

**QUALIblood**

Your expert in blood testing



## WE OFTEN HEAR...

*“ BIOLOGICAL INVESTIGATIONS IN HUMAN SAMPLES IS SO COMPLEX AND A PLENTY OF TESTS ARE SO CONSTANTLY RENEWED THAT A CONTINUOUS EXPERTISE IN THIS FIELD IS HARD TO REACH”*










*“I RISK MY TIME AND MONEY IN INEFFICIENT OR OUTDATED BLOOD TESTING. COULD A SPECIFIC LABORATORY HELP ME IN THE CHOICE OF THE MOST APPROPRIATE BIOMARKERS TO ENSURE TRANSFERABILITY TO REAL-WORLD APPLICATIONS?”*




# BECAUSE EVERY RESULT MATTERS, **QUALIBLOOD** IS OFFERING TO PHARMACEUTICAL AND IVD COMPANIES A TAILOR-MADE ACCOMPANIMENT IN BLOOD ANALYSES




## ASSISTANCE IN PROTOCOL DEFINITION

-  DETERMINATION OF TESTING STRATEGY TO AVOID UNNECESSARY TESTING
-  ADVICE OF (PRE)ANALYTICAL ISSUES ACCORDING TO LABORATORY STANDARDS
-  ELABORATION AND WRITING OF LABORATORY MANUAL TO ENSURE SAMPLE INTEGRITY
-  PROVISION OF INDIVIDUAL COLLECTION DEVICES IN ALL STUDY SITES ACCORDING TO STUDY DESIGN
-  SHIPPING, REPATRIATION AND HANDLING OF BIOLOGICAL SAMPLES
-  STORAGE OF BIOLOGICAL SAMPLES
-  ANALYSES OF BIOLOGICAL SAMPLES ACCORDING TO GCLP GUIDELINES



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 FOLLOW-UP AND CLOSE INTERACTIONS AFTER THE DELIVERY OF THE RESULTS

-  RESULTS INTERPRETATIONS
-  REPORTS ACCORDING REGULATORY REQUIREMENTS
-  PRESENTATION OF RESULTS FOR PUBLICATIONS AND MARKETING PURPOSE



# FACILITIES FOR PHARMACEUTICAL COMPANIES

## DEFINITION OF ANALYTICAL PROTOCOLS

- ACCOMPANIMENT IN THE REDACTION OF THE PROTOCOL
- ADVICE AND LITERATURE REVIEW FOR DETERMINATION OF MOST APPROPRIATE BIOMARKERS

## PRE-CLINICAL INVESTIGATIONS – GRP AND/OR GLP ENVIRONMENT

- IN VITRO INVESTIGATION ON PLASMA, SERUM, WHOLE BLOOD OR CELL LINES
- TAILOR-MADE SOLUTIONS AND TESTING VALIDATED ACCORDING TO REGULATORY REQUIREMENTS
- BLOOD TESTING IN SAMPLES FROM ANIMAL STUDIES

## CLINICAL INVESTIGATIONS – GCLP ENVIRONMENT – TESTING ACCORDING TO ISO 17025

- BLOOD TESTING OF SAMPLES FROM HUMAN STUDIES – PHASE I TO IV
- CENTRAL LABORATORY SERVICES – LAB KITS MANUFACTURING, SAMPLE MANAGEMENT, TESTING, BIOBANKING

## BATCH RELEASE – GLP ENVIRONMENT

- EVALUATION OF THE QUALITY OF PLASMA-DERIVED MEDICINAL PRODUCT ACCORDING TO REGULATORY REQUIREMENTS



# MORE THAN 400 DIFFERENT TESTS AVAILABLE IN THE FIELD OF

 HEMOSTASIS

 CYTOKINES AND CHEMOKINES

 CHEMISTRY

 ENZYME SPECIFICITY TESTING

 SEROLOGY

 HEMATO-ONCOLOGY

 COMPLEMENT SYSTEM

For more information on available analyses –  
contact us at

<http://www.qualiblood.eu/contact>



# WHO ARE WE?

- SPIN-OFF FROM THE UNIVERSITY OF NAMUR IN CLOSE COLLABORATIONS WITH LOCAL AND FOREIGN HOSPITALS
- 10 YEARS OF RESEARCH IN HEMATOLOGY WITH A FOCUS ON HEMOSTASIS AND HEMOCOMPATIBILITY – MORE THAN 100 PUBLICATIONS IN THE FIELD
- GATHERING OF EXPERTISE FROM LABORATORY, CLINICAL AND REGULATORY SCIENTISTS



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# WHO ARE WE?

 CEO & CSO

JONATHAN DOUXFILS

 CFO & HR DIRECTOR

BERNARD MASEREEL

 QUALITY MANAGER

SABRINA MELCHIONDA

 LABORATORY MANAGER

CÉLINE BOUVY

 PROJECT LEADERS & ASSAY DIRECTORS LAURE MORIMONT

CLARA DAVID

 LABORATORY TECHNOLOGISTS

ÉLISE MODAFFARI

+ 7 PHD STUDENTS



# WHO ARE WE?

- MEMBERS AND CHAIRMAN OF DIFFERENT SUBCOMMITTEES AT THE INTERNATIONAL SOCIETY OF THROMBOSIS AND HEMOSTASIS
- EXPERT AT THE INTERNATIONAL COUNCIL FOR STANDARDISATION IN HAEMATOLOGY
- EXPERT AT THE ECAT FOUNDATION
- REGULATORY EXPERTISE AND NETWORKING



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# EXAMPLES AND SUCCESSFUL PROJECTS

DEVELOPMENT OF DRUGS CANDIDATES



GEDEON RICHTER



EVALUATION OF DIAGNOSTIC TOOLS



ADVICE AND REGULATORY CONSULTANCY



EUROPEAN MEDICINES AGENCY  
SCIENCE MEDICINES HEALTH



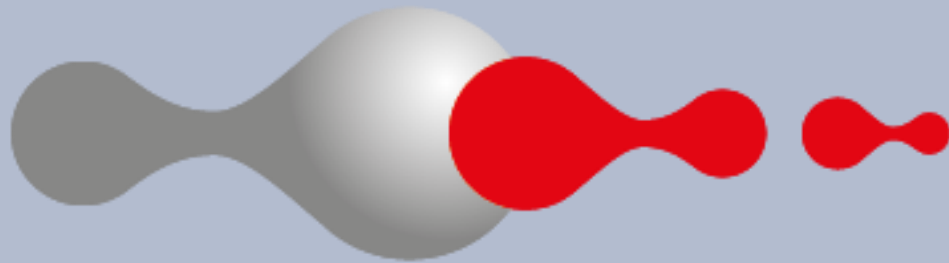
Daiichi-Sankyo

IN BUSINESS WITH PHARMACEUTICAL AND DIAGNOSTIC COMPANIES









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