



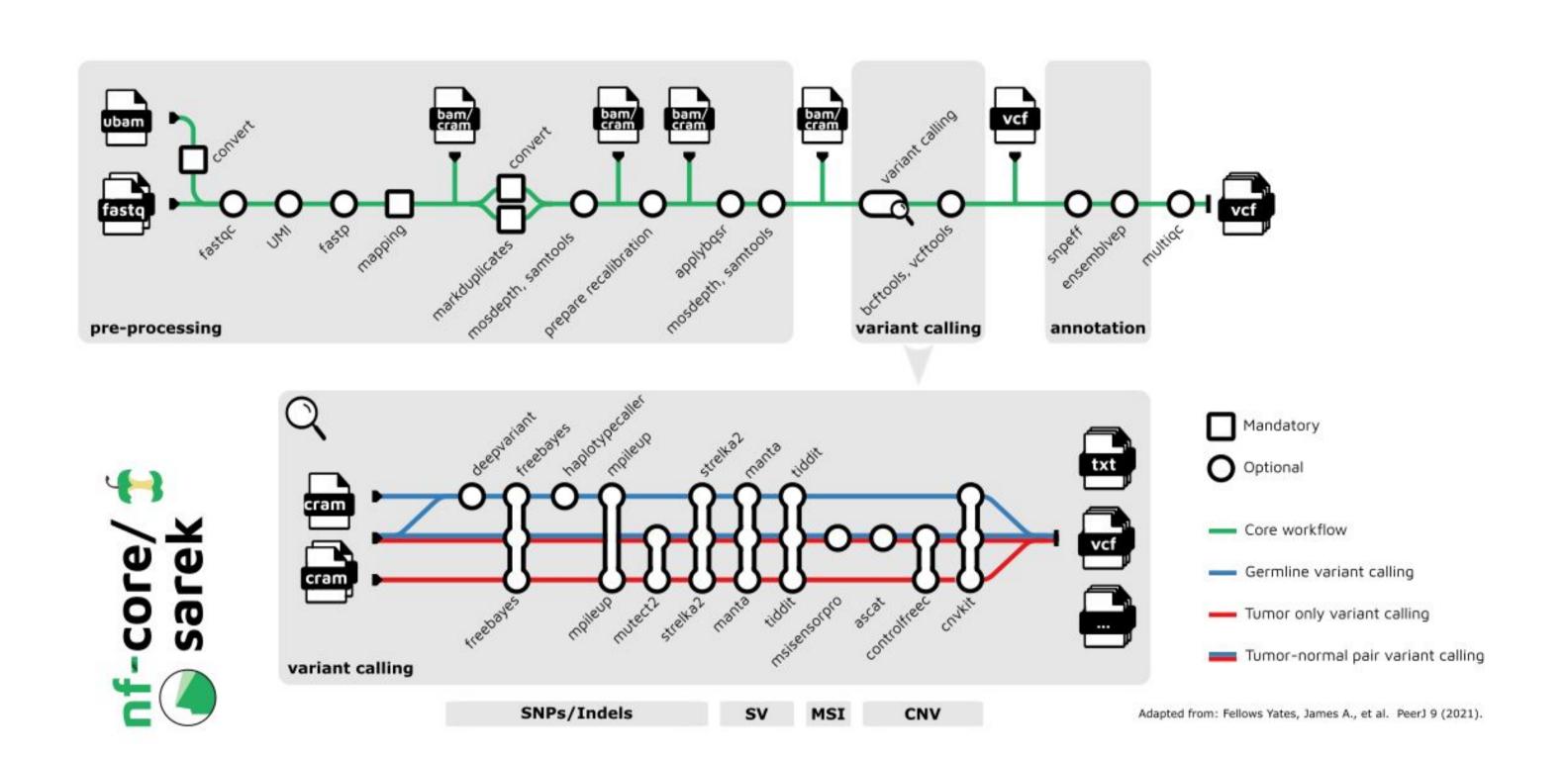
nf-core/sarek: a pipeline for efficient germline, tumor-only, and somatic analysis of NGS data on different compute infrastructures

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Overview

Somatic variant calling studies often include many patients with dataset sizes varying widely between oncopanel, whole-exome, and whole-genome sequencing data. nf-core/sarek¹ is an established pipeline for exploring single-nucleotide variants, structural variation, microsatellite instability, and copy-number alterations of germline, tumor-only, and paired tumor-normal short-reads nf-core/sarek is part of nf-core², a community project which provides an infrastructure to create reproducible, scalable, and portable open-source Nextflow³-based pipelines. Here, we show the latest updates including improvements to the data flow and tool selection reducing time, compute resources, and doud computing costs, as well as modularization improving code maintainability. Preprint available at: biorxiv.org/content/10.1101/2023.07.19.549462v1



Pipeline metro map showing a high-level view of the different analysis steps. The pipeline can be started from six different entry points and run through all subsequent tasks. All optional tools can be selected in any combination. This allows to recompute and extend the results throughout a project's duration.

Scatter/gathering speeds up analysis

