

Obesity is a risk factor for acute promyelocytic leukemia: evidence from population and cross-sectional studies and correlation with FLT3 mutations and polyunsaturated fatty acid metabolism



Luca Mazzarella,^{1,2} Edoardo Botteri,³ Anthony Matthews,⁴ Elena Gatti,¹ Davide Di Salvatore,⁵ Vincenzo Bagnardi,⁶ Massimo Breccia,⁷ Pau Montesinos,⁸ Teresa Bernal,⁹ Cristina Gil,¹⁰ Timothy J Ley,¹¹ Miguel Sanz,⁸ Krishnan Bhaskaran,⁴ Francesco Lo Coco^{12*} and Pier Giuseppe Pelicci^{1,13}

¹Department of Experimental Oncology, European Institute of Oncology IRCCS, Milan, Italy; ²Division of Early Drug Development for Innovative Therapies, European Institute of Oncology IRCCS, Milan, Italy; ³Cancer Registry of Norway, Oslo, Norway; ⁴London School of Hygiene and Tropical Medicine, London, UK; ⁵IFOM-FIRC Institute of Molecular Oncology, Milan, Italy; ⁶University of Milan-Bicocca, Milan, Italy; ⁷Division of Hematology, Department of Cellular Biotechnologies and Hematology, Sapienza University, Rome, Italy; ⁸Hematology Department, Hospital Universitario y Politécnico La Fe, Valencia, Spain; ⁹Hospital Central de Asturias, Oviedo, Spain; ¹⁰Hospital General, Alicante, Spain; ¹¹Department of Medicine, Division of Oncology, Washington University in St. Louis, Saint Louis, MO, USA; ¹²Department of Hematology, University Tor Vergata, Roma, Italy and ¹³Department of Oncology and Hemato-Oncology, University of Milan

Haematologica 2020
Volume 105(6):1559-1566

ABSTRACT

Obesity correlates with hematologic malignancies including leukemias, but risk of specific leukemia subtypes like acute promyelocytic leukemia and underlying molecular mechanisms are poorly understood. We explored multiple datasets for correlation between leukemia, body mass index (BMI) and molecular features. In a population-based study (n=5.2 million), we correlated BMI with promyelocytic leukemia, and other acute myeloid, lymphoid or other leukemias. In cross-sectional studies, we tested BMI deviation in promyelocytic leukemia trial cohorts from that expected based on national surveys. We explored The Cancer Genome Atlas for transcriptional signatures and mutations enriched in promyelocytic leukemia and/or obesity, and confirmed a correlation between body mass and FLT3 mutations in promyelocytic leukemia cohorts by logistic regression. In the population-based study, hazard ratio per 5 kg/m² increase was: promyelocytic leukemia 1.44 (95% CI: 1.0-2.08), non-promyelocytic acute myeloid leukemias 1.17 (95% CI: 1.10-1.26), lymphoid leukemias 1.04 (95% CI: 1.0-1.09), other 1.10 (95% CI: 1.04-1.15). In cross-sectional studies, body mass deviated significantly from that expected (Italy: $P < 0.001$; Spain: $P = 0.011$; USA: $P < 0.001$). Promyelocytic leukemia showed upregulation of polyunsaturated fatty acid metabolism genes. Odds of FLT3 mutations were higher in obese acute myeloid leukemias (odds ratio=2.4, $P = 0.007$), whether promyelocytic or not, a correlation confirmed in the pooled promyelocytic leukemia cohorts (OR=1.22, 1.05-1.43 per 5 kg/m²). These results strengthen the evidence for obesity as a *bona fide* risk factor for myeloid leukemias, and in particular APL. FLT3 mutations and polyunsaturated fatty acid metabolism may play a previously underappreciated role in obesity-associated leukemogenesis.

Introduction

The etiology of acute myeloid leukemia (AML) remains poorly understood. Genetic predisposition or clear exposure to environmental mutagenic agents (smoking, benzene, radiation, prior chemotherapy) can be demonstrated only in a minority of cases.¹ Age is an independent risk factor, probably linked to the pro-

This paper is dedicated to the memory of our wonderful colleague, Prof. Francesco Lo Coco, who recently passed away

Correspondence:

LUCA MAZZARELLA
luca.mazzarella@ieo.it

PIER GIUSEPPE PELICCI
piergiuseppe.pelicci@ieo.it

Received: April 5, 2019.

Accepted: September 11, 2019.

Pre-published: September 12, 2019.

doi:10.3324/haematol.2019.223925

Check the online version for the most updated information on this article, online supplements, and information on authorship & disclosures: www.haematologica.org/content/106/6/1559

©2020 Ferrata Storti Foundation

Material published in *Haematologica* is covered by copyright. All rights are reserved to the Ferrata Storti Foundation. Use of published material is allowed under the following terms and conditions:

<https://creativecommons.org/licenses/by-nc/4.0/legalcode>. Copies of published material are allowed for personal or internal use. Sharing published material for non-commercial purposes is subject to the following conditions: <https://creativecommons.org/licenses/by-nc/4.0/legalcode>, sect. 3. Reproducing and sharing published material for commercial purposes is not allowed without permission in writing from the publisher.



gressive mutation accumulation and clonal stem cell expansion that accompanies aging.² Although obesity has recently emerged as a prominent risk factor for a variety of solid tumors,³ its impact on hematologic neoplasms has received less attention. A moderate but consistently positive correlation between body mass index (BMI) and incidence of leukemias has been identified in observational studies,⁴⁻⁶ yet none of the collected evidence has been considered sufficiently strong to consider obesity as a *bona fide* risk factor for AML.^{3,7} Most studies did not distinguish between myeloid/lymphoid and acute/chronic forms, nor between genetic subtypes within each form. AML is recognized as a highly heterogeneous disease with genetically diverse subtypes.⁸ Subtypes have radically different outcomes and, similarly, their risk may be differentially affected by environmental factors. Identification of subtype-specific risk associations, however, is made difficult by their rarity.

A genetic subset of AML, acute promyelocytic leukemia (APL), is characterized by a specific chromosomal translocation (t15;17), homogeneous biology and response to clinical agents all-trans retinoic acid (ATRA) and arsenic trioxide, which have made it the most curable form of AML to date.⁹ We previously demonstrated that the risk of relapse after ATRA/idarubicin is significantly increased in overweight/obese APL patients.¹⁰ In the present report, we investigated the association of overweight/obesity with the risk of developing APL and other leukemias. We describe the results of multiple studies across four western populations with significantly different dietary regimens and prevalence of obesity. All the studies demonstrated increased risk of developing APL in overweight/obesity subjects. In an effort to generate mechanistic hypotheses to explain this relationship, we analyzed transcriptomic and mutational data from the AML project in The Cancer Genome Atlas (TCGA)¹¹ and identified alterations selectively associated with obesity and/or APL which may be involved in obesity-associated leukemogenesis.

Methods

UK population-based study: data collection and statistical methods

Full details of the methods for the UK population study were described previously.⁶ The study was approved by the London School of Hygiene and Tropical Medicine Ethics Committee. To identify outcomes of specific leukemia sub-types, Clinical Practice Research Datalink (CPRD) clinical records were searched for codes relating to specific leukemia subgroups. We controlled for multiple co-variables at time of the BMI record(s): age, smoking status, alcohol use, previous diabetes diagnosis, index of multiple deprivation, calendar period, and stratified by gender. We excluded people with missing smoking [49,206 of 5.24 million (0.9%)] and alcohol [394,196 of 5.24 million (7.5%)] status. Confidence intervals (CI) in Figure 1 are presented at the 99% level; all other CIs are presented at the 95% level.

Cross-sectional studies: data collection and statistical methods

Acute promyelocytic leukemia cases from Spain were extracted from the PETHEMA database to include 414 cases diagnosed between 1998 and 2012. APL cases from Italy, where 134 adult patients were treated under the AIDA protocol, were included in the previously described cohort.¹⁰ APL cases from the USA includ-

ed the entire cohort of the published AML The Cancer Genome Atlas (TCGA) project¹¹ (n=20) plus 22 additional APL cases, unselected for any clinical variable, diagnosed at Washington University, St. Louis, MO, USA (Expanded TCGA cohort). For all case cohorts, BMI was measured at the time of diagnosis.

Data collection was approved by the Research Ethics Board of each participating institution, as referenced.¹¹⁻¹⁴ Data sources for expected BMI in the local population are described in the *Online Supplementary Appendix*.

We compared the distribution of BMI observed in the three APL case cohorts to the distribution of BMI expected in the general population of the same countries. Specifically, to calculate the expected distribution of BMI in Italy, we used data from the Italian National Institute of Statistics,¹⁴ and we selected the area of Lazio, where the APL cases were diagnosed, in the years 2000-2010. For Spain, we used data from the Eurostat,¹⁵ and we selected the general population of Spain in the year 2008, the only year available. For both Italy and Spain, the expected BMI distribution was calculated using the available age- and gender-specific BMI distribution of the general population classified into three categories (<25, 25-29.9, ≥30). For the USA, we used the 2009-2010 data from the American National Health and Nutrition Examination Survey.¹⁶ The expected BMI distribution was calculated using the available race-, age-, and gender-specific BMI distribution of the general population classified into four categories (<25, 25-29.9, 30.0-34.9, ≥35).

The global null hypothesis that the observed counts did not differ from the expected ones across the BMI categories was tested in a null Poisson regression model, where the observed counts were considered as dependent variable and the expected counts as the offset. BMI was included in the model as an ordinal variable to test the log-linear relationship between BMI and the observed to expected ratio (i.e. to test for linear trend). Pearson's χ^2 goodness of fit test *P*-value was reported.

Expression data analysis

Expression data (RPKM matrix) were downloaded from the AML TCGA data portal. Cases with available RNAseq, BMI and French-American-British (FAB) classification data (177 of 200) were used in the present study. Cases were classified by FAB in "APL" (FAB="M3") and "non-APL" (FAB ≠ "M3"), and by BMI in "obese" (BMI ≥ 30) and "non-obese" (BMI < 30). Genes with < 0.2 reads per kilo base per million mapped reads (RPKM) in at least 75% of patients were removed.¹¹ The Quantitative Set Analysis for Gene Expression method, as implemented in the quSAGE package¹⁵ in the R programming language (v3.2.3), was used to conduct supervised gene set enrichment analysis. For each expressed gene, the quSAGE algorithm calculates a probability density function (PDF) of differential expression between two groups of samples. For each gene set, it then calculates "activity", i.e. the mean difference in log-expression of individual genes included in a gene set. Gene sets with False Discovery Rate (FDR) < 0.05 were considered significant. We focused on the Kyoto Encyclopedia of Genes and Genomes (KEGG) and chemical and genetic perturbations (CGP) gene set collections, downloaded from MSigDB (<http://software.broadinstitute.org/gsea/msigdb/>). The CGP collection was used to confirm enrichment of previously identified APL-specific gene signatures¹⁶ (*Online Supplementary Table S1*). We focused on the KEGG collection, as it is enriched for metabolism-associated gene annotations.¹⁶ The script to generate the present results is available on request.

Mutational data analysis

For the analysis of the TCGA data, mutational data were retrieved from the TCGA AML paper,¹¹ and AML driver genes (restricted to those with at least 2 mutations in the dataset) were

down-loaded from IntOGen.¹⁷ For each gene, different mutations were conflated so that gene status in each patient was either "mutated" or "wild-type". For each gene, we then calculated the number of mutated or wild-type patients in the obese or non-obese groups, and calculated odds ratios (OR), 95% confidence intervals (CI), and *P*-values by Fisher's test with Benjamini-Hochberg correction. Only genes with >1 mutation in the dataset were considered, using the *fdsm* package in R.

For the analysis of the retrospective cohort, FLT3 Internal Tandem Duplication (ITD) mutational data were provided by the referring centers. Logistic regression was used to calculate OR with 95% CI.

Further details of the methods used are provided in the *Online Supplementary Appendix*.

Results

Population-based cohort study in the UK

Overall characteristics of the 5.24 million UK adults included in this study have been described previously.⁶ A total of 5,833 subjects with a diagnosis of "leukemia" over the observational time were included in the present analy-

sis. These events were further classified in the following groups: "APL" (n=26), "non APL-AML" (n=1,012), lymphoid leukemias ("LL"; n=2,823), and "other" (n=1,972). Median time lapse between BMI measurement and diagnosis was similar across subgroups (APL: 1,810 days; AML: 2,280 days; LL: 1,928 days; other: 1,894 days).

We fit BMI as a three-knot cubic spline and as a linear term. There was no evidence of non-linearity (*P*=0.94), suggesting that the relationship was best described by the linear model. After adjusting for co-variates, per each 5 kg/m² increase we obtained hazard ratios (HR) of 1.44 for

Table 1. Hazard ratios from the UK population study.

Disease	N. of events	Adjusted HR (per 5 kg/m ² increase in BMI)	95% CI
APL	26	1.44	1.00, 2.08
AML	1,012	1.17	1.10, 1.26
LL	2,823	1.04	1.00, 1.09
Other	1,972	1.10	1.04, 1.15

N: number; HR: hazard ratio; BMI: body mass index; CI: confidence interval; APL: acute promyelocytic leukemia; AML: acute myeloid leukemia; LL: lymphoid leukemias.

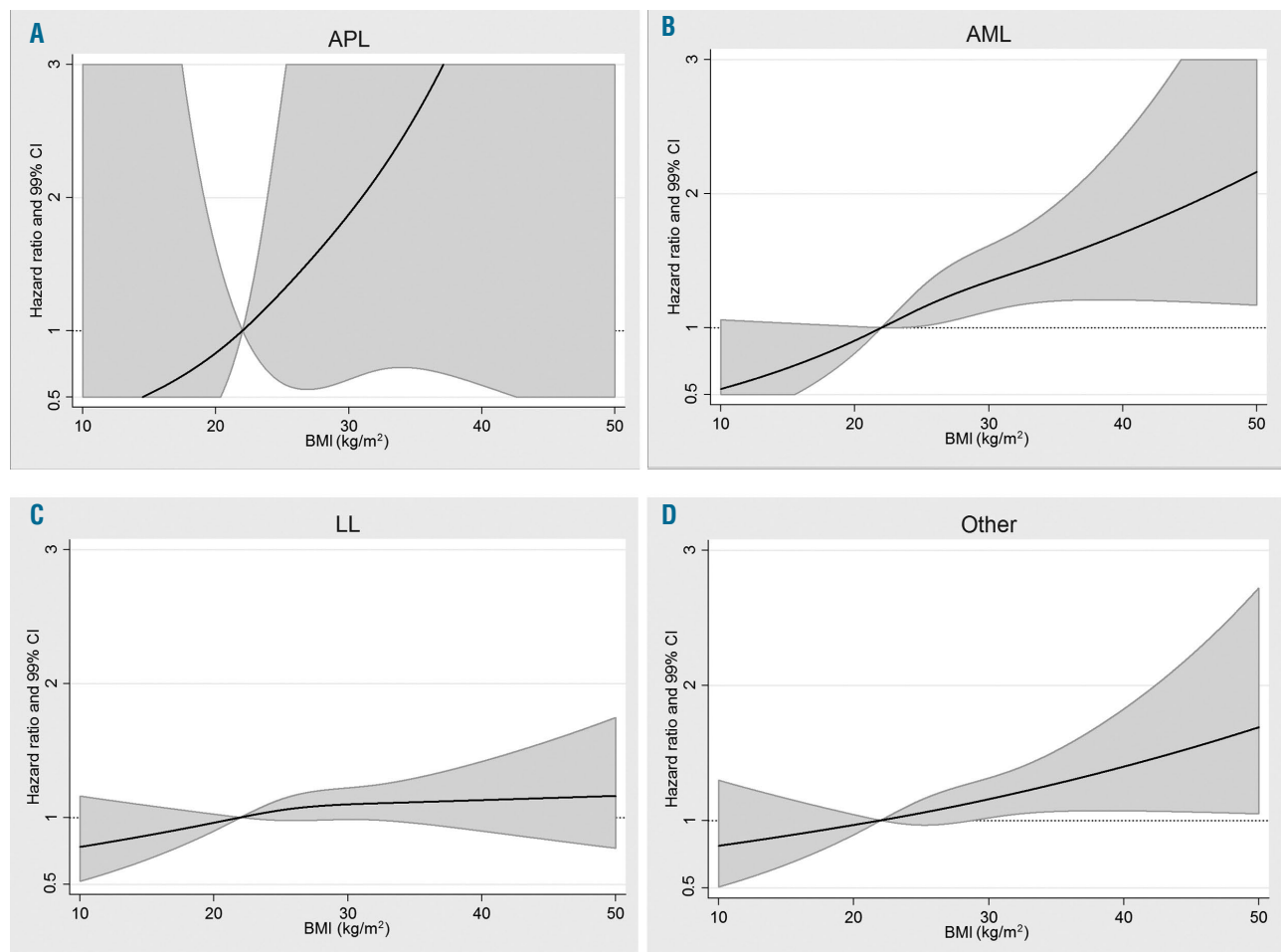


Figure 1. Relationship between body mass index (BMI) and log-hazard ratio (HR) for leukemias in the UK population. (A) Acute promyelocytic leukemia (APL); (B) other acute myeloid leukemia (AML); (C) lymphoid leukemias; (D) all leukemias. Plots show mean (dark line) \pm 95% confidence intervals (shaded area).

APL (95%CI: 1.0-2.08), 1.17 for non APL-AML (95%CI: 1.10-1.26), 1.04 for LL (95%CI: 1.0-1.09), and 1.10 for other leukemias (95%CI: 1.04-1.15) (Table 1 and Figure 1). Stratification by gender suggested a stronger effect for male gender in APL (HR 1.82, 95%CI: 1.10-3.00 vs. female HR 1.19, 95%CI: 0.67-1.98), although the sample size becomes very small (n=13 each). Together, these results suggest that higher BMI is associated with increased risk of all sub-types of leukemia, particularly APL.

Cross-sectional studies in Italian, Spanish and US trial cohorts

Though APL showed the strongest association with higher BMI in the cohort analysis described above, results were not conclusive due to the small number of cases identified (n=26) and the consequently wide confidence intervals. To strengthen the evidence, we carried out retrospective case-control studies using cohorts of APL

patients from national registries of clinical trials from Spain (PETHEMA) and Italy (GIMEMA), and patients from the US-based AML genome sequencing study (the AML TCGA cohort with 22 additional cases characterized at Washington University, St Louis, MO). In all three groups, APL diagnosis was established using gold standard diagnostic procedures.

Demographic characteristics of the three case cohorts (Italy n=134, Spain n=414 and USA n=42) are described in Table 2. Gender (female 53.0%, 55.2%, 50%, respectively) and age (median of 45, 45, 47 years, respectively) were similarly represented. Information on ethnicity was unavailable for the Spanish and Italian cohorts, whereas white, black and hispanic ethnicities were represented in the US cohort.

To generate control groups for comparison, we obtained anthropometric data from epidemiological surveys of the general population in the different countries. As the prevalence of obesity has increased dramatically in most coun-

Table 2. Description of the cross-sectional cohorts.

		Italy n=134	Spain n=414	USA n=42
Age	18 - 35	46 (34.3%)	113 (27.3%)	13 (31.0%)
	36 - 50	34 (25.4%)	145 (35.0%)	12 (28.6%)
	51 - 65	40 (29.9%)	102 (24.6%)	11 (26.2%)
	> 65	14 (10.4%)	54 (13.0%)	6 (14.3%)
	Median (IQR)	45 (31-57)	45 (34-57)	47 (33-60)
Gender	Male	63 (47.0%)	227 (54.8%)	21 (50.0%)
	Female	71 (53.0%)	187 (45.2%)	21 (50.0%)
Year of diagnosis	Median (range)	2002 (1997-2010)	2003 (1996-2012)	2007 (2001-2011)
Race	White	-	-	36 (85.7)
	Black	-	-	5 (11.9)
	Hispanic	-	-	1 (2.4)
BMI	Median (IQR)	26 (23-28)	26 (23-29)	34 (28-39)

IQR: interquartile range; BMI: body mass index; n: number.

Table 3. Observed body mass index (BMI) distribution in acute promyelocytic leukemia (APL) cases and expected BMI distribution in general population (percentages in brackets).

Italy	BMI	Obs	All Exp ^a	P	Obs	Males Exp ^b	P	Obs	Females Exp ^d	P
	<25.0	48 (35.8%)	77.8 (58.0%)	<0.001	16 (25.4%)	29.0 (46.0%)	<0.001	32 (45.1%)	48.8 (68.7%)	<0.001
	25.0-29.9	71 (53.0%)	44.8 (33.4%)		42 (66.7%)	28.0 (44.5%)		29 (40.8%)	16.8 (23.6%)	
	≥30.0	15 (11.2%)	11.4 (8.5%)		5 (7.9%)	6.0 (9.5%)		10 (14.1%)	5.5 (7.7%)	
	Total	134	134		63	63		71	71	
Spain	BMI	Obs	Exp ^a	P	Obs	Exp ^b	P	Obs	Exp ^d	P
	<25.0	172 (41.5%)	189.9 (45.9%)	0.011	79 (34.8%)	85.0 (37.4%)	0.130	93 (49.7%)	104.9 (56.1%)	0.033
	25.0-29.9	156 (37.7%)	158.1 (38.2%)		99 (43.6%)	103.2 (45.5%)		57 (30.5%)	55.9 (29.4%)	
	≥30.0	86 (20.8%)	66.0 (15.9%)		49 (21.6%)	38.8 (17.1%)		37 (19.8%)	27.2 (14.6%)	
	Total	414	414		227	227		187	187	
USA	BMI	Obs	Exp ^a	P	Obs	Exp ^b	P	Obs	Exp ^d	P
	<25.0	2 (4.8%)	12.8 (30.6%)	<0.001	1 (4.8%)	5.3 (25.4%)	0.002	1 (4.8%)	7.5 (35.7%)	0.003
	25.0-29.9	13 (31.0%)	13.7 (32.6%)		5 (23.8%)	7.9 (37.7%)		8 (38.1%)	5.8 (27.6%)	
	30.0-34.9	12 (28.6%)	8.6 (20.5%)		9 (42.9%)	4.8 (23.0%)		3 (14.3%)	3.8 (18.0%)	
	≥35.0	15 (35.7%)	6.9 (16.3%)		6 (28.6%)	2.9 (13.9%)		9 (42.9%)	3.9 (18.8%)	
	Total	42	42		21	21		21	21	

Expected frequencies were obtained from the body mass index (BMI) distribution in the general population of the area of the acute promyelocytic leukemia (APL) cases, period of APL diagnosis and in addition: ^aage class and gender; ^bage class, gender and race; ^cage class; ^dage class and race. Obs: observed.

tries in the last decades (especially in the USA), we obtained data that were as close as possible to the median year of diagnosis (2002 for Italy, 2003 for Spain, 2007 for the USA) (Table 3; see also Methods section).

In all three cohorts, there was strong evidence that the observed BMI distribution for cases across World Health Organization (WHO) BMI classes was different from that expected under the null hypothesis of no association (Italy: $P < 0.001$; Spain: $P = 0.011$; USA: $P < 0.001$) (Table 3) in gender-, age-, and ethnicity- (for USA) matched controls. In particular, in all three datasets, there were more cases than expected in the higher BMI groups, irrespective of gender in all cohorts apart from Spain, in which significance was not reached for males ($P = 0.130$) despite a similar trend (Table 3).

Correlation of TCGA transcriptomics data with body mass index and acute myeloid leukemia subtype

The availability of the TCGA dataset prompted us to search for signatures that could suggest a mechanistic rationale for the association between APL and obesity. We interrogated available AML transcriptomes with supervised gene set enrichment analysis using quSage.¹⁵ Focusing on the KEGG gene set collection, APL was associated with increased activity of 13 and decreased activity of 64 out of 186 gene sets (Table 4, Figure 2A and *Online Supplementary Table S2*). Intriguingly, among significantly up-regulated gene sets, we found pathways associated with the metabolism of long-chain unsaturated fatty acids (linoleic and arachidonic), which are precursors of eicosanoids mediating inflammation-associated cancers.¹⁸

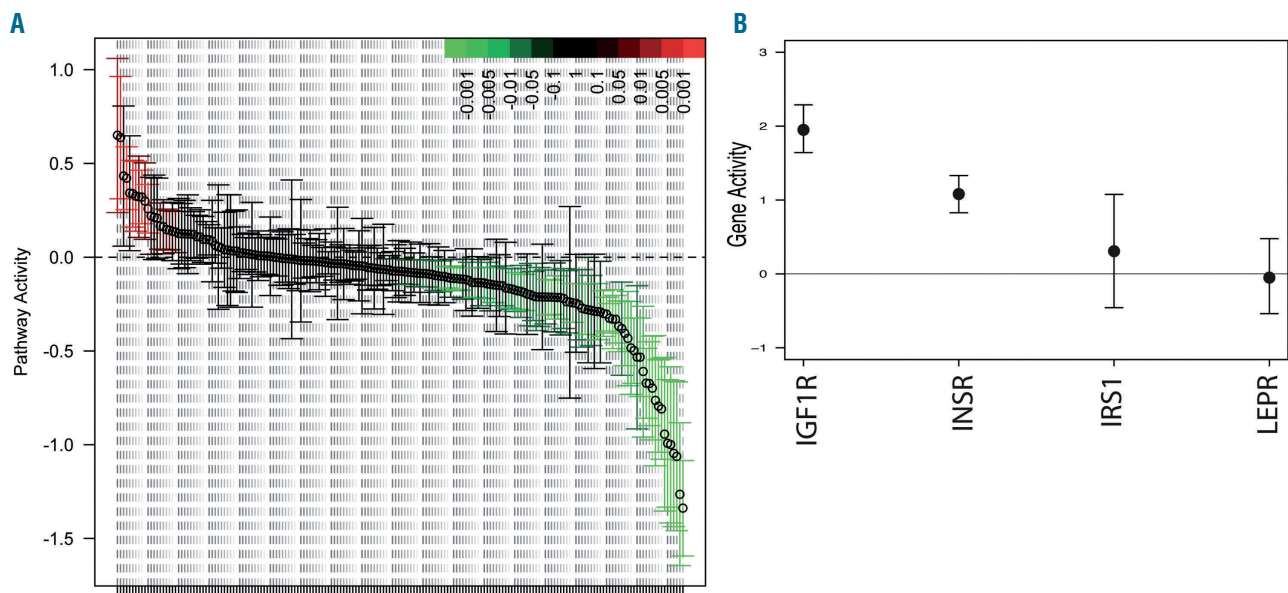


Figure 2. Differential activities of Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways and insulin/leptin receptors in the M3 versus non-M3 quSage comparison in The Cancer Genome Atlas (TCGA). (A) Activity score with 95% Confidence Intervals (CI) of 186 KEGG gene sets; significant gene sets are color-coded in red (if up-regulated) or green (if down-regulated). (B) Insulin/IGF1 receptor pathway and leptin receptors. Mean \pm 95% confidence interval are plotted.

Table 4. Significantly up-regulated Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways in acute promyelocytic leukemia versus acute myeloid leukemia in The Cancer Genome Atlas (TCGA).

pathway.name	log.fold.change	P	FDR
KEGG_RENIN_ANGIOTENSIN_SYSTEM	0.6503	0.0023	0.0093
KEGG_LINOLEIC_ACID_METABOLISM	0.6381	0.0002	0.0010
KEGG_GLYCOSAMINOGLYCAN_BIOSYNTHESIS_HEPARAN_SULFATE	0.4217	0.0000	0.0000
KEGG_GLYCOSPHINGOLIPID_BIOSYNTHESIS_LACTO_AND_NEOLACTO_SERIES	0.3391	0.0003	0.0017
KEGG_ALANINE_ASPARTATE_AND_Glutamate_METABOLISM	0.3258	0.0009	0.0039
KEGG_ARACHIDONIC_ACID_METABOLISM	0.3221	0.0037	0.0130
KEGG_GLYCOSAMINOGLYCAN_DEGRADATION	0.3208	0.0000	0.0001
KEGG_HISTIDINE_METABOLISM	0.2996	0.0044	0.0148
KEGG_ARGININE_AND_PROLINE_METABOLISM	0.2582	0.0001	0.0008
KEGG_LIMONENE_AND_PINENE_DEGRADATION	0.1662	0.0087	0.0250
KEGG_CARDIAC_MUSCLE_CONTRACTION	0.1475	0.0084	0.0245
KEGG_PROTEIN_EXPORT	0.1439	0.0066	0.0204
KEGG_PATHWAYS_IN_CANCER	0.1346	0.0169	0.0428

Also noticeable was the APL-associated upregulation of insulin and insulin-like growth factor (IGF1) receptors, but not leptin receptor (Figure 2B); insulin signaling-associated pathways were also specifically up-regulated in obese *versus* non-obese APL patients ("type II diabetes mellitus" and "insulin signaling") (Online Supplementary Table S2).

No pathway was significantly enriched in obese *versus* non-obese patients among non-M3 cases.

Correlation of mutational data with body mass index

We then asked whether obesity is associated with specific driver mutations in AML in the TCGA cohort. Out of 23 established driver genes mutated at least twice in the cohort, mutations in FLT3 were positively associated with obesity (33 of 88 obese *vs.* 22 of 110 non-obese; OR=2.4, FDR=0.16, $P=0.007$) (Figure 3 and Online Supplementary Table S3). The correlation remained statistically significant both in non-APL AML (27 of 49 obese *vs.* 22 of 102 non-obese; OR=2, $P=0.04$), and in APL (6 of 12 obese *vs.* 0 of 8 non-obese; $P=0.04$). When we analyzed the two main classes of FLT3 mutations separately [tyrosine kinase domain (TKD) and internal tandem duplication (ITD)], the association held statistically significant for ITD (24 of 88 obese *vs.* 14 of 110 non-obese; OR=2.6, $P=0.01$) but not for TKD (9 of 88 obese *vs.* 8 of 110 non-obese; $P=0.6$). In APL, where all FLT3 mutations were ITD, the correlation remained statistically significant (6 of 12 obese *vs.* 0 of 8 non-obese; $P=0.04$). In non-APL AML, with 32 ITD and 17 TKD, overall FLT3 mutations were still significantly enriched in obese patients (27 of 49 obese *vs.* 22 of 102 non-obese; OR=2, $P=0.04$) but not when analyzed separately ($P=0.11$ for ITD and 0.44 for TKD).

We then attempted to validate this finding in the APL cohorts, for which data on the most representative FLT3ITD mutation (ITD) were available (Table 5). In the pooled analysis (163 mutated patients, of a total 569), OR of having a FLT3 ITD was 1.22 (95%CI: 1.05-1.43) per each 5 kg/m² increase. In the individual cohorts, results were significant in the Italian (30 of 114 mutated, OR=2.35, 95%CI: 1.25-4.42) and US (14 of 41 mutated, OR=1.44, 95%CI: 0.93-2.24) cohorts, but not in the Spanish (119 of 414 mutated, OR=1.09, 95%CI: 0.89-1.33).

Discussion

Here we provide substantial evidence for an association between elevated BMI and risk of developing AML. The risk was particularly high with the APL subtype, with an estimated 44% HR increase per each 5 kg/m². This was qualitatively confirmed by comparing expected *versus* observed BMI distributions in APL cohorts across three western countries (the USA, Spain and Italy) with different

obesity prevalence and dietary habits. In addition, we provide hypothesis-generating evidence for molecular mechanisms underlying such an association, in particular, the possible involvement of pro-inflammatory fatty acid metabolism and mutations of the tyrosin kinase FLT3.

Our epidemiological results expand a growing body of literature identifying overweight/obesity as a *bona fide* risk factor for leukemias. The most recent meta-analysis reported an adjusted relative risk of 1.14 (95%CI: 1.04-1.26; $P=0.008$) for AML overall.⁵ Despite the growing evidence, the notion of obesity as a risk factor for leukemia remains widely overlooked.⁷ Among the highly heterogeneous AML subtypes, APL is the most clinically and biologically coherent. We and others previously showed that in APL, but not in other AML, an elevated BMI significantly affects outcome.^{10,19} This is also in line with the few retrospective studies that have assessed APL as a separate disease entity.^{20,21} No study had addressed this question prospectively, a task made difficult by the rarity of the disease, but made possible in our case by the very large study

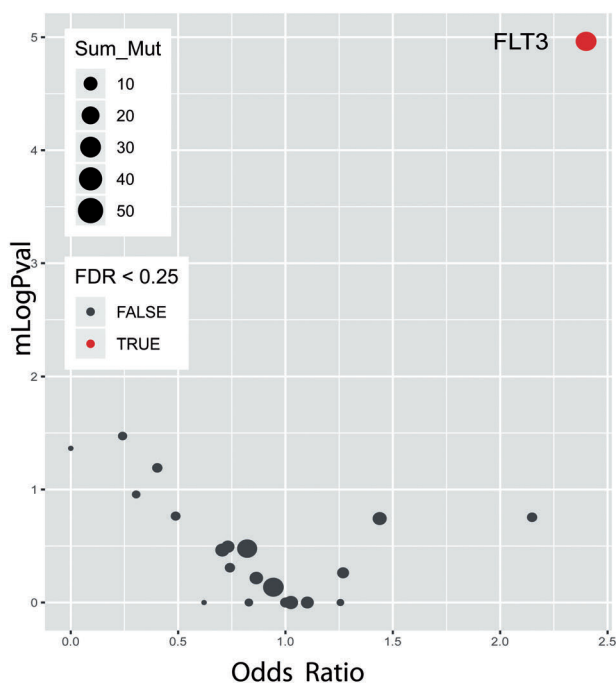


Figure 3. Association between obesity and FLT3 mutations. (A) Bubble plot representing Odds Ratio versus $-\log P$ value (mLogPval) of any mutation in 23 driver genes in The Cancer Genome Atlas (TCGA) acute myeloid leukemia cohort. FLT3 (in red) is the only gene with False Discovery Rate (FDR) < 0.25. Bubble size reflects the number of obese patients with a mutation. Data are tabulated in Online Supplementary Table S3.

Table 5. Logistic regression of body mass index (BMI) and FLT3 ITD mutations.

BMI	All 3 cohorts 163/569 ^a OR (95% CI)	ITALY 30/114 ^a OR (95% CI)	SPAIN 119/414 ^a OR (95% CI)	USA 14/41 ^a OR (95% CI)
5 unit increase	1.22 (1.05-1.43)	2.35 (1.25-4.42)	1.09 (0.89-1.33)	1.44 (0.93-2.24)
≥ 25 <i>vs.</i> <25 ^b	-	4.40 (1.63-11.9)	1.15 (0.75-1.78)	-
≥ 30 <i>vs.</i> <30 ^b	-	-	-	6.46 (1.21-34.5)

^aMutations / All patients. ^bGiven the small number of obese patients in Italy and Spain, we compared overweight/obese patients *versus* normal weight patients (i.e. BMI ≥ 25 *vs.* <25). Given the small number of normal weight patients in the USA, we compared obese patients *versus* non-obese patients (i.e. BMI ≥ 30 *vs.* <30). OR: odds ratios; CI: confidence intervals.

population (5.2 million). The largest prospective study to date (EPIC), which revealed a statistically significant higher risk only in female AML, but not in other gender and biological subgroups,⁴ was based on a relatively small number of incident cases: only 671 out of 375,021 participants over 11.5 years of median follow up. The use of orthogonal epidemiological approaches is a strength of the study, as it attempts to mitigate some weaknesses of each design. Registry-based studies have little patient selection bias, providing results that are more comparable to real-life scenarios. However, the quality of case identification is likely to be sub-optimal; erroneous assignment of APL to the general AML ICD code might "deplete" incident cases and further reduce statistical power. Case-control studies in the context of clinical trials, on the other hand, offer the advantage of gold standard diagnosis but might be affected by significant patient selection biases. This may have counter-selected obese patients in the present study, since the correlated comorbidities may be associated with limited access to clinical trials.

Another limitation of the study is that we could not provide the same degree of geographical homogeneity for control subjects in the case-control studies. This may be particularly relevant for the USA, known to have wide state-specific differences in BMI distribution. However, this variation is mainly due to demographic parameters,²² such as age, gender and race, and is, therefore, at least partly accounted for in our multivariate analysis. We also note that our US APL cohort includes a single patient of hispanic ethnicity. Hispanics are considered at higher incidence of APL, although some large studies based on Surveillance, Epidemiology, and End Results (SEER) data dispute this commonly held conclusion.²³

Understanding the molecular mechanism causing increased cancer risk in obese subjects is crucial for adequate nutritional management in disease prevention, given the sustained rise of obesity worldwide, particularly in emerging economies. The possibility of matching transcriptional and mutational profiles from TCGA to patient clinical and BMI data provided an opportunity to generate hypotheses grounded on actual data. However, extracting biological significance from large molecular datasets remains challenging. Shifting the analytical focus from single genes to gene sets or pathways may allow signals to be captured even when the changes affecting individual genes are minimal, provided they are coherent. The gene set-based method we used here for transcriptional analysis does not assume equal variances, resulting in improved sensitivity and specificity over similar competing methods.¹⁵ Our main finding is the upregulation of several genes involved in the metabolism of pro-inflammatory ω -6 polyunsaturated fatty acids (PUFA, linoleic and arachidonic) in APL. These molecules are increased in the plasma of metabolically impaired subjects, including the obese,²⁴

and may lead to elevated production of derivative molecules with multiple effects in signaling and inflammation, enhancing leukemogenesis through several independent mechanisms: direct growth promotion, generation of genotoxic oxidative stress, immune modulation,^{18,25} and generation of endogenous agonists for Peroxisome proliferator-activated receptors (PPAR).²⁶ PPAR are known insulin sensitizers²⁷ and their transcriptional targets are up-regulated in APL (*Online Supplementary Table S2*); APL expressed higher levels of insulin and IGF1 receptors, and its growth may thus be favored by the increased insulin/IGF1 levels in obese subjects.^{3,28} Elevated generation of PUFA-derived eicosanoids by APL cells may also explain the association between obesity and ATRA differentiation syndrome (DS),¹⁰ as eicosanoids strongly promote leukocyte adhesion and chemokine release in the lungs.²⁹

Finally, the association between FLT3 mutations and a higher BMI, although unconfirmed in the larger Spanish cohort, is an intriguing finding that we think deserves additional research. FLT3 mutations are associated with specific metabolic dependencies which may be differentially affected according to the systemic nutritional status.³⁰ It cannot be entirely ruled out that geographical differences in dietary composition may account for the discrepancies in the association between BMI and APL risk (weakest in Spain) and FLT3 mutations (null in Spain). Consistent with this highly speculative view, a recent EPIC substudy revealed marked differences in nutritional patterns between European nations. Despite sharing a theoretical propensity for "Mediterranean" diets, Italy and Spain were highly polarized, especially in terms of average polyunsaturated fatty acid consumption (3% vs. 38% of the participants in the highest quintile, respectively).³¹ More mechanistic studies are needed to clarify whether FLT3 mutations are favored by specific nutritional components.

In conclusion, based on evidence provided here, we propose including obesity among environmental factors increasing risk for myeloid neoplasms and in particular APL. Additional studies with experimental models will clarify the molecular determinants of this relationship, and test whether and how specific nutritional components like PUFA can determine specific mutational and transcriptional alterations able to influence the natural history of the disease.

Acknowledgments

AIRC-Cariplo Foundation, Italian Ministry of Health, European Hematology Association, Wellcome Trust and Royal Society

Role of the funding source

Funding sources had no role in study design, collection, analysis, and interpretation of data; report writing nor decision to submit the paper for publication.

References

- Mazzarella L, Riva L, Luzi L, Ronchini C, Pelicci PG. The genomic and epigenomic landscapes of AML. *Semin Hematol*. 2014;51(4):259-272.
- Shlush LI, Zandi S, Itzkovitz S, Schuh AC. Aging, clonal hematopoiesis and preleukemia: not just bad luck? *Int J Hematol*. 2015;102(5):513-522.
- Lauby-Secretan B, Ph D, Scoccianti C, Ph D, Loomis D, Ph D. Body Fatness and Cancer — Viewpoint of the IARC Working Group. *N Engl J Med*. 2016;375(8):794-798.
- Saber Hosnijeh F, Romieu I, Gallo V, et al. Anthropometric characteristics and risk of lymphoid and myeloid leukemia in the European Prospective Investigation into Cancer and Nutrition (EPIC). *Cancer Causes Control*. 2013;24(3):427-438.
- Li S, Chen L, Jin W, et al. Influence of body mass index on incidence and prognosis of acute myeloid leukemia and acute promyelocytic leukemia: A meta-analysis. *Sci Rep*. 2017;7(1):17998.
- Bhaskaran K, Douglas I, Forbes H, dos-

- Santos-Silva I, Leon DA, Smeeth L. Body-mass index and risk of 22 specific cancers: a population-based cohort study of 5.24 million UK adults. *Lancet*. 2014;384(9945):755-765.
7. American Cancer Society. Leukemia–Acute Myeloid (Myelogenous). <http://www.cancer.org/cancer/leukemia-acutemyeloidaml/detailedguide/leukemia-acute-myeloid-myelogenous-risk-factors>.
 8. Komanduri K V, Levine RL. Diagnosis and Therapy of Acute Myeloid Leukemia in the Era of Molecular Risk Stratification. *Annu Rev Med*. 2016;67:59-72.
 9. Coombs CC, Tavakkoli M, Tallman MS. Acute promyelocytic leukemia: where did we start, where are we now, and the future. *Blood Cancer J*. 2015;5:e304.
 10. Breccia M, Mazzarella L, Bagnardi V, et al. Increased BMI correlates with higher risk of disease relapse and differentiation syndrome in patients with acute promyelocytic leukemia treated with the AIDA protocols. *Blood*. 2012;119(1):49-54.
 11. Ley TJ, Miller C, Ding L, et al. Genomic and epigenomic landscapes of adult de novo acute myeloid leukemia. *N Engl J Med*. 2013;368(22):2059-2074.
 12. Avvisati G, Lo-Coco F, Paoloni FP, et al. AIDA 0493 protocol for newly diagnosed acute promyelocytic leukemia: very long-term results and role of maintenance. *Blood*. 2011;117(18):4716-4725.
 13. Lo-Coco F, Avvisati G, Vignetti M, et al. Front-line treatment of acute promyelocytic leukemia with AIDA induction followed by risk-adapted consolidation for adults younger than 61 years: results of the AIDA-2000 trial of the GIMEMA Group. *Blood*. 2010;116(17):3171-3179.
 14. Adès L, Sanz MA, Chevret S, et al. Treatment of newly diagnosed acute promyelocytic leukemia (APL): a comparison of French-Belgian-Swiss and PETHEMA results. *Blood*. 2008;111(3):1078-1084.
 15. Yaari G, Bolen CR, Thakar J, Kleinstein SH. Quantitative set analysis for gene expression: A method to quantify gene set differential expression including gene-gene correlations. *Nucleic Acids Res*. 2013;41(18):1-11.
 16. Morishima K, Tanabe M, Furumichi M, Kanehisa M, Sato Y. KEGG: new perspectives on genomes, pathways, diseases and drugs. *Nucleic Acids Res*. 2016;45(D1):D353-D361.
 17. Gonzalez-Perez A, Perez-Llamas C, Deu-Pons J, et al. IntOGen-mutations identifies cancer drivers across tumor types. *Nat Methods*. 2013;10(11):1081-1082.
 18. Wang D, Dubois RN. Eicosanoids and cancer. *Nat Rev Cancer*. 2010;10(3):181-193.
 19. Castillo JJ, Mulkey F, Geyer S, et al. Relationship between obesity and clinical outcome in adults with acute myeloid leukemia: A pooled analysis from four CALGB (alliance) clinical trials. *Am J Hematol*. 2016;91(2):199-204.
 20. Wong O, Harris F, Yiyang W, Hua F. A hospital-based case-control study of acute myeloid leukemia in Shanghai: Analysis of personal characteristics, lifestyle and environmental risk factors by subtypes of the WHO classification. *Regul Toxicol Pharmacol*. 2009;55(3):340-352.
 21. Estey E, Thall P, Kantarjian H, Pierce S, Kornblau S, Keating M. Association between increased body mass index and a diagnosis of acute promyelocytic leukemia in patients with acute myeloid leukemia. *Leukemia*. 1997;11(10):1661-1664.
 22. Flegal KM, Carroll MD, Kit BK, Ogden CL. Prevalence of obesity and trends in the distribution of body mass index among US adults, 1999-2010. *JAMA*. 2012;307(5):491-497.
 23. Matasar MJ, Ritchie EK, Consedine N, Magai C, Neugut AI. Incidence rates of acute promyelocytic leukemia among Hispanics, blacks, Asians, and non-Hispanic whites in the United States. *Eur J Cancer Prev*. 2006;15(4):367-370.
 24. Caspar-Bauguil S, Fioroni A, Galinier A, et al. Pro-inflammatory phospholipid arachidonic acid/eicosapentaenoic acid ratio of dysmetabolic severely obese women. *Obes Surg*. 2012;22(6):935-944.
 25. TrabANELLI S, Chevalier MF, Martinez-Usatorre A, et al. Tumour-derived PGD2 and NKp30-B7H6 engagement drives an immunosuppressive ILC2-MDSC axis. *Nat Commun*. 2017;8(1):593.
 26. Kliewer SA, Lenhard JM, Willson TM, Patel I, Morris DC, Lehmann JM. A prostaglandin J2 metabolite binds peroxisome proliferator-activated receptor γ and promotes adipocyte differentiation. *Cell*. 1995;83(5):813-819.
 27. Peters JM, Shah YM, Gonzalez FJ. The role of peroxisome proliferator-activated receptors in carcinogenesis and chemoprevention. *Nat Rev Cancer*. 2012;12(3):181-195.
 28. Poloz Y, Stambolic V. Obesity and cancer, a case for insulin signaling. *Cell Death Dis*. 2015;6:e2037.
 29. Dahlén SE, Björk J, Hedqvist P, et al. Leukotrienes promote plasma leakage and leukocyte adhesion in postcapillary venules: in vivo effects with relevance to the acute inflammatory response. *Proc Natl Acad Sci U S A*. 1981;78(6):3887-3891.
 30. Stockard B, Garrett T, Guingab-Cagmat J, Meshinchi S, Lamba J. Distinct Metabolic features differentiating FLT3-ITD AML from FLT3-WT childhood Acute Myeloid Leukemia. *Sci Rep*. 2018;8(1):5534.
 31. Moskal A, Pisa PT, Ferrari P, et al. Nutrient Patterns and Their Food Sources in an International Study Setting: Report from the EPIC Study. *PLoS One*. 2014;9(6):e98647.