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PRODUCT REGISTRATION

(Phase I-IIb)

SERVICES

EARLY

Through our harmonized clinical and laboratory facilities in The Netherlands, the United States,

DEVELOPMENT

and Central and Eastern Europe, PRA offers over 220 beds worldwide.

SERVICES

Your Full Service Clinical Research Partner

(Phase II-III)

Our teams successfully deliver complex, global, multi-centre trials and have supported more than a dozen drug approvals across a range of therapeutic areas.

LATE PHASE SERVICES

(Phase IIIb-IV & Registries)

Combining scientific expertise and operational execution to achieve regulatory and sponsor objectives for post approval studies.



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