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► **To cite this version:**

Candice Trocmé, Hubert Marotte, Athan Baillet, Béatrice Pallot-Prades, Jérôme Garin, et al.. Apolipoprotein A-I and platelet factor 4 are biomarkers for infliximab response in rheumatoid arthritis.. *Annals of the Rheumatic Diseases*, BMJ Publishing Group, 2009, 68 (8), pp.1328-33. 10.1136/ard.2008.093153 . inserm-00382264

HAL Id: inserm-00382264

<https://www.hal.inserm.fr/inserm-00382264>

Submitted on 2 Jun 2009

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Apolipoprotein A-I and platelet factor 4 are biomarkers for Infliximab response in Rheumatoid Arthritis

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Keywords: rheumatoid arthritis, SELDI-TOF-MS, infliximab, biomarkers, treatment response

Word count: 3000

ABSTRACT

Objectives: The use of biologics such as infliximab has dramatically improved the treatment of rheumatoid arthritis (RA). However, factors predictive of therapeutic response need to be identified. A proteomic study was performed prior to infliximab therapy to identify a panel of candidate protein biomarkers of RA predictive of treatment response.

Methods: Plasma profiles of 60 RA patients (28 non-responders ACR 20 negative and 32 responders ACR 70 positive to infliximab) were studied by SELDI-TOF-MS technology on two types of arrays, an anion exchange array (SAX2) and a nickel affinity array (IMAC3-Ni). Biomarker characterization was carried out using classical biochemical methods (purification by ammonium sulfate precipitation or metal affinity chromatography) and identification by MALDI-TOF analysis.

Results: Two distinct protein profiles were observed on both arrays and several proteins were differentially expressed in both patient populations. Five proteins at 3.86, 7.77, 7.97, 8.14 and 74.07 kDa were overexpressed in the non-responder group, whereas one at 28 kDa was increased in the responder population (sensitivity > 56%, specificity > 77.5%). Moreover combination of several biomarkers improved both the sensitivity and specificity of the detection of patient response to over 97%. The 28 kDa protein was characterized as apolipoprotein A-I and the 7.77 kDa biomarker was identified as platelet factor 4.

Conclusions: We characterized six plasma biomarkers, enabling the detection of patient response to infliximab with high sensitivity and specificity. Apolipoprotein A-1 was predictive of a good response to infliximab, whereas platelet factor 4 was associated with non-responders.

Prevalence of rheumatoid arthritis (RA) varies between 0.3 and 0.8% in European countries.[1] Joint invasion and destruction are mainly mediated by inflammatory cells, cytokines and proteolytic enzymes. The heterogeneity of pathological manifestations, disease course, and response to treatments suggests that several subtypes of RA exist, especially at a molecular level.

Early referral, diagnosis and initiation of treatments to control tissue damage and structural progression are highly recommended, considering the disease's high propensity towards joint destruction, even during periods of clinical remission.[2] Although classical treatments are still used [3, 4], a radical new approach to RA treatment is now available with the use of the biological therapies, such as tumor necrosis factor α (TNF α) inhibitors (etanercept, infliximab, adalimumab), anti-CD20 antibodies (rituximab), interleukin-1 (IL-1) receptor antagonists (anakinra) and an inhibitor of T lymphocyte costimulation (abatacept). Almost 70% of treated patients respond to these therapies, but for unknown reasons, approximately one-third of RA patients do not respond [5]. Indeed, the currently used markers reflecting bone and cartilage turnover [6] are few and inaccurate, and no predictive factor of treatment resistance has been identified. Discovery of biomarkers for diagnosis, prognosis and disease monitoring [7] is necessary.

In order to identify proteins predictive of a treatment response, we chose to study the plasma proteome (i.e. the whole protein content) of RA patients by the technology SELDI-TOF-MS (Surface Enhanced Laser Desorption/Ionization – Time Of Flight Mass Spectrometry) developed by Ciphergen Biosystems, Inc..[8, 9] This method has allowed characterization of numerous biomarkers for many diseases [10-12], particularly cancer.[13-16]

Concerning RA, a first SELDI-based study on 12 synovial fluid samples characterized an RA biomarker absent in osteoarthritis patients, the pro-inflammatory S100A8 protein.[17] More recently the same protein was identified in a larger survey comparing serum protein profiles of RA and non-inflammatory control samples.[18]

In our study, we used this SELDI-TOF-MS technology to characterize biomarkers predictive of an anti-TNF α therapy response in RA patients. Protein profiles were performed on plasma specimens taken prior to the start of infliximab treatment, from patients that were further evaluated as non-responders (ACR 20 negative) or responders (ACR 70 positive) to the therapy. Several biomarkers were isolated and were highly correlated with the infliximab response both independently or when associated in a decision tree algorithm. Moreover, corresponding proteins were identified using classical biochemistry techniques.

PATIENTS AND METHODS

Patients and study protocol

Patients with a RA diagnosis according to ACR (American College of Rheumatology) criteria were recruited in four university hospitals from the Rhône-Alpes area. All patients had disease duration of at least 6 months, had never received anti-TNF α therapy before inclusion and had no active or latent bacterial, fungal or viral infection at the time of enrolment. Blood was withdrawn on a spray-coated K₂EDTA tube before the first infliximab infusion and plasma aliquots were stored at -80°C. Patients then received the same infliximab treatment as previously described [19] and clinical outcome was assessed at week 30. Among the 60 recruited patients, 28 were ACR 20 negative (non-responders), whereas 32 were ACR 70 positive (responders). Differences between both populations of each classical parameter measured at inclusion were evaluated with the Mann-Whitney U-test or the Chi² test.

Proteomic analysis

The plasma protein profile was investigated with the SELDI-TOF-MS method developed by Ciphergen Biosystems, Inc (Fremont, CA), that associates a first phase of protein fractionation on chips with distinct chromatographic surfaces (cationic, anionic, hydrophobic ...) to a second detection step of protein molecular weight by mass spectrometry after laser desorption. We used an anion exchange array (SAX2) and a nickel affinity chip (IMAC3-Ni). Firstly, the sample protein concentration was evaluated using the Bradford method.[20] For the SAX2 array, 5 μ l of plasma diluted to a final concentration of 0.7 μ g/ μ l in a washing / loading buffer (10 mM Tris, 0.1% Triton X100 pH 7) was loaded twice in duplicate for a 30 min incubation period at room temperature. Then spots were washed twice with 5 μ l of washing buffer, twice with 5 μ l of the same buffer without Triton X100 and finally with 5 μ l of 10 mM Hepes pH 7. For the IMAC-Ni array, 20 μ l of plasma was mixed with 30 μ l of a denaturation buffer (50 mM Tris, 8 M urea, 1% CHAPS pH 7.4) for 15 min at +4°C with frequent agitations, and 5 μ l of this protein solution diluted in the washing / loading buffer (PBS, 0.5 M NaCl, 0.1% Triton X100 pH 7.3) to 0.7 μ g/ μ l was applied on each spot previously activated with 100 mM NiSO₄. Incubation and washing steps were performed as previously described, using the IMAC3 buffer with (first two washes) or without Triton X100 (last two washes). Then 0.8 μ l of saturated sinapinic acid in 0.5% trifluoroacetic acid, 50% acetonitrile was applied twice on each spot and chips were analyzed on the ProteinChip[®] reader (model PBSII). Mass data processing was performed with the Ciphergen ProteinChip[®] software 3.1 and intensities, (height from baseline), of characterized proteins were compared in each group using the Mann-Whitney U-test. ROC (Receiver Operating Characteristic) curve analyses were performed to measure the diagnostic potential of each biomarker and correlations between variables (biomarkers and clinical data) were evaluated using the Spearman rank test. Finally, CART (Classification and Regression Tree) analysis allowed data classification in a tree-like structured decision diagram.[21]

Protein purification and identification

Apolipoprotein A-I was purified from 300 μ l of responder plasma by protein precipitation with 50% ammonium sulfate. After centrifugation at 14000 g for 10 min, 5 μ l of supernatant (6.35 μ g) was loaded on a 15% SDS-PAGE gel which was further stained with Coomassie blue R-250. The only band revealed at about 28 kDa was excised from the gel, trypsin-digested and sequenced by LC-MSMS (liquid chromatography-tandem mass spectrometry).

Platelet factor 4 (PF4) was purified from 5.2 ml of non-responder plasma on a cobalt-based affinity resin. After washings, the matrix-bound proteins were desorbed with 6 ml PBS 1X containing 20 mM imidazole and 0.5 M NaCl. This specific eluate was dialyzed overnight in distilled water, lyophilized and the resuspended powder was finally loaded on an 18% SDS-PAGE gel. A thick Coomassie blue R-250 stained band between 6 kDa and 14 kDa was sequenced by LC-MSMS as previously described. As results were not conclusive, we set up an ELISA for platelet factor 4 in order to deplete the column specific eluate from the 7,769 Da biomarker. Then 100 μ l of eluate diluted twice in 2% bovine serum albumin (BSA) was incubated on a 96-well plate for 2 h at room temperature in wells previously coated with 1, 2 or 10 μ g of monoclonal anti-human PF4 antibody (R&D Systems, Lille, France). Next 2 μ l of supernatant was loaded on a normal phase NP20 chip which was analyzed on the ProteinChip[®] reader. Results were expressed as a ratio of the intensity of the 7,769 Da peak to the BSA peak value in order to overcome possible loading discrepancies and significance was assessed using the Wilcoxon test.

RESULTS

Blood protein profiles of RA patients before the start of infliximab treatment

Characteristics of enrolled patients are summarized in table 1. Both populations were similar in terms of patient identity (age and sex), RA characteristics (disease duration, number of swollen and tender joints) and biological parameters such as rheumatoid factor (RF), C-reactive protein (CRP) and erythrocyte sedimentation rate (ESR), demonstrated that in this panel as well as in general RA population, no classical marker of RA severity and aggressiveness could predict the patient response to an anti-TNF α therapy.

Table 1: Demographic characteristics of patients at inclusion

Group characteristics	ACR 20 negative	ACR 70 positive
Number	28	32
Age (years)	53 (37.7-61.3)	58 (31-78.15)
Women (%)	78	90
Disease duration (years)	11,5 (5-31.45)	12.5 (4.95-28.9)
RF positive (%)	75	95
CRP level (mg/liter)	25.65 (0.85-58.05)	13.5 (2.85-45.1)
ESR (mm first hour)	34 (7.4-76.35)	34 (15.9-78.4)
Number tender joints	11 (3.55-21)	13 (5.75-27)
Number swollen joints	4.5 (0.85-10)	6 (0.95-14)

Values correspond to medians + 5th and 95th percentiles, except for sex and RF expressed as percentages. RF = Rheumatoid Factor; CRP = C-Reactive Protein; ESR = Erythrocyte Sedimentation Rate.

Two distinct protein profiles were obtained through plasma analysis on the SAX2 and IMAC-Ni arrays (fig 1A&B). Moreover, duplicate loadings of samples showed a similar protein pattern, characterizing the reproducibility of our experimental procedure (data not shown). Several proteins were found to be differentially expressed in both patient populations: only one

small protein at 3.86 kDa was characterized on the SAX2 anion exchange array whereas five distinct markers at 7.77, 7.97, 8.21, 28.0 and 74.07 kDa were identified from the IMAC-Ni affinity array (table 2). All biomarkers were significantly increased in the non-responder population, except the 28.0 kDa protein which was overexpressed in responders, and they discriminated both populations with an overall high specificity (>77.5%), a lower sensitivity (>56%) and an AUC ranging from 0.761 to 0.846, depending on the biomarkers. Correlation analyses between the six biomarkers showed that only the 3 proteins around 8 kDa isolated on IMAC-Ni were strongly positively related: 7.77 and 7.97 ($r=0.587$, $p<0.0001$), 7.77 and 8.21 ($r=0.597$, $p<0.0001$), and 7.97 and 8.14 ($r=0.869$, $p<0.0001$).

Table 2: Characterization of plasma biomarkers for infliximab treatment response in RA

Marker (kDa)	Peak intensity (mean \pm SD)		p	Cut-off	AUC	Spec (%)	Sens (%)
	NR group	R group					
SAX 2							
3.86	17.22 \pm 7.23	8.37 \pm 3.30	0.0000004	13.14	0.846	82.5	77
IMAC3-Ni							
7.77	20.50 \pm 8.69	11.05 \pm 6.23	0.000005	16.51	0.809	80	71
7.97	15.20 \pm 5.94	9.82 \pm 4.44	0.00002	12.73	0.785	80	71
8.21	17.38 \pm 6.31	11.81 \pm 4.76	0.00002	15.94	0.761	83	56
28.0	1.88 \pm 0.94	3.74 \pm 2.16	0.000005	2.39	0.796	77.5	82
74.07	8.55 \pm 4.34	4.45 \pm 3.01	0.000009	7.18	0.800	87.5	62

NR = Non-Responder; R = Responder; SD = Standard Deviation; Spec = Specificity; Sens = Sensitivity; AUC = Area Under the Curve; p was calculated using the Mann-Whitney test

The entire protein profiles of both populations were then compared in a multivariate CART analysis. The best tree, obtained with the IMAC3-Ni proteomic profile, comprises five nodes: following the path according to the node value (protein peak intensity) leads to a terminal node classifying the patient in either of the two populations. This classification tree predicts infliximab response in this population with a specificity of 97.5% and a sensitivity of 97.1% (fig 2). Combining proteomic profiles with clinical data (CRP, ESR, disease duration and number of swollen or tender joints) or using both the SAX2 and IMAC3 profiles did not improve specificity and sensitivity. Moreover three proteins of the tree at 7.77, 8.21 and 28 kDa are individually highly discriminating (table 2). Box plot representation of individual peak intensities of these proteins, showed a tiny overlap of values between both populations (fig 3A, B&C), especially for the 7.77 and 28 kDa biomarkers. We then focused on the characterization of these two biomarkers.

Purification and identification of the 7.77 and 28.0 kDa biomarkers

The 28 kDa biomarker was first purified by ammonium sulfate precipitation: although part of the protein was precipitated in the pellet, a nearly pure band was recovered at about 28 kDa in the supernatant as shown on an SDS-PAGE gel (fig 4A). Loading supernatant on an IMAC3-Ni chip confirmed this protein to be the previous identified biomarker at m/z 27,976 (data not shown). Finally the sequencing of this protein by LC-MSMS identified it undoubtedly as apolipoprotein A-I (on-line table 1). Correlation analysis showed that this biomarker was independent of clinical variables.

Identification of the 7.77 kDa required more steps. A first purification step on an IMAC-based column allowed the isolation in the specific eluate of one main protein at m/z 7,769 contaminated by faint amounts of 3 proteins between 8 and 9 kDa, as characterized on the IMAC-Ni and NP20 arrays (data not shown and fig 4B). Fractioning this sample on an SDS-

PAGE revealed a thick band around 8 kDa that was sequenced by LS-MSMS: several proteins were characterized, among which platelet factor 4 (PF4, or CXCL4) was the most represented (on-line table 2). Based on previous studies identifying this 7,769 Da protein as mature PF4 (without the first 31 amino acids of the signal peptide) [22, 23], we tried to deplete the column eluate from the biomarker with a homemade PF4 ELISA. The decrease of the peak intensity in the supernatant after incubation with anti-PF4 antibodies, evaluated on an NP20 array, was proportional to the increase of the amount of antibodies coated in the wells (fig 4B), with a significant 44% reduction of the signal for the highest antibody concentration (fig 4C). This last experiment formally identified the 7.77 kDa biomarker as platelet factor 4. As previously, this biomarker was not correlated to the clinical parameters.

DISCUSSION

In this study with SELDI-TOF-MS we characterized several biomarkers closely correlated to the response to infliximab therapy in RA patients. Although other clinical and biological parameters such as number of inflamed joints, HLA haplotypes or CRP level, can partially predict treatment efficiency [7], this is the first time to our knowledge that biomarkers with specificity as high as 87.5% independently and 97.5% when associated in a classification tree, have been determined. Indeed, these prediction characteristics were obtained for the discrimination of two extreme patient groups (ACR 20 negative and ACR 70 positive) and it is likely that specificity and sensitivity of these biomarkers will be lower when used on a more homogeneous RA population. Six biomarkers between 3.86 and 74.07 kDa were isolated on two chromatographic surfaces, suggesting different reactivities of these proteins. The smallest one at 3.86 kDa was characterized on an anion array, meaning that this protein is negatively

charged at pH 7. The other 5 biomarkers isolated on an IMAC-Ni chip have a specific affinity for this metallic ion. Three of these proteins were also recovered in the classification tree built from the IMAC-Ni plasma profile, emphasising their importance in the determination of the infliximab response.

We focused initially on the 28 kDa protein that is overexpressed in the responder population and is also the first node of the decision tree, and we identified it as apolipoprotein A-I. Apolipoprotein A-I is the major protein component of HDL (High Density Protein) particles and the primary acceptor for cholesterol in extra-hepatic tissues. In RA, several studies have reported a decrease of circulating levels of apolipoprotein A-I and HDL-cholesterol in patients compared to the general population [24-26], and concentration of both parameters increases significantly in patients responding to a DMARD (Disease-Modifying Anti-Rheumatic Drug) treatment compared to non-responders.[27] On the other hand, apolipoprotein A-I was highly expressed in inflamed RA synovial tissues particularly in perivascular areas containing infiltrated T cells and macrophages, but was not detected in normal tissue [28] or in the synovium of non-inflammatory RA.[29] Moreover, increased levels of apolipoprotein A-I and cholesterol have been measured in RA synovial fluid.[30] As apolipoprotein A-I was reported to inhibit the synthesis of the major inflammatory cytokines TNF α and interleukin-1 β by blocking direct contact between T lymphocytes and monocytes [31], increased levels of this protein in synovial tissue could modulate inflammation and disease evolution by controlling interactions between immune cells and then cytokine production.[28, 32] This specific function as a “negative” acute phase protein is also emphasised in a study reporting that patients with lower levels of apolipoprotein A-1 develop more systemic inflammatory response syndrome criteria.[33]

Finally, it has been recently shown in a retrospective study among blood donors (n=1078) who later developed RA (n=79) that the lipid profile was perturbed at least 10 years prior to the

onset of symptoms, especially the HDL-cholesterol value that was significantly decreased.[34] The authors suggested that this specific lipid profile may be favourable to the onset of RA or the presence of a constant high inflammatory state due to low levels of HDL-cholesterol. In our study, we demonstrated that high pre-treatment levels of apolipoprotein A-I were predictive of a good response to infliximab. This increased initial concentration may favour the response to anti-TNF α treatment, via the decrease of chronic inflammation, by inhibiting cytokine release and potentiating the efficiency of therapy in a synergistic way.

The second biomarker we focused on was overexpressed in non-responder patients and was identified as platelet factor 4 (PF4) (or CXCL4). This chemokine specific for platelet α -granules exerts multiple functions in several physiological processes.[35] In RA, several studies reported an increase of PF4 in patient synovial fluid [36, 37] and elevated plasma levels in patients with cutaneous vasculitis.[38] In our study, the increase in plasma PF4 was correlated to a non-response to infliximab, and the same rise was recently shown in sera of Crohns disease patients who had not responded to infliximab therapy.[39] Two of its coagulation-unrelated functions may explain this result: PF4 was reported to induce degranulation of TNF α -primed neutrophils [40] and to inhibit monocyte apoptosis.[41] A high PF4 concentration may increase the inflammatory response by potentiating the pro-inflammatory functions of neutrophils and monocytes, that may counteract the efficiency of an anti-TNF α therapy.

In this proteomic study that investigated the plasma protein profile of RA patients who were either responders or non-responders to infliximab therapy, we identified two biomarkers (apolipoprotein A-I and PF4) that may be key elements for RA treatment monitoring. Firstly, these proteins are involved in RA physiopathology and may be associated with specific clinical situations such as inflammation or vasculitis and secondly, both are strongly correlated with

and without clinical response to infliximab treatment. However, these biomarkers need to be validated in a larger cohort of patients and to be evaluated for determination of intermediate treatment response assessed by the ACR score (ACR50) or more linearly by the Disease Activity Score (DAS) 28 index. The identification of the other biomarkers is currently under investigation.

ACKNOWLEDGMENTS

This work was supported by grants from the “Ministère de l’Enseignement Supérieur et de la Recherche”, the “Région Rhône-Alpes”, Joseph Fourier University, the GEFLUC (department of Grenoble), the “Fondation pour la Recherche Médicale”, the “Direction Régionale de la Recherche Clinique” (CHU Grenoble) and the “Ligue Nationale contre le Cancer”.

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LEGENDS

Figure 1: Representative SELDI-TOF-MS plasma profiles on SAX2 and IMAC3-Ni arrays

Plasma proteome of RA patients was analyzed by SELDI-TOF-MS with different laser intensities to characterize large (A) and smaller (B) proteins. Spectra showed relative intensity of each protein according to its mass/charge (m/z) value. Some peak intensities are marked in italic typing.

Figure 2: Decision tree built from IMAC3-Ni protein profile

CART analysis was performed on the whole proteome peaks characterized on the IMAC-Ni array. The intensity (I) of each node corresponding to a protein will draw a path for each patient leading to a terminal node, classifying the patient as a responder (R) or a non-responder (NR)

Figure 3: Relative peak intensities of the 7,769 Da (A) the 8,210 (B) and the 27,976 Da biomarkers in the responder (R) and non-responder (NR) populations

The box represents the interquartile range, the line across the box is the median and the whiskers represent the 5th and 95th percentiles.

Figure 4: Identification of the 27,976 Da and the 7,769 Da biomarkers

(A) Proteins from a responder plasma were precipitated with 50% ammonium sulfate and after centrifugation, 1 µl of pellet (lane 2 – 42.63 µg), 5 µl of desalted supernatant (lane 3 - 6.35 µg) and 1 µl of pure plasma (lane 1 – 77.43 µg) were loaded on a 15% SDS-PAGE gel. The 28 kDa band in the supernatant (circled on the gel) was identified by LC-MSMS as apolipoprotein A-I.
(B) The 7,769 Da biomarker purified on a cobalt-based IMAC was incubated with coated anti-

PF4 monoclonal antibodies (1 to 10 μg / well) and after a 2 h incubation period, supernatant was loaded on an NP20 array. (C) The intensity of this protein peak (in parenthesis on spectra) normalized to the BSA signal decreased while amount of coated antibody increased (* $p=0.022$ calculated with the Wilcoxon's test).

APPENDIX

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Fig. 1

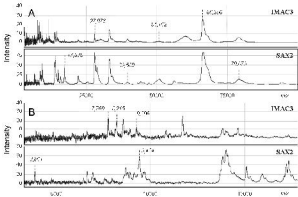


Fig 2

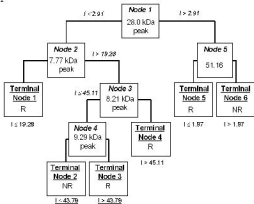


Fig 3

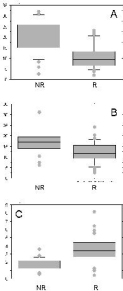


Fig.4

