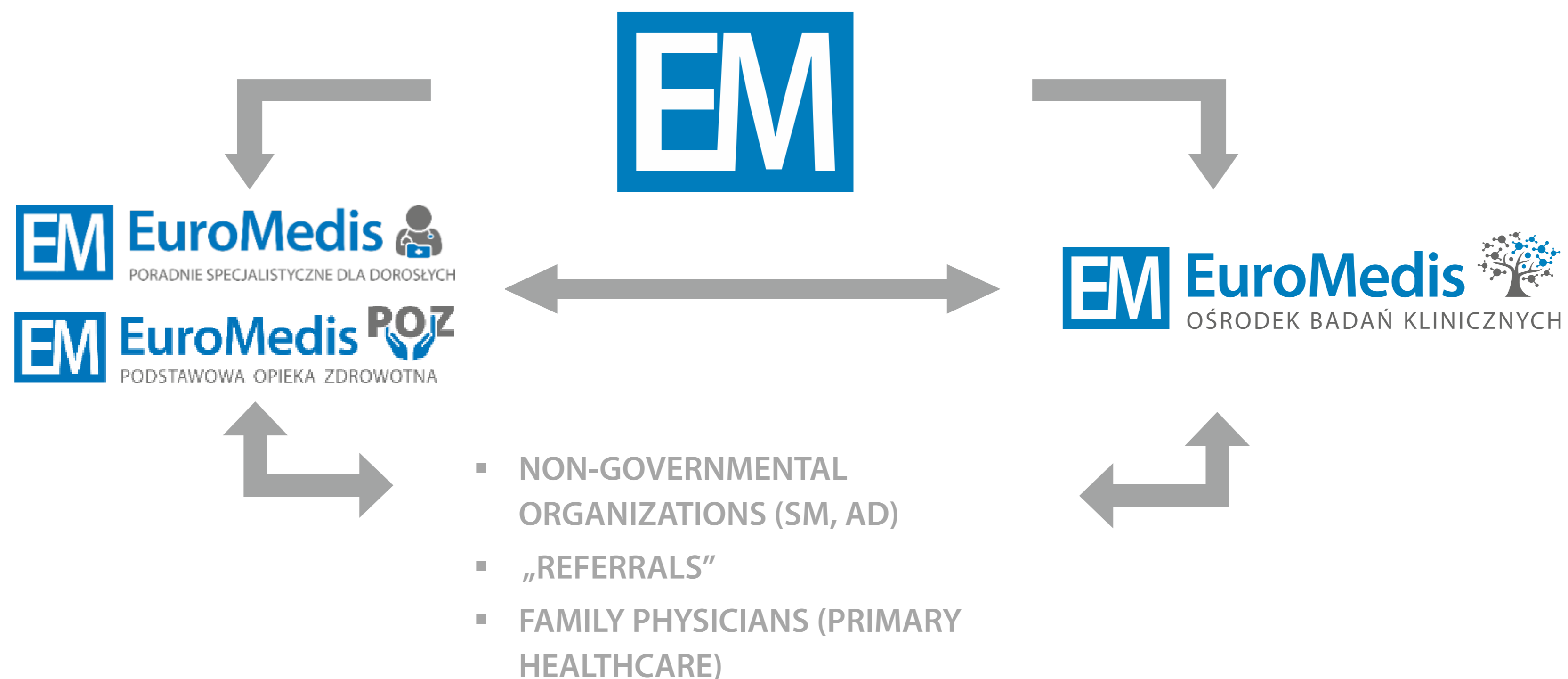


- DISCUSSION OF EVERY INDIVIDUAL PATIENT'S CASE AND RELATING ISSUES
- OVERVIEW OF THE COURSE OF EACH CLINICAL TRIAL AND THE RELATED PROBLEMS
- TRAININGS

STRATEGIES IN RECRUITMENT

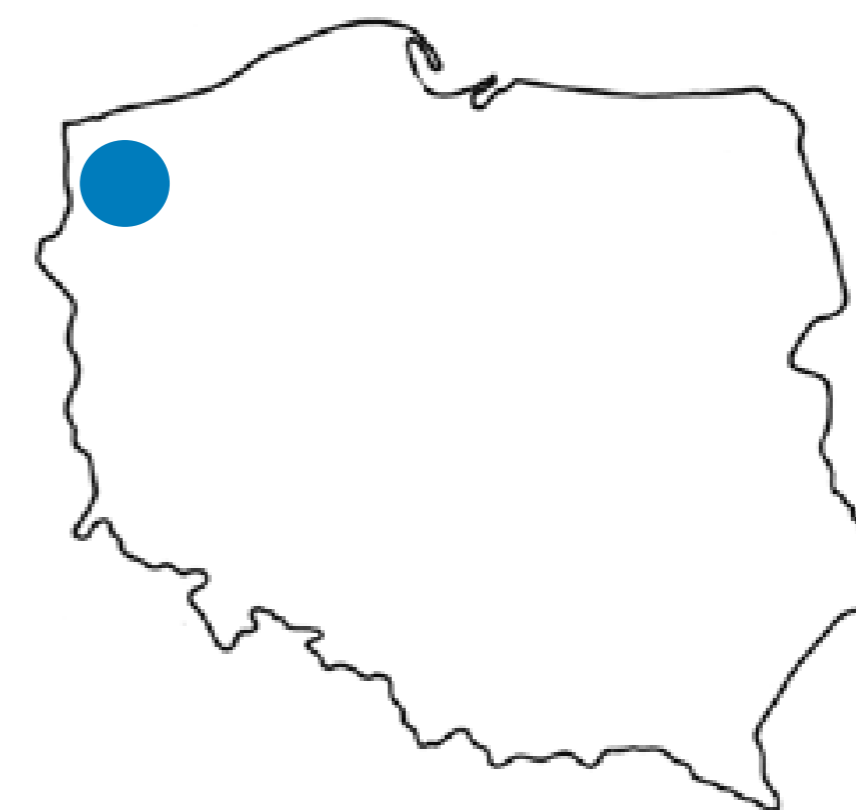
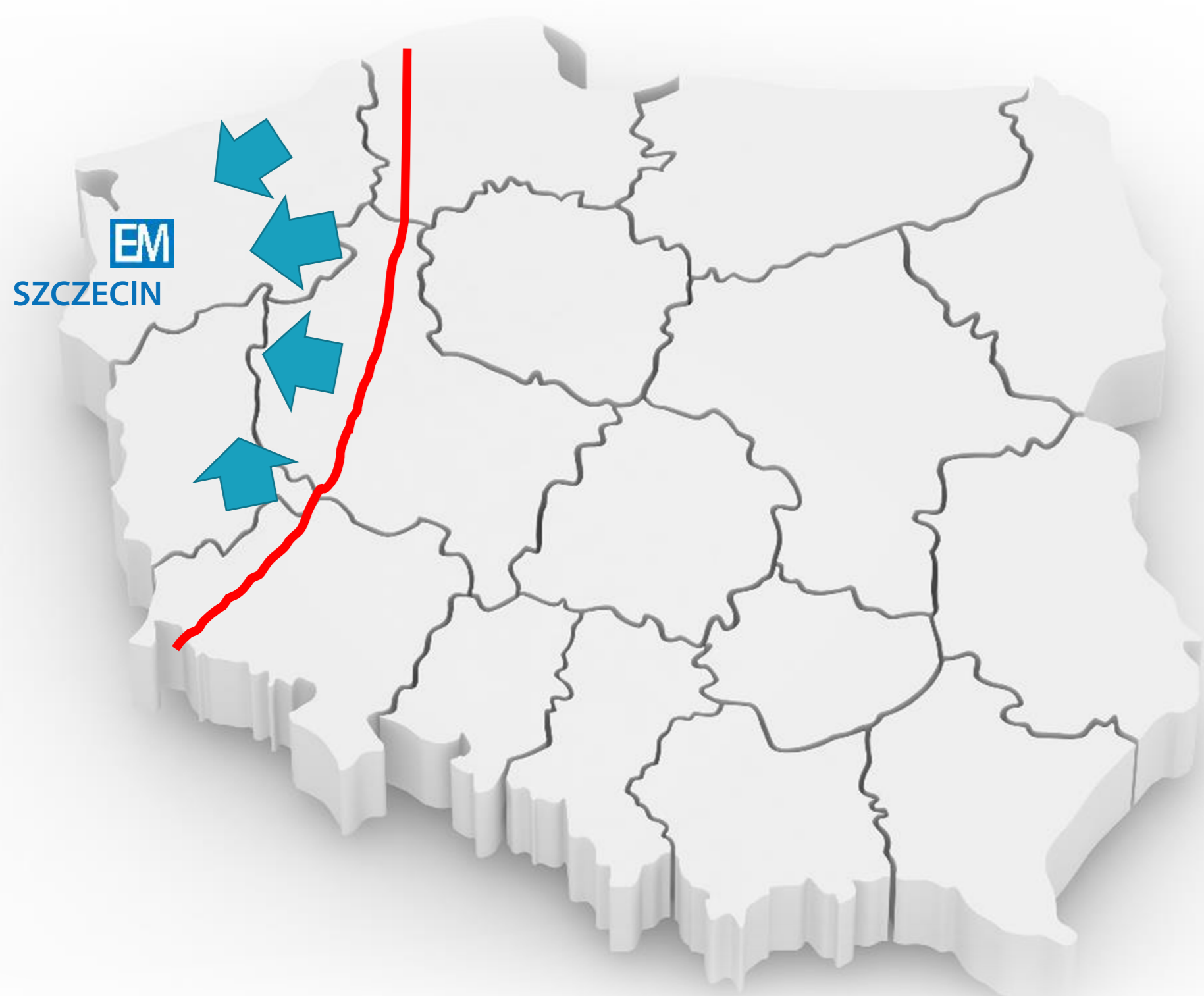
Connecting the medical activity provided by Specialistic Outpatient Clinics and Clinical Trial Centre in one facility enables constant flow of patients, it includes also GP's, 'referrals' and associations dealing with diseases on which they conduct clinical trials. It considerably increases recruitment possibilities from own database of patients, however, it also enables further care provided by the very same specialists once their participation in the trial has ended.

All patients are informed about the possibility of participating in clinical trials as a treatment option. The patients who meet the criteria of being included in the trial and still are willing to take part, are invited to a special visit where all procedures and ICF – Informed Consent form are discussed. After the visit and patient's consent, we start the procedures connected with performing the given examination



POPULATION

- Region of north-west Poland: **3.000.000 people**
- Szczecin and adjacent poviats: **600.000 people**



STRATEGIES IN RECRUITMENT

EuroMedis runs a website devoted to clinical trials as well as numerous information campaigns in social media. We also prepare our own information materials regarding clinical trials.

EuroMedis is also an organizer of scientific seminars for patients and their guardians as well as scientific and information meetings for doctors.



www.EuroMedis.pl /
portal for doctors /
social media

information
materials

direct meetings/
conferences

FUTURE PLAN

- ELEKTRONIC MEDICAL DOCUMENTATION WITH REMOTE ACCESS TO SD VERIFICATION FOR CRA
- OWN MRI
- OWN WŁASNA ENDOSCOPY





WHY EUROMEDIS CLINICAL TRIAL CENTER?

- OVER 15 YEARS OF EXPERIENCE ON THE MARKET
- SPECIALIZED MEDICAL STAFF
- EXPERIENCED CLINICAL TRIAL COORDINATORS
- DEDICATED TEAM FOR CONTACT WITH REFERRALS
- FULL DIAGNOSTIC AND LABORATORY FACILITIES
- 24/7 DIAGNOSTIC DEPARTMENT FOR 1 PHASE OF CLINICAL TRIALS
- MODERN TECHNOLOGICAL SOLUTIONS – 24h TEMPERATURE MONITORING SYSTEM
- TELEPHONE CONTACT - 24h AND COMPANY TRANSPORT FOR THE PATIENTS OF THE FACILITY
- QUALITY DEPARTMENT

BADANIA KLINIC



C O N T A C T U S

WWW.EUROMEDIS.PL



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