

1). FISH analysis showed the initial *MYC* amplification in 1% of uncultured BM cells, however a new trisomy 8 involving the *MYC* locus (8q24) occurred. We initiated salvage therapy with azacitidine and venetoclax (aza-ven)³ and prepared the patient for allogeneic stem cell transplantation. However, BM aspiration after the second cycle revealed refractory disease with 60% blasts and persistent *NPM1* mutational load. Another salvage therapy with high-dose cytarabine and mitoxantrone (HAM) was initiated, but the disease was refractory with novel extramedullary skin lesions. Considering the poor prognosis of this chemo-refractory disease, as well as the patient's age and wish a best supportive care therapy was initiated. The patient died 2 months after the appearance of the skin lesions.

In order to gain a deeper understanding of the clonal composition and evolution of this unusual course of the disease, we performed single-cell DNA sequencing (scSeq) at four time points indicated in Figure 1 and Table 1 using the Mission Bio Tapestry single-cell DNA and protein sequencing platform investigating genes covered in Mission Bio's myeloid panel⁴ with v2 of the protocol. Additionally, we used antibodies against four surface proteins to simultaneously generate a single-cell protein library. Libraries were sequenced to a DNA/protein read depth of 324/31 M (ID), 267/31 M (CRi-1), 292/37 M (CRi-2) and 225/37 M (Rel-1) on an Illumina NextSeq2000 sequencer (P3, 2x150bp). Sequencing details are available upon request. Using scSeq, we were able to confirm the *MYC* amplification at initial diagnosis in a large fraction of CD34⁺ blasts.

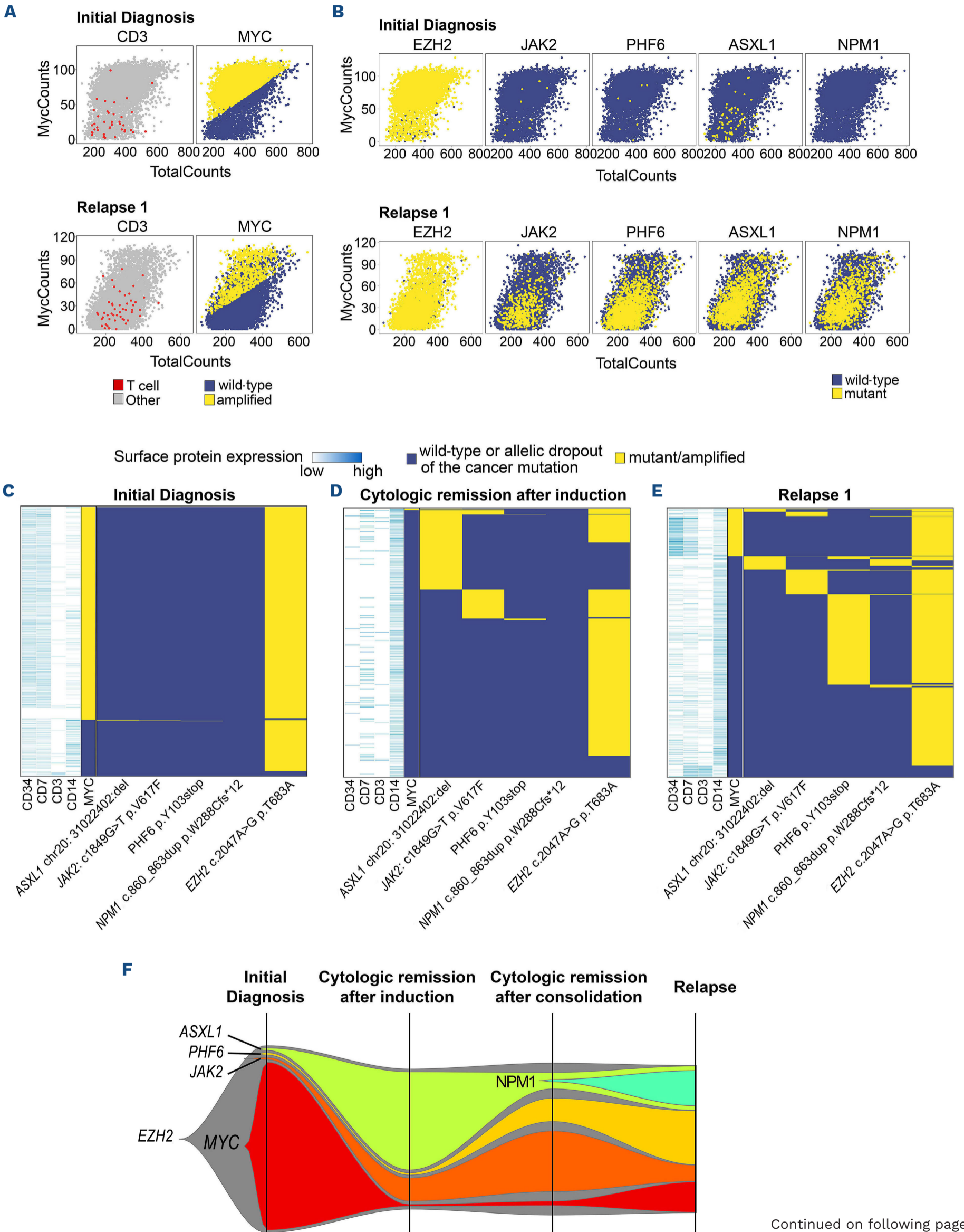
In order to discern *MYC*-amplified from wild-type cells, we used CloneTracer to assign clonal probabilities to each cell.⁵ Those cells being assigned to the *MYC* clone were termed *MYC*-amplified (*MYC*amp) (Figure 2A). Notably, healthy T cells were reliably identified as non-*MYC*amp as expected (Figure 2A). We identified an *EZH2* missense mutation p.T683A in CD34⁺ cells with aberrant CD7 expression, which seemed to be present in the *MYC*amp and also non-*MYC*amp CD34⁺CD7⁺ cells, but not in healthy CD3⁺CD34⁻ T cells. This suggested that the *EZH2* mutation occurred in an ancestral committed progenitor cell giving rise to the *MYC*amp leukemic clone (Figure 2B, C). Moreover, an *ASXL1* p.E635Rfs mutation was detected in around 1% of the cells not harboring *MYC*amp (Figure 2B, C).

In the remission sample post induction (CRi-1), which was considered measurable residual disease (MRD)-negative by flow cytometry, the *EZH2* mutation was present in the majority of CD3⁻ BM cells, but *MYC*amp was detectable in a minority of cells (Figure 2D). However, we identified a *JAK2* V617F mutation typically found in myeloproliferative neoplasms (MPN) and AML,⁶ in 9% of the cells, and the *ASXL1* mutation had increased in frequency. Notably, we observed a high allelic dropout rate of *EZH2*, suggesting that many cells called as *EZH2* wild-type were actually mutant, and that the overall fraction of *EZH2* mutant myeloid cells was very high. Considering also that the false-positive rate of the applied technology is very low,^{7,8} the abundant co-occurrence of the *ASXL1* and *EZH2* mutations in the first remission time point

Table 1. Clonal composition and evolution of the disease.

	Clinical time point	Karyotype	Blast count bone marrow %	<i>MYC</i> amp (FISH) %	<i>MYC</i> amp scSeq	<i>ASXL1</i> p.Glu635Rfs	<i>ASXL1</i> p.Glu635fs COSM36165	<i>EZH2</i> p.Thr683Ala %	<i>EZH2</i> p.Tyr646Cys	<i>JAK2</i> p.Val617Phe COSM12600	<i>NPM1</i> p.Trp288fs COSM17559	<i>PHF6</i> p.Tyr103stop	<i>TET2</i> p.Try1245fs COSM87129 %
Clinical diagnostics	Diagnosis (ID)	46,XY,del(16)(p13.1),1~19dmin[13]/46,XY,del(16)(p13.1)[3]/46,XY,2~15dmin[3]/46,XY[1]	80	71	-	-	-	48*	-	-	-	-	49
	Relapse (Rel-1)	48,XY,+8,+19,2~4dmin[11]/49,sl,+8[2]/46,XY,inv(16)(p11.2q22)[4]/46,XY[3]	60	2	-	3	-	34	-	9	11	-	48
	Progressive disease		65	18+8	-	-	-	20	27	10	31	-	49
scSeq	Diagnosis (ID)		67 PB	-	72	-	1	80	-	-	-	-	-
	CRi-1		4	-	0	-	42	47	-	8	-	-	-
	CRi-2		<5	-	1	-	6	64	-	23	-	10	-
	Relapse (Rel-1)		60	-	12	-	18	51	-	6	15	19	-

*Not reported initially by clinic, probably because no COSMIC initial diagnosis (ID) and near 50% so considered single-nucleotide polymorphism. *MYC*amp: *MYC*-amplified; FISH: fluorescence *in situ* hybridization; Rel-1: relapse one; PB: peripheral blood; CRi-1: complete remission with incomplete hematological recovery one; CRi-2: complete remission with incomplete hematological recovery two; scSeq: single-cell sequencing.



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Figure 2. Single-cell DNA sequencing analysis. (A) Dot plots showing total counts on the x-axis and *MYC* counts on the y-axis, as well as highlighted in color the cell type (left panels: red: T cells, gray: other cells) or mutational status (right panels: blue: wild-type, yellow: *MYC*-amplified) at the indicated time points initial diagnosis (top) and relapse (bottom). (B) Dot plots showing total counts on the x-axis and *MYC* counts on the y-axis, as well as highlighted in color the mutational status (blue: wild-type, yellow: mutant) at the indicated time points initial diagnosis (top) and relapse (bottom). (C-E) Heatmaps showing the mutational status for the indicated genes at first diagnosis (C), remission 1 (D), and at relapse (E). Cells with dropout in any of the genes were removed, and the cells were additionally annotated with the expression of surface proteins (CD34, CD7, CD3, CD14) measured with cellular indexing of transcriptomes and epitopes by sequencing (CITE-seq). (F) Schematic of the clonal evolution as detected through single-cell DNA sequencing.

(Figure 2D) strongly suggested that *ASXL1* occurred downstream of the *EZH2* mutation.

In the second remission sample post consolidation two (CRi-2) *MYC*amp was detectable at a very low rate by scSeq, while FISH was still negative. The *JAK2* clone increased in abundance (25%), continued to co-occur with the *EZH2* p.T683A variant, but did not overlap with *MYC*amp cells (Table 1). In addition, a new *PHF6* mutation (p.Y103stop) was detected in 10% of cells. These cells shared the *EZH2* p.T683A, but they were devoid of the *JAK2* mutation and *MYC*amp. Of note, blasts were below 5% at this stage and flow cytometry did not detect the leukemia-associated immunophenotype from initial diagnosis, so that the patient was considered flow MRD-negative.

In the cytologic relapse sample, which was acquired 6 months later, *MYC*amp was detectable again by scSeq (12%; Figure 2E, F). The *JAK2* clone had decreased from 25% to 7%, while the *PHF6*-mutated cells represented the second largest clone at relapse downstream of the *EZH2* mutation (22%). An *NPM1* TCTG-insertion was found in 15% of cells. Interestingly, the *NPM1*-mutated cells harbored the *ASXL1* mutation, which was already present at diagnosis independently of the other three clones (Figure 2E, F).

Several observations are remarkable in this scSeq study: an *EZH2* missense mutation p.T683A was detectable already at initial diagnosis and was shared by the initial *MYC*amp clone, but also by two distinct *JAK2*- and *PHF6*-mutated clones emerging during cytologic remission. Based on the in-depth scSeq information this AML would have been classified as high-risk AML with myelodysplasia-related gene mutations (AML-MR) according to the new World Health Organization (WHO) and ELN 2022 classification, so that a therapy with CPX-351 would have been recommended.¹

While the *JAK2*-mutated clone decreased from remission to relapse, the *PHF6*-mutated clone became the second largest subclone downstream of the *EZH2* mutation at relapse potentially pointing towards clonal selection during the maintenance therapy with glasdegib. *PHF6* mutations in AML are often frameshift or nonsense mutations.⁹ As *PHF6* is an X-linked gene, these alterations are predominantly found in males as in our patient. *PHF6* mutations are associated with immature AML (FAB subtypes M0–M2).⁹ Moreover, *PHF6* along with other epigenetic regulators such as *DNMT3A*, *TET2*, and *ASXL1*, is frequently altered in patients with clonal hematopoiesis.⁹ Genomic amplifications, such as *dmin*, homogeneously stain-

ing regions (*hsr*), are rare in leukemia, accounting for less than 1% of cytogenetically abnormal hematological malignancies and are associated with elderly patients and poor prognosis.¹⁰ While we found *MYC*amp in form of *dmin* at initial diagnosis by FISH, the relapse sample revealed a novel trisomy 8 involving the *MYC* locus confirming the hypothesis that different *MYC*amp mechanisms can co-exist within the same leukemic cell population as previously described.¹¹ scSeq was able to detect *MYC*amp in both conditions, and the fraction of cells highly corresponded to the fractions detected by FISH or cytogenetics. *MYC* mutations have been reported to coincide with other mutations in AML, such as *FLT3*, *NPM1*, and *DNMT3A* mutations.¹² The interesting observation in this case is, that the additional clones developing during therapy did not occur in the *MYC*amp clone as expected. This might suggest, that the *MYC*amp clone facilitated development of the other AML clones by paracrine mechanisms, e.g., via BM niche remodeling.

Another striking observation in this case is the occurrence of a therapy-associated *NPM1* mutation. Approximately 15% of therapy-related AML cases carry *NPM1* mutations, frequently accompanied by a normal karyotype and *DNMT3A* mutations, and rarely associated with chromosome aberration.¹³ Cases of *NPM1*-mutated AML following chemotherapy for previous lymphoid malignancies appear to arise from a background of *DNMT3A*- or *TET2*-driven clonal hematopoiesis (CH) rather than being a direct result of cytotoxic therapy.¹⁴ It is therefore remarkable, that the *NPM1* mutation in this case occurred in cells harboring mutations in two other epigenetic regulators, *ASXL1* and *EZH2*, suggesting a common mechanism. The *ASXL1*-mutated background and the co-existence of *NPM1* wild-type clones might explain the non-response to aza-ven in this case.

In summary we present a very unusual AML case with emergence of multiple parallel genetic clones during remission after treatment with intensive chemotherapy and targeted drugs including the hedgehog inhibitor glasdegib and the CD33-directed antibody “GO”. The complex clonal evolution observed in this case resulted in a very aggressive and multi-refractory disease, which raises the question, whether and how the agents applied might have contributed to clonal selection or exit from dormancy. These findings underscore the importance of employing scSeq to dissect individual cases, which can offer valuable mechanistic insights and help optimize therapeutic strategies.

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Disclosures

No conflicts of interest to disclose.

Contributions

CP and SJ designed the study, performed clinical data analysis and wrote the manuscript. MS, LV, CST, and SBC performed single cell sequencing and data analysis. MH performed the clinical diagnostics. SJ, RS, and CMT designed the clinical study. All authors contributed to writing the manuscript.

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Data-sharing statement

The authors welcome requests for the original data related to the laboratory analysis results.