Description of Additional Supplementary File

File name: Supplementary Data 1

Description: Targeted sequencing data of 800 patients enrolled in plasmaMATCH

Sheet 1: Summary matrix. Mutation positive (‘1’) or negative (‘0’) for all patients (rows) and genes (columns). Includes pathogenic SNVs and indels.

Sheet 2: SNVs and indels. Includes the judgement of whether the mutation call was considered pathogenic (‘pathogenic’, see “Methods”).

Sheet 3: CNVs.

Sheet 4: Fusions.