



Genome Build GRCh37 (hg19)

Gene	GenBank Accession Number*	Exons
<i>ANKRD26</i>	NM_014915.2	1–4, 5'UTR c.-172
<i>ASXL1</i>	NM_015338.5	10–13
<i>BCOR</i>	NM_001123385.1	4–15
<i>CALR</i>	NM_004343.3	9
<i>CBL</i>	NM_005188.3	8–9, intron 7 100bp before exon 8 and intron 8
<i>CEBPA</i>	NM_004364.4	1
<i>CSF3R</i>	NM_000760.3	14 and 17
<i>DDX41</i>	NM_016222.2	1–17
<i>DNMT3A</i>	NM_022552.4	8–23
<i>ELANE</i>	NM_001972.2	1–5
<i>ETNK1</i>	NM_018638.4	2–5
<i>ETV6</i>	NM_001987.4	3–8
<i>EZH2</i>	NM_004456.4	2–20
<i>FLT3</i>	NM_004119.2	14–20
<i>GATA1</i>	NM_002049.3	2 and 4
<i>GATA2</i>	NM_001145661.1	3–7, intron 5 c.1017+1 to 1017+730
<i>IDH1</i>	NM_005896.3	4
<i>IDH2</i>	NM_002168.3	4
<i>JAK2</i>	NM_004972.3	12–16
<i>KDM6A (UTX)</i>	NM_021140.3	1–29
<i>KIT</i>	NM_000222.2	8–11 and 17
<i>KRAS</i>	NM_033360.3	2–3
<i>MPL</i>	NM_005373.2	10–12
<i>NPM1</i>	NM_002520.6	9–11, intron 10 30bp before exon 11
<i>NRAS</i>	NM_002524.4	2 and 3
<i>PHF6</i>	NM_001015877.1	2–10
<i>PTPN11</i>	NM_002834.3	3–4 and 12–13
<i>RAD21</i>	NM_006265.2	1, 2, 4–7, 9–11, 13, 14
<i>RUNX1</i>	NM_001001890.2	1–6, intron 4 c.725–13T>A and intron 5 c.886+1–4del
<i>SETBP1</i>	NM_015559.2	partial exon 4; amino acids 400–950
<i>SH2B3 (LNK)</i>	NM_005475.2	2–8
<i>SF3B1</i>	NM_012433.2	13–16

<i>SRP72</i>	NM_006947.3	6, 10
<i>SMC3</i>	NM_005445.3	7, 8, 13, 17, 19, 21, 29
<i>SRSF2</i>	NM_003016.4	1 and 2
<i>STAG2</i>	NM_001042750.1	4–34
<i>TERT</i>	NM_198253.2	2–16
<i>TET2</i>	NM_001127208.2	3–11
<i>TP53</i>	NM_000546.4	4–9
<i>U2AF1</i>	NM_001025203.1	2, 6, 8
<i>WT1</i>	NM_024426.2	1–10
<i>ZRSR2</i>	NM_005089.3	1–11

Default offset is +/- 10bps around each exon with exceptions for *CBL*, *GATA2* and *RUNX1*, which have certain intron regions of interest. For *SETBP1*, only part of exon 4 is evaluated. *NPM1* has coverage set to -30bps before exon 11 because of a downstream polymorphic region. *ANKRD26* has a region of interest in the 5' untranslated region (5'UTR). *RAD21* exon 10 and *STAG2* exons 12, 17, and 22 have coverage set to -15bps before the exons due to homopolymer regions.

*Reference transcript numbers may have been updated due to database re-versioning. Refer to the patient report for the most updated gene transcript information.