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Heinrich Hofmann et al.

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Significance of surface charge and shell material of superparamagnetic iron oxide nanoparticle (SPION) based core/shell nanoparticles on the composition of the protein corona†

Usawadee Sakulkhu, a Morteza Mahmoudi, b,c Lionel Maurizi, a Geraldine Coullerez, a Margarethe Hofmann-Amtenbrink, Marcel Vries, Mahdi Motazacker, Marcel Vries, Mahdi Motazacker, Mahdi Mahdi Mahdi Motazacker, Mahdi Mahd Farhad Rezaee^e and Heinrich Hofmann*^a

As nanoparticles (NPs) are increasingly used in many applications their safety and efficient applications in nanomedicine have become concerns. Protein coronas on nanomaterials' surfaces can influence how the cell "recognizes" nanoparticles, as well as the in vitro and in vivo NPs' behaviors. The SuperParamagnetic Iron Oxide Nanoparticle (SPION) is one of the most prominent agents because of its superparamagnetic properties, which is useful for separation applications. To mimic surface properties of different types of NPs, a core-shell SPION library was prepared by coating with different surfaces: polyvinyl alcohol polymer (PVA) (positive, neutral and negative), SiO₂ (positive and negative), titanium dioxide and metal gold. The SPIONs with different surfaces were incubated at a fixed serum: nanoparticle surface ratio, magnetically trapped and washed. The tightly bound proteins were quantified and identified. The surface charge has a great impact on protein adsorption, especially on PVA and silica where proteins preferred binding to the neutral and positively charged surfaces. The importance of surface material on protein adsorption was also revealed by preferential binding on TiO2 and gold coated SPION, even negatively charged. There is no correlation between the protein net charge and the nanoparticle surface charge on protein binding, nor direct correlation between the serum proteins' concentration and the proteins detected in the coronas

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Introduction

The SuperParamagnetic Iron Oxide Nanoparticle (SPION) has promising biomedical applications. The biomedical applications of coated SPION can be divided into 3 major groups: separation (e.g. cellular proteomics, cell sorting, purification), therapy (e.g. hyperthermia,2 drug delivery3) and diagnosis (e.g. Magnetic Resonance Imaging (MRI),^{4,5} cell tracking⁶⁻⁹).

Coating of SPION with a variety of polymers has been approved for some clinical applications. The nanoparticles coated with dextran i.e. ferucarbotran have been used as contrast agents in MRI.¹⁰ Coating with polyelectrolyte PEI-PEG-chitosan copolymer was used for gene delivery.11 Moreover, coating of NPs with other hydrophilic polymers such as polyethylene glycol (PEG), polyvinyl alcohol (PVA) or poly (acrylic acid) has been used for imaging and drug delivery. 12 In addition to polymeric coating, inorganic coatings of SPION with silica or gold have interested researchers and industry for many years towards further surface derivatization of NPs and because of their influences on colloidal stability and the biological behaviour of SPION in biomedical applications. 13,14 SPION is also of interest in the discovery of biomarker proteins for example in blood plasma, serum or urine, which are easily accessible in the acquisition of proteins secreted or released from cells and various interconnecting tissues. These proteins may indicate a disease status if detected and by this serve as biomarkers. On the other hand, as the majority of the nanoparticle formulation, SPION must also be safe for any biomedical applications. The safety of the nanoparticles used in biomedical

^aLaboratory of Powder Technology, Ecole Polytechnique Fédérale de Lausanne, Lausanne, Switzerland. E-mail: heinrich.hofmann@epfl.ch

^bNanotechnology Research Center and Department of Nanotechnology, Faculty of Pharmacy, Tehran University of Medical Sciences, Tehran, Iran

^cDivision of Cardiovascular Medicine, Stanford School of Medicine, Stanford University, CA, USA

^dMat Search Consulting Hofmann, Ch. Jean Pavillard 14, 1009 Pully, Switzerland ^eUniversity Medical Center Groningen (UMCG)University of Groningen, Groningen,

 $[^]f$ Department for Experimental and Molecular Medicine, Academic Medical Center, University of Amsterdam, Amsterdam, The Netherlands

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applications mainly relates to NP stability, biodistribution and toxicity.15 Without coating, naked SPIONs are not stable and become aggregated at physiological pH. In order to overcome this shortcoming, various materials are employed as coatings on the surface of SPION. Coating of SPION does not only improve the colloidal stability but also increases the opportunity to functionalize molecules (e.g. fluorescent dyes, polymers, radiotracers, drugs or targeting biomolecules such as antibodies) onto the SPION surface.

Once the NPs are exposed to biological environments, biomolecule adsorption immediately occurs. If NPs were injected into the body, blood proteins are the biomolecules that have the highest chance to interact with the nanoparticles. 16 Protein adsorption forms a protein corona on the nanoparticles. Protein adsorptions can be categorized by their physiological location into 2 groups: (i) extracellular and (ii) inter-/ intracellular (effects between cell-cell junctions and within cells). Extracellular protein adsorption depends on the nanoparticle characteristics (e.g. size, shape, surface area, surface charge, roughness and porosity, functional groups, ligands, crystallinity and hydrophobicity-hydrophilicity) and on the suspending media.¹⁷ The largest factors of influence on the adsorption arise from acids and bases, salts and multivalent ions, and natural or synthetic organic matter (e.g. proteins, lipids, surfactants, polymers, and polyelectrolytes). All of these factors determine nanoparticle stability and behavior in biological fluids. 18 The extracellular protein adsorption will consequently influence the protein adsorption at the inter-/ intracellular level.

Protein adsorption is one of the major driving forces in the selection of a nanoparticle destination for theranostic applications, which will further affect cells, tissues and finally a body system. Common proteins like albumin, fibrinogen, IgG, complement C3, apolipoprotein A-I and apolipoprotein E¹⁹⁻²² bind to various semi-solid nanoparticles, for instance liposomes, and solid nanoparticles of polymeric or inorganic nature, such as iron oxide, silica, titanium oxide, etc.; some of them also covered by polymer coatings or carbon based nanoparticles such as nanotubes. All of the proteins mentioned are highly abundant in blood plasma, in which at least 99% of the protein mass is covered by 22 of the most abundant proteins. ^{23–25} It is now a challenge of biomarker development to find the specific but very rare protein disease markers (tissue-derived proteins) within the 1% of the remaining proteins of the blood plasma, already diluted on their way to the blood stream.

The constitution of the protein corona depends on two main parameters: (i) composition of the biological milieu and (ii) the surface property of SPION. Further cellular/tissue responses arise due to the composition of the corona. The uptake of nanoparticles is determined by particle-protein and corona-membrane interactions, receptor-ligand binding interactions, membrane wrapping, biomolecule interaction and conformational change in the biomolecules.¹⁷ It is reported that the protein coating reduces the targeting capability of surface engineered NPs by blocking the active sites of the tar-

geting ligands.²⁶ It was revealed that the presence of the corona (protein layer) inhibits the formation of AB fibrils (which is the main cause of Alzheimer's diseases)²⁷ for all of the tested nanomaterials (e.g. carbon nanotubes and silica nanoparticles). All of these factors finally determine either the activation of or damage to physiological responses towards, for example, oxidant injury or mitochondrial and lysosomal damage.²⁸ If the cellular damage is severe, it might cause an adverse effect on the body at the system level. This is the reason why the synthesis and surface modification of NPs to eliminate or minimize the negative effects on cell behavior, as well as to reduce the clearance of NPs in vivo, has become one of the most interesting topics for medical applications. Recently Giri et al.29 have presented a detailed study of the composition of the protein coronas of gold nanoparticles at incubation times between 15 and 24 h. They showed that proteins adsorbed in the first 15 min have a significantly higher isoelectric point than the mean theoretical isoelectric point of all proteins present in the incubation media. Regarding the influence of the protein secondary structure on adsorption, Fleischer et al. 30 showed that BSA adsorbed onto negatively charged (carbocylate modified) and positively charged (amine) polystyrene particles. Circular dichroism spectra show no changes for BSA on negatively charged particles whereas a significant change was observed for BSA adsorbed on positively charged NPs. As a consequence, the amount of adsorbed proteins and interaction with receptors at the cell surface is different for negatively and positively charged particles.

Research on the interactions between plasma proteins and nanoparticles has been increasingly reported. Many ex situ methods have been applied to investigate the protein coronas on NPs, all showing advantages and limitations, such as centrifugation, gel filtration, or membrane-based microfiltration.31,32 Centrifugation assays have been reported as an efficient way to retrieve enough protein for safe identification of protein adsorption patterns, if conducted with care and accompanied with other methods to avoid false positive results due to insufficient washing of high abundance proteins. Moreover, sedimentation of large proteins, protein aggregates, and co-precipitation may further complicate the picture. SPION-containing nanoparticles offer the possibility to be rapidly isolated from biofluids by magnetic separation, therefore preventing contamination or degradation during the purification process and with significant improvement in the recovery yield of proteins and particles.22 A magnetic separation technique with a high gradient magnetic separation column offers in particular the greatest advantage in the simplicity of operation, in the possibility of optimizing the separation efficiency upon flow rate under reduced shear forces and in controlled elution conditions.33,34

Nonspecific protein adsorption on the surface of nanoparticles and the formation of protein coronas are widely seen as negative side effects, screening the targeting ability of functional nanoparticles towards specific receptors on cells in biological fluids in vitro or in vivo.35 On the other hand the formation of a biomolecular corona is also relevant in the cre-

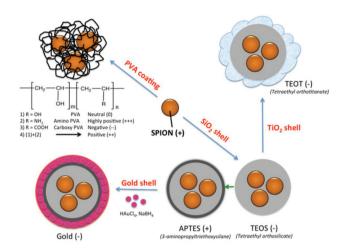


Fig. 1 Outlining of the various coatings employed on the surface of SPION

ation of bionanoconjugates that can stabilize the nanoparticle, 36 promote interactions with biological systems 37,38 or as a way to capture therapeutic biomarkers through an engineered nanoparticle-protein interface.³⁹ The advantages and disadvantages of the protein-nanoparticle interaction, with the corresponding in vitro or in vivo biological impacts, are strongly dependent on the type and amount of corona protein content. In the current study, SPION coated with PVA polymer, varying the surface charge from highly positive (amine modified), to positive (mixture of amine modified PVA copolymer and unmodified plain PVA), neutral (unmodified plain PVA), and negative (carboxylic acid modified PVA), as well as SPION coated with inorganic coatings such as silica shell using tetraethyl orthosilicate (TEOS), further modified with 3-aminopropyltriethoxysilane (APTES) and with gold layer, as well as with TiO2, were synthesized (Fig. 1). The different surface properties of the NPs were expected to influence the formation of the protein corona in fetal bovine serum (FBS) commonly used in in vitro cell culture. As it is essential to identify the protein corona that may ultimately play a biological role, a high resolution LC-MS/MS analysis was applied to allow a precise quantification of all adsorbed proteins. The aim of this study was to determine whether the different surface coatings influence the preferential binding of certain serum proteins (low and high abundance proteins) and to determine the relative abundance of the adsorbed proteins.

Results and discussion

Coated SPION characterization (sizes and zeta potentials)

The 9 different core/shell iron oxide nanoparticles were produced to investigate FBS protein adsorption. The characteristics of the produced nanoparticles are shown in Fig. 2 and Table 1. The particle diameter size of naked SPION, inorganic and metallic core-shell NPs were measured by Transmission Electron Microscopy (TEM), while the hydrodynamic diameter of PVA coated NPs and the zeta potential of all NPs were measured in deionized water by Photon Correlation Spectroscopy (PCS). The nanoparticles used in this study were all less than 150 nm in diameter. The variation in the size of highly positively charged PVA(NH2)-SPION, positively charged PVA(NH2)/(OH)-SPION, neutral PVA(OH)-SPION and negatively

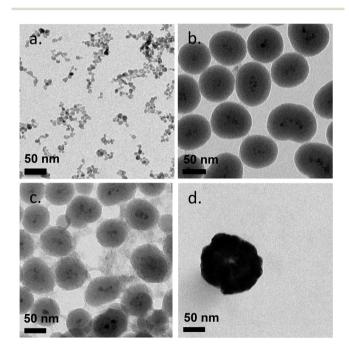


Fig. 2 Transmission electron microscopy (TEM) images of naked SPION (a), SiO₂-SPION (b), TiO₂-SPION (c) and gold-SPION (d).

Table 1 Summary of size and zeta potential of polymeric, inorganic and metallic coated SPION measured in DI water

Sample	Hydrodynamic mean diameter (by number) (nm)	TEM (nm)	Zeta potential (mV)	Point of zero charge (pzc)
Naked SPION	28 ± 3	7.7 ± 2.2	27.5 ± 3.0	7.0
PVA _(NH2) -SPION (++++)	71.8 ± 0.143	_	34.6 ± 2.1	9.0
PVA _{(NH₂)/(OH)} -SPION (+++)	46.8 ± 0.088	_	13.6 ± 1.3	8
PVA _(OH) -SPION (0)	112.7 ± 0.155	_	5.6 ± 0.9	7
PVA _(COOH) -SPION (-)	78.5 ± 0.006	_	-15.5 ± 1.0	5
SiO _{2(TEOS)} -SPION	_	102.1 ± 25.2	-36.6 ± 1.4	3
SiO _{2(APTES)} -SPION	_	100.7 ± 28.2	36.8 ± 1.9	9
TiO ₂ -SPION	_	102.1 ± 25.2	-24.9 ± 0.5	4
Gold-SPION	_	143.9 ± 18.4	-48.2 ± 1.7	6

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charged PVA(COOH)–SPION (71, 47, 113, and 79 nm respectively), suggested that the coverage of the polymer on the surface of the NPs and the differing hydration degree of the PVA layer should be considered in protein adsorption. The polydispersity index was between 0.2 and 0.25. The mean diameter of SiO_2 , TiO_2 and Au coated SPION, measured by TEM, varied from 50 to 144 nm (with the iron oxide core ~8 nm).

Coating of an initial naked SPION (zeta potential: ~27 mV) with different materials led to NPs with different surface charges. SPION coated with PVA containing amino and carboxylic acid groups showed positive and negative charges of ~13 and ~-15 mV, respectively. SPION coated with a plain PVA (-OH) showed a slightly positive charge of 6 mV and was used as representative of a neutral particle. The increasing size and the different surface charges of these 3 particles suggested successful coating of polymer onto a SPION surface when compared to a naked SPION core. The inorganic SiO2 shell nanoparticles showed negative and positive charges after coating with TEOS and APTES, respectively. The particles were spherical in shape with multiple core SPIONs (bead form). TiO2 and gold coated SPION showed strongly negatively charged surfaces with zeta potentials of -24.9 and -48 mV, respectively, confirming successful coating of these two materials onto SPION as reported in previous works. 40,41 In addition, the presence of the inorganic shell was confirmed by TEM images (Fig. 2). The SPION core is represented by dark 8 nm nanoparticles (a) inside a lighter shell of silica SiO₂ (b) and TiO₂ (c), while the gold shell was confirmed by a larger nanoparticle size with a highly electron-dense shell (d).

Nanoparticle-protein adsorption

Serum is a complex fluid that contains for human about 3700 different proteins with concentrations up to 70 mg ml⁻¹; in which the most abundant proteins such as albumin, immunoglobulin G (IgG) and α₂-macroglobulin represent 97% of the total protein content whereas the remaining 3% is a mixture composed of low abundance proteins. Identification of all the proteins is therefore a challenging task where proteomic techniques, e.g. electrophoretic methods, chromatography or mass spectrometry, can provide qualitative and quantitative analysis of protein patterns and enable the detection of potentially relevant biomarkers at very low concentrations in protein coronas when combined with nanomaterials (<1 ng cm⁻²). 42-44 Based on the available proteomic information of fetal bovine serum (FBS) reported by Zheng et al.45 the table of abundant proteins was adapted (Table 2). Fetal bovine serum proteins were therefore categorized, as a percentage relative concentration of proteins, into 3 groups: high (\geq 6%), medium (\geq 3%) and low (\leq 3%). Tables 3–5 show the list of proteins adsorbed on the polymer, inorganic and metallic coated SPION with increased specificity for nanoparticle surface, i.e. proteins common to several surface types (>2), common to 2 surface types and common to only 1 surface type, respectively. A color code was used to show the adsorbed protein given their relative abundance in FBS, e.g. red (high), green (medium) and yellow (low abundance).

Table 2 Level of protein abundance in fetal bovine serum

Level of abundance	Protein	% Relative concentration of protein
High (>6%)	Serum albumin	18.7
0 ()	Alpha-1-antiproteinase	7.9
	Plasminogen	6.4
	Cone cGMP-specific 3',5'-cyclic phosphodiesterase alpha-subunit	6.1
	Lactoperoxidase	6.1
Medium $(3\% < X \le 6\%)$	NADH-ubiquinone oxidoreductase 75	5.2
	Alpha-2-HS-glycoprotein	5.2
	Kininogen, LMW II	4.7
	Hemiferrin	4.7
	Integrin beta-1	4.7
	Prothrombin	4.4
	Apolipoprotein A-I	3.8
	Antithrombin-III	3.8
	Beta-2-glycoprotein I	3.5
	Alpha-2-antiplasmin	3.2
Low (≤3%)	Alpha-1-1-microglobulin and inter alpha-trypsin inhibitor light chain	2.9
	Hemoglobin beta fetal chain	2.9
	Alpha 1 antichymotrypsin	2.3
	Apolipoprotein A-II	1.7
	Hemoglobin R chain	1.7

Effect of nanoparticle surface charge on the protein corona

As depicted in Table 3, based on the theoretical protein isoelectric point from http://web.expasy.org/compute_pi/, no clear correlation between protein isoelectric point (IEP) and protein adsorption on nanoparticles with different surface charge could be observed. For instance, alpha-2-HS-glycoprotein (IEP 5.26) tends to have a negative charge at physiological pH 7.4, so according to electrostatic binding the protein should preferentially adsorb onto the positively charged nanoparticles. However, alpha-2-HS glycoprotein bound on all types of nanoparticles irrespective of their surface charge. The electrostatic effect alone, between protein and NPs of opposite surface charge, is therefore not the only driving force that can explain the protein adsorption. It is well known that proteins have an inhomogeneous distribution of charges at their surfaces. Therefore, even if the overall net charge of the protein is negative, positive charge domains may allow an electrostatic interaction with the particle surface. In addition, the formation of the protein corona will involve simultaneously a combination of protein-particle and protein-protein interactions. Adsorption models of coronas consisting of sequential attachment and multiple layers of proteins are also proposed, which consist of primary binding proteins on the NP surface followed by secondary binders mediated by protein-protein interactions, 46 e.g. oppositely charged anionic to cationic proteins, or by specific recognition of molecular cues on the primary protein layer. Protein conformational change and denaturation could additionally occur after adsorption onto a solid interface driven by a favorable protein-surface interaction and an due to loss of ordered gain structure. M. Rankl et al. (2006) have proven that immuno-

 Table 3
 List of protein adsorption on polymeric, inorganic and metallic coated SPION (protein bound to >2 surface types). The color indicates the relative abundance of the protein in the serum (red: high, green: medium, yellow: low and white: very low)

				Core	Organic s	Organic shell (PVA)			Inorganic shell	shell		Metallic shell	
		Protein		Positive	Highly positive	Positive	Neutral	Negative	Positive	Negative	Negative	Negative	Minho
Protein	Accession	weight (Da)	Theoretical Protein IEP	SPION	-NH ₂	-NH ₂ / OH	НО	Н000	$\begin{array}{c} SiO_2 \\ (APTES) \end{array}$	SiO_2 (TEOS)	${ m TiO}_2$	Au	of NP type
Alpha-2-HS-olvooprotein	FFTIIA	38.410	963										σ
omplement C3	CO3		6.41										0
Serum albumin	ALBU		5.82	٠									000
Alpha-1-antiproteinase	ATAT		6.05										0
Apolipoprotein A-I	APOA1		5.71										0 00
Actin. extonlasmic 2	ACTG		5.31										0 00
Actin extonlasmic 1	ACTB		5.29										000
Apolipoprotein E	APOE		5.55			T							0 00
Alpha-2-macroolohilin	A2MG		5 71										·
Complement factor B	CFAR		7.87										
Fibrinogen alpha chain	FIRA		6 73										
Pioment enithelium-derived factor	PEDE		6.57										
Hemoolohin suhunit alaha	HRA		8 07										
Drothrombin	THDE	70 506	7 0.7										. 4
Kininogen-2	KNG2		6.09										9
Alpha-2-antiplasmin	AZAP		5.45										9
Clusterin	CLUS	51114	5.73										9
Hemoglobin subunit beta	HBB		7.02										9
Hemoglobin fetal subunit beta	HBBF		6.51										9
Thrombospondin-1	TSP1		4.74										5
Inter-alpha-trypsin inhibitor heavy chain H4	ITIH4		6.22										5
Inter-alpha-trypsin inhibitor heavy chain H3	ITIH3	99 551	5.59										5
Actin, alpha skeletal muscle	ACTS		5.23										2
Actin, alpha cardiac muscle 1	ACTC		5.23										2
Actin, aortic smooth muscle	ACTA		5.24										2
Actin, gamma-enteric smooth muscle	ACTH		5.31										5
Plasminogen	PLMN	91 216	7.68										4
Myosin-10	MYH10	229 097	5.43										4
Gelsolin	GELS	80 731	5.54										4
Kininogen-1	KNG1	068 89	6.14										4
Apolipoprotein A-IV	APOA4	43 018	5.3										4
Coagulation factor V	FA5	248 981	5.53										3
Heat shock protein HSP 90-alpha	HS90A	84 731	4.92										3
78 kDa glucose-regulated protein	GRP78	72 400	5.07										3
Tetranectin	TETN	22 144	5.47										3
Apolipoprotein A-II	APOA2		7.8										3
	200021	02000	7 06			_							1

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Table 4 List of protein adsorption on polymeric, inorganic and metallic coated SPION (protein bound to 2 surface types). The color indicates the relative abundance of the protein in the serum (red: high, green: medium, yellow: low and white: very low)

Highly Protein Protein					Core	Organic s	Organic shell (PVA)			Inorganic shell	shell		Metallic shell	
Public P			Protein		Positive	Highly positive	Positive	Neutral	Negative	Positive	Negative	Negative	Negative	M
FIBB	Protein	Accession	molecular weight (Da)	Theoretical protein IEP	SPION	-NH ₂	-NH ₂ / OH		СООН	SiO_2 (APTES)	SiO ₂ (TEOS)	TiO_2	Au	of NP type
CO4 3101551 FINC 272151 CCAH 140 374 CO1A1 138 939 CO7 93 090 CO7 93 090 CO7 93 090 CO7 71 241 KLKB1 70 294 A HS71A 70 259 B HS71B 68 588 FETA 68 588 FETA 68 588 FA10 54 510 VTDB 53 342 HEMO 52 209 F12AI 51 723 CMGA 50 015 TBA4A 49 924 ain LDHB 36 724 adin LDHB 36 724 ading protein 2 IBP2 34 015 oOSTR 30 969 OOSTR 30 904 TTHY 15 227 B2MG 13 677 APOC3 10 692	Fibrinogen beta chain	FIBB	53.340	8.45							0			2
FINC 272 151 CFAH 140 374 CO1A1 138 939 CO7 93 090 TO7 93 090 TO 94 A HS71A 70 294 A HS71A 70 299 FETA 68 588 FA10 54 510 VTDB 53 342 HEMO 52 209 F12AI 51 723 CMGA 50 015 TRAA 49 924 ain LDHB 36 724 ading protein 2 IBP2 34 015 nding protein 3 IBP3 31 570 OSTR 30 969 OSTR 23 1344 TTHY 15 27 B2MG 13 677 APOC3 10 692	Complement C4 (Fragments)	CO4	3 101 551	6.68										7
CFAH 140 374 CO1A1 138 939 CO7 93 090 TOTAL 138 939 CO7 93 090 MKLKB1 70 994 A HS71A 70 229 B HS71B 70 229 FETA 68 588 FA10 54 510 VTDB 53 342 HEMO 52 209 F12AI 51 723 CMGA 50 015 TBA4A 49 924 pitor IPSP 45 297 atin LDHB 36 724 adding protein 2 IBP2 34 015 nding protein 2 IBP2 34 015 nding protein 3 IBP3 31 570 OSTR 9094 1433Z 27 745 SPP24 13 377 BEMG 13 677	Fibronectin	FINC	272 151	5.32										2
CO1A1 138 939 CO7 93 090 TOTAL 138 939 CO7 93 090 A HSP7C 71 241 KLKB1 70 994 A HS71A 70 259 B HS71B 70 259 FETA 68 588 FA10 54 510 VTDB 53 342 HEMO 52 209 FT2AI 54 209 FT2AI 54 209 TRAAA 49 924 Ditor IPSP 45 297 atin LDHB 36 724 adding protein 2 IBP2 34 015 adding protein 2 IBP2 34 015 adding protein 3 IBP3 31 570 OSTP 9069 OSTP 1433Z 27 745 SPP24 13 377 SPP24 13 377 SPP24 13 377 SPP24 13 377 APOC3 10 692	Complement factor H	CFAH	140 374	6.43										2
CO7 93 090 Protein HSP7C 71 241 A HS71A 70 259 B HS71B 70 259 B HS71B 70 259 FF10 54 510 VTDB 53 342 HEMO 52 209 F12AI 51 23 CMGA 50 015 TBAAA 49 924 Ditor IPSP 45 297 atin LDHB 36 724 uding protein 2 IBP2 34 015 uding protein 3 IBP3 31 570 OSTP 30 969 OSTP 30 969 OSTP 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677	Collagen alpha-1(I) chain	CO1A1	138 939	5.6										2
A HSP7C 71 241 KIKB1 70 994 H HS71A 70 259 B HS71B 70 229 FETA 68 588 FA10 54 200 VTDB 53 342 HEMO 52 209 F12AI 54 220 F12AI 51 723 CMGA 50 015 TBA4A 49 924 ain LDHB 36 724 ading protein 2 IBP2 34 015 ading protein 3 IBP3 31 570 OSTP 30 969 OSTP 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677	Complement component C7	CO7	93 090	6.91										2
A KLKB1 70 994 B HS71A 70 259 B FETA 68 588 FA10 54 510 VTDB 53 42 HEMO 52 209 F12AI 51 723 CMGA 50 015 TBA4A 49 924 Sitor IPSP 45 297 adin LDHB 36 724 nding protein IBP2 34 015 nding protein IBP2 34 015 nding protein IBP2 34 015 OSTK 30 969 OSTP OSTP 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	Heat shock cognate 71 kDa protein	HSP7C	71 241	5.37										2
A HS71A 70 259 B HS71B 70 229 FETA 68 588 FA10 54 510 VTDB 53 342 HEMO 52 209 F12AI 51 723 CMGA 50 015 TBA4A 49 924 sitor IPSP 45 297 adm LDHB 36 724 adding protein 2 IBP2 34 015 adding protein 3 IBP2 34 015 adding protein 3 IBP2 34 015 AGTH 30 909 OSTP 30 909 OSTP 7145 SPP24 23 134 TTHY 15 77 B2MG 13 677 APOC3 10 692	lasma kallikrein	KLKB1	70 994	8.64										2
B HS71B 70 229 FETA 68 588 FA10 54 510 VTDB 53 342 HEMO 52 209 F12AI 51 723 CMGA 50 015 TBA4A 49 924 sitor IPSP 45 297 ain LDHB 36 724 adding protein 2 IBP2 34 015 adding protein 3 IBP3 31 570 OSTY 30 969 OSTY 30 904 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	Heat shock 70 kDa protein 1A	HS71A	70 259	6.57										2
FETA 68 588 FA10 54 510 VTDB 53 342 HEMO 52 209 F12AI 51 723 CMGA 50 015 TBA4A 49 924 nitor IPSP 45 297 ain LDHB 36 724 nding protein IBP2 34 015 nding protein IBP2 34 015 OSTK 30 969 OSTY 30 904 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	Feat shock 70 kDa protein 1B	HS71B	70 229	5.67										2
FA10 54 510 VTDB 53 342 HEMO 52 209 F12AI 51 723 CMGA 50 015 TBA4A 49 924 ain LDHB 36 724 ading protein 2 IBP2 34 015 oostrk 30 969 Oostrk 30 969 Oostrk 30 969 Oostrk 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	Jpha-fetoprotein	FETA	68 588	5.92										2
VTDB 53 342 HEMO 52 209 F12AI 51 723 CMGA 50 015 TBA4A 49 924 ain LDHB 36 724 nding protein IBP2 34 015 nding protein IBP3 34 015 nding protein IBP3 30 969 OSTK 30 969 OSTP 30 904 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	loagulation factor X	FA10	54510	5.35										2
HEMO 52 209 F12AI 51723 CMGA 50 015 TBA4A 49 924 ain IDHB 36 724 ading protein 2 IBP2 34 015 ading protein 3 IBP3 31 570 OSTR 30 969 OSTP 30 904 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	itamin D-binding protein	VTDB	53 342	5.36										2
F12AI 51 723 CMGA 50 015 TBA4A 49 924 alitor IPSP 45 297 admin LDHB 36 724 ading protein 3 IBP2 34 015 ading protein 3 IBP3 31 570 OSTR 30 969 OSTR 30 969 OSTP 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	Jemopexin	HEMO	52209	7.9										2
CMGA 50 015 TBA4A 49 924 Ditor IPSP 45 297 Rain LDHB 36 724 Inding protein 2 IBP2 34 015 Inding protein 3 IBP3 30 969 OSTR 30 969 OSTR 30 904 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	actor XIIa inhibitor	F12AI	51723	6.19										2
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oitor IPSP 45 297 tain LDHB 36 724 nding protein 2 IBP2 34 015 nding protein 3 IBP3 31 570 OSTR 30 969 OSTP 30 904 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	ubulin alpha-4A chain	TBA4A	49 924	4.93										2
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nding protein 3 IBP3 31 570 OSTK 30 969 OSTP 30 904 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	nsulin-like growth factor-binding protein 2		34 015	7.13										2
OSTK 30 969 OSTP 30 904 1433Z 27 745 SPP24 23 134 TTHY 15 727 BZMG 13 677 APOC3 10 692	nsulin-like growth factor-binding protein 3		31 570	9.03										2
OSTP 30 904 1433Z 27 745 SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	Osteopontin-K		30 969	4.57										2
1433Z 27.745 SPP24 23.134 TTHY 15.727 B2MG 13.677 APOC3 10.692	Osteopontin	OSTP	30 904	4.49										2
SPP24 23 134 TTHY 15 727 B2MG 13 677 APOC3 10 692	4-3-3 protein zeta/delta	1433Z	27 745	4.73										2
TTHY 15 727 B2MG 13 677 APOC3 10 692	secreted phosphoprotein 24	SPP24	23 134	8.3										2
B2MG 13 677 APOC3 10 692	ransthyretin	TTHY	15727	5.9										2
APOC3 10 692	3eta-2-microglobulin	B2MG	13 677	7.79										2
	Apolipoprotein C-III	APOC3	10692	5.02										2

Table 5 List of protein adsorption on polymeric, inorganic and metallic coated SPION (protein bound to 1 surface types). The color indicates the relative abundance of the protein in the serum (red: high, green: medium, yellow: low and white: very low)

				Core		Organic s	Organic shell (PVA)		In	Inorganic shell	ell	Metallic shell	
		Protein		Positive	Highly positive	Positive	Neutral	Negative	Positive	Negative	Negative	Negative	N.
Protein	Accession	weight (Da)	Theoretical Protein IEP	SPION	$-NH_2$	-NH ₂ / OH	НО	Н000	SiO ₂ (APTES)	SiO ₂ (TEOS)	TiO2	Au	of NP type
Serotransferrin	TRFE	77 753	6.75										+
Antithrombin-III	ANT3	52 347	7.01										1
Beta-2-glycoprotein 1	APOH	38 252	8.53										1
Versican core protein	CSPG2	369991	4.47										1
Collagen alpha-2(I) chain	CO1A2	129 064	9.23										1
Thrombospondin-4	TSP4	105 974	4.44										1
Inter-alpha-trypsin inhibitor heavy chain H1	ITIH1	101 237	86.9										1
Neural cell adhesion molecule 1	NCAM1	93 894	4.87										₩,
Caunerin-5	CADHS	8/46/	5.5										- T
Cartilage ongoment matrix protein Heat shook 70 kDs arotein 1-like	COMP HS711	20 3 3 6 2	4.3/										- -
Moesin	MOFS	67 975	0.0										٠.
Glucosidase 2 subunit beta	GLUZB	60 151	4.36										·
4-trimethylaminobutyraldehyde dehydrogenase	AL9A1	53 977	5.84										
Carboxypeptidase N catalytic chain	CBPN	52 669	8.75										1
Integrin-linked protein kinase	ILK	51 447	8.3										1
Fibrinogen gamma-B chain	FIBG	50 244	5.53										1
Tubulin alpha-1B chain	TBA1B	50 152	4.94										1
Elongation factor 1-alpha 1	EF1A1	50141	9.1										1
Tubulin beta-5 chain	TBB5	49 671	4.78										1
Calreticulin	CALR	48 039	4.31										1
Coagulation factor IX (Fragment)	FA9	46 785	5.47										1
Fetuin-B	FETUB	42 663	5.59										₩,
Pentraxin-related protein PTX3	PTX3	42021	5.08										₩,
Na(+)/H(+) exchange regulatory coractor NHE-KF1	AMED	39 603	5.29										·
FIUCIII AMBE	LIIM	39 756	7.01										- T
Correted frigaled related protein 2	CED D2	26.724	0.75										٦.
Tronomyosin alnha-3 chain	TPM3	37 819	6.79										- -
Tronomyosin alpha-1 chain	TPM1	32.695	4 69										٠.
14-3-3 protein heta/alpha	1433B	28 081	8.4										· —
Complement factor D	CFAD	27 878	7.64										. +
Brain acid soluble protein 1	BASP1	23 011	4.53										1
Transgelin	TAGE	22 599	8.87										1
Transgelin-2	TAGL2	22 426	8.4										1
Histone H1.3	H13	22154	10.97										1
Flavin reductase (NADPH)	BLVRB	22 132	6.58										-
Ras-related protein Rap-1b	RAP1B	20 825	5.65										· -
Beta-lactoglobulin	LACB	19 883	4.93										· —
Profilin-1	PROF1	15 057	8.46										т т
Fatty acid-binding protein, liver	FABPL	14 227	7.78										1
Apolipoprotein C-II	APOC2	11 061	5.67										1
Histone H1.2 (Fragment)	H12	10365	11.01										1
Thymosin beta-4	TYB4	5053	5.02										1
Thymosin beta-10	TYB10	4805	6.18										1

globulin G (IgG) undergoes conformational changes during non-specific binding once it is exposed to different surface models.⁴⁷ Particle size, i.e. surface curvature, can also have a significant effect on the protein conformational change 48,49 and ultimately on the protein corona with variability in the surface chemistry and surface charge. Size dependent protein adsorption was for example shown on polystyrene (PS) nanoparticles of two different sizes (50 nm, 200 nm) with a more pronounced trend in amine- and carboxylic acid-containing particles compared to neutral surfaces.⁵⁰ Moreover, protein adsorption on NP surfaces also depends on the length of time of the protein-particle incubation. The adsorption/desorption process could happen via either reversible or irreversible protein conformational change during incubation.⁵¹

The protein corona appears to follow a general structure with few proteins adsorbed at high abundance and many more at low abundance although the composition in the corona does not necessarily correlate with the relative abundances of the proteins in the biological milieu. The proteins compete for the surface through a dynamic process ("Vroman Effect") based on protein abundances, affinities, and incubation time with the nanoparticles. Upon exposure of the NPs to the serum the most abundant and smaller molecular weight proteins first cover the NPs, e.g. albumin, IgG, or fibrinogen in plasma and are replaced by proteins with slower adsorption rates but higher affinity, e.g. apolipoproteins although the phenomenon is still debated for nanomaterials.⁵² Protein adsorption is a dynamic system. The adsorption/desorption process could happen via either reversible or irreversible protein conformational change during incubation⁵¹ and the protein corona is therefore hypothesized to consist of a long-lived hard shell of proteins assumed to be irreversibly bound. The corona is not immediately established but changes over time until equilibrium is reached. 53,54 Temporal studies showed that equilibrium takes place in few minutes for nanoparticles incubated with full protein serum.⁵⁵

Based on competitive binding between serum proteins, high abundance proteins were generally expected to bind to the NPs with higher probability than low abundance proteins and to all nanoparticle surfaces. Surprisingly, from the 5 listed high abundance proteins, 2 proteins were not detected on any particle surfaces, i.e. cone cGMP-specific 3',5'-cyclic phosphodiesterase alpha-subunit and lactoperoxidase. The highly abundant plasminogen binds to only 4 of the 9 investigated NPs. The surface specific binding of high abundance proteins reveals that the need of a specific surface character for protein adsorption is not ignorable. Only serum albumin and alpha-1-antiproteinase were found on 8 of the 9 different particles; both did not adsorb onto negatively charged silica coated SPION.

It is obvious that proteins preferred to bind onto positively charged NPs rather than on negatively charged NPs. Positively charged PVA coated and positively charged SiO_{2(APTES)} coated NPs showed similar patterns of adsorbed proteins indicating an influence of surface charge on protein adsorption. However, the presence of alpha-2-antiplasmin and plasminogen on highly positively charged PVA coated SPION (-NH₂), but

not on SiO_{2(APTES)}-SPION is clear evidence for the effect of the surface materials. Previous studies showed that the main proteins that associated with large particles (and hydrophobic surfaces) were albumin, IgG and antibodies, complement proteins, and apolipoproteins.^{22,56} Notably, these proteins belong to the group of high abundance plasma proteins (some being complement factors), with apolipoproteins as hydrophobic proteins. Surprisingly, none of the high abundance proteins, e.g. serum albumin, alpha-1-antiproteinase and plasminogen, bound onto negatively charged silica coated SPION (SiO_{2(TEOS)}-SPION), while these high abundance proteins (except plasminogen) bound to all PVA coated nanoparticles. One of the reasons could be that albumin (IEP 4.7-4.8) is negatively charged at physiological pH. Although TiO2 and gold coated SPION had negative charges, more than 50 proteins bound onto these particles. This indicates that the surface materials play an important role in protein adsorption. There are some proteins bound specifically to certain nanoparticles (Table 5). This knowledge would be useful in further experimental design for a biomarker study.

Effect of different surface charge of the polymer coating on protein adsorption

PVA has been widely used in biomedicine for biomedical devices and pharmaceutical applications because of suitable properties, e.g. it is hydrophilic, biocompatible, nontoxic, non carcinogenic, non immunogenic, inert in body fluids and confers biopassive properties to surfaces. Although those surfaces show reduced protein adsorption, a large number of proteins can still adsorb from full serum and vary with the polymer physicochemical properties and coating.^{57,58} The total number of proteins that bind to PVA-SPION decreased upon changing the PVA coating on SPION in the order PVA(NH2) > PVA(OH) > PVA(NH₂/OH) > PVA(COOH). Neutral to highly positively charged NPs bound a wide range of proteins and showed similar protein binding patterns whereas negatively charged PVA coated SPION bound a lower number of proteins (Table 3). Only 5 protein types bound onto the negatively charged PVA coated SPION while 24, 55, 31 and 34 proteins were found on naked SPION, highly positive, positive and neutral PVA coated SPION, respectively. Serum albumin, alpha-1-antiproteinase, alpha-2-HS-glycoprotein were high abundance proteins which bound to naked SPION and to all PVA coated SPION. In addition, complement C3, a low abundance protein also binds on all types of PVA surfaces. Serum albumin is a highly abundant protein which plays a role in opsonization and increasing nanoparticle half-life in blood.⁵⁹ Alpha-1-antiproteinase is an inhibitor of serine proteases. Its primary target is elastase, but it also has a moderate affinity for plasmin and thrombin and inhibits trypsin, chymotrypsin and plasminogen activators.⁶⁰ Alpha-2-HS glycoprotein is more abundant in fetal than in adult blood. It is involved in several functions, such as endocytosis, brain development and the formation of bone tissue. 61 Hirsch et al.³⁴ also found these three abundant proteins on positively, neutral and negatively charged PVA coated SPION. Deviation in the total amount of proteins found on all investi**Biomaterials Science**

gated PVA-SPION surfaces described in the work by Hirsch et al. could be due to different protein content fluids performed at a lower concentration of serum (10% v/v in PBS) versus full FBS serum in our study and sample preparation for MS analysis.

The protein coronas on polymer coated particles did not reflect the relative abundance of proteins in the serum. Albumin, the most abundant protein in serum, is well-detected on all the surfaces but the composition of the protein coronas also revealed enrichment in apolipoprotein versus albumin in all of the PVA coated SPION. Albumin can indeed be displaced by other proteins in serum and results in a different protein corona, e.g. around positive and negative charged particles.³⁴ Apolipoprotein A-I has furthermore been detected in the coronas of many other silica, lipid and polymeric nanoparticles, e.g. PS and polyNIPAM copolymers, 20,31,50 suggesting that lipid coating is a general feature of nanoparticles in physiological conditions.⁶² Apolipoprotein might be affected in different ways by the surface characters of the particles and may undergo in particular conformational changes on surfaces of different charges⁶³ that can explain the clear difference of protein composition between negatively charged PVA-SPION and other PVA-SPION. Multiple studies have however also reported that surface hydrophobicity played an important role in protein adsorption onto nanoparticles e.g. mediated by interaction with the lipid binding domain. Difference in the relative abundance of apolipoprotein might reflect the relative abundance of amino, alcohol, and carboxylic acid functional groups and the relative hydrophobicity/hydrophilicity of the different polymer layers. The central role of lipoproteins was also shown in the fouling of protein plasma on various polymeric biomaterials on flat surfaces.⁶⁴

The absence of kininogen on positively charged nanoparticles was already observed for amine-modified silica nanoparticles in the work from Lundqvist et al. 50 Interestingly, kininogen (high molecular weight) has been reported as a surface binding protein on iron oxide through histidine-rich sequences and is a possible marker of an incompletely masked iron oxide core as shown with loose dextran coating of ferumoxides.⁴⁶

The negatively charged PVA-SPION is the only particle type that adsorbs almost no proteins, or surprisingly specifically those high abundance proteins showing an isoelectric point (IEP) < 7 and therefore net negatively charged under physiological conditions, e.g. the pH present in 100% serum. These findings are consistent with previous reports that found that the protein coronas of negatively charged silica NPs were preferentially composed of negatively charged proteins with IEP < 7.65 Interestingly, the low protein adsorption onto negatively charged PVA-SPION is supported by a greater colloidal stability and formation of smaller aggregates than for other PVA-SPION in FBS-supplemented cell culture medium.³⁴ The reduced fouling of PVA(COOH) coated SPION suggests an enhanced stability of the PVA polymer layer, indeed mediated by carboxylate-iron coordination in addition to multiple electrostatic and hydrogen bonding interactions with the iron oxide surface. The exchange of the PVA dispersant by proteins

might therefore be reduced and the SPION core less accessible for protein binding. The decrease in total protein adsorption for carboxylated PVA was also demonstrated for PVA membranes exposed to human plasma as a result of negative surface potentials and anionic substitution on PVA.⁵⁷

Given that many abundant proteins bind to all different PVA-SPION independently of the surface charges and that the protein coronas of the negatively charged particles did not correlate with protein charge either, electrostatic interaction is likely not the only effect in modulating the protein adsorption on PVA-SPION. Difference in protein corona composition might also be induced by variation in the effective size of the particle (hydrodynamic diameter), such that it was not possible to effectively decouple in our study the net surface charge effect from size variation. 50,66 To date, very few surfaces were found to be resistant to protein adsorption onto nanoparticles exposed to full serum, since the polymer surface density and the conformation might additionally influence the pattern of adsorbed proteins. 67,68 The adsorption patterns will therefore result in a combination of physicochemical properties: in this case, the charge, size, the functional groups, molecular structures and the polymer conformation on the SPION surface.

Effect of different surface charge of inorganic and metallic coated SPION on protein adsorption

Here we investigated the adsorption of proteins on SPION coresolid shell nanoparticles, i.e. silica, titanium dioxide and gold, with special focus on the surface charge and effect of functionalities, e.g. amino, on the total amount of adsorbed proteins. Metal oxide nanoparticles were also selected as a model system as they are widely used for life science applications and are exposed to the general public in many commercial applications. Numerous high to medium abundance serum proteins were absorbed onto the oxide and metal nanoparticles.

There were 40, 19, 53 and 50 proteins bound on SiO_{2(APTES)}-SPION, SiO_{2(TEOS)}-SPION, TiO₂-SPION and gold-SPION, respectively. For SiO2 coated NPs, as observed in PVA-SPION, proteins preferentially bound onto positively charged silica coated nanoparticles rather than the negatively charged NPs, indicating the influence of the surface charge of SiO2 on protein adsorption. Alpha-2-HS-glycoprotein, apolipoprotein A-I, complement 3, fibrinogen alpha chain, cytoplasmic actin 1 & 2, apolipoprotein E, hemoglobin subunit alpha and complement factor B were proteins that were found on all investigated inorganic and metallic nanoparticles. The presence of kininogen-2 on SiO_{2(TEOS)}-SPION, but not on SiO_{2(APTES)}-SPION showed a surface charge specific binding of this protein on the silica surface. Kininogen-2 is an inhibitor of thiol proteases which plays an important role in blood coagulation, inhibiting the thrombin- and plasmin-induced aggregation of thrombocytes.⁶⁹ Protein adsorption on silica NPs studied by Monopoli et al. showed some of the same proteins as in our work, such as alpha-2-SH-glycoprotein, apolipoprotein E, apolipoprotein AI, kininogen-1 and thrombospondin-1.70 In Monopoli's work, serum albumin was observed on the particles after washing 3 times with PBS. It could be inter-

preted that serum albumin is not a loosely bound protein. In our work, however, serum albumin was eluted out after washing with high ionic strength solutions. This indicated that serum albumin is neither a loosely bound protein nor a tightly bound protein; it is an intermediate.

Interestingly, TiO2 coated and gold coated NPs which also have highly negative charges showed binding of 53 and 50 proteins, respectively. The high numbers of proteins observed on both positive and highly negatively charged particles indicated a requirement of sufficient surface charge for protein adsorption. In addition, protein adsorption markedly depends on the surface materials. Qualitatively, high and low abundance adsorbed serum proteins were generally common to all negatively and positively charged surfaces. However a lower number of proteins were common to 2 types of surfaces (Table 4) and few low abundant proteins were found on only one material surface (Table 5) suggesting that the different particle surfaces, bare metal and metal oxide, bound proteins in a more specific manner and not simply in proportion of their abundance in the serum. Although most studies investigated the adsorption of single proteins on nanoparticle surfaces, it was shown that the protein amount adsorbed at physiological pH correlated well with the zeta-potentials and IEPs of surfaces.⁷¹ For Si, Ti and Fe oxide NPs, the points of zero charge are in the order of SiO_2 (pH ~ 2) < TiO_2 (pH ~ 5) < Fe_2O_3 (pH ~ 7) also reflecting the order of acid strength of the hydroxylated surfaces, e.g. at physiological pH silica and titanium dioxide are negatively charged while maghemite is positively charged (Table 1). However, in a complex composition of FBS, adsorption ability of each inorganic particle does not correlate to the IEP and decreases in the order of TiO₂ > Fe₂O₃ > SiO₂. This was also confirmed by work of Horie et al.72 The difference of protein adsorption results from FBS and one protein system might be explained by the influence of protein-protein interaction and other biomolecule interference such as lipids on proteinnanoparticle complex formation.

Tedja et al., reported the influence of serum protein adsorption on TiO2 increasing nanoparticle stability and cellular uptake by A549 cells. TiO2 was found to be taken up inside the cell by an endocytosis mechanism.⁷³ Although in our work, the crystalline type of TiO2 was not studied. The TiO2 surface area has been reported to play a more important role than crystallinity.⁷² Gold NPs directly interact with lipid membranes, enhancing internalization of the particles⁷⁴ and revealing the possibility of gold NPs binding onto the lipid related compartment. Gold NPs are also able to induce protein conformational changes.⁷⁵ A high number of proteins bound on gold coated NPs can also be explained by preferential binding of the -SH group of proteins on the gold NPs.

Based on the results presented herein and in the recent publication from Giri²⁹ it seems that the surface charge of the particles and of the proteins could play an important role in the formation of the protein corona. Because the surface charge of proteins is inhomogeneously distributed over the surface of the protein, we have chosen the isoelectric point (IEP) as a characteristic parameter to investigate the interaction in detail. The difference between the IEP value for a protein and the pH of the media determines the charge (positive if IEP < pH, negative if IEP > pH) but is also in a first estimation proportional to the difference of charges. The same is valid for the IEP of the nanoparticles. In the case of opposed charges of proteins and NP we expect attraction as vice versa to NPs and proteins with the same sign of charge. Therefore we propose an electrostatic interaction index (ESII) which is defined as follows:

$$ESII = (IEP_{protein} - pH_{media})(IEP_{Nanoparticle} - pH_{media})$$

ESII is positive in the case of both differences being negative or positive, meaning the particles have the same sign of charge (positive or negative); ESII is negative if we have a positive and negative difference. In the first case we can expect repulsion between protein and NP in the second case attraction (this is analogous to colloidal chemistry where attraction has a negative and repulsion a positive potential). In the supplement[†], the ESII for all investigated particles is given, taking 7.4 as the pH in the media and the molecular weight of the proteins as a further parameter. The calculation of ESII allows in a further step to determine to what degree the adsorption of proteins is driven by the electrostatic interaction. When the percentage of adsorbed proteins with a negative and positive ESII per particle type taking into account all detected proteins was taken as 100%, it is evident that with increasing IEP the amount of proteins adsorbed increases (Fig. S1†). At low IEP (COOH coated particles), PVA coated and naked SPION (all with IEP < 7.4), it is not specific what has been adsorbed, meaning the electrostatic interaction is not significantly important. If the IEP > 7.4 and/or the particles are functionalized with amino groups, the electrostatic interaction is predominant and the amount of adsorbed proteins is increased. Interestingly, a very similar behavior could be observed with particles with inorganic coating. Negatively charged particles (IEP < 7.4) have a low number of adsorbed proteins and the IEP is not relevant, whereas particles with IEP near to the pH of the media show increased adsorption of proteins but this adsorption is indifferent regarding the ESII. Finally, the particles with NH2 groups at the surface clearly attract proteins with a negative charge (IEP < 7.4) and negative ESII.

Experimental

Materials section is detailed in the ESI.†

Methods

Polyvinyl alcohol coated SPION. SPION was produced by alkaline co-precipitation as described in previous works. 76,77

In order to obtain highly positively, positively, neutral and negatively charged PVA coated SPION, the different polymer solutions (i.e. 0.2% w/v M12, a mixture of Mowiol®3-85 and M12 at a mass ratio of 45, 10% w/v Mowiol®3-85 and 6% w/v KL-506 solutions: see ESI†) were mixed with 10 mg_{Fe} ml⁻¹ SPION suspension at a v/v ratio of 1. The particle suspensions

were stored for at least one week and kept at 4 °C until further use. The particle suspension was adjusted to pH 7.4 by 1 M NaOH at least one day before used.

TEOS silica and APTES-TEOS silica coated SPION. Silica coated NPs were prepared according to the Stöber sol-gel process⁷⁸ (see ESI†). The obtained particles (SiO_{2(TEOS)}-SPION) were washed twice and dispersed in DI water (final concentration approx. $0.1 \text{ mg}_{Fe} \text{ ml}^{-1}$).

The silica particles were functionalized with amines to produce positively charged silica coated SPION (see ESI†). The obtained particles (SiO_{2(APTES)}-SPION) were washed twice and dispersed in DI water (final concentration approx. 0.1 mg_{Fe} ml⁻¹). The nanoparticles were stored at 4 °C.

TiO₂ coated SPION. SiO_{2(TEOS)}-SPION were produced by hydrolysis and condensation as mentioned before. 79,80 1 ml of SiO_{2(TEOS)}-SPION was added into a 50 ml ethanol solution. Titanium(IV) tetraethoxide (TEOT) as calculated to obtain 5 nm TiO2 shell thickness was then added into a nanoparticle suspension. The reaction was allowed to proceed at room temperature for 3 h. The obtained nanoparticles were then centrifuged at 30 000g for 45 min. The pellet was washed 3 times with absolute ethanol before being resuspended in DI water and kept at 4 °C.

Gold coated SPION. Because it is difficult to precipitate a homogeneous gold layer on SPION, an indirect method was used. Inspired by the work of Rasch et al., 81 gold nanoparticles were deposited as seeds on the positively charged amino silica coated SPION (SiO_{2(APTES)}-SPION) and these seeds were grown by adding additional Au precursor.

Particle characterization by photon correlation spectroscopy (PCS). Particle sizes weighted in number were measured by dynamic laser scattering measurements carried out at 90° using a ZetaPALS (Brookhaven instruments corporation, USA) equipped with a BI-9000AT digital autocorrelator. The CONTIN method was used for data processing. Nanoparticles were suspended in distilled water at 100 µg_{Fe} ml⁻¹ and sonicated for a few seconds. The theoretical refractive index of magnetite (2.42)⁷ was used to calculate the number-weighted distribution from the raw intensity-weighted data. Viscosity, refractive index and dielectric constant of DI water were used as characteristic of the solvent.

Zeta potential measurements (ζ -potential) by ZetaPALS. Zeta potential measurements were performed by using phase analysis light scattering with the same instrument equipped with a platinum electrode. ζ-potentials were calculated from electrophoretic mobility by using theoretical models (Smoluchowski method) in ZetaPALS software packages (Brookhaven Instruments Corporation, USA) for data processing. Nanoparticles were diluted to 100 μg_{Fe} ml⁻¹ in DI water.

Size measurement by transmission electron microscopy (TEM). The size and morphology of naked SPION, silica-, gold- and TiO2-coated SPIONs were performed using a transmission electron microscope at 120 kV accelerating voltage (Philips/FEI CM12). Nanoparticle suspensions were diluted in DI water, and then sonicated for a few seconds. The diluted nanoparticle suspensions were dropped onto carbon-coated

copper grids and were allowed to dry at room temperature. TEM pictures were analyzed using image analysis software (ImageJ). The mean diameter of the nanoparticles and the thickness of the shell of more than a hundred particles were measured.

Incubation of nanoparticles with FBS. Different surface types of nanoparticles were incubated in the same batch of fetal bovine serum at a constant serum volume to particle surface ratio of 2.8 ml m⁻² as previously reported by Dawson's group in similar studies of protein corona from human serum proteins adsorbed onto nanoparticles.⁵⁰ The serum proteins are in excess to the particle surface to more likely reflect a true biological situation in body fluids.32 To reduce the interference of the dilution effect on protein adsorption, all serum-NP incubation fractions were adjusted to the same final volume (77.8% serum). Particle suspensions were incubated for 1 h at 37 °C. (Remark: due to the unidentified and highly rough surface of TiO2-SPION, the surface area of SiO2(TEOS)-SPION was used for calculation of the serum volume to particle surface ratio for TiO2-SPION).

Protein separation by a magnetic fixed bed reactor. After incubation with FBS serum the NPs were loaded into a column in a magnetic reactor. In order to elute the proteins adsorbed on the surface of the NPs, the NPs trapped on a Ni-Fe wire were washed sequentially with buffers at increased ionic strengths i.e. phosphate buffered saline (PBS) and supplemented with a high salt gradient from 0.5-2.0 M NaCl, at the flow rate of 0.5 ml min⁻¹. During each washing step, trapped nanoparticle-protein complexes were equilibrated with each elution solution for 5 min before the elution fractions were collected. Finally, the column with Ni-Fe wire was removed from the magnetic field of the reactor and the trapped NPs were collected in 1000 µl of the final elution buffer. The collected fractions are as follows: flow through, PBS1, PBS2, PBS(0.5 M NaCl), PBS(1.0 M NaCl), PBS(2.0 M NaCl) and eluted NPs. All collection fraction volumes were 1 column volume (400 µl) except for the last NP fraction (1000 µl). Tightly bound proteins on the NP fraction were then identified by MS analysis.

Semi-quantitative analysis of proteins associated with nanoparticles by nLC-MS/MS. Proteins associated with NPs were analysed following a method detailed in the ESI† and in previous works^{82,83} in order to obtain normalized Spectral Count (SpC) amounts of each protein, identified in the LC-MS/MS study of smooth and jagged surfaces, and this was calculated by applying the following equation:

$$\mathrm{NpSpC}_k = \left(\frac{(\mathrm{SpC}/M_{\mathrm{w}})_k}{\sum\limits_{i=1}^n{(\mathrm{SpC}/M_{\mathrm{w}})_i}}\right) \times 100$$

where $NpSpC_k$ is the normalized percentage of spectral count for protein k, SpC is the spectral count identified, and $M_{\rm w}$ is the molecular weight (in kDa) of the protein k. The MS analysis including the $NpSpC_k$ of tightly bound protein eluted from

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each type of nanoparticle was represented in the supplementary information (Tables S1 to S9†).

Conclusions

This study provides important information about SPIONs with different nanomaterial coating and their link with the protein composition of coronas derived from a biological fluid (i.e. serum). Also, our results shed more light on the application of SPION with a specific surface meant for the binding of a specific cluster of proteins (in vitro and in vivo) in nanomedicine and nanobiology. The results presented in this work indicated that the composition of the protein corona is very difficult to predict. Highly abundant serum proteins are not always the most abundant proteins in the nanoparticleprotein corona. Some highly abundant serum proteins are even missing in the tightly bound protein corona. Also the charge of the particles is not the only decisive factor. It is likely to be a combination of surface chemistry and charge, which determines protein adsorption. One of the reasons for the absence of correlation between the net charge of adsorbed proteins and the surface charge of the nanoparticles is the complexity of protein nature such as protein conformation, charge distribution and the hydrophobicity/hydrophilicity grade of the protein. We showed clearly that nanoparticleprotein binding strongly depends on the exact surface chemistry of the nanoparticles more than the abundancy level of protein in the biological system. The surface charges, especially on polymeric coated SPION, play a dominant role in protein adsorption. The importance of the material can clearly be seen by comparing polymeric to inorganic and metallic coatings. To quantify this observed relationship between the charge of the proteins and that of the nanoparticles at a given pH of the media, an electrostatic interaction index was introduced and applied with success to the investigated particles. The results presented in this work indicated that the corona composition could be generated for specific biomedical engineering, using NPs with specific surface properties. Moreover understanding of nanoparticle-protein adsorption could finally lead to the prediction of nanoparticle behavior in vitro.

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