

10 reasons to choose quantitative TCR sequencing in immuno-oncology

02

Seamless TCR-sequencing analysis platform, from extraction of the genetic material, library preparation, sequencing and bio-IT analysis, to the (optional) interpretation by specialised clinical immuno-oncologists



01



Delivery of accurate absolute quantifications of the T-lymphocyte receptor repertoires (technology based on RNA sequencing)

03

Platform developed and implemented by clinical oncologists from the Laboratory of Translational Oncology IPG / GHdC in Charleroi (Belgium) to answer research and clinical questions from medical end-users



04



Flexibility to use starting material from multiple sample types (PBMC blood samples, frozen tissue samples, cultured cells, sorted cells...)

05

Technical approach (using unique molecular indexes) to overcome the well-known pitfalls of other types of TCR-sequencing analyses, such as library amplification bias and polymerase errors



06

Sensitivity of the analysis down to a few cells in case of solid tumours



07

Possibility to combine quantitative TCR-sequencing analysis with anatomo-pathology, immuno-histochemistry analysis (e.g. routine PDL-1 staining, CD8 & PD-1) and spatial biology



08

Comparable quality in benchmarking against other marketed solutions for TCR-sequencing analyses



09

On-demand delivery of more comprehensive in-depth reports with additional tailor-made scientific analysis, such as T-cell clonotype tracking, multiple samples and time-points comparisons and the corresponding medical interpretation



10

Ideal support of quantitative TCR sequencing for target discovery, pre-clinical and clinical projects (uncovering predictive and companion biomarkers) in immuno-oncology



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