

We are IntechOpen, the world's leading publisher of Open Access books Built by scientists, for scientists

4,800

Open access books available

122,000

International authors and editors

135M

Downloads

Our authors are among the

154

Countries delivered to

TOP 1%

most cited scientists

12.2%

Contributors from top 500 universities



WEB OF SCIENCE™

Selection of our books indexed in the Book Citation Index
in Web of Science™ Core Collection (BKCI)

Interested in publishing with us?
Contact book.department@intechopen.com

Numbers displayed above are based on latest data collected.

For more information visit www.intechopen.com



Genetic Alterations and Their Clinical Implications in Acute Myeloid Leukemia

Hsin-An Hou, Wen-Chien Chou and Hwei-Fang Tien
 Division of Hematology, Department of Internal Medicine, National Taiwan University
 Hospital, College of Medicine, National Taiwan University, Taipei,
 Taiwan

1. Introduction

Acute myeloid leukemia (AML) is a hematologic malignancy with great variability in the pathogenesis, clinical features, and treatment outcomes. Advances in molecular research have greatly improved our understanding of the leukemogenesis in AML. A two-hit model proposes that the development of AML requires the cooperation between at least two classes of gene mutations. (Frohling, *et al* 2005, Gilliland 2002) Class I mutations, such as *RAS*, *FLT3*, *KIT*, *PTPN11* and *JAK2* mutations, involve genes in the kinase signaling pathways leading to cell survival and proliferation and Class II mutations, such as *t(15;17)/PML-RARA*, *inv(16)/CBFB-MYH11* and *t(8;21)/RUNX1-RUNX1T1* fusions, and *MLL/PTD*, and *CEBPA* and *AML1/RUNX1* mutations, involve transcription factors or cofactors resulting in impaired hematopoietic differentiation. In addition to genetic abnormalities, increasing evidences show that epigenetic deregulations are also critical to the pathogenesis of AML. (Chen, *et al* 2010) Compatible with these findings, several novel mutations involving genes related to epigenetic modifications, such as *isocitrate dehydrogenase 1 (IDH1)*, *IDH2*, *ten-eleven translocation 2 (TET2)*, *additional sex comb-like 1 (ASXL1)*, and DNA methyltransferase 3A (*DNMT3A*) were detected in AML recently. (Chou, *et al* 2010b, Delhommeau, *et al* 2009, Gelsi-Boyer, *et al* 2009, Ley, *et al* 2010, Mardis, *et al* 2009, Metzeler, *et al* 2011)

Risk-adapted treatment may not only improve the prognosis, but also reduce the toxicity from the therapy in patients with AML. In addition to the conventional risk factors, such as age, white blood cell (WBC) counts and cytogenetics, molecular genetic alterations, such as mutations of *NPM1*, *CEBPA*, *AML1/RUNX1*, *WT1*, *FLT3*, *TET2*, and *DNMT3A* etc., are also important prognostic factors in AML patients. Furthermore, the gene mutations which are stable during treatment courses can also be used as biomarkers to monitor minimal residual disease (MRD). Herein, we will review the gene mutations in AML and discuss their clinical implications.

2. Class I mutations that lead to cell survival and proliferation

2.1 *FLT3* mutations

FMS-like tyrosine kinase 3 (*FLT3*), mapped at 13q12, encodes a receptor tyrosine kinase. (Kiyoi, *et al* 1998) *FLT3*-internal tandem duplication (*FLT3-ITD*) mutation, one of the

most common mutations in AML, was found by Nakao et al in 1996.(Nakao, *et al* 1996) The mutation occurs as a duplication of nucleotide sequences of variable lengths in exons 14 and 15, leading to addition of repeated peptide in the juxtamembrane domain in the cytoplasm. Another activating *FLT3* mutation occurs in tyrosine kinase domain (*FLT3*-TKD), causing point mutations, small deletions or insertions mainly at codon 835 or 836 within the activation loop of the second kinase domain.(Bacher, *et al* 2008, Yamamoto, *et al* 2001) The *FLT3* mutant protein constitutively activates the cascade of *FLT3* signaling in the absence of *FLT3* ligand promoting cell proliferation and decreased apoptosis.

FLT3-ITD occurs in about 25% of adult AML and shows association with normal karyotype and *NPM1* mutation. The patients with this mutation have higher WBC counts, shorter disease-free survival (DFS) and overall survival (OS), and increased relapse rate.(Kottaridis, *et al* 2001, Kottaridis, *et al* 2002) While mutant size may not be related to prognosis, higher mutant levels are associated with higher relapse rate and shorter survival.(Gale, *et al* 2008) Absence of *FLT3*-ITD combined with *NPM1* mutation is regarded as a favorable prognostic genotype.(Gale, *et al* 2008, Schlenk, *et al* 2008) Up to one third of AML patients with *FLT3*-ITD can lose the mutation at disease relapse, indicating that this mutation is much less stable than *NPM1* mutation, and is not a good marker for disease monitoring.(Chou, *et al* 2011b, Kottaridis, *et al* 2002, Palmisano, *et al* 2007, Shih, *et al* 2002) *FLT3*-TKD occurred in about 4%-10% of AML patients.(Yamamoto, *et al* 2001, Bacher, *et al* 2008) AML with this mutation also shows specific clinical and biologic features, such as elevated WBC counts at diagnosis, higher frequency of normal karyotype and mutations in *NPM1*, *CEBPA*, and *NRAS*. However, the prognostic significance is still inconclusive.(Bacher, *et al* 2008, Whitman, *et al* 2008)

2.2 RAS mutations

The RAS proteins are a large superfamily of low molecular-weight guanine nucleotide-binding proteins, which are activated by cytokine receptors in response to ligand stimulation and therefore control cell proliferation and survival of hematopoietic progenitors.(Downward 2003, Reuther and Der 2000, Shields, *et al* 2000, Wittinghofer 1998) Three members of the RAS family, HRAS, KRAS and NRAS, are found to be activated by mutations in human cancers.(Bos 1989, Downward 2003) Almost all RAS mutations occur by single nucleotide substitutions in codons 12, 13 and 61.(Bos, *et al* 1987, Farr, *et al* 1988, Senn, *et al* 1988, Toksoz, *et al* 1989) *NRAS* and *KRAS* mutations are found in approximately 12-30% and 9-14%, respectively, of AML patients. In a large cohort study of 2502 AML patients, the mutations were found much prevalent in patients with *inv*(16)/*t*(16;16) and *inv*(3)/*t*(3;3), but seldom found in those with *t*(15;17) and complex karyotype.(Bacher, *et al* 2006)

The prognostic relevance of *RAS* mutation in AML has not been firmly established. Some studies showed *RAS* mutation predicted poor prognosis(De Melo, *et al* 1997, Kiyoi, *et al* 1999, Meshinchi, *et al* 2003), some showed no impact on clinical outcome,(Bacher, *et al* 2006, Bowen, *et al* 2005, Radich, *et al* 1990, Ritter, *et al* 2004) whereas others found *RAS* mutations were associated with a favorable prognosis.(Coghlan, *et al* 1994, Neubauer, *et al* 1994) However, higher dose of cytarabine may decrease the relapse rate in *RAS*-mutated AML patients.(Neubauer, *et al* 2008)

2.3 KIT mutations

KIT, a member of type III receptor tyrosine kinase family, is important for the development of hematopoietic progenitor cells and also crucial in leukemogenesis.(Blume-Jensen and

Hunter 2001, Bowen, *et al* 2005, Radich, *et al* 1990) High expression of *KIT* (CD117) is a common finding in AML, (Ikeda, *et al* 1991, Reuss-Borst, *et al* 1994) and activation mutations of *KIT*, most commonly affecting exons 8 and 17 can be identified in 20–45% of AML with *inv* (16) and 12.8–46.8% of AML with *t*(8;21), but infrequently found in other AML types. (Beghini, *et al* 2000, Beghini, *et al* 2004, Care, *et al* 2003) Most, though not all, studies, (Boissel, *et al* 2006, Care, *et al* 2003, Schnittger, *et al* 2006) showed *KIT* mutation was associated with inferior outcome in the core binding factor (CBF) AML, especially *KIT*-D816 mutations in *t*(8;21)/*RUNX1-RUNX1T1*-positive AML.

2.4 *JAK2* mutations

JAK2 is a nonreceptor tyrosine kinase. The *JAK2* V617F mutation induces the activation of *JAK2*-*STAT5* signal transduction pathway and then substantially alters the proliferation and self-renewal of hematopoietic precursors. (Liu, *et al* 1999, Walz, *et al* 2006) Although the *JAK2* V617F mutation is a common genetic event in the patients with myeloproliferative neoplasms (MPN), (Baxter, *et al* 2005, Goldman 2005, Kralovics, *et al* 2005, Levine, *et al* 2005b) it is seldom found (<1%-2%) in *de novo* AML patients. (Frohling, *et al* 2006, Illmer, *et al* 2007, Lee, *et al* 2006, Levine, *et al* 2005a) Illmer *et al* showed 3.6% of patients with CBF AML had *JAK2* V617F mutation and these patients had an aggressive clinical course and poor outcome. (Illmer, *et al* 2007)

2.5 *PTPN 11* mutations

SHP-2 is encoded by *PTPN11* which is located on chromosome 12q24. The protein is a non-receptor tyrosine phosphatase participating in intracellular signaling elicited by a number of growth factors, cytokines, hormones and adhesion molecules. (Neel, *et al* 2003, Tartaglia, *et al* 2004) Germline *PTPN11* mutations were first reported by Tartaglia *et al* in patients afflicted with Noonan syndrome. (Tartaglia, *et al* 2002, Tartaglia, *et al* 2001) Subsequently, somatic *PTPN11* mutations were also found in patients with juvenile myelomonocytic leukemia, and myelodysplastic syndrome (MDS). (Chen, *et al* 2006, Loh, *et al* 2004b, Tartaglia, *et al* 2003) The *PTPN11* mutation is not a frequent molecular event (4-5%) in AML. (Hou, *et al* 2008, Loh, *et al* 2004a, Tartaglia, *et al* 2005) In a study of 272 primary AML patients, we found this gene mutation was closely associated with older age, French-American-British (FAB) M4/M5 subtype, CD14 expression, normal karyotype and *NPM1* mutation. (Hou, *et al* 2008) Loh *et al* and Tartaglia *et al* revealed that the *PTPN11* mutation had no prognostic implication for pediatric patients with AML; (Loh, *et al* 2004a, Tartaglia, *et al* 2005) however, this mutation may be a poor-risk factor for OS in adult AML patients without *NPM1* mutations. (Hou, *et al* 2008)

3. Class II mutations that impair hematopoietic differentiation

3.1 *CEBPA* mutations

CCAAT/enhancer binding protein α (C/EBP α) is a 42-kDa transcription factor that possesses a DNA-binding basic leucine zipper domain (bZIP) in the COOH terminus and two transactivation domains TAD 1 and TAD 2 in the NH2 terminus. (Friedman and McKnight 1990) As a transcription factor, it plays a crucial role in granulocytic differentiation and diminished C/EBP α activity contributes to myeloid progenitor transformation. (Cammenga, *et al* 2003, Oelgeschlager, *et al* 1996, Smith, *et al* 1996) *CEBPA*

mutations are observed in 7% to 18% of patients with AML.(Frohling, *et al* 2004, Lin, *et al* 2005, Pabst, *et al* 2001b, Preudhomme, *et al* 2002) Two major types of *CEBPA* mutations have been identified; one alters COOH terminal bZIP of *CEBPA*, resulting in decreased DNA-binding and/or dimerization activity and the other disrupts translation of the C/EBP α NH2 terminus, thereby up-regulates an alternative 30-kDa isoform with dominant-negative effect on the full-length wild-type C/EBP α .(Koschmieder, *et al* 2009, Lin, *et al* 1993, Pabst, *et al* 2001b) Most patients with *CEBPA* mutations harbored biallelic mutations involving both the NH2-terminal TAD region and the COOH-terminal bZIP domain.(Hou, *et al* 2009, Lin, *et al* 2005, Pabst, *et al* 2009, Renneville, *et al* 2009a) *CEBPA* mutations occur most frequently in patients with FAB subtype M2, and are closely associated with CD7, CD15, CD34, and HLA-DR expression on the leukemic cells, higher circulatory blasts and normal cytogenetics.(Frohling, *et al* 2004, Lin, *et al* 2005, Pabst, *et al* 2001a, Zhang, *et al* 1997) This mutation seems quite stable during AML evolution and may be a potential marker to monitor MRD. The fact that none of the AML patients who do not have *CEBPA* mutations at diagnosis acquire the mutation at relapse suggests that this mutation may not play a major role in the progression of AML.(Lin, *et al* 2005, Tiesmeier, *et al* 2003) Several studies have shown mutant *CEBPA* predicts favorable outcome in AML patients with intermediate or normal cytogenetics.(Barjesteh van Waalwijk van Doorn-Khosrovani, *et al* 2003, Bienz, *et al* 2005, Frohling, *et al* 2004, Preudhomme, *et al* 2002) The favorable impact of *CEBPA* mutations in the AML patients is only observed in the absence of *FLT3/ITD* or other associated cytogenetic abnormalities.(Renneville, *et al* 2009a) Moreover, only double *CEBPA* mutations, but not single *CEBPA* mutation, are associated with better prognosis and define a distinct genetic entity.(Dufour, *et al* 2010, Hou, *et al* 2009, Pabst, *et al* 2009, Wouters, *et al* 2009)

3.2 *MLL* -PTD

The *MLL* partial tandem duplication (*MLL*-PTD) most commonly results from a duplication of a genomic region encompassing exon 5 through exon 11/12 and insertion of the duplicated segment into intron 4 of the full-length *MLL* gene.(Caligiuri, *et al* 1994, Schichman, *et al* 1994, Whitman, *et al* 2005) The duplication involves a portion of the gene corresponding to the amino terminus of the *MLL* protein which contains the AT hook and a region of homology to DNA methyltransferase motifs.(Schichman, *et al* 1995) The mechanism by which *MLL*-PTD contributes to leukemic phenotype is not clear, but may be through silencing of the wild-type *MLL* by epigenetic mechanisms.(Dimartino and Cleary 1999, Whitman, *et al* 2005) This mutation is found in 5-12% of patients with cytogenetically normal AML (CN-AML),(Dohner, *et al* 2002, Munoz, *et al* 2003, Schnittger, *et al* 2000, Shiah, *et al* 2002) and up to 54% of AML patients with trisomy 11.(Rege-Cambrin, *et al* 2005) Compared with patients without *MLL*-PTD, patients with this mutation more often have FAB M2 subtype, CD11b expression, wild-type *NPM1* and high *BAALC* expression, but lower WBC counts, less frequently extramedullary involvement and FAB M4/M5 subtype at diagnosis.(Shiah, *et al* 2002, Whitman, *et al* 2007) The presence of *MLL*-PTD predicts shorter remission duration and worse OS;(Dohner, *et al* 2002, Munoz, *et al* 2003, Shiah, *et al* 2002) however, more intensive consolidation therapy that includes hematopoietic stem cell transplantation (HSCT) during first complete remission (CR) may reverse the poor prognosis conferred by this mutation.(Whitman, *et al* 2007)

3.3 AML1/RUNX1

The *AML1/RUNX1* gene (Ito 2008), consisting of 10 exons (exons 1-6, 7A, 7B, 7C and 8), is one of the most frequently deregulated genes in leukemia through chromosomal translocations and point mutations. (Friedman 2009, Niebuhr, *et al* 2008, Osato 2004, Yamagata, *et al* 2005) Monoallelic germ-line mutation of the *RUNX1* gene occurs in rare cases of familial platelet disorder with predisposition to AML (FPD/AML). (Michaud, *et al* 2002) Acquired *RUNX1* mutation was frequently reported in therapy-related MDS or MDS/AML. (Harada, *et al* 2004) The incidence of *RUNX1* mutation in AML varies from 2.9% to 46% depending on the population selected, the regions of *RUNX1* screened, and the methods used. (Dicker, *et al* 2007, Preudhomme, *et al* 2000, Tang, *et al* 2009) In a large cohort study of 470 adult patients with *de novo* non-M3 AML, we detected *RUNX1* mutation in 13.2% of cases. The *RUNX1* mutation is closely associated with older age, immature FAB subtypes (M0/M1) and specific cytogenetic abnormalities such as trisomy 8 (+8), +13, or +21. (Dicker, *et al* 2007, Schnittger, *et al* 2011, Tang, *et al* 2009) None of the patients with t(8;21), inv(16), t(15;17) or 11q23 translocation shows *RUNX1* mutation. (Tang, *et al* 2009) One half of *RUNX1*-mutated patients have concurrently other gene mutations, mostly (83.9%) Class I mutations, especially *FLT3/ITD*, *FLT3/TKD* and *N-RAS* mutations (Tang, *et al* 2009) which all result in hyperactivation of the receptor tyrosine kinase (RTK)-RAS signalling pathways. (Niimi, *et al* 2006) This finding is consistent with the two-hit model of leukemogenesis. (Frohling, *et al* 2005, Gilliland 2002) Further, the *RUNX1* mutation is mutually exclusive with *CEBPA* and *NPM1* mutations, but closely associated with *MLL/PTD*. (Schnittger, *et al* 2011, Tang, *et al* 2009) The mutation may be lost at relapse in *RUNX1*-mutated patients, but none of the patients who do not harbor *RUNX1* mutation at diagnosis acquire novel mutation at relapse. (Tang, *et al* 2009) *RUNX1* mutation is an independent poor-risk factor for DFS and OS in *de novo* AML patients. (Gaidzik, *et al* 2011, Schnittger, *et al* 2011, Tang, *et al* 2009) In addition, HSCT seems to ameliorate the poor survival impact of *RUNX1* mutations. (Gaidzik, *et al* 2011, Tang, *et al* 2009)

4. Other mutations

4.1 NPM1 mutations

NPM1 mutation in AML was first identified by Dr. Falini's group, who noticed that some AML patients' leukemia cells exhibited aberrant cytoplasmic localization of *NPM1* protein, which normally located in nucleoli in non-mitotic cells. (Falini, *et al* 2005) Subsequent investigation revealed a tetra-nucleotide insertion near the C-terminal end of the coding sequence of *NPM1*. The most frequent form of mutation is duplication of TCTG (type A, c.860_863dupTCTG), resulting in alteration of the peptide sequence from DLWQWRKSL* to DLCLAVEEVSLRK*. *NPM1* mutation occurs in about 30% of AML, more frequently in elder patients, (Falini, *et al* 2011, Falini, *et al* 2005) and is highly associated with normal karyotype, and *FLT3/ITD*, but significantly exclusive with *CEBPA* mutation, favorable karyotype, and expression of CD34 and HLA-DR. (Boissel, *et al* 2005, Chou, *et al* 2006, Dohner, *et al* 2005, Falini, *et al* 2005, Schnittger, *et al* 2005, Suzuki, *et al* 2005, Verhaak, *et al* 2005) *NPM1* mutation generally renders better prognosis, (Falini, *et al* 2005) especially when *FLT3-ITD* is absent. (Schlenk, *et al* 2008, Thiede, *et al* 2006) Further refinement of patient groups disclosed 3 groups with distinct prognosis: good (*NPM1*⁺/*FLT3-ITD*⁻), intermediate (*NPM1*⁺/*FLT3-ITD*⁺ or *NPM1*⁻/*FLT3-ITD*⁻), and poor (*NPM1*⁻/*FLT3-ITD*⁺). (Gale, *et al* 2008) *NPM1* mutation seems quite consistent with disease status. (Chou, *et al* 2007, Schnittger, *et al*

2009) Serial analyses of *NPM1* mutations showed the mutation disappeared at CR, but the same mutation usually reappeared at relapse. This feature makes *NPM1* mutation an ideal marker for MRD monitoring. Studies have shown *NPM1* mutant levels reflect disease status, predict impending relapse, and bring prognostic implication. (Chou, *et al* 2007, Gorello, *et al* 2006, Kronke, *et al* 2011, Schnittger, *et al* 2009)

4.2 *WT1* mutations

The *Wilms' Tumor 1* (*WT1*) gene, encoding a zinc-finger transcription factor, is physiologically expressed in hematopoietic stem cells and involved in regulation of cellular growth and differentiation. (Baird and Simmons 1997, Ellisen, *et al* 2001) *WT1* was initially identified as a tumor suppressor gene. (Haber, *et al* 1990) but was later found to be overexpressed in AML as well as other cancers and thus was suggested to be an oncogene. (Bergmann, *et al* 1997, King-Underwood, *et al* 1996, Miwa, *et al* 1992) Mutations in *WT1* gene are found in about 7-13% of CN-AML patients with hotspots in the four Cys-His zinc finger domains on exons 7 and 9. (Gaidzik, *et al* 2009, Hou, *et al* 2010, King-Underwood, *et al* 1996, Paschka, *et al* 2008, Virappane, *et al* 2008) The precise role of *WT1* mutations in the leukemogenesis remains to be defined. The majority of *WT1* mutations are frame-shift mutations occurring in exon 7, followed by single amino acid substitutions in exon 9; whereas frame-shift mutations in exon 9 are rare. *WT1* mutations occur with similar frequencies in patients with normal karyotype and those with abnormal cytogenetics. (Hou, *et al* 2010) Chromosomal abnormality t(7;11)(p15;15), a translocation resulting in *NUP98/HOXA9* fusion, is closely associated with *WT1* mutation. (Hou, *et al* 2010) *WT1* mutations are positively associated with *FLT3/ITD* and *CEBPA* mutations. (Gaidzik, *et al* 2009, Renneville, *et al* 2009b) Paschka *et al* showed patients with *WT1* mutations had higher expression of *ERG* and *BAALC* than patients without. (Paschka, *et al* 2008) *WT1* mutation is an independent poor prognostic factor in CN-AML as well as total AML patients. (Hou, *et al* 2010, Paschka, *et al* 2008, Renneville, *et al* 2009b, Virappane, *et al* 2008), though different results have been reported. (Gaidzik, *et al* 2009, Santamaria, *et al* 2009)

5. Mutations of genes that involve epigenetic modifications

Different from genetic abnormalities which result in DNA sequence changes, epigenetic dysregulation causes aberrant gene expression without alteration of gene sequences. (Baylin and Ohm 2006, Chen, *et al* 2010, Jones and Baylin 2002) Epigenetic regulation includes DNA methylation, histone modifications, such as methylation, acetylation and phosphorylation, etc, and microRNA expression. (Baylin and Ohm 2006, Chen, *et al* 2010, Jones and Baylin 2002) The recent findings that mutations of genes related to epigenetic modifications, such as *IDH1*, *IDH2*, *TET2*, *ASXL1* and *DNMT3A*, are detected in AML patients provide new insights into mechanisms of epigenetic deregulation in the leukemogenesis.

5.1 *TET2* mutations

TET2 protein can catalyze the conversion of 5-methylcytosine (5-mC) of DNA to 5-hydroxymethylcytosine (5-hmC), with ferrous iron and α -ketoglutarate (α -KG) as cofactors, indicating a role of *TET2* in DNA methylation. (Ito, *et al* 2010) Mutations of *TET2* result in global DNA hypermethylation. (Figuroa, *et al* 2010) *TET2* mutation was originally identified in myeloid malignancies via single nucleotide polymorphism and comparative genomic-

hybridization array, which revealed common deletion of this gene in chromosome 4q.(Delhommeau, *et al* 2009) Subsequent studies confirmed this mutation in MDS, MPN, MDS/MPN, and secondary AML, with frequencies around 10% to 26%, 7% to 13%, 22% to 58% and 24% to 32%, respectively.(Bacher, *et al* 2010, Couronne, *et al* 2010, Flach, *et al* 2010, Jankowska, *et al* 2009, Kosmider, *et al* 2009a, Kosmider, *et al* 2009b, Langemeijer, *et al* 2009, Saint-Martin, *et al* 2009, Schaub, *et al* 2010, Smith, *et al* 2010, Tefferi, *et al* 2009a, Tefferi, *et al* 2009b) *TET2* mutation occurs in 18.0% to 23% of CN-AML patients.(Chou, *et al* 2011a, Metzeler, *et al* 2011) It is closely associated with older age, higher WBC count, but mutually exclusive with *IDH* mutation.(Chou, *et al* 2011a, Metzeler, *et al* 2011) In our study of AML patients with and without chromosomal abnormalities, *TET2* mutation was also found to be positively associated with normal karyotype, intermediate-risk cytogenetics, isolated trisomy 8, *NPM1* mutation, and *ASXL1* mutation.(Chou, *et al* 2011a) In European LeukemiaNet (ELN) favorable-risk group (patients with CN-AML with mutated *CEBPA* and/or mutated *NPM1* without *FLT3-ITD*),(Dohner, *et al* 2010) but not intermediate-1 risk group (CN-AML with wild-type *CEBPA* and wild-type *NPM1* and/or *FLT3-ITD*), *TET2*-mutated patients were found to have a lower CR rate, shorter DFS and OS, compared with *TET2*-wild type patients.(Metzeler, *et al* 2011) However, we did not have the same finding, but found that *TET2* mutation was an unfavorable prognostic factor in patients with intermediate-risk cytogenetics, and its negative impact was further enhanced when the mutation was combined with *FLT3-ITD*, *NPM1*-wild, or unfavorable genotypes (other than ELN favorable-risk group).(Chou, *et al* 2011a) More studies are needed to clarify the prognostic implication of *TET2* mutations in AML.

5.2 *IDH* mutations

IDH1 and *IDH2* genes encode two isoforms of isocitrate dehydrogenase which catalyzes the carboxylation of isocitrate to α -KG. *IDH1* and *IDH2* mutations were first detected in patients with brain tumors.(Parsons, *et al* 2008) Later, *IDH1* mutations (Mardis, *et al* 2009) and then *IDH2* mutations were discovered in AML patients, too.(Abbas, *et al* 2010, Marcucci, *et al* 2010, Ward, *et al* 2010) *IDH1* mutations affect arginine residue in position 132 (R132) and *IDH2* mutations, in R140 and R172 of exon 4. *IDH* mutations occur at low frequencies (3.6% to 5%) in MDS,(Kosmider, *et al* 2010) and in chronic-phase MPN (about 1.8%)(Pardanani, *et al* 2010), but obviously increased as these diseases progress to AML (7.5% to 21%),(Kosmider, *et al* 2010, Pardanani, *et al* 2010) indicating a role of *IDH* mutations in leukemogenesis. In *de novo* AML, *IDH2* mutations occur more frequently than *IDH1* mutations, with frequencies of 11% vs. 6% in patients younger than 60 years,(Abbas, *et al* 2010) 15.4% vs. 7.7% in total patients,(Ward, *et al* 2010) and 19% vs. 14% in adults with normal karyotype.(Marcucci, *et al* 2010) The underlying mechanism of *IDH* mutations in the leukemogenesis of AML remains to be determined, but several implications of *IDH1/2* mutations in AML have been generated. First, *IDH* mutations are loss-of-function mutations, as mutant *IDH* proteins show decreased enzyme activities,(Zhao, *et al* 2009) and have dominant-negative effects on wild type *IDH* upon homodimerization.(Zhao, *et al* 2009) Secondly, *IDH* mutations are also gain-of-function mutations because the mutant proteins can convert α -KG to 2-hydroxyglutarate (2-HG), a metabolite that may contribute to tumor growth through activating hypoxia-inducing factor-1 α (HIF-1 α).(Dang, *et al* 2009, Reitman, *et al* 2010, Ward, *et al* 2010) Thirdly, *IDH* mutations reduce production of α -KG, a cofactor of *TET2*, thus impair catalytic function of *TET2* resulting in global DNA hypermethylation,

similar to the effect of *TET2* mutations. 2-HG converted from α -KG in *IDH*-mutated cells is also shown to inhibit *TET2*-mediated hydroxymethylation of cytosine, indicating overlapping effects of these two mutations.(Xu, *et al* 2011) Compatible with this, *IDH* and *TET2* mutations are mutually exclusive in AML patients.(Figueroa, *et al* 2010, Metzeler, *et al* 2011)

Studies have shown similar clinical features between AML with *IDH1* and *IDH2* mutations, including strong association of both mutations with normal karyotype and isolated monosomy 8, but inverse correlation with expression of HLA-DR. However, some differences exist. *IDH1* mutation shows strong correlation with *NPM1* mutation, and FAB M1 subtype, but is inversely associated with FAB M4 subtype and expression of CD13 and CD14. On the other hand, mutation of *IDH2* is associated with higher platelet counts, but is inversely correlated with expression of CD34, CD15, CD7, and CD56, and is mutually exclusive with *WT1* mutation and chromosomal translocations involving CBF. While there is no impact of *IDH1* mutation on patient survival, multivariate analysis reveals *IDH2* mutation as an independent favorable prognostic factor,(Chou, *et al* 2010a, Chou, *et al* 2011c,) but different results have also been reported.(Marcucci, *et al* 2010, Thol, *et al* 2010) More intriguing are the differences of clinical presentations between patients with R140 and R172 mutations. Compared with *IDH2* R140 mutation, *IDH2* R172 mutation is associated with younger age, lower WBC count and LDH level, and is mutually exclusive with *NPM1* mutation. Recent studies also reported worse prognosis in AML patients bearing *IDH2* R172Q,(Boissel, *et al* 2010, Marcucci, *et al* 2010) while *IDH2* R140Q, in the contrary, conferred a better prognosis.(Green, *et al* 2010) Why mutations in different isoforms or loci of the same gene render distinct clinical and prognostic features remains to be investigated. Serial analyses of *IDH1/2* mutations at both diagnosis and relapse confirmed high stability of these two mutations.(Chou, *et al* 2010a, Chou, *et al* 2011c)

5.3 *ASXL1* mutations

Recently, mutations in exon 12 of *Additional sex comb-like 1 (ASXL1)* gene were found in various types of myeloid malignances, including MDS, MPN, MDS/MPN, and AML.(Abdel-Wahab, *et al* 2010, Boultonwood, *et al* 2010, Carbuccia, *et al* 2009, Carbuccia, *et al* 2010, Gelsi-Boyer, *et al* 2009) *ASXL1*, a human homologue of the *Additional sex combs (Asx)* gene of *Drosophila*, mapped to chromosome 20q11, a region commonly involved in cancers.(Fisher, *et al* 2003) It consists of an N-terminal ASX Homology (ASXH) domain and a C-terminal plant homeodomain (PHD) zinc finger region.(Fisher, *et al* 2003, Fisher, *et al* 2006) In human, the exact function of *ASXL1* mutation remains to be defined, but it is involved in regulation of histone methylation by cooperation with heterochromatin protein-1 (HP1) to modulate the activity of LSD1, a histone demethylase for H3K4 and H3K9.(Dange and Colman 2010, Wang, *et al* 2009) *ASXL1* mutations in exon 12 are found with an incidence of 10.8%, 8.9% and 12.9% among total cohort of patients with *de novo* AML, those with normal karyotype and abnormal cytogenetics, respectively.(Chou, *et al* 2010b) Most of the mutations appear to be either non-sense or frame-shift mutations, leading to disruption of the plant homeodomain (PHD) at the C terminal of *ASXL1*, which is well conserved among different species and can recognize methylated H3K4.(Abdel-Wahab, *et al* 2010, Fisher, *et al* 2003, Pena, *et al* 2006, Shi, *et al* 2006, Wysocka, *et al* 2006) Up to two thirds of mutations occurred at c.1934dupG (an extra G after 1934th nucleotide of the coding sequence of *ASXL1*) causing G646WfsX12 (change of glycine to tryptophan at amino acid 646, with a

premature stop codon after another 11 amino acid).(Chou, *et al* 2010b) The mutation was closely associated with older age, male gender, isolated trisomy 8, *RUNX1* mutation, and expression of HLA-DR and CD34, but inversely associated with t(15;17), complex cytogenetics, *FLT3*-ITD, *NPM1* mutations, *WT1* mutations, and expression of CD33 and CD15.(Chou, *et al* 2010b) Patients with *ASXL1* mutations had a shorter OS than those without, but the mutation was not an independent adverse prognostic factor in multivariate analysis. Sequential analyses showed that the original *ASXL1* mutations could disappear at relapse and/or refractory status in some patients. Moreover, two out of the 109 *ASXL1*-wild patients acquired a novel *ASXL1* mutation at relapse.(Chou, *et al* 2010b) Thus, the *ASXL1* mutation status can change during disease evolution in a few patients.

5.4 DNMT3A mutations

By whole genome sequencing on a single patient with normal cytogenetics, Ley and his colleagues found a mutation in *DNMT3A* gene, which encodes the enzyme DNA methyltransferase 3A which belongs to the family of DNMTs that catalyze the addition of methyl group to cytosine of CpG dinucleotide.(Ley, *et al* 2010) In this seminal study, *DNMT3A* mutation was detected in 22.1% of AML patients. Most of the mutations occurred at R882 amino acid. Others included mis-sense, non-sense and frame-shift mutations. Although *DNMT3A* is directly related to DNA methylation, the real significance of this mutation to leukemogenesis remains unknown. First, the wide spreading of mutation spots in *DNMT3A* suggests a loss-of-function mutation, but the remarkable aggregate of mutation at R882 implies a gain of function. Reduction of DNA methylation in 182 genomic areas was noted in R882 mutation-harboring AML cells, however, the methylation patterns of vast majority of cytosine methylation regions are the same as wild type.(Ley, *et al* 2010)

DNMT3A mutations are associated with intermediate or normal cytogenetics, higher WBC counts, FAB M4/M5 subtypes, and *FLT3*-ITD, *NPM1*, and *IDH1* mutations but mutually exclusive with favorable karyotypes.(Ley, *et al* 2010, Thol, *et al* 2011) In our study of 500 AML patients, *DNMT3A* mutations were identified in 14% of total patients and 22.9% of patients with CN-AML. (Hou, *et al* 2011) In addition to the findings shown in previous reports, (Ley, *et al* 2010, Thol, *et al* 2011) we for the first time identified the *DNMT3A* mutation was positively associated with *PTPN11* and *IDH2* mutations, but negatively associated with *CEBPA* mutation.(Hou, *et al* 2011) Intriguingly, the majority (97.1%) of the *DNMT3A*-mutated patients showed additional molecular alterations at diagnosis. This mutation renders poor OS among all AML patients, patients with a normal karyotype, and those with *FLT3*-ITD.(Hou, *et al* 2011, Ley, *et al* 2010, Thol, *et al* 2011) Importantly, *DNMT3A* mutation is an independent poor prognostic factor. Further, *DNMT3A* mutation is rather stable during disease progression and can be a potential biomarker for monitoring of MRD.(Hou, *et al* 2011)

6. Gene mutations as markers to monitor Minimal Residual Disease (MRD)

Since gene mutations are theoretically absent in healthy people and restricted in leukemia cells, it is reasonable to monitor MRD by detection of gene mutations. This is an advantage of leukemia over solid tumors in that leukemia cells are indigenous to blood and marrow, which are easy for access. There are two critical considerations of MRD monitoring by gene mutations: one is the stability of the mutations, and the other is the pattern of mutation. An

ideal MRD marker should be very consistent with disease status, while those that may disappear after disease evolution are not suitable for this purpose. Also, if the mutation appears as a point mutation, probably only qualitative rather than absolute quantitative measurement can be achieved because of inevitable background signals due to minimal sequence differences between wild-type and mutant alleles. Moreover, if the mutation occurs sporadically across the whole coding sequence without a hot spot, the absolute quantification techniques (usually fluorescence-based real-time PCR) would become very cumbersome.

Among the mutations in AML, *NPM1* mutation is perhaps the most useful and intensively studied marker of MRD because this mutation is quite stable, relatively prevalent, highly concentrated at a hot spot, and has 4 nucleotide insertion, which can be clearly discriminated from the wild-type allele in quantitative real-time PCR.(Chou, *et al* 2007, Schnittger, *et al* 2009) Studies have shown *NPM1* mutant levels reflect disease status, predict impending relapse, and bring prognostic implication.(Chou, *et al* 2007, Gorello, *et al* 2006, Kronke, *et al* 2011, Schnittger, *et al* 2009) Another stable marker is *IDH* mutation. *IDH1* and *IDH2* mutations are stable and highly consistent with disease status.(Chou, *et al* 2010a, Chou, *et al* 2011c) We have developed a single-tube, highly sensitive and specific PCR method to detect all *IDH1* mutations at R132 residue.(Chou, *et al* 2010c) However, the *IDH* mutation is not a good marker for MRD monitoring because the minimal difference between the point mutation and normal allele.

Other gene mutations are not readily applicable in MRD monitoring. *FLT3*-ITD is not stable. This mutation can disappear at disease relapse in a significant proportion of patients,(Chou, *et al* 2011b, Shih, *et al* 2002) although this length mutation can be readily and sensitively detected by GeneScan-based method.(Stirewalt and Radich 2003) *DNMT3A* mutation at R882, which occurs at a frequency of up to 60% of all *DNMT3A* mutation, can be a potential marker for qualitative assessment of MRD, but awaits for further testing.(Ley, *et al* 2010, Thol, *et al* 2011, Hou, *et al* 2011) *ASXL1* and *TET2* mutations do not have hot spots and are not stable during AML evolution. Other mutations have lower incidences and have not been well investigated in MRD monitoring.

7. Risk-adapted treatment according to gene mutations in AML patients

The ultimate goal of risk stratification according to molecular alterations is to explore personalized therapy, thereby reduce the risk of relapse and treatment-related side effects. How to integrate gene mutations into clinical management is a crucial issue. The choice between high-dose Cytarabine (HDAC) and allogeneic HSCT as the post-remission therapy is traditionally based on the cytogenetic risks and the patients' condition. The meta-analysis showed that allogeneic HSCT resulted in better clinical outcome in younger AML patients with intermediate- and unfavorable-risk cytogenetics in first CR.(Cornelissen, *et al* 2007, Koreth, *et al* 2009) Although allogeneic HSCT reduces the risk of relapse and is a curative approach for AML patients, the higher rate of transplantation related morbidity and mortality counterbalances its beneficial effect. Thus, allogeneic HSCT is currently recommended only in those patients with acceptable benefit-risk ratio. Given that AML is a heterogenous disease especially in intermediate-risk cytogenetics and CN-AML, increasing understanding of novel molecular genetic markers in AML leukemogenesis can further help to reassess the value of HSCT in different prognostic groups.

Recently, ELN proposed a new classification to stratify AML patients into different risk groups according to cytogenetics and genetic alterations.(Dohner, *et al* 2010) In addition to CBF AML, CN-AML with mutated *NPM1* without *FLT3*-ITD and those with mutated *CEBPA* are categorized as favorable-risk groups; the regimen using repetitive cycles of HDAC as postremission therapy is considered beneficial for this group of patients. Allogeneic HSCT in first CR is not beneficial for CN-AML patients with mutated *NPM1* without *FLT3*-ITD,(Schlenk, *et al* 2008) and probably neither for those with mutated *CEBPA*. Allogeneic HSCT is generally not considered in patients with CBF AML in first CR, but may be indicated in those who harbor *KIT* mutations because such patients did poorly with chemotherapy. For the patients with adverse-risk genotype (other than mutated *NPM1* without *FLT3*/ITD or mutated *CEBPA*), an allogeneic HSCT from a matched related donor or even unrelated donor in first CR is suggested.(Basara, *et al* 2009, Cornelissen, *et al* 2007, Slovak, *et al* 2000, Suciu, *et al* 2003, Tallman, *et al* 2007) Recent studies showed that allogeneic HSCT may be considered in patients with *FLT3*-ITD even if definite results of prospective trials are not available.(Bornhauser, *et al* 2007, Gale, *et al* 2005, Schlenk, *et al* 2008) Besides, allogeneic HSCT also ameliorates the poor survival impact of *RUNX1* mutations on AML patients.(Gaidzik, *et al* 2011, Tang, *et al* 2009) The treatment of choice for patients with other recently documented poor-risk mutations, such as *WT1*, *TET2* and *DNMT3A* mutations is currently unclear.

In addition to chemotherapy and transplantation, targeted therapies aiming to specific molecular pathway are evolving as an adjunctive treatment in AML patients. *FLT3*/ITD and *FLT3*/TKD occur in about 20-35% of AML patients. Since *FLT3* is a receptor tyrosine kinase and promote cancer phenotypes, it is an ideal target for therapy. Several *FLT3* inhibitors, such as sorafenib, PKC-412 (midostaurin), sunitinib, semaxanib, tandutinib, AC220, KW-2449, and CEP701 (lestaurtinib) have been used in clinical trials and some effects were noticed in relapse/refractory setting.(Levis, *et al* 2002, Metzelder, *et al* 2009, Stone, *et al* 2005, Zhang, *et al* 2008) An ongoing international intergroup trial (10603 RATIFY), incorporating midostaurin into induction, consolidation or maintenance setting is currently underway. All-trans retinoic acid in combination with chemotherapy was found to be beneficial for *NPM1*-mutated patients (Burnett, *et al* 2010); however this preliminary result was not confirmed by the other study done on younger patients.(Schlenk, *et al* 2009) Tyrosine kinase inhibitor, such as imatinib, might be of clinical value in treatment of patients with *KIT* mutations.(Kindler, *et al* 2004, Kindler, *et al* 2003, Kohl, *et al* 2005) Epigenetic modification through demethylation agent azacitidine or decitabine may play a role in the treatment of patients with *MLL* rearrangement,(Altucci and Minucci 2009) and those with genetic alterations relating to epigenetic changes, such as *TET2* mutations.(Itzykson, *et al* 2011) Besides, recent report demonstrated that inhibition of glutaminase preferentially killed *IDH1*-mutated glial cells, which were more dependent on glutaminolysis pathway to supply α -KG, so glutaminase itself could be a potential therapeutical target.(Seltzer, *et al* 2010) Eventually, it may be reasonable to use combinations of molecularly targeted therapies and chemotherapy to improve the clinical outcome in AML patients.

8. Acknowledgment

This work was partially sponsored by grants from the National Science Council (NSC 97-2314-B002-015-MY3, 98-2314-B-002-033-MY3, 100-2325-B-002-032, 100-2325-B-002-033 and 100-2628-B-002-003-MY3), National Health Research Institute (NHRI-EX97-9731BI),

Department of Health (DOH100-TD-C-111-001), and Department of Medical Research (NTUH.99P14), National Taiwan University Hospital, Taiwan, Republic of China

9. References

- Abbas, S., et al. (2010) Acquired mutations in the genes encoding IDH1 and IDH2 both are recurrent aberrations in acute myeloid leukemia: prevalence and prognostic value. *Blood*, 116, 2122-2126.
- Abdel-Wahab, O., et al. (2010) Genetic analysis of transforming events that convert chronic myeloproliferative neoplasms to leukemias. *Cancer Res*, 70, 447-452.
- Altucci, L. & Minucci, S. (2009) Epigenetic therapies in haematological malignancies: searching for true targets. *Eur J Cancer*, 45, 1137-1145.
- Bacher, U., et al. (2006) Implications of NRAS mutations in AML: a study of 2502 patients. *Blood*, 107, 3847-3853.
- Bacher, U., et al. (2008) Prognostic relevance of FLT3-TKD mutations in AML: the combination matters--an analysis of 3082 patients. *Blood*, 111, 2527-2537.
- Bacher, U., et al. (2010) Mutations of the TET2 and CBL genes: novel molecular markers in myeloid malignancies. *Ann Hematol*, 89, 643-652.
- Baird, P.N. & Simmons, P.J. (1997) Expression of the Wilms' tumor gene (WT1) in normal hemopoiesis. *Exp Hematol*, 25, 312-320.
- Barjesteh van Waalwijk van Doorn-Khosrovani, S., et al. (2003) Biallelic mutations in the CEBPA gene and low CEBPA expression levels as prognostic markers in intermediate-risk AML. *Hematol J*, 4, 31-40.
- Basara, N., et al. (2009) Early related or unrelated haematopoietic cell transplantation results in higher overall survival and leukaemia-free survival compared with conventional chemotherapy in high-risk acute myeloid leukaemia patients in first complete remission. *Leukemia*, 23, 635-640.
- Baxter, E.J., et al. (2005) Acquired mutation of the tyrosine kinase JAK2 in human myeloproliferative disorders. *Lancet*, 365, 1054-1061.
- Baylin, S.B. & Ohm, J.E. (2006) Epigenetic gene silencing in cancer - a mechanism for early oncogenic pathway addiction? *Nat Rev Cancer*, 6, 107-116.
- Beghini, A., et al. (2000) C-kit mutations in core binding factor leukemias. *Blood*, 95, 726-727.
- Beghini, A., et al. (2004) KIT activating mutations: incidence in adult and pediatric acute myeloid leukemia, and identification of an internal tandem duplication. *Haematologica*, 89, 920-925.
- Bergmann, L., et al. (1997) High levels of Wilms' tumor gene (wt1) mRNA in acute myeloid leukemias are associated with a worse long-term outcome. *Blood*, 90, 1217-1225.
- Bienz, M., et al. (2005) Risk assessment in patients with acute myeloid leukemia and a normal karyotype. *Clin Cancer Res*, 11, 1416-1424.
- Blume-Jensen, P. & Hunter, T. (2001) Oncogenic kinase signalling. *Nature*, 411, 355-365.
- Boissel, N., et al. (2005) Prevalence, clinical profile, and prognosis of NPM mutations in AML with normal karyotype. *Blood*, 106, 3618-3620.
- Boissel, N., et al. (2006) Incidence and prognostic impact of c-Kit, FLT3, and Ras gene mutations in core binding factor acute myeloid leukemia (CBF-AML). *Leukemia*, 20, 965-970.

- Boissel, N., et al. (2010) Prognostic impact of isocitrate dehydrogenase enzyme isoforms 1 and 2 mutations in acute myeloid leukemia: a study by the Acute Leukemia French Association group. *J Clin Oncol*, 28, 3717-3723.
- Bornhauser, M., et al. (2007) Improved outcome after stem-cell transplantation in FLT3/ITD-positive AML. *Blood*, 109, 2264-2265; author reply 2265.
- Bos, J.L., et al. (1987) Mutations in N-ras predominate in acute myeloid leukemia. *Blood*, 69, 1237-1241.
- Bos, J.L. (1989) ras oncogenes in human cancer: a review. *Cancer Res*, 49, 4682-4689.
- Boulwood, J., et al. (2010) Frequent mutation of the polycomb-associated gene ASXL1 in the myelodysplastic syndromes and in acute myeloid leukemia. *Leukemia*, 24, 1062-1065.
- Bowen, D.T., et al. (2005) RAS mutation in acute myeloid leukemia is associated with distinct cytogenetic subgroups but does not influence outcome in patients younger than 60 years. *Blood*, 106, 2113-2119.
- Burnett, A.K., et al. (2010) The impact on outcome of the addition of all-trans retinoic acid to intensive chemotherapy in younger patients with nonacute promyelocytic acute myeloid leukemia: overall results and results in genotypic subgroups defined by mutations in NPM1, FLT3, and CEBPA. *Blood*, 115, 948-956.
- Caligiuri, M.A., et al. (1994) Molecular rearrangement of the ALL-1 gene in acute myeloid leukemia without cytogenetic evidence of 11q23 chromosomal translocations. *Cancer Res*, 54, 370-373.
- Cammenga, J., et al. (2003) Induction of C/EBPalpha activity alters gene expression and differentiation of human CD34+ cells. *Blood*, 101, 2206-2214.
- Carbuccia, N., et al. (2009) Mutations of ASXL1 gene in myeloproliferative neoplasms. *Leukemia*, 23, 2183-2186.
- Carbuccia, N., et al. (2010) Mutual exclusion of ASXL1 and NPM1 mutations in a series of acute myeloid leukemias. *Leukemia*, 24, 469-473.
- Care, R.S., et al. (2003) Incidence and prognosis of c-KIT and FLT3 mutations in core binding factor (CBF) acute myeloid leukaemias. *Br J Haematol*, 121, 775-777.
- Chen, C.Y., et al. (2006) Acquisition of JAK2, PTPN11, and RAS mutations during disease progression in primary myelodysplastic syndrome. *Leukemia*, 20, 1155-1158.
- Chen, J., et al. (2010) Leukaemogenesis: more than mutant genes. *Nat Rev Cancer*, 10, 23-36.
- Chou, W.C., et al. (2006) Nucleophosmin mutations in de novo acute myeloid leukemia: the age-dependent incidences and the stability during disease evolution. *Cancer Res*, 66, 3310-3316.
- Chou, W.C., et al. (2007) Clinical implications of minimal residual disease monitoring by quantitative polymerase chain reaction in acute myeloid leukemia patients bearing nucleophosmin (NPM1) mutations. *Leukemia*, 21, 998-1004.
- Chou, W.C., et al. (2010a) Distinct clinical and biologic characteristics in adult acute myeloid leukemia bearing the isocitrate dehydrogenase 1 mutation. *Blood*, 115, 2749-2754.
- Chou, W.C., et al. (2010b) Distinct clinical and biological features of de novo acute myeloid leukemia with additional sex comb-like 1 (ASXL1) mutations. *Blood*, 116, 4086-4094.
- Chou, W.C., et al. (2010c) A single-tube, sensitive multiplex method for screening of isocitrate dehydrogenase 1 (IDH1) mutations. *Blood*, 116, 495-496.
- Chou, W.C., et al. (2011a) TET2 mutation is an unfavorable prognostic factor in acute myeloid leukemia patients with intermediate-risk cytogenetics. *Blood*.

- Chou, W.C., et al. (2011b) Sensitive measurement of quantity dynamics of FLT3 internal tandem duplication at early time points provides prognostic information. *Ann Oncol*, 22, 696-704.
- Chou, W.C., et al. (2011c) The prognostic impact and stability of Isocitrate dehydrogenase 2 mutation in adult patients with acute myeloid leukemia. *Leukemia*, 25, 246-253.
- Coghlan, D.W., et al. (1994) The incidence and prognostic significance of mutations in codon 13 of the N-ras gene in acute myeloid leukemia. *Leukemia*, 8, 1682-1687.
- Cornelissen, J.J., et al. (2007) Results of a HOVON/SAKK donor versus no-donor analysis of myeloablative HLA-identical sibling stem cell transplantation in first remission acute myeloid leukemia in young and middle-aged adults: benefits for whom? *Blood*, 109, 3658-3666.
- Couronne, L., et al. (2010) Analyses of TET2 mutations in post-myeloproliferative neoplasm acute myeloid leukemias. *Leukemia*, 24, 201-203.
- Dang, L., et al. (2009) Cancer-associated IDH1 mutations produce 2-hydroxyglutarate. *Nature*, 462, 739-744.
- Dange, M. & Colman, R.F. (2010) Each conserved active site Tyr in the three subunits of human isocitrate dehydrogenase has a different function. *J Biol Chem*.
- De Melo, M.B., et al. (1997) N-ras gene point mutations in Brazilian acute myelogenous leukemia patients correlate with a poor prognosis. *Leuk Lymphoma*, 24, 309-317.
- Delhommeau, F., et al. (2009) Mutation in TET2 in myeloid cancers. *N Engl J Med*, 360, 2289-2301.
- Dicker, F., et al. (2007) Trisomy 13 is strongly associated with AML1/RUNX1 mutations and increased FLT3 expression in acute myeloid leukemia. *Blood*, 110, 1308-1316.
- Dimartino, J.F. & Cleary, M.L. (1999) Mll rearrangements in haematological malignancies: lessons from clinical and biological studies. *Br J Haematol*, 106, 614-626.
- Dohner, H., et al. (2010) Diagnosis and management of acute myeloid leukemia in adults: recommendations from an international expert panel, on behalf of the European LeukemiaNet. *Blood*, 115, 453-474.
- Dohner, K., et al. (2002) Prognostic significance of partial tandem duplications of the MLL gene in adult patients 16 to 60 years old with acute myeloid leukemia and normal cytogenetics: a study of the Acute Myeloid Leukemia Study Group Ulm. *J Clin Oncol*, 20, 3254-3261.
- Dohner, K., et al. (2005) Mutant nucleophosmin (NPM1) predicts favorable prognosis in younger adults with acute myeloid leukemia and normal cytogenetics: interaction with other gene mutations. *Blood*, 106, 3740-3746.
- Downward, J. (2003) Targeting RAS signalling pathways in cancer therapy. *Nat Rev Cancer*, 3, 11-22.
- Dufour, A., et al. (2010) Acute myeloid leukemia with biallelic CEBPA gene mutations and normal karyotype represents a distinct genetic entity associated with a favorable clinical outcome. *J Clin Oncol*, 28, 570-577.
- Ellisen, L.W., et al. (2001) The Wilms tumor suppressor WT1 directs stage-specific quiescence and differentiation of human hematopoietic progenitor cells. *Embo J*, 20, 1897-1909.
- Falini, B., et al. (2005) Cytoplasmic nucleophosmin in acute myelogenous leukemia with a normal karyotype. *N Engl J Med*, 352, 254-266.
- Falini, B., et al. (2011) Acute myeloid leukemia with mutated nucleophosmin (NPM1): is it a distinct entity? *Blood*, 117, 1109-1120.

- Farr, C.J., et al. (1988) Analysis of RAS gene mutations in acute myeloid leukemia by polymerase chain reaction and oligonucleotide probes. *Proc Natl Acad Sci U S A*, 85, 1629-1633.
- Figueroa, M.E., et al. (2010) Leukemic IDH1 and IDH2 mutations result in a hypermethylation phenotype, disrupt TET2 function, and impair hematopoietic differentiation. *Cancer Cell*, 18, 553-567.
- Fisher, C.L., et al. (2003) A human homolog of Additional sex combs, ADDITIONAL SEX COMBS-LIKE 1, maps to chromosome 20q11. *Gene*, 306, 115-126.
- Fisher, C.L., et al. (2006) Characterization of *Asx11*, a murine homolog of Additional sex combs, and analysis of the *Asx*-like gene family. *Gene*, 369, 109-118.
- Flach, J., et al. (2010) Mutations of JAK2 and TET2, but not CBL are detectable in a high portion of patients with refractory anemia with ring sideroblasts and thrombocytosis. *Haematologica*, 95, 518-519.
- Friedman, A.D. (2009) Cell cycle and developmental control of hematopoiesis by Runx1. *J Cell Physiol*, 219, 520-524.
- Friedman, A.D. & McKnight, S.L. (1990) Identification of two polypeptide segments of CCAAT/enhancer-binding protein required for transcriptional activation of the serum albumin gene. *Genes Dev*, 4, 1416-1426.
- Frohling, S., et al. (2004) CEBPA mutations in younger adults with acute myeloid leukemia and normal cytogenetics: prognostic relevance and analysis of cooperating mutations. *J Clin Oncol*, 22, 624-633.
- Frohling, S., et al. (2005) Genetics of myeloid malignancies: pathogenetic and clinical implications. *J Clin Oncol*, 23, 6285-6295.
- Frohling, S., et al. (2006) Rare occurrence of the JAK2 V617F mutation in AML subtypes M5, M6, and M7. *Blood*, 107, 1242-1243.
- Gaidzik, V.I., et al. (2009) Prognostic impact of WT1 mutations in cytogenetically normal acute myeloid leukemia: a study of the German-Austrian AML Study Group. *Blood*, 113, 4505-4511.
- Gaidzik, V.I., et al. (2011) RUNX1 mutations in acute myeloid leukemia: results from a comprehensive genetic and clinical analysis from the AML study group. *J Clin Oncol*, 29, 1364-1372.
- Gale, R.E., et al. (2005) No evidence that FLT3 status should be considered as an indicator for transplantation in acute myeloid leukemia (AML): an analysis of 1135 patients, excluding acute promyelocytic leukemia, from the UK MRC AML10 and 12 trials. *Blood*, 106, 3658-3665.
- Gale, R.E., et al. (2008) The impact of FLT3 internal tandem duplication mutant level, number, size, and interaction with NPM1 mutations in a large cohort of young adult patients with acute myeloid leukemia. *Blood*, 111, 2776-2784.
- Gelsi-Boyer, V., et al. (2009) Mutations of polycomb-associated gene ASXL1 in myelodysplastic syndromes and chronic myelomonocytic leukaemia. *Br J Haematol*, 145, 788-800.
- Gilliland, D.G. (2002) Molecular genetics of human leukemias: new insights into therapy. *Semin Hematol*, 39, 6-11.
- Goldman, J.M. (2005) A unifying mutation in chronic myeloproliferative disorders. *N Engl J Med*, 352, 1744-1746.

- Gorello, P., et al. (2006) Quantitative assessment of minimal residual disease in acute myeloid leukemia carrying nucleophosmin (NPM1) gene mutations. *Leukemia*, 20, 1103-1108.
- Green, C.L., et al. (2010) The prognostic significance of IDH1 mutations in younger adult patients with acute myeloid leukemia is dependent on FLT3/ITD status. *Blood*, 116, 2779-2782.
- Haber, D.A., et al. (1990) An internal deletion within an 11p13 zinc finger gene contributes to the development of Wilms' tumor. *Cell*, 61, 1257-1269.
- Harada, H., et al. (2004) High incidence of somatic mutations in the AML1/RUNX1 gene in myelodysplastic syndrome and low blast percentage myeloid leukemia with myelodysplasia. *Blood*, 103, 2316-2324.
- Hou, H.A., et al. (2008) Characterization of acute myeloid leukemia with PTPN11 mutation: the mutation is closely associated with NPM1 mutation but inversely related to FLT3/ITD. *Leukemia*, 22, 1075-1078.
- Hou, H.A., et al. (2009) Reply to 'Heterogeneity within AML with CEBPA mutations; only CEBPA double mutations, but not single CEBPA mutations are associated with favorable prognosis'. *Br J Cancer*, 101, 738-740.
- Hou, H.A., et al. (2010) WT1 mutation in 470 adult patients with acute myeloid leukemia: stability during disease evolution and implication of its incorporation into a survival scoring system. *Blood*, 115, 5222-5231.
- Hou, H.A., et al. (2011) DNMT3A Mutations in Acute Myeloid Leukemia-Stability during Disease Evolution and the Clinical Implication. *Blood*, in revision.
- Ikeda, H., et al. (1991) Expression and functional role of the proto-oncogene c-kit in acute myeloblastic leukemia cells. *Blood*, 78, 2962-2968.
- Illmer, T., et al. (2007) Tyrosine kinase mutations of JAK2 are rare events in AML but influence prognosis of patients with CBF-leukemias. *Haematologica*, 92, 137-138.
- Ito, S., et al. (2010) Role of Tet proteins in 5mC to 5hmC conversion, ES-cell self-renewal and inner cell mass specification. *Nature*, 466, 1129-1133.
- Ito, Y. (2008) RUNX genes in development and cancer: regulation of viral gene expression and the discovery of RUNX family genes. *Adv Cancer Res*, 99, 33-76.
- Itzykson, R., et al. (2011) Impact of TET2 mutations on response rate to azacitidine in myelodysplastic syndromes and low blast count acute myeloid leukemias. *Leukemia*, 25, 1147-1152.
- Jankowska, A.M., et al. (2009) Loss of heterozygosity 4q24 and TET2 mutations associated with myelodysplastic/myeloproliferative neoplasms. *Blood*, 113, 6403-6410.
- Jones, P.A. & Baylin, S.B. (2002) The fundamental role of epigenetic events in cancer. *Nat Rev Genet*, 3, 415-428.
- Kindler, T., et al. (2003) Sustained complete hematologic remission after administration of the tyrosine kinase inhibitor imatinib mesylate in a patient with refractory, secondary AML. *Blood*, 101, 2960-2962.
- Kindler, T., et al. (2004) Efficacy and safety of imatinib in adult patients with c-kit-positive acute myeloid leukemia. *Blood*, 103, 3644-3654.
- King-Underwood, L., et al. (1996) Mutations in the Wilms' tumor gene WT1 in leukemias. *Blood*, 87, 2171-2179.
- Kiyoi, H., et al. (1998) Internal tandem duplication of the FLT3 gene is a novel modality of elongation mutation which causes constitutive activation of the product. *Leukemia*, 12, 1333-1337.

- Kiyoi, H., et al. (1999) Prognostic implication of FLT3 and N-RAS gene mutations in acute myeloid leukemia. *Blood*, 93, 3074-3080.
- Kohl, T.M., et al. (2005) KIT exon 8 mutations associated with core-binding factor (CBF)-acute myeloid leukemia (AML) cause hyperactivation of the receptor in response to stem cell factor. *Blood*, 105, 3319-3321.
- Koreth, J., et al. (2009) Allogeneic stem cell transplantation for acute myeloid leukemia in first complete remission: systematic review and meta-analysis of prospective clinical trials. *JAMA*, 301, 2349-2361.
- Koschmieder, S., et al. (2009) Dysregulation of the C/EBPalpha differentiation pathway in human cancer. *J Clin Oncol*, 27, 619-628.
- Kosmider, O., et al. (2009a) TET2 mutation is an independent favorable prognostic factor in myelodysplastic syndromes (MDSs). *Blood*, 114, 3285-3291.
- Kosmider, O., et al. (2009b) TET2 gene mutation is a frequent and adverse event in chronic myelomonocytic leukemia. *Haematologica*, 94, 1676-1681.
- Kosmider, O., et al. (2010) Mutations of IDH1 and IDH2 genes in early and accelerated phases of myelodysplastic syndromes and MDS/myeloproliferative neoplasms. *Leukemia*, 24, 1094-1096.
- Kottaridis, P.D., et al. (2001) The presence of a FLT3 internal tandem duplication in patients with acute myeloid leukemia (AML) adds important prognostic information to cytogenetic risk group and response to the first cycle of chemotherapy: analysis of 854 patients from the United Kingdom Medical Research Council AML 10 and 12 trials. *Blood*, 98, 1752-1759.
- Kottaridis, P.D., et al. (2002) Studies of FLT3 mutations in paired presentation and relapse samples from patients with acute myeloid leukemia: implications for the role of FLT3 mutations in leukemogenesis, minimal residual disease detection, and possible therapy with FLT3 inhibitors. *Blood*, 100, 2393-2398.
- Kralovics, R., et al. (2005) A gain-of-function mutation of JAK2 in myeloproliferative disorders. *N Engl J Med*, 352, 1779-1790.
- Kronke, J., et al. (2011) Monitoring of Minimal Residual Disease in NPM1-Mutated Acute Myeloid Leukemia: A Study From the German-Austrian Acute Myeloid Leukemia Study Group. *J Clin Oncol*, 29, 2709-2716.
- Langemeijer, S.M., et al. (2009) Acquired mutations in TET2 are common in myelodysplastic syndromes. *Nat Genet*, 41, 838-842.
- Lee, J.W., et al. (2006) The JAK2 V617F mutation in de novo acute myelogenous leukemias. *Oncogene*, 25, 1434-1436.
- Levine, R.L., et al. (2005a) The JAK2V617F activating mutation occurs in chronic myelomonocytic leukemia and acute myeloid leukemia, but not in acute lymphoblastic leukemia or chronic lymphocytic leukemia. *Blood*, 106, 3377-3379.
- Levine, R.L., et al. (2005b) Activating mutation in the tyrosine kinase JAK2 in polycythemia vera, essential thrombocythemia, and myeloid metaplasia with myelofibrosis. *Cancer Cell*, 7, 387-397.
- Levis, M., et al. (2002) A FLT3-targeted tyrosine kinase inhibitor is cytotoxic to leukemia cells in vitro and in vivo. *Blood*, 99, 3885-3891.
- Ley, T.J., et al. (2010) DNMT3A mutations in acute myeloid leukemia. *N Engl J Med*, 363, 2424-2433.

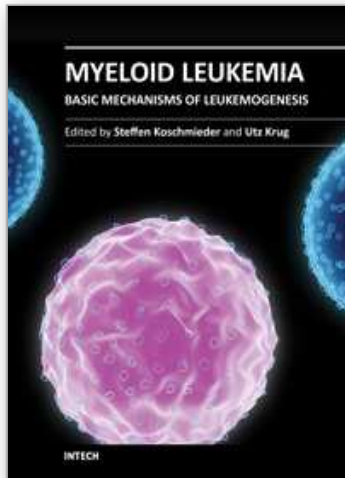
- Lin, F.T., et al. (1993) A 30-kDa alternative translation product of the CCAAT/enhancer binding protein alpha message: transcriptional activator lacking antimetabolic activity. *Proc Natl Acad Sci U S A*, 90, 9606-9610.
- Lin, L.I., et al. (2005) Characterization of CEBPA mutations in acute myeloid leukemia: most patients with CEBPA mutations have biallelic mutations and show a distinct immunophenotype of the leukemic cells. *Clin Cancer Res*, 11, 1372-1379.
- Liu, R.Y., et al. (1999) Constitutive activation of the JAK2/STAT5 signal transduction pathway correlates with growth factor independence of megakaryocytic leukemic cell lines. *Blood*, 93, 2369-2379.
- Loh, M.L., et al. (2004a) PTPN11 mutations in pediatric patients with acute myeloid leukemia: results from the Children's Cancer Group. *Leukemia*, 18, 1831-1834.
- Loh, M.L., et al. (2004b) Mutations in PTPN11 implicate the SHP-2 phosphatase in leukemogenesis. *Blood*, 103, 2325-2331.
- Marcucci, G., et al. (2010) IDH1 and IDH2 gene mutations identify novel molecular subsets within de novo cytogenetically normal acute myeloid leukemia: a Cancer and Leukemia Group B study. *J Clin Oncol*, 28, 2348-2355.
- Mardis, E.R., et al. (2009) Recurring mutations found by sequencing an acute myeloid leukemia genome. *N Engl J Med*, 361, 1058-1066.
- Meshinchi, S., et al. (2003) Activating mutations of RTK/ras signal transduction pathway in pediatric acute myeloid leukemia. *Blood*, 102, 1474-1479.
- Metzelder, S., et al. (2009) Compassionate use of sorafenib in FLT3-ITD-positive acute myeloid leukemia: sustained regression before and after allogeneic stem cell transplantation. *Blood*, 113, 6567-6571.
- Metzeler, K.H., et al. (2011) TET2 mutations improve the new European LeukemiaNet risk classification of acute myeloid leukemia: a Cancer and Leukemia Group B study. *J Clin Oncol*, 29, 1373-1381.
- Michaud, J., et al. (2002) In vitro analyses of known and novel RUNX1/AML1 mutations in dominant familial platelet disorder with predisposition to acute myelogenous leukemia: implications for mechanisms of pathogenesis. *Blood*, 99, 1364-1372.
- Miwa, H., et al. (1992) Expression of the Wilms' tumor gene (WT1) in human leukemias. *Leukemia*, 6, 405-409.
- Munoz, L., et al. (2003) Acute myeloid leukemia with MLL rearrangements: clinicobiological features, prognostic impact and value of flow cytometry in the detection of residual leukemic cells. *Leukemia*, 17, 76-82.
- Nakao, M., et al. (1996) Internal tandem duplication of the flt3 gene found in acute myeloid leukemia. *Leukemia*, 10, 1911-1918.
- Neel, B.G., et al. (2003) The 'Shp'ing news: SH2 domain-containing tyrosine phosphatases in cell signaling. *Trends Biochem Sci*, 28, 284-293.
- Neubauer, A., et al. (1994) Prognostic importance of mutations in the ras proto-oncogenes in de novo acute myeloid leukemia. *Blood*, 83, 1603-1611.
- Neubauer, A., et al. (2008) Patients with acute myeloid leukemia and RAS mutations benefit most from postremission high-dose cytarabine: a Cancer and Leukemia Group B study. *J Clin Oncol*, 26, 4603-4609.
- Niebuhr, B., et al. (2008) Gatekeeper function of the RUNX1 transcription factor in acute leukemia. *Blood Cells Mol Dis*, 40, 211-218.
- Niimi, H., et al. (2006) Hyperactivation of the RAS signaling pathway in myelodysplastic syndrome with AML1/RUNX1 point mutations. *Leukemia*, 20, 635-644.

- Oelgeschlager, M., et al. (1996) C/EBP, c-Myb, and PU.1 cooperate to regulate the neutrophil elastase promoter. *Mol Cell Biol*, 16, 4717-4725.
- Osato, M. (2004) Point mutations in the RUNX1/AML1 gene: another actor in RUNX leukemia. *Oncogene*, 23, 4284-4296.
- Pabst, T., et al. (2001a) AML1-ETO downregulates the granulocytic differentiation factor C/EBPalpha in t(8;21) myeloid leukemia. *Nat Med*, 7, 444-451.
- Pabst, T., et al. (2001b) Dominant-negative mutations of CEBPA, encoding CCAAT/enhancer binding protein-alpha (C/EBPalpha), in acute myeloid leukemia. *Nat Genet*, 27, 263-270.
- Pabst, T., et al. (2009) Heterogeneity within AML with CEBPA mutations; only CEBPA double mutations, but not single CEBPA mutations are associated with favourable prognosis. *Br J Cancer*, 100, 1343-1346.
- Palmisano, M., et al. (2007) NPM1 mutations are more stable than FLT3 mutations during the course of disease in patients with acute myeloid leukemia. *Haematologica*, 92, 1268-1269.
- Pardanani, A., et al. (2010) IDH1 and IDH2 mutation analysis in chronic- and blast-phase myeloproliferative neoplasms. *Leukemia*, 24, 1146-1151.
- Parsons, D.W., et al. (2008) An integrated genomic analysis of human glioblastoma multiforme. *Science*, 321, 1807-1812.
- Paschka, P., et al. (2008) Wilms' tumor 1 gene mutations independently predict poor outcome in adults with cytogenetically normal acute myeloid leukemia: a cancer and leukemia group B study. *J Clin Oncol*, 26, 4595-4602.
- Pena, P.V., et al. (2006) Molecular mechanism of histone H3K4me3 recognition by plant homeodomain of ING2. *Nature*, 442, 100-103.
- Preudhomme, C., et al. (2000) High incidence of biallelic point mutations in the Runt domain of the AML1/PEBP2 alpha B gene in Mo acute myeloid leukemia and in myeloid malignancies with acquired trisomy 21. *Blood*, 96, 2862-2869.
- Preudhomme, C., et al. (2002) Favorable prognostic significance of CEBPA mutations in patients with de novo acute myeloid leukemia: a study from the Acute Leukemia French Association (ALFA). *Blood*, 100, 2717-2723.
- Radich, J.P., et al. (1990) N-ras mutations in adult de novo acute myelogenous leukemia: prevalence and clinical significance. *Blood*, 76, 801-807.
- Rege-Cambrin, G., et al. (2005) Trisomy 11 in myeloid malignancies is associated with internal tandem duplication of both MLL and FLT3 genes. *Haematologica*, 90, 262-264.
- Reitman, Z.J., et al. (2010) IDH1 and IDH2: not your typical oncogenes. *Cancer Cell*, 17, 215-216.
- Renneville, A., et al. (2009a) The favorable impact of CEBPA mutations in patients with acute myeloid leukemia is only observed in the absence of associated cytogenetic abnormalities and FLT3 internal duplication. *Blood*, 113, 5090-5093.
- Renneville, A., et al. (2009b) Wilms tumor 1 gene mutations are associated with a higher risk of recurrence in young adults with acute myeloid leukemia: a study from the Acute Leukemia French Association. *Cancer*, 115, 3719-3727.
- Reuss-Borst, M.A., et al. (1994) AML: immunophenotypic heterogeneity and prognostic significance of c-kit expression. *Leukemia*, 8, 258-263.
- Reuther, G.W. & Der, C.J. (2000) The Ras branch of small GTPases: Ras family members don't fall far from the tree. *Curr Opin Cell Biol*, 12, 157-165.

- Ritter, M., et al. (2004) Prognostic significance of N-RAS and K-RAS mutations in 232 patients with acute myeloid leukemia. *Haematologica*, 89, 1397-1399.
- Saint-Martin, C., et al. (2009) Analysis of the ten-eleven translocation 2 (TET2) gene in familial myeloproliferative neoplasms. *Blood*, 114, 1628-1632.
- Santamaria, C.M., et al. (2009) Molecular stratification model for prognosis in cytogenetically normal acute myeloid leukemia. *Blood*, 114, 148-152.
- Schaub, F.X., et al. (2010) Clonal analysis of TET2 and JAK2 mutations suggests that TET2 can be a late event in the progression of myeloproliferative neoplasms. *Blood*, 115, 2003-2007.
- Schichman, S.A., et al. (1994) ALL-1 partial duplication in acute leukemia. *Proc Natl Acad Sci U S A*, 91, 6236-6239.
- Schichman, S.A., et al. (1995) Self-fusion of the ALL1 gene. A new genetic mechanism for acute leukemia. *JAMA*, 273, 571-576.
- Schlenk, R.F., et al. (2008) Mutations and treatment outcome in cytogenetically normal acute myeloid leukemia. *N Engl J Med*, 358, 1909-1918.
- Schlenk, R.F., et al. (2009) Gene mutations and response to treatment with all-trans retinoic acid in elderly patients with acute myeloid leukemia. Results from the AMLSG Trial AML HD98B. *Haematologica*, 94, 54-60.
- Schnittger, S., et al. (2000) Screening for MLL tandem duplication in 387 unselected patients with AML identify a prognostically unfavorable subset of AML. *Leukemia*, 14, 796-804.
- Schnittger, S., et al. (2005) Nucleophosmin gene mutations are predictors of favorable prognosis in acute myelogenous leukemia with a normal karyotype. *Blood*, 106, 3733-3739.
- Schnittger, S., et al. (2006) KIT-D816 mutations in AML1-ETO-positive AML are associated with impaired event-free and overall survival. *Blood*, 107, 1791-1799.
- Schnittger, S., et al. (2009) Minimal residual disease levels assessed by NPM1 mutation-specific RQ-PCR provide important prognostic information in AML. *Blood*, 114, 2220-2231.
- Schnittger, S., et al. (2011) RUNX1 mutations are frequent in de novo AML with noncomplex karyotype and confer an unfavorable prognosis. *Blood*, 117, 2348-2357.
- Seltzer, M.J., et al. (2010) Inhibition of glutaminase preferentially slows growth of glioma cells with mutant IDH1. *Cancer Res*, 70, 8981-8987.
- Senn, H.P., et al. (1988) Mutation analysis of the N-ras proto-oncogene in active and remission phase of human acute leukemias. *Int J Cancer*, 41, 59-64.
- Shi, X., et al. (2006) ING2 PHD domain links histone H3 lysine 4 methylation to active gene repression. *Nature*, 442, 96-99.
- Shiah, H.S., et al. (2002) Clinical and biological implications of partial tandem duplication of the MLL gene in acute myeloid leukemia without chromosomal abnormalities at 11q23. *Leukemia*, 16, 196-202.
- Shields, J.M., et al. (2000) Understanding Ras: 'it ain't over 'til it's over'. *Trends Cell Biol*, 10, 147-154.
- Shih, L.Y., et al. (2002) Internal tandem duplication of FLT3 in relapsed acute myeloid leukemia: a comparative analysis of bone marrow samples from 108 adult patients at diagnosis and relapse. *Blood*, 100, 2387-2392.

- Slovak, M.L., et al. (2000) Karyotypic analysis predicts outcome of preremission and postremission therapy in adult acute myeloid leukemia: a Southwest Oncology Group/Eastern Cooperative Oncology Group Study. *Blood*, 96, 4075-4083.
- Smith, A.E., et al. (2010) Next-generation sequencing of the TET2 gene in 355 MDS and CMML patients reveals low-abundance mutant clones with early origins, but indicates no definite prognostic value. *Blood*, 116, 3923-3932.
- Smith, L.T., et al. (1996) PU.1 (Spi-1) and C/EBP alpha regulate the granulocyte colony-stimulating factor receptor promoter in myeloid cells. *Blood*, 88, 1234-1247.
- Stirewalt, D.L. & Radich, J.P. (2003) The role of FLT3 in haematopoietic malignancies. *Nat Rev Cancer*, 3, 650-665.
- Stone, R.M., et al. (2005) Patients with acute myeloid leukemia and an activating mutation in FLT3 respond to a small-molecule FLT3 tyrosine kinase inhibitor, PKC412. *Blood*, 105, 54-60.
- Suciu, S., et al. (2003) Allogeneic compared with autologous stem cell transplantation in the treatment of patients younger than 46 years with acute myeloid leukemia (AML) in first complete remission (CR1): an intention-to-treat analysis of the EORTC/GIMEMAAML-10 trial. *Blood*, 102, 1232-1240.
- Suzuki, T., et al. (2005) Clinical characteristics and prognostic implications of NPM1 mutations in acute myeloid leukemia. *Blood*, 106, 2854-2861.
- Tallman, M.S., et al. (2007) Impact of cytogenetics on outcome of matched unrelated donor hematopoietic stem cell transplantation for acute myeloid leukemia in first or second complete remission. *Blood*, 110, 409-417.
- Tang, J.L., et al. (2009) AML1/RUNX1 mutations in 470 adult patients with de novo acute myeloid leukemia: prognostic implication and interaction with other gene alterations. *Blood*, 114, 5352-5361.
- Tartaglia, M., et al. (2001) Mutations in PTPN11, encoding the protein tyrosine phosphatase SHP-2, cause Noonan syndrome. *Nat Genet*, 29, 465-468.
- Tartaglia, M., et al. (2002) PTPN11 mutations in Noonan syndrome: molecular spectrum, genotype-phenotype correlation, and phenotypic heterogeneity. *Am J Hum Genet*, 70, 1555-1563.
- Tartaglia, M., et al. (2003) Somatic mutations in PTPN11 in juvenile myelomonocytic leukemia, myelodysplastic syndromes and acute myeloid leukemia. *Nat Genet*, 34, 148-150.
- Tartaglia, M., et al. (2004) SHP-2 and myeloid malignancies. *Curr Opin Hematol*, 11, 44-50.
- Tartaglia, M., et al. (2005) Somatic PTPN11 mutations in childhood acute myeloid leukaemia. *Br J Haematol*, 129, 333-339.
- Tefferi, A., et al. (2009a) Detection of mutant TET2 in myeloid malignancies other than myeloproliferative neoplasms: CMML, MDS, MDS/MPN and AML. *Leukemia*, 23, 1343-1345.
- Tefferi, A., et al. (2009b) Mutation in TET2 in myeloid cancers. *N Engl J Med*, 361, 1117; author reply 1117-1118.
- Thiede, C., et al. (2006) Prevalence and prognostic impact of NPM1 mutations in 1485 adult patients with acute myeloid leukemia (AML). *Blood*, 107, 4011-4020.
- Thol, F., et al. (2010) Prognostic impact of IDH2 mutations in cytogenetically normal acute myeloid leukemia. *Blood*, 116, 614-616.
- Thol, F., et al. (2011) Incidence and Prognostic Influence of DNMT3A Mutations in Acute Myeloid Leukemia. *J Clin Oncol*.

- Tiesmeier, J., et al. (2003) Evidence for allelic evolution of C/EBPalpha mutations in acute myeloid leukaemia. *Br J Haematol*, 123, 413-419.
- Toksoz, D., et al. (1989) Ras genes and acute myeloid leukaemia. *Br J Haematol*, 71, 1-6.
- Verhaak, R.G., et al. (2005) Mutations in nucleophosmin (NPM1) in acute myeloid leukemia (AML): association with other gene abnormalities and previously established gene expression signatures and their favorable prognostic significance. *Blood*, 106, 3747-3754.
- Virappane, P., et al. (2008) Mutation of the Wilms' tumor 1 gene is a poor prognostic factor associated with chemotherapy resistance in normal karyotype acute myeloid leukemia: the United Kingdom Medical Research Council Adult Leukaemia Working Party. *J Clin Oncol*, 26, 5429-5435.
- Walz, C., et al. (2006) Activated Jak2 with the V617F point mutation promotes G1/S phase transition. *J Biol Chem*, 281, 18177-18183.
- Wang, J., et al. (2009) The lysine demethylase LSD1 (KDM1) is required for maintenance of global DNA methylation. *Nat Genet*, 41, 125-129.
- Ward, P.S., et al. (2010) The common feature of leukemia-associated IDH1 and IDH2 mutations is a neomorphic enzyme activity converting alpha-ketoglutarate to 2-hydroxyglutarate. *Cancer Cell*, 17, 225-234.
- Whitman, S.P., et al. (2005) The MLL partial tandem duplication: evidence for recessive gain-of-function in acute myeloid leukemia identifies a novel patient subgroup for molecular-targeted therapy. *Blood*, 106, 345-352.
- Whitman, S.P., et al. (2007) Long-term disease-free survivors with cytogenetically normal acute myeloid leukemia and MLL partial tandem duplication: a Cancer and Leukemia Group B study. *Blood*, 109, 5164-5167.
- Whitman, S.P., et al. (2008) FLT3 D835/I836 mutations are associated with poor disease-free survival and a distinct gene-expression signature among younger adults with de novo cytogenetically normal acute myeloid leukemia lacking FLT3 internal tandem duplications. *Blood*, 111, 1552-1559.
- Wittinghofer, A. (1998) Signal transduction via Ras. *Biol Chem*, 379, 933-937.
- Wouters, B.J., et al. (2009) Double CEBPA mutations, but not single CEBPA mutations, define a subgroup of acute myeloid leukemia with a distinctive gene expression profile that is uniquely associated with a favorable outcome. *Blood*, 113, 3088-3091.
- Wysocka, J., et al. (2006) A PHD finger of NURF couples histone H3 lysine 4 trimethylation with chromatin remodelling. *Nature*, 442, 86-90.
- Xu, W., et al. (2011) Oncometabolite 2-hydroxyglutarate is a competitive inhibitor of alpha-ketoglutarate-dependent dioxygenases. *Cancer Cell*, 19, 17-30.
- Yamagata, T., et al. (2005) Runx1/AML1 in normal and abnormal hematopoiesis. *Int J Hematol*, 82, 1-8.
- Yamamoto, Y., et al. (2001) Activating mutation of D835 within the activation loop of FLT3 in human hematologic malignancies. *Blood*, 97, 2434-2439.
- Zhang, D.E., et al. (1997) Absence of granulocyte colony-stimulating factor signaling and neutrophil development in CCAAT enhancer binding protein alpha-deficient mice. *Proc Natl Acad Sci U S A*, 94, 569-574.
- Zhang, W., et al. (2008) Mutant FLT3: a direct target of sorafenib in acute myelogenous leukemia. *J Natl Cancer Inst*, 100, 184-198.
- Zhao, S., et al. (2009) Glioma-derived mutations in IDH1 dominantly inhibit IDH1 catalytic activity and induce HIF-1alpha. *Science*, 324, 261-265.



Myeloid Leukemia - Basic Mechanisms of Leukemogenesis

Edited by Dr Steffen Koschmieder

ISBN 978-953-307-789-5

Hard cover, 484 pages

Publisher InTech

Published online 14, December, 2011

Published in print edition December, 2011

The current book comprises a series of chapters from experts in the field of myeloid cell biology and myeloid leukemia pathogenesis. It is meant to provide reviews about current knowledge in the area of basic science of acute (AML) and chronic myeloid leukemia (CML) as well as original publications covering specific aspects of these important diseases. Covering the specifics of leukemia biology and pathogenesis by authors from different parts of the World, including America, Europe, Africa, and Asia, this book provides a colorful view on research activities in this field around the globe.

How to reference

In order to correctly reference this scholarly work, feel free to copy and paste the following:

Hsin-An Hou, Wen-Chien Chou and Hwei-Fang Tien (2011). Genetic Alterations and Their Clinical Implications in Acute Myeloid Leukemia, *Myeloid Leukemia - Basic Mechanisms of Leukemogenesis*, Dr Steffen Koschmieder (Ed.), ISBN: 978-953-307-789-5, InTech, Available from:
<http://www.intechopen.com/books/myeloid-leukemia-basic-mechanisms-of-leukemogenesis/genetic-alterations-and-their-clinical-implications-in-acute-myeloid-leukemia>

INTECH

open science | open minds

InTech Europe

University Campus STeP Ri
Slavka Krautzeka 83/A
51000 Rijeka, Croatia
Phone: +385 (51) 770 447
Fax: +385 (51) 686 166
www.intechopen.com

InTech China

Unit 405, Office Block, Hotel Equatorial Shanghai
No.65, Yan An Road (West), Shanghai, 200040, China
中国上海市延安西路65号上海国际贵都大饭店办公楼405单元
Phone: +86-21-62489820
Fax: +86-21-62489821

© 2011 The Author(s). Licensee IntechOpen. This is an open access article distributed under the terms of the [Creative Commons Attribution 3.0 License](#), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

IntechOpen

IntechOpen