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A transcriptomic signature predicting septic outcome in patients undergoing autologous stem cell transplantation

Running title: Predictive transcriptomic signature of sepsis

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Highlights :

- 1) A transcriptomic signature predicts infection in neutropenic patients
- 2) Expression of eleven genes can be used routinely for sepsis prediction
- 3) Early anti-infectious treatment may improve sepsis outcome
- 4) Identification of a predictive transcriptomic signature is ongoing in leukemia

1 **Abstract**

2 Autologous hematopoietic stem cell transplantation (auto-HSCT) is a standard treatment in multiple
3 myeloma and relapsed or refractory lymphomas. After auto-HSCT, hematologic reconstitution and
4 infectious complications are the main two critical issues. Though many patients develop infectious
5 complications after therapeutic intensification, it remains impossible to predict infection for each
6 individual. The goal of this work was to determine and identify a predictive transcriptomic signature
7 of systemic inflammatory response syndrome (SIRS) and/or sepsis in patients receiving auto-HSCT.

8 High throughput transcriptomic and bioinformatics analysis were performed to analyze gene
9 expression modulation in peripheral blood mononuclear cells (PBMCs) in 21 patients undergoing
10 auto-HSCT for hematological malignancies (lymphoma or multiple myeloma [MM]).

11 Transcriptomic analysis of PBMCs samples collected just after conditioning regimen identified an
12 eleven genes signature (*CHAT, CNN3, ANKRD42, LOC100505725, EDAR, GPAT2, ENST00000390425,*
13 *MTRM8, C6orf192 and LOC10289230 and XLOC-005643*) that was able to early predict (at least 2 to 7
14 days before its occurrence) the development of SIRS or sepsis.

15 The possibility of SIRS or sepsis occurrence early prediction (2–7 days before occurrence) opens up to
16 new therapeutic strategies based on pre-emptive antibiotic and/or antifungal prophylaxis adapted to
17 the specific risk profile of each patient.

18

19 **Key words:** Systemic Inflammatory Response Syndrome (SIRS); Sepsis; Autologous Hematopoietic
20 Stem Cell Transplantation (Auto-HSCT); Transcriptomic Analysis; Treatment related mortality (TRM)

21 **Abstract word count: 195**

22 **Introduction**

23 Auto-HSCT is based on the administration of myelosuppressive high-dose chemotherapy, followed by
24 infusion of autologous hematopoietic stem cells to obtain hematologic reconstitution. Hematopoietic
25 stem cells (HSCs) infusion reduces chemotherapy-induced myelosuppression period and procedure-
26 related mortality rate below 3% [1–3]. With few exceptions (solid tumors, autoimmune diseases),
27 auto-HSCT is essentially indicated for selected hematological malignancies and considered as a
28 standard treatment in young patients with MM and for relapsed or refractory lymphoma.

29 Besides direct toxicity of conditioning regimens, deep (<0.5 G/L neutrophils) and prolonged (usually
30 7–12 days) neutropenia exposes patients to significant risks of infection. The saprophytic gram-
31 negative bacilli (such as *Escherichia coli*) are the most common cause of septic shock [4] and chronic
32 immunosuppression exposes to the risk of fungal infection. An anti-fungal prophylaxis is usually
33 administered [5] but antimicrobial prophylaxis is less often given because its effectiveness is not
34 clearly established and increases the risk of *Clostridium difficile* diarrhea [6]. Unfortunately, it is not
35 possible to foresee which patients will develop a SIRS and/or sepsis. Therefore, it remains impossible
36 to adjust the antibiotic or antifungal prophylaxis to the specific risk profile of each patient.

37 The main objective of this work was to determine and identify a predictive transcriptomic signature
38 of the SIRS and/or sepsis in patients receiving auto-HSCT, leading to the possibility of a pre-emptive
39 anti-infectious treatment.

40

41 ***Patients, materials and methods***

42 ***Patients***

43 The **prospective** study was approved by the institutional review board of the Assistance Publique des
44 Hôpitaux de Marseille (AP-HM – AORC2012 2012-08). Written informed consent was obtained from
45 each patient. Patients were admitted in the hematology department of the Conception university
46 hospital for undergoing auto-HSCT. All patients were under 65 years and were already followed in
47 the hematology department for MM or high-grade lymphoma. Before auto-HSCT, patients were in
48 complete remission (CR) or in partial remission (PR) after conventional chemotherapy. Inclusion
49 criteria were the same as required for being eligible to auto-HSCT. Twenty four patients were
50 included in this study protocol. Twenty one were analyzed. Blood samples were collected for each
51 patient at three moments: before the conditioning regimen (T1), after the conditioning regimen and
52 before the graft infusion (T2) and at the end of the neutropenic phase (T3).

53 **All patients (after written informed consent) hospitalized in the Hematology and Cellular Therapy**
54 **Department of the university hospital of Marseille between August 2012 and December 2015 have**
55 **been included. Only the patients (21 in all) in whom the RNA extraction was a success (after**
56 **quantification and quality control) have been analyzed.**

57

58 ***SIRS and sepsis definitions***

59 SIRS and sepsis definition are based on the American College of Chest Physicians society of Critical
60 Care Medicine consensus. SIRS is defined as the systemic inflammatory response to a variety of
61 severe clinical symptoms with at least two of the following criteria: a) Temperature higher than 38 °C
62 or lower than 36 °C b) Heart rate higher than 90 beats/min c) Respiratory rate higher than 20
63 breaths/min or PaCO₂ lower than 32 mmHg d) White blood cell counts higher than 12,000 cells/mm³
64 or lower than 4,000 cells/mm³, or the presence of more than 10% immature neutrophils. The last
65 criterion cannot be considered in auto-HSCT context because of the aplasia phase following HSC

66 infusion [7, 8]. Sepsis is defined as SIRS secondary to documented or suspected infection. Patients
67 with severe sepsis are patients with sepsis and at least one organ dysfunction. Septic choc is defined
68 by severe sepsis associated with refractory hypotension [8].

69

70 ***Conditioning regimens***

71 Conditioning regimen for therapeutic intensification was high dose melphalan (200 mg/m²) for
72 patients with MM and BEAM for patients with lymphoma (carmustin 300 mg/m² at day -6,
73 etoposide 150 mg/m² from day -5 to -2 twice daily, cytarabin 200mg/m² from day -5 to -2 twice daily
74 and melphalan 140mg/m² day -1, with auto-HSCT on day 0).

75

76 ***Hematopoietic stem cell collection***

77 HSCs mobilization required hematopoietic growth factors (G-CSF in most cases). Two procedures
78 have been used to mobilize HSCs. For steady-state collection, G-CSF was injected at the dose of
79 10µg/kg-a-day. Mobilization chemotherapy consisted in high doses cyclophosphamide (1.5 to 4g/m²)
80 followed by 5µg/kg-a-day of G-CSF. When HSCs collection failed with these usual procedures,
81 plerixafor (SDF-1/CXCR4 interaction inhibitor) was used at 240 µg/kg.

82

83 ***PBMCs isolation***

84 Blood samples were centrifuged to separate plasma and other blood components. Concentrate blood
85 was diluted and PBMCs were collected by Ficoll-Hystopaque density-gradient centrifugation [9].

86

87 ***RNA extraction, quantification quality controls***

88 Depending on cell number, two kits were used for RNA extraction, the RNeasy Mini Kit Qiagen™
89 (Qiagen, Valencia, California), which accepts 10 million of cells and the RNeasy Midi Kit Qiagen™,
90 which accepts up to 100 million cells. RNAs were extracted according to the Qiagen protocol. RNAs

91 were quantified by NanoDrop 1000 (Nano Drop Technologies, San Diego, CA). Optical density was
92 measured at 260 and 280 nm and the ratio 260/280 (> 1.8) indicates its purity. Extracted RNA quality
93 was checked with Agilent 2100 Bioanalyzer™ (Agilent Technologies, Santa Clara, California). A score
94 on a scale of 0 to 10 was automatically attributed to each sample and corresponded to RNA Integrity
95 Number (RIN). Sample with a RIN under 7 was discarded.

96

97 ***Pangenomic gene expression assay***

98 One hundred nanograms of total RNA was labelled using One-Color Microarray-Based Gene
99 Expression Analysis: Low Input Quick Amp (LIQA) labelling protocol. 0.6 µg of the purified Cy3
100 labelled cRNA were hybridized for 17h at 65 °C, at 60 rpm, using the SurePrint G3 human GE 8x60K
101 V2 chips (Agilent Technologies, Santa Clara, California). Microarrays were composed of 62 928
102 features. Probes synthesized on chips had a size of 60 nucleotides. Microarrays were washed using
103 Gene Expression Wash Buffer Kit (Agilent Technologies) and scanned through the standard Agilent
104 protocol. Data were processed using Feature Extraction software.

105

106 ***Differential gene expression analysis***

107 The library AgiND is implemented in R software in order to analyze and visualize data. AgiND was
108 developed on Bioconductor library model (tagc.univ-mrs.fr/ComputationalBiology/AgiND/) and is
109 used to diagnose data quality and data-microarray normalization. Quantile method was used to
110 normalize data; the objective was to homogenize distribution of microarray intensity [10]. A filter
111 was applied on row data to delete controls, then a second filter was applied to delete genes which
112 were expressed under the background in at least 80% of samples in each group (SIRS–, SIRS+,
113 Sepsis+).

114 To test co-factors effects (gender, treatment, infection) on gene expression, GeneANOVA software
115 was used to perform ANalysis Of Variance (ANOVA) on normalized data to determine an estimation

116 of the contribution of each factor (genes, gender, treatment, infection) in gene expression variation.
117 Global ANOVA model is given in the following formula: $Y = \mu + \beta_1 G + \beta_2 T + \beta_3 I + \epsilon$, where Y is
118 explained variable, μ is global mean, β_1 , β_2 , β_3 , are model coefficients, and G, T, I are the quantitative
119 variables; $\beta_1 G$, is the gene effects, $\beta_2 T$, is the treatment effects, $\beta_3 I$, the infection effects, and ϵ is
120 the error term [11]. Differential gene expression analysis was performed using linear Models for
121 Microarray Data (Limma). Limma is a multivariate analysis and takes into account co-factors-effect
122 tested by ANOVA analysis (treatment and gender) (**Table 2**). In order to determine infection impact
123 on gene expression and on the heat map, new gene expression was calculated after subtracting
124 gender and treatment co-factors effects. Unsupervised hierarchical clustering was applied on
125 adjusted gene expression median adjusted data to group genes and samples, according to their
126 expression using 'TMeV' (Tigr MultiExperiment Viewer) MeV: MultiExperiment Viewer | Part of the
127 TM4 Microarray Software Suite [<http://www.tm4.org/mev/>]. Pearson correlation was used, and
128 clusters were grouped on the basis of average linkage method.

129

130 ***Support Vector Machines (SVMs)***

131 SVMs were applied to predict classification of patients according to the predictive signature. SVMs
132 use training set in which genes known to be related to each other by function or samples related to a
133 group are described as positive examples and genes or samples known not to be members of that
134 class are labelled as negative examples. In this study, samples were attributed to two groups: SIRS
135 and sepsis patients in the first group and patients with no temperature in the second. They were
136 combined into a set of training examples used by SVM to distinguish class members from non-
137 members on the basis of expression data. After learning the class expression features, the SVM can
138 be used to recognize and classify each sample on the basis of their expression [11, 12]. SVMs tool is
139 implemented in TMeV software MeV: MultiExperiment Viewer | Part of the TM4 Microarray
140 Software Suite. [<http://www.tm4.org/mev/>].

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High-throughput quantitative PCR

RT-PCR was performed using SuperScript™ VILO™ MasterMix protocol described by Invitrogen. Sixty nanogram of RNA of each sample was retrotranscribed, using 4µL of SuperScript™ VILO™ MasterMix, and RNase DNase Free water for a final volume of 20µL. Mix was incubated 10 min at 25 °C, 60min at 42 °C and 5min at 85°C. Quantitative PCR was performed with FlexSix array on BioMark™ HD (Fluidigm). A pre-amplification of each sample was applied according to the protocol provided by Fluidigm. The pre-amplification step was followed by an Exonuclease I treatment (BioLegend) to remove unincorporated primers. The final product was diluted 5-fold using 18µl of TE Buffer (10 mM Tris-HCl, 1mM EDTA). In a second part, 12 x 12 (samples x primers) qRT-PCR reactions are performed for each primer pair on each sample on the 12 x 12 array (FlexSix). We used the EvaGreen detection assay for following standard Fluidigm protocols. Ct values were calculated from the system software Biomark Real-Time PCR Analysis (Fluidigm). The primers used were:

Gene	R	F
SLC18B1	5'-GCAGGACAGCTTTTCAGTATCAC-3'	5'-CCTGGCCTTAAACATCACCG-3'
CNN3	5'-GGCTGGCTCCTTTATTAGTGC-3'	5'-AAGCTGGCCAAAGTGAATTG-3'
MTMR8	5'-ACTTAAGTGGGGAGAGGGGT-3'	5'-GAGGCAGACTACTCCAAGCA-3'
GPAT2	5'-GCCCAGAGAAGCCTACATCA-3'	5'-CATCAGAAGCTCCTGGGGGA-3'
PLCG1	5'-ACAGGAATCTTGGTGCTTCAGT-3'	5'-CAGGGAGGTACATGGCCAAT-3'
ANKRD42	5'-CGCCCATGAAAAGCTGCATA-3'	5'-GGAGCAAATCTGACAGCCCA-3'
GAPDH	5'-CCACCACCCTGTTGCTGTGA-3'	5'-CCCCTCCTCCACCTTTGAC-3'
EDAR	5'-ACATGAGCTGACTGGCTG-3'	5'-TGAGTGTGCCATTCCAGGAT-3'
TRAV3 -	5'-GCCTCGGTTGGGGTATTGAA-3'	5'-CTCAGCCGGAAGATCAGGTC-3'

CHAT	5'-CAGCAGAACATCTCCGTCGT-3'	5'-ATGGCCATTGACAACCACCT-3'
LOC100289230	5'-GTGCTCCGGAAAACGTAAA-3'	5'-TCCTGTGCCCGTAATTTCTC-3'

157

158 **Results**

159 ***Patient's analysis***

160 Twenty-four patients were included in the study, among these patients; 21 validated the molecular
161 criteria for transcriptomic analysis. Among these 21 patients, 6 patients did not develop a fever or
162 SIRS (28.6%), nine patients developed SIRS (42.6%), 5 a sepsis (23.8%) and 1 a severe sepsis (5%).
163 Patients' clinical data are summarized in the **Table 1**.

164

165 ***Pangenomic gene expression***

166 The 21 samples were analyzed on Agilent microarrays 'SurePrint G3 GE 8x60K human.' After 17 hours
167 of hybridization, the chips were washed and scanned. Results passed microarrays quality controls.
168 Raw data were transformed into *log2* and normalized with quantile method. 24 046 probes
169 expressing a higher signal than background (in at least one group) have been selected. Two methods
170 were used for statistical analysis: ANOVA analysis to measure the impact of each factor on gene
171 expression variation (infection, gender, treatment) and Limma analysis on three groups (SIRS-,
172 SIRS+, sepsis +) with the following co-factors: gender and treatment, to define differential gene
173 expression (**Figure 1**). ANOVA analysis was performed on the 24 046 probes to estimate impact of
174 each factor (infection, gender, treatment) in gene expression variation. P-value was calculated for
175 each factor. For all factors (genes, infections, gender, and treatment) p-values were $<10^{-4}$ (**Table 2**).
176 Infection F score was the highest, suggesting that 'infection' had the greatest impact on the gene
177 expression. Genes F score was of 115.45 (p-value $<10^{-4}$) and explains genes fluctuations. Treatment
178 and gender had an impact on the variation of gene expression with a F score of respectively 176.21
179 and 156.07 (p-value $<10^{-4}$). Unsupervised hierarchical clustering method was used to classify the
180 differently expressed genes identified by Limma analysis. Expression similarity profiles of the genes
181 were grouped on the horizontal axis and samples on the vertical axis. Gene expression profiles are

182 shown in heat map (**Figure 2**). With a FDR (False Discovery Rate) fixed to 5%, 11 genes differentially
183 expressed were identified between the patients who did not have temperature (SIRS-) and the
184 patients who developed SIRS and sepsis (taking account effect of gender and treatment). The eleven
185 differentially expressed genes were: *CHAT*, *CNN3*, *ANKRD42*, *LOC100505725*, *EDAR*, *GPAT2*,
186 *ENST00000390425*, *MTRM8*, *LOC10289230* and *XLOC-005643* (**Table 3**). Gene expression profiling
187 distinguished two groups: patients who did not develop temperature, on the one hand, and patients
188 with SIRS or sepsis, on the other hand. *CHAT*, *CNN3*, *C6orf192*, *ANKRD42*, *LOC100505725*, *EDAR*,
189 *GPATZ*, *ENST00000390425*, *MTRM8*, *LOC10289230* were over expressed and *XLOC-005643* was
190 under-expressed in patient samples with SIRS or Sepsis in contrast to patients who did not develop
191 fever and SIRS.

192

193 **SVM**

194 SVM and leave-one-out cross-validation were used to classify patients according to their gene
195 expression. 21 samples were separated into two groups, the first group was composed of 15 SIRS and
196 septic patients and the second of the 6 patients who did not have temperature. SIRS and sepsis
197 patients were considered as positive experiments, and the other as negative. All 15 patients of the
198 positive experiments were classified as positive, staying in positive class (true positives=15) and none
199 was transferred from negative class to positive (False negatives=0). On the 6 patients of the negative
200 experiments, all were classified as negative, all retained in negative class (true negatives=15) and
201 none was recruited into negative class from positives (false positives=0). **We performed a Fisher**
202 **exact test on our results; the p-value = 1.84 E-5.**

203

204 **RT-qPCR**

205 In order to confirm the microarray gene expression results, RT-qPCR was performed on 9 genes that
206 composed the transcriptomic signature. The *LOC 100289230* gene was differently expressed (p-value

207 = 0.003) in patients who developed SIRS or sepsis compared with patients who did not. *TRAV3*, *EDAR*,
208 *PLCG1-AS1-001*, *GPAT2*, *MTRM8*, *CNN3*, and *SLC18B1* were also differentially expressed with p-values
209 of 0.004, 0.01, 0.01, 0.005, 0.009, 0.09 and 0.01, respectively (Figure 3).

210

211

212 ***Discussion and conclusion***

213 This clinical study identified eleven genes significantly and differentially expressed in patients who
214 developed SIRS or sepsis after the conditioning regimen for auto-HSCT. Ten of them were up-
215 regulated (*CHAT*, *CNN3*, *ANKRD42*, *LOC100505725*, *EDAR*, *GPATZ*, *ENST00000390425*, *MTRM8*,
216 *C6orf192* and *LOC10289230*) while only one was down-regulated (*XLOC-005643*). All patients with
217 this specific transcriptomic signature developed a SIRS or sepsis within 48 hours (range 48 hours-7
218 days) following conditioning regimen. All patients were classified in the right group according to their
219 gene expression and based on SVMs analysis. After a wide scientific literature review, genes
220 composing our predictive signature are not directly involved in sepsis or infection pathways. In this
221 cohort of 21 patients, 9 developed a SIRS and 5 a sepsis. Only one patient developed a severe sepsis,
222 and no patient had septic shock, thus impeding the possibility to identify a specific transcriptomic
223 signature predictive for these life-threatening conditions. In addition, since a SIRS/sepsis predictive
224 signature before the conditioning regimen was not identified, the transcriptomic signature was not
225 linked to the patient pre-auto HSCT status but depended on the conditioning regimen patient's
226 response.

227 We wondered about the definition of sepsis and SIRS and the clustering of sirs and sepsis;
228 first, white blood cells count criteria is not relevant in patients in deep aplasia. Furthermore, in the
229 same way, patients in deep aplasia present grade II-III anemia especially after BEAM conditioning
230 regimen. In patients with no cardiovascular and/or pulmonary comorbidity, hemoglobin until 8g/dl is

231 tolerated. However, that often results in an increase of the heart rate higher than 90 beats/min
232 and/or respiratory rate higher than 20 breaths/min, especially in patients with fever.
233 Fever is neither sensitive or specific in conventional patients. But, in our severely
234 immunocompromised patients fever is more sensitive and specific than in immunocompetent
235 patients. Nevertheless, it is not unusual to start an antibiotic therapy in patients with no fever but
236 with a microbiological documentation and/or a major increase of the C reactive protein only.
237 At last, we have been very drastic on the definition of our sepsis patients group. Only the patients
238 with fever and a microbiological documentation have been considered in sepsis. Anyway, our
239 immunocompromised patients are paucisymptomatic and very few infections were clinically
240 probable.

241 Our transcriptomic signature predicts SIRS/sepsis profiles and is more robust than the main
242 confounding factors, such as conditioning regimen, type or gender. BEAM conditioning regimen was
243 more myelosuppressive than melphalan alone and men have had a more significant risk to develop
244 infectious complications than women [13]. Auto-HSCT patients affected by MM are conditioned by
245 high dose of melphalan while patients with lymphoma by high dose BEAM chemotherapy, meaning
246 that treatment and pathology are confounding factors. SAM and ANOVA analyses were performed
247 on the data of patient samples before chemotherapy, and no differentially expressed genes were
248 found between patients affected by lymphoma or MM.

249 This work proposes a transcriptomic approach of the sepsis issue during auto-HSCT
250 neutropenic phase. Vanska et al. have shown that high pentraxin 3 level predicted septic shock and
251 bacteremia at the onset of febrile neutropenia after intensive chemotherapy of hematologic patients
252 [14]. Nonetheless, high pentraxin 3 level had only a predictive value for septic shock in patients who
253 already had a febrile neutropenia. In contrast, our predictive transcriptomic signature identifies
254 patients who have a major risk to develop SIRS and/or sepsis at least 48 hours (range 48 hours–7
255 days) before onset of fever. In order to confirm our results, our transcriptomic signature has been

256 validated on a 10 patient's prospective validation cohort (data not shown). Since total gene
257 expression analysis by microarrays is not possible in clinical routine for time and cost-effective
258 reasons, RT-qPCR analysis on a blood sample of the eleven deregulated genes will have to be
259 developed in order to assess the possibility of a routine use.

260

261 Prophylactic anti-fungal and/or anti-microbial treatments have not proven a high efficiency
262 regarding the outcome in patients undergoing highly myeloablative chemotherapy. For example, the
263 use of ciprofloxacin induces a 20% reduction of infection rates but lead to 70% increase in the
264 development of ciprofloxacin resistance [15]. Early identification of patients who will develop SIRS
265 and/or sepsis could perhaps contribute to a better use of anti-infectious agents as preemptive
266 treatment instead of a prophylactic treatment, although this is only an hypothesis by should be
267 tested in a prospective clinical trial. A limitation of our study, directly related to the small number of
268 patients and of infectious events, is the impossibility to identify a specific profile for microbial or
269 fungal or viral infections, in order to precisely orientate the preemptive treatment. Finally,
270 identification of high vs low infectious risk patients after myeloablative chemotherapy, could help to
271 select patients with a safe septic profile and allows an early hospital discharge or even outpatient
272 based blood stem cell transplantation, in order to develop a more rational utilization of hospital
273 resources but still maintaining optimal safety conditions [16-19]. A validation of this transcriptomic
274 signature in a prospective and larger cohort of patients is necessary to have a stronger clinical
275 impact. Another strategy (work in progress) is to validate in RT-qPCR only the eleven genes of
276 interest in a prospective and larger cohort.

277

278 ***Conflict of interest***

279 All authors declare no financial conflict of interest

280

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285

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296

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348 **Table legends:**

349 **Table 1: Patients, pathologies and auto-HSCT characteristics**

350 **Table 2: co-factors effect on gene expression.**

351 ANOVA analysis was performed to test the co-factors impact (infection, treatment and gender) on
352 gene expression for samples taken at T2 (after chemotherapy). With p-values <0, 05 infection,
353 treatment and gender have a significant effect on the gene expression. DF: Degrees of Freedom

354 **Table 3: Differentially expressed gene characteristics**

355 Ch: Chromosome, +1: Forward strand, from 3' to 5', -1: Reverse strand, from 5' to 3'

356

357

358 **Figure legends:**

359 **Figure 1: Differential gene expression flow chart**

360 Pangenomic array for the transcriptomic analysis have been used, data were filtered and the controls
361 were suppressed, cofactors effects were tested based on their implication on variation of gene
362 expression. A multivariate analysis – LIMMA – using the cofactors already tested was performed to
363 define the gene differentially expressed using Bioconductor library. The LIMMA model is given in the
364 following formula: $Y = \alpha + \beta_1 \cdot T + \beta_2 \cdot G + \beta_3 \cdot I + \epsilon$, where Y is the explained variable, α is the global
365 mean, $\beta_1 \cdot T$, $\beta_2 \cdot G$, $\beta_3 \cdot I$, are the model coefficients, T, G, I, are the quantitative variables, $\beta_1 \cdot T$ is the
366 treatment effect, $\beta_2 \cdot G$ is the gene effect, $\beta_3 \cdot I$ is the infection effect and ϵ is the error term. Then, we
367 performed multi-testing correction by fixing the threshold to 5%. Finally, we adjusted the expression
368 data based on the co-factors before performing the hierarchical clustering *: Linear Models for
369 Microarray data.

370

371 **Figure 2: Gene expression profile adjusted on the effects of the cofactors**

372 Microarray analysis revealed 11 genes differentially expressed. This genes cluster distinguished SIRS
373 or sepsis patients from patients who didn't develop any fever or SIRS.

374

375 **Figure 3: Transcriptomic validation signature by RT-qPCR**

376 Relative gene expression composed the transcriptomic signature of patients who developed SIRS or
377 sepsis compared to patients who did not develop fever and SIRS

378 * = 0.05; ** = 0.01; *** ≤0.01, NS = not significant

379

Table 1 : Patients, pathologies and auto-HSCT characteristics

Baseline and Demographic Characteristics	n = 21
Patients age (median)	58 years Range [32-68]
Patient gender	
Male	14 (67%)
Female	7 (33%)
Hematologic malignancy	
Lymphoma	2
Multiple Myeloma	19
Conditioning Regimen	
BEAM	2
Melphalan high dose	19
Injected CD 34⁺ (x 10⁶ / kg)	2.82 Range [0.751-3.64]
Mean duration of fever	3 days SEM [-4.30 - 4.30]
Mean duration of neutropenia	9 days SEM [-2.90 – 2.90]
Documented bacterial infections	6 (30%)
Gram-positive bacteria	75%
<i>Staphylococcus hominis</i>	1
<i>Staphylococcus epidermidis</i>	2
<i>Staphylococcus aureus</i>	1
<i>Streptococcus mitis</i>	1
<i>Clostridium difficile</i>	1
Gram-negative bacteria	25%
<i>Escherichia coli</i>	1
<i>Klebsiella pneumonia</i>	1
Documented fungal infections	0 (0%)
Patient classification	
SIRS-	6 (29%)
SIRS+	9 (42%)
Sepsis+	5 (24%)
Severe sepsis+	1 (5%)
Septic shock+	0 (0%)

Table 2: Co-factors effect on gene expression.

ANOVA analysis was performed to test the co-factors impact (infection, treatment and gender) on genes expression for samples taken at T2 (after chemotherapy). With p-values < 0, 05 infection, treatment and gender have a significant effect on gene expression. DF: Degrees of Freedom

Factors	Sum of squares	DF	Variance	F score	P-value
Genes	3476905.49	28114	123.67	115.45	< 10 ⁻⁴
Infection	954.39	1	954.39	890.96	< 10 ⁻⁴
Treatment	188.75	1	188.75	176.21	< 10 ⁻⁴
Gender	167.18	1	167.18	156.07	< 10 ⁻⁴
Residual	602328.16	562297	1.07	-	-
Total	4080543.97	590414	6.91	-	-

Table 3: Differentially expressed gene characteristics.

Ch : Chromosome, +1 : Forward strand, from 3' to 5', -1 : Reverse strand, from 5' to 3'

Gene	Complete name	Ensembl ID	Location	Sens
SLC18B1 / C6ORF192	solute carrier family 18 member B1	ENSG00000146409	Chr 6: 132,769,370-132,798,553	-1
CNN3	calponin 3	ENSG00000117519	Chr 1: 94,896,949-94,927,278	-1
MTMR8	myotubularin related protein 8	ENSG00000102043	Chr X: 64,268,081-64,395,431	-1
LOC100505725 PLCG1-AS1-001 / TOP1-AS1 /	PLCG1 antisense RNA 1	ENSG00000226648	Chr 20: 41,098,329-41,138,003	-1
CHAT	choline O-acetyltransferase	ENSG00000070748	Chr 10: 49,609,095-49,665,104	1
EDAR	ectodysplasin A receptor	ENSG00000135960	Chr 2: 108,894,471-108,989,372	-1
ANKRD42	ankyrin repeat domain 42	ENSG00000137494	Chr 11: 83,193,739-83,260,694	1
GPAT2	glycerol-3-phosphate acyltransferase 2, mitochondrial	ENSG00000186281	Chr 2: 96,021,946-96,039,451	-1
ENST00000390425/ TRAV3	T cell receptor alpha variable 3 (gene/pseudogene)	ENSG00000211777	Chr 14: 21,723,713-21,724,321	1
LOC100289230	NS	NS	Chr 5: 98,929,134-98,931,009	1
XLOC_005643	linc-CMAHP-1:1 / linc-FAM65B- 1/RP3-425P12.2	ENSG00000230372	Chr 6: 25,061,853-25,063,735	-1

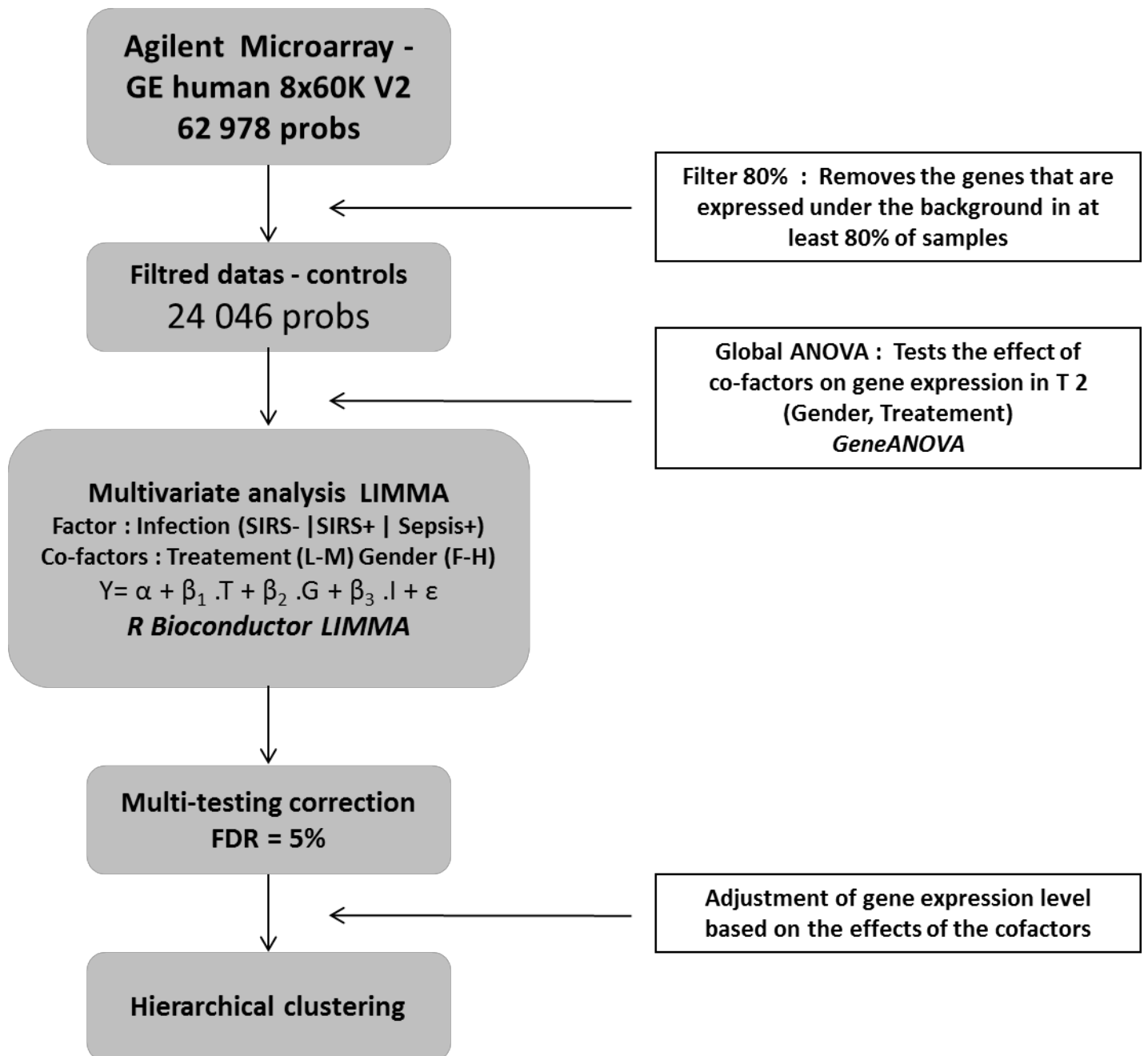


Figure 1: Differential genes expression flow-chart

Figure2

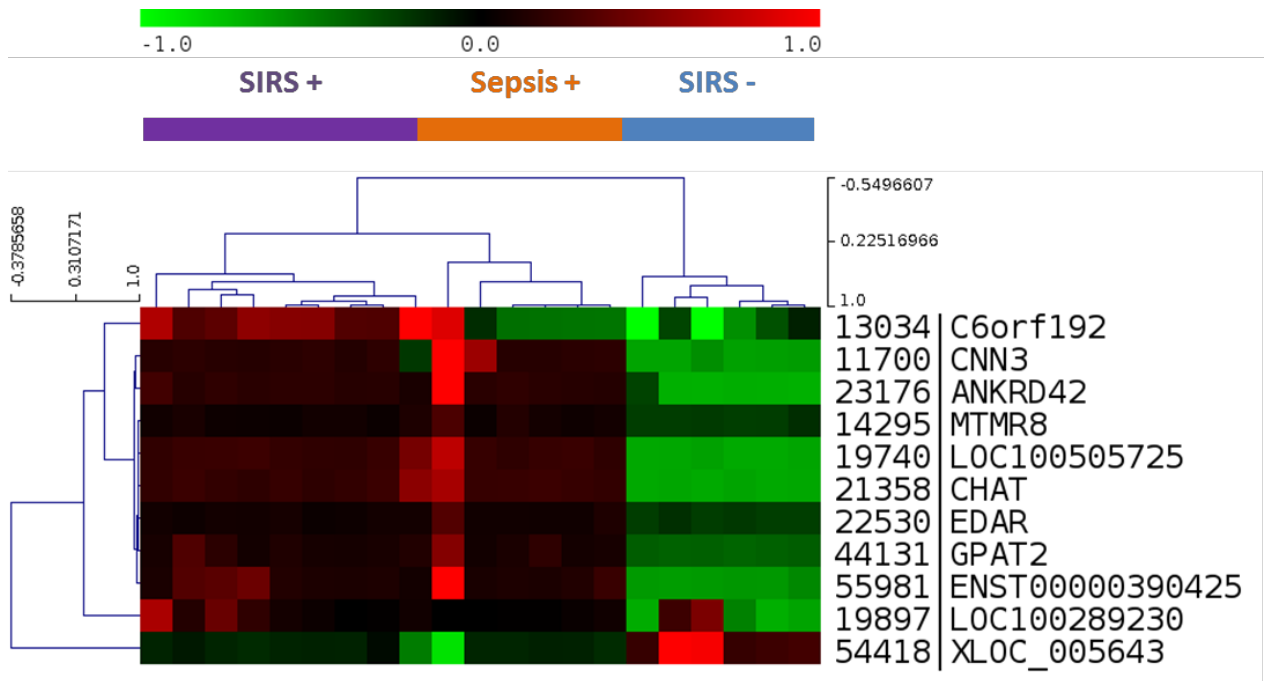


Figure 2: Gene expression profile adjusted on the effects of the cofactors

Color should be used

Figure3

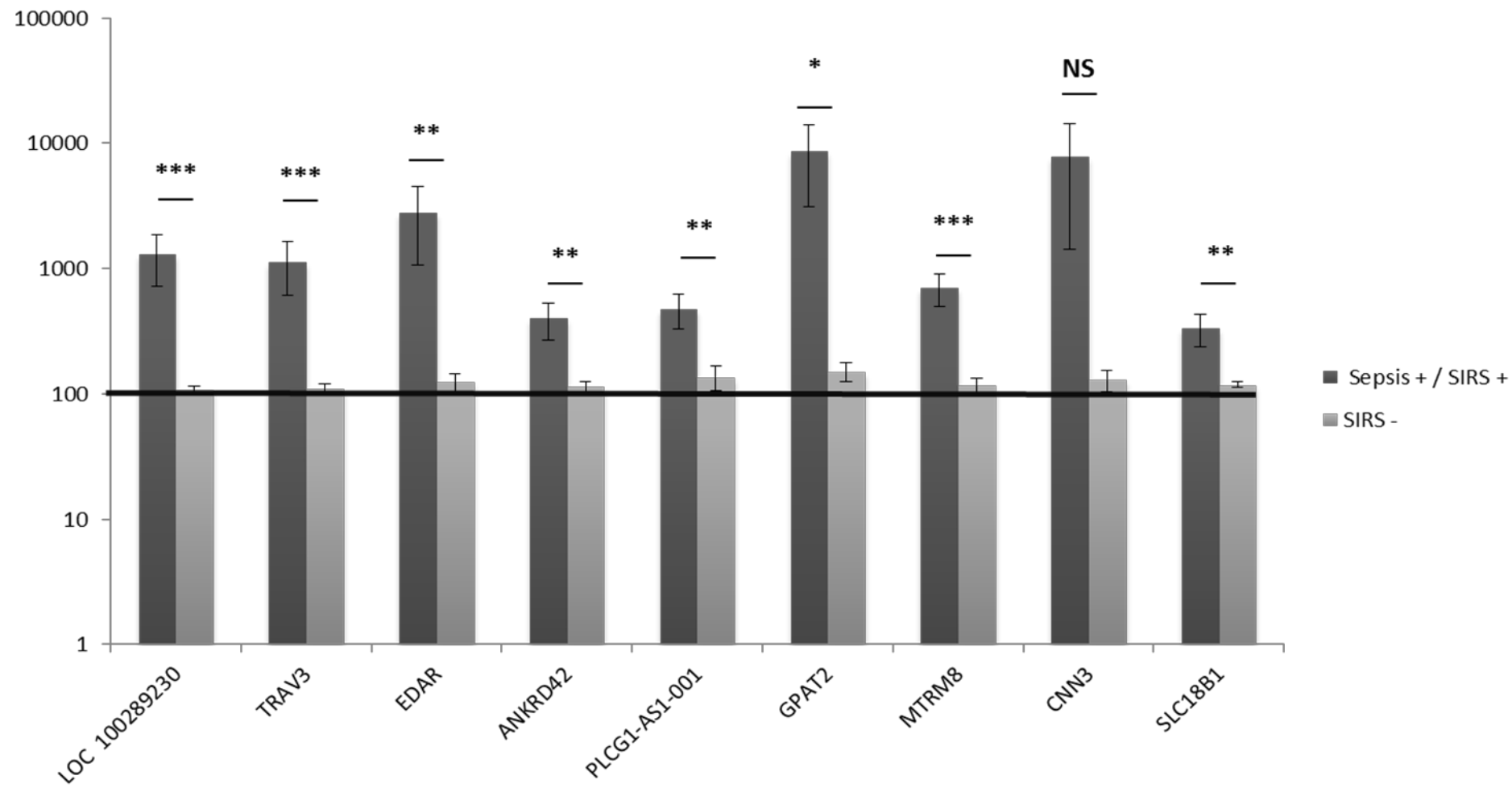


Figure 3 : Transcriptomic validation signature by RT-qPCR