



Berlin Longterm Observation of Vascular Events

STEERING COMMITTEE

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SHORT SUMMARY

The objective of BeLOVE is to retinopathy, kidney dysfunction) participation of basic and clinical identify novel pathophysiological on risk and prognosis of these scientists as well as clinical cross-disease mechanisms and vascular diseases as well as epidemiologists from BIH, Charité, treatment targets that determine effects of specific/individual and MDC will promote the short and long-term vascular events on remote organs identification of cross-disease both outcome of high-risk patients with (e.g., neurocardiogenic damage mechanisms (e.g., contribution of acute vascular disease (i.e., acute after stroke, cognitive deficits gut microbiome, immune system, cerebrovascular disorder, acute after acute cardiovascular events. metabolic factors, stress, vascular coronary syndrome, acute heart BeLOVE will provide a uniform function, (epi)genetic factors).standardized backbone BeLOVE serves as a unique failure, and acute kidney injury) and an phenotyping of all patients, but resource of integrated diabetes. With high and interdisciplinary approach and enables different levels of quality biomedical data and deep phenotyping, BeLOVE will participation and combination biomaterials open to the BIH investigate the impact of systemic with modules for additional community for testing of current disease-specific testing. Broad and future research hypotheses. factors and co-morbidities (e.g.

G. Rauch/ K. Schmidt-Ott /

J. Schulz-Menger / B. Singerink / J. Spranger

Recognize risks. Understanding interrelations.



THE GENERAL CONCEPT

Aims

Improve prediction and mechanistic understanding of cardiovascular disease progression and outcomes in patients in very high risk in order to provide the basis for improved, individualized management.

BERLIN

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Strategy

Combining cutting edge phenotyping technologies with short- and long-term observation in patients with distinct cardiovascular disease manifestations to identify

Figure 1: General concept of BeLOVE. (Acute heart failure (AHF), Acute coronary syndrome (ACS), Acute cerebrovascular disorders (stroke), Acute kidney injury stage 2 or 3 lasting ≥72h (AKI), Type 2 diabetes mellitus (T2DM)

- a) unifying and divergent mechanisms of disease progression
- b) endogenous protective mechanisms
- c) risk factors
- d) novel therapeutic targets
- e) cross-organ interactions
- f) methods of improved risk prediction based on complex data analysis

INNOVATION

BeLOVE is open to new research hypotheses developed by the broad scientific community of the BIH, Charité, and MDC. However, it focuses on three overarching themes:

 Disease overarching mechanisms (e.g., microbiome & immune homeostasis, body composition & metabolic function)

METHOD



- 2. Risk prediction & target identification (e.g., early vs. late, risk stratification, cross-over risk, comprehensive risk models, machine-learning techniques)
- Interaction between diseases and organ systems
 (systemic effects of vascular events, organinteraction, effects on remote organs)

AKI CONTROLETOR						
T2DM		Questionaires Endpoints	Deep Phenotyping	Questionaires Endpoints	Deep Phenotyping	Questionaires Endpoints
Acute/Short-Term Follow-Up			Chronic/Long-Term Follow-Up			
Acute Events Day 0 Chronic (T2DM)	Baseline 3–7 days	Day 60	Day 90	1 Year	2 Years	3 Years to 10 Years

Figure 2: Study design. Follow-up timeline and procedures of the BeLOVE initiative







