

RWE in Europe Paper IV:

Engaging pharma in the RWE Roadmap

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Abstract

This paper summarises discussions held at the fourth round-table

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... tangible outputs going forward.

... around data generation and collaborations discussed here are applicable to all ...

(...)

... mean that data has to be aggregated, high quality, complete and longitudinal with ... and patient-level data linkage. *A priori* ... study objectives and analysis plans as well as careful cohort selection.

... improving treatment and accelerating research in the cancer arena, has partnered ... enable it to be used for regulatory purposes.

In the oncology arena trials are both timely and expensive, whilst treatment protocols are dynamic. This means that by the time lengthy trials have been completed treatment options have developed further and those on trial have become irrelevant. RWD can ... periodic. Furthermore, oncologists are often highly time stretched and have a limit on the data variables they are able to capture. In order to develop regulatory grade ... data. Their databases, which encompass two million active patient records, 2500 ... are collected and standardised using a technology platform that involves human data ... room for a more automated system that can limit levels of human error, reduce any bias and improve the speed at which data can be made available.



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the product lifecycle.

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References

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