Your expert partner for

Dermatological Research

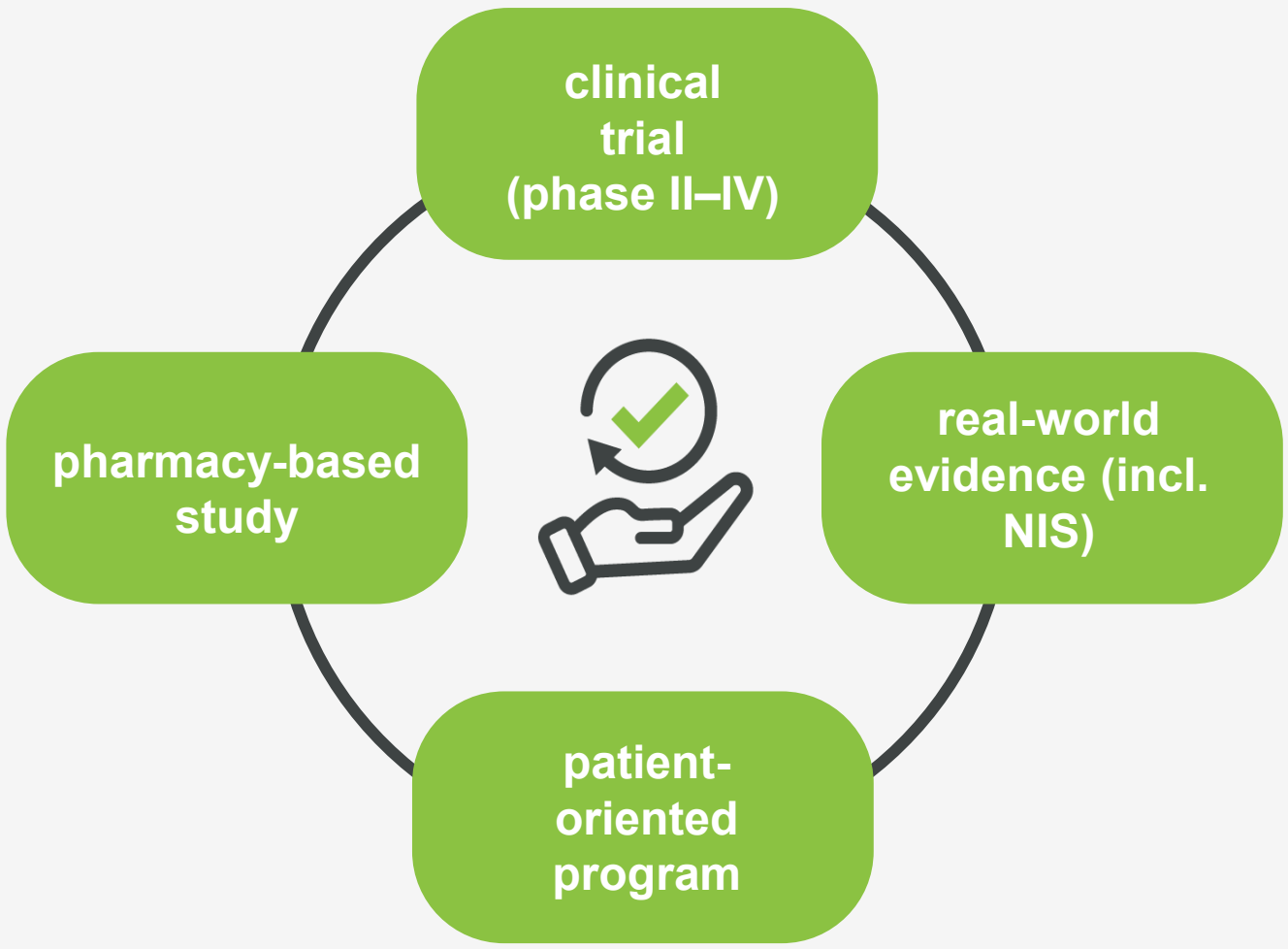
Winicker Norimed GmbH
Clinical Research

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Dermatology

Studies that turn data into value

You want to conduct a dermatology study? We got you covered.



Who are we?

We are Winicker Norimed – a CRO with more than 30 years of expertise in patient surveys and extensive experience in clinical research (phase II to IV as well as non-interventional studies) with a strong focus on dermatology and its manifold indications.

Expertise you can trust

With years of specialized experience in dermatology research, Winicker Norimed delivers high-quality study design, execution and evaluation. Our team combines scientific precision with a deep understanding of regulatory and clinical requirements as well as biostatistics and safety management.

Innovative systems & modern documentation tools

We integrate the latest technology into every study. We continuously develop new concepts for study methodology, participant engagement and data quality enhancement – making your dermatological studies more dynamic, efficient and valuable.

A strong network of specialists

We collaborate with an extensive network of dermatologists, clinical specialists, and study sites - ensuring fast recruitment, reliable data, and consistently high study standards.

Dermatological Study Experience

Dermatology studies — where expertise meets innovation



Plaque psoriasis:

- Non-interventional studies with 70-150 sites and up to 950 patients
- Paper-based surveys
- Use of questionnaires for patient-reported outcomes (e.g. DLQI, ActiPso, PBI, itch and skin pain NRS) incl. licensing
- Photo documentation according to clinical practice to record the change of skin lesions throughout the study



Psoriasis arthritis (PsA):

- Patient-oriented programs with ~70 sites and up to 600 patients
- Non-interventional studies with 130 sites and up to 3.400 patients
- Clinical trials (among others phase III) with ~50 sites and about 300 patients
- Complex eCRF documentation of multiple study arms for over 6 years
- Diagnostic sonography with portable ultrasound devices (incl. logistics and provision of tablets)

Atopic dermatitis:

- Non-interventional studies with ~60 sites and up to 500 patients
- Analysis of a wide range of patient-reported outcomes (e.g. POEM, RECAP, PP-NRS, TSQM, WHO-5) and physician assessments (e.g. EASI, BSA, vIGA-AD)

Dermatological Study Experience

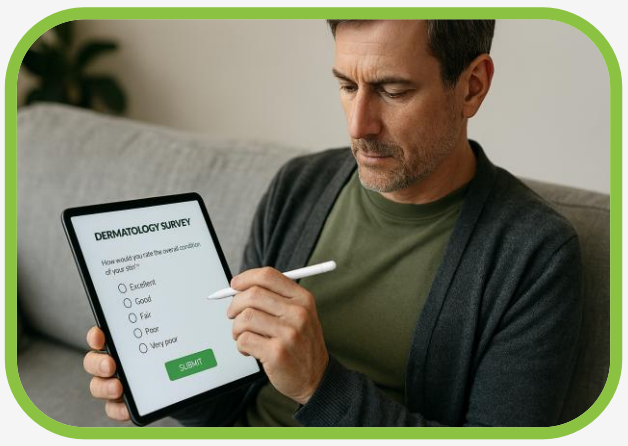
Inflammatory skin disease:

- Pharmacy-based studies with over 1.200 pharmacies and up to 3.000 patients included
- Direct-to-patient-approach: Easy access via link / QR code on own device to complete questionnaires
- Strong and fast enrollment using online advertisements



Infantile haemangioma:

- Retrospective drug utilisation study (PASS) with ~20 sites and around 250 patients
- Straightforward paper-based documentation at sites and data entry into the analysis database by our experienced team



Vitiligo:

- Non-interventional studies with ca. 50 sites and 600 patients
- State-of-the-art documentation options for sites through our long-standing partners in the field of eCRF services

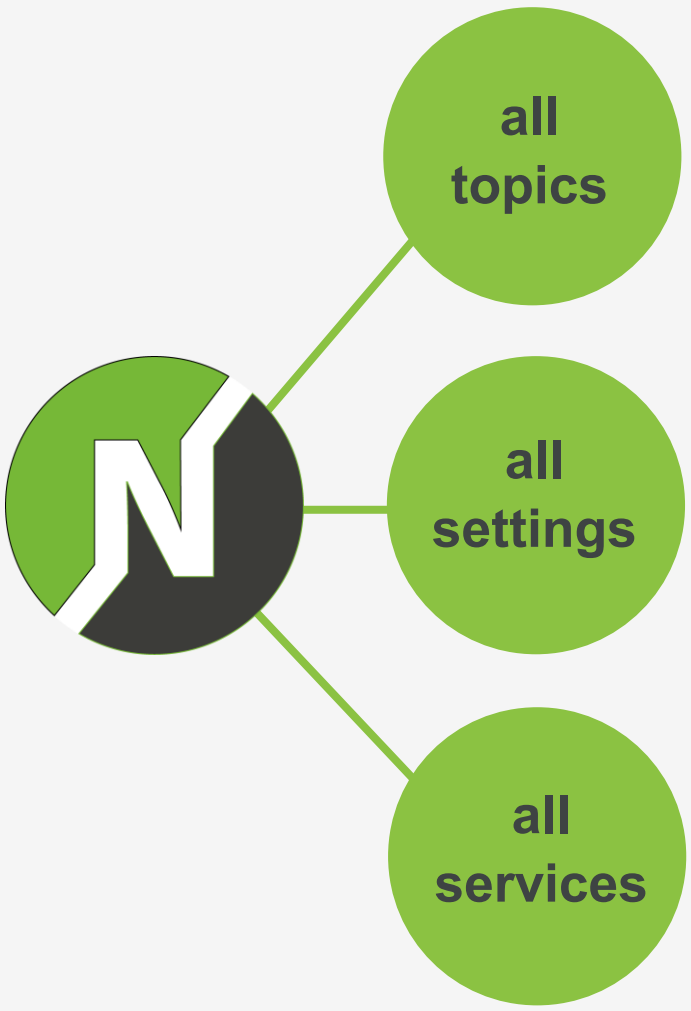
Scabies:

- Clinical trials (phase III and IV) with about 20 sites and up to 300 patients
- Patient diaries for real-time documentation of current state of health enabling prompt evaluation of safety signals

Actinic keratosis:

- Clinical trials (phase III) with around 10 sites and 400 patients

What we offer



- Pharmaceuticals
- Medical devices
- General studies

- Pre- & post-marketing authorization
- Pre- & post-CE marketing
- Hospitals, physicians, pharmacies, patients

- Full service
- Individual services
- Collaborations
- Certified GCP and MPDG trainings

We are a full-service CRO for your projects from start to finish and also provide highly specialized single services like:

Biostatistics:

- Biostatistical consulting and analysis
- Processing of secondary health and tracking data
- Informative tables and figures for presentations and publications
- Statistical review of inputs for protocols, manuscripts and clinical study reports

Medical writing:

- Support and consulting on the structure and content design of study documents
- Interim and final reports (incl. detailed methodology, results, and interpretation)
- AMNOG dossiers
- Safety writing (e.g. ASRs, DSURs, PSUR, PBRER, RMP)
- Posters, abstracts and publications

Safety & vigilance:

- Support in all areas (pre- and post-authorisation / CE marking)
- Creation & maintenance of safety-relevant documents acc. to regulatory requirements
- Case processing including follow-up & authority submission
- Data entry and maintenance in safety databases (e.g. EudraVigilance, ARGUS)
- Signal detection & evaluation

In addition to our expert knowledge in terms of content and our many years of experience, we offer you the advantages of a medium-sized full-service CRO.

You can count on us – just like various clients from the areas of pharma, biotechnology and medical devices already do!

What we offer

- A highly qualified and experienced team
- Independent work in close coordination with the client
- Strict adherence to timelines − we know this is essential
- Willingness to take on new challenges

We look forward to your requests!



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