Coronary Intraplaque Hemorrhage Evokes a Novel Atheroprotective Macrophage Phenotype

Intraplaque hemorrhage accelerates atherosclerosis via oxidant stress and contributes to lesion development and destabilization. Normally, macrophages scavenge hemoglobin-haptoglobin (HbHp) complexes via CD163, and this process provokes the secretion of the anti-inflammatory atheroprotective cytokine interleukin (IL)-10. We therefore tested the hypothesis that HbHp complexes may drive monocytic differentiation to an atheroprotective phenotype. Examination of the macrophage phenotype in hemorrhaged atherosclerotic plaques revealed a novel hemorrhage-associated macrophage population (HA-mac), defined by high levels of CD163, but low levels of human leukocyte antigen-DR. HA-mac contained more iron, a pro-oxidant catalyst, but paradoxically had less oxidative injury, measured by 8-oxoguanosine content. Differentiating monocytes with HbHp complexes reproduced the CD163-high human leukocyte antigen-DR-low HA-mac phenotype in vitro. These in vitro HA-mac cells cleared Hb more quickly, and consistently showed less hydrogen peroxide release, highly reactive oxygen species and oxidant stress, and increased survival. Differentiation to HA-mac was prevented by neutralizing IL-10 antibodies, indicating that IL-10 mediates an autocrine feedback mechanism in this system. Nonlinear dynamic modeling showed that an IL-10/CD163-positive feedback loop drove a discrete HA-mac lineage. Simulations further indicated an all-or-none switch to HA-mac at threshold levels of HbHp, and this conversion was experimentally verified. These data demonstrate the creation of a novel atheroprotective (HA-mac) macrophage subpopulation in response to intraplaque hemorrhage and raise the possibility that therapeutically reproducing this macrophage phenotype may be cardio-protective in cases of atherosclerosis. (Am J Pathol 2009, 174:1097–1108; DOI: 10.2353/ajpath.2009.080431)

Atherosclerotic intraplaque hemorrhage is an important contributor to lesion development and destabilization. In the carotid artery, hemorrhage promotes progression and clinical symptoms. Furthermore, coronary intraplaque hemorrhages are tightly associated with thrombosis. Mechanisms for atherogenicity of hemorrhage include the delivery of cholesterol-rich erythrocyte membranes and hemoglobin-derived iron, which can catalyze hydrogen peroxide conversion into highly reactive oxygen species (hROS). Thus a clearer understanding of macrophage responses to intraplaque hemorrhage is critical.

Hemoglobin (Hb) is principally cleared by complexing with haptoglobin (Hp), followed by uptake via CD163, a macrophage scavenger receptor. The association of Hb with Hp is normally crucial to its binding to CD163, although in diabetes there is evidence for Hp-independent CD163 binding of glycosylated Hb. The importance of Hp is strongly supported by the association of Hp genotypes with many forms of vascular disease. As a consequence of CD163 binding, HbHp induces interleukin 10 (IL10) and heme oxygenase-1 (HO-1), which are anti-inflammatory cytokines that promote atheroprotection.
inflammatory and atheroprotective. However, it is possible that CD163 functions primarily as an endocytic receptor, with down-stream signaling being induced by internalized heme. Taken together, experimental studies favor a protective role of CD163, and CD163 expression in atherosclerotic tissues may therefore represent an adaptation limiting the atherogenicity of plaque hemorrhage.

Hypercholesterolaemia-sensitive monocyte subsets (CD14^{hi}CD16^{ull} & CD14^{low}CD16^{pos}) have been described in blood. However, macrophage differentiation to separate phenotypes occurs primarily in response to tissue micro-environmental influences, which in plaques include oxidatively modified low density lipoproteins (OxLDL). Indeed, Kruth has recently published evidence that macrophage subsets defined by the presence or absence of CD14 can be found in human atherosclerotic tissue.

We tested the hypothesis that intraplaque hemorrhage results in monocyte differentiation to macrophages specialized for safe hemoglobin disposal. We show that culprit atherosclerotic plaques contain a novel anti-oxidant hemorrhage-associated macrophage subset (HA-mac) defined by high CD163 and low HLA-DR, which are distinct from pro-inflammatory lipid core macrophages. HA-mac could be generated in vitro by culturing monocytes with HbHp complexes through a mechanism centered on an IL-10 autocrine feedback loop. Our data indicate that monocytes entering atherosclerotic plaques may be adaptively modeled by plaque hemorrhage. Understanding this differentiation pathway may allow intervention to induce protective macrophage differentiation therapeutically.

**Materials and Methods**

**Pathology, Immunohistochemistry, Confocal, Image Analysis**

Human plaques were from a series of paraffin-embedded plaques that derive from consecutive West of Scotland hospital autopsies and have already been fully described. Tissues were studied with consent of next of kin for autopsy and for use of tissues for research. Local Research Ethics Committee and Central Office for Research Ethics Committees approved the research, and the tissues and site were registered under the UK Human Tissue Act. Exclusion criteria were severe sepsis or hematological malignancy.

To study the effects of hemorrhage on macrophages, we selected for study one lesion per patient with both evident hemorrhage and macrophage infiltrates on H&E sections. Consistent with the literature, these lesions were thrombosed due either to rupture (six cases) or intimal erosion (two cases). Crosschecking with clinical information indicated that the selected plaques corresponded to a tightly defined group of fatal myocardial infarction, occurring within 48 hours of event onset. As a consecutive hospital series, this cohort is very different to studies of sepsis or individual hospital autopsies and have already been fully described. Tissues were studied with consent of next of kin for autopsy and for use of tissues for research. Local Research Ethics Committee and Central Office for Research Ethics Committees approved the research, and the tissues and site were registered under the UK Human Tissue Act. Exclusion criteria were severe sepsis or hematological malignancy.

For immunostaining, sections were deparaffinized, microwaved for 20 minutes in citrate buffer (600W), blocked in serum, and peroxidase-blocked. Sections were incubated with anti-CD163 (clone 10D6, Novocastra, Newcastle on Tyne, UK), then with biotinylated anti-mouse secondary (1:200, Dako Ltd, Ely, UK) and streptavidin-alkaline phosphatase (1:100, Roche Diagnostics Ltd., Lewes, UK). Color was developed with Vector Blue (Vector Laboratories Ltd., Peterborough, UK). Sections were then incubated with anti-HLA-DR (clone CR3/43, Dako), which was detected with polymer-peroxidase kit (Menarini Diagnostics UK, Wokingham, UK) followed by color development with amino-ethyl-carmine (Vector Laboratories). CD68 was probed with antibody clone PG-M1 (Dako) and detected with polymer-peroxidase kit (Menarini) and color was developed with 3’-5’-di-aminobenzidine (DAB, Sigma-Aldrich, Poole, UK) and hematoxylin counterstaining (Pioneer Research Chemicals, Colchester, UK). For CD163/Iron dual staining, sections were probed for CD163 as above with polymer immunoperoxidase-aminocarbazole-benzidine (DAB, Sigma-Aldrich, Poole, UK) and hematoxylin counterstaining (Pioneer Research Chemicals, Colchester, UK). For CD163/Haemoglobin dual staining, sections were probed for CD163 as above with polymer immunoperoxidase-aminocarbazole-benzidine (DAB, Sigma-Aldrich, Poole, UK) and hematoxylin counterstaining (Pioneer Research Chemicals, Colchester, UK) followed by detection with polymer-peroxidase/DAB/hematoxylin as above. Negative controls were without brown staining.

Counting of CD68-positive macrophages (in total 9326) on CD163/HLA-DR dual-stained sections was performed manually. We measured the shortest straight line distance from the macrophages to the specified feature (lipid core, hemorrhage, or CD34-positive neovessels) with an eyepiece graticule. Neovessels were defined as CD34-bounded spaces of any size. Iron was assessed semiquantitatively on a standard hemochromatosis histology scale of 0 (no iron) to 3+ (maximum iron), where 3+ is equivalent to the images shown.

Quantification of HO-1 and 8-oxo-G intensity was by minor adaptation of validated image morphometry protocols, using Image J (a PC-compatible version of NIH-Image), rather than NIH-Image (Macintosh version). Digitally captured RGB images (Olympus Camedia CS050, Olympus, Watford, UK) were RGB split, after which the blue channel was subtracted from the red, producing a black and white image of brown density. This was measured by the Image J intensity histogram function to generate the arithmetic mean intensity. A region of interest was manually drawn to measure only parts of the image that included macrophages, and this was repeated for ≥5 representative images per area. Mean density units of intensity were then expressed as a continuous variable. Since normality testing by Kolmogorov-Smirnov indicated a non-Gaussian distribution, significance testing was by Mann-Whitney.

**Confocal Microscopy**

For confocal detection, we substituted fluorescent secondary antibodies for the enzyme-labeled second layers,
with staining and storage in the dark. Primary antibodies, blocking and microwaving were as above. Anti-CD163 (clone 10D6) was detected with goat-anti-mouse AlexaFluor 647 (Molecular Probes, Invitrogen, Paisley, UK; excitation [Ex.] 633 nm, emission [Em.] 650 to 700 nm); anti-HLA-DR with goat-anti-mouse AlexaFluor 568 (Molecular Probes; Ex. 568 nm, Em. 580 to 610 nm); and anti-CD68 with fluorescein-isothiocyanate (FITC)-labeled PK1 (Dako; Ex. 488 nm, Em. 500 to 535 nm). Sections were counterstained for 5 minutes in nuclear dye 7-actinomycin D (Molecular Probes) and mounted in 80% glycerol/20% PBS (both Sigma-Aldrich). We used a Zeiss LSM 510 Meta inverted confocal microscope, with three laser lines. Pinhole and unvorable filter settings were at defaults. One scan was performed using differential interference contrast to define general toponography. A further confocal scan was then performed at FITC excitation and Cy5 emission wavelengths (Ex. 488 nm, Em. 650 to 700 nm), yielding a selective image of 7-actinomycin D due to its characteristic wide Stoke’s shift. Scan and photomultiplier settings were set to optimize signal/noise ratio for each emission wavelength. Processing was with Zeiss LSM Image Browser, and comprised addition of scalebars, and adjustment of brightness and contrast, before import into Microsoft Powerpoint for assembly of montage Figures.

**In Vitro Macrophage Cultures and Reagents**

Human monocytes were isolated as described from venous blood of healthy consenting volunteers, following Local Research Ethics Committee approval. In brief, 50 ml of donor blood was collected aseptically, immediately citrated, and centrifuged to separate plasma from cells. Plasma was then recalciﬁed to produce autologous serum. Mononuclear cells were puriﬁed by centrifugation through Histopaque 1077 (10771, Sigma-Aldrich) at 500 × g for 20 minutes. In some experiments, erythrocytes were removed by dextran sedimentation (Phar- macia, GE Health care, Amersham, UK) and Percoll (Phar- macia) density gradient centrifugation. Macrophages were puriﬁed on the culture plastic by adherence incubation for 1 hour followed by three washes of warm Iscove’s Modiﬁed Dulbecco’s Medium (IMDM, Cat. No. 21056, Invitrogen). These cells had >95% purity by ﬂow cytometry for forward scatter, side scatter, and CD14 staining. Unless otherwise stated, macrophages were cultured for 7 days at approximately 105 cells per well in 24 well plates in 10% autologous donor serum, IMDM (10% AHS IMDM), with penicillin (100 IU/ml) and streptomycin (100 µg/ml). Hemoglobin (lyophilized stabilized purified Aα, ferrous hemoglobin, H0267, Sigma-Aldrich) and haptoglobin phenotype 1-1 (lyophilized puriﬁed, H0138, Sigma-Aldrich) were reconstituted in PBS at 1 mg/ml. HbHp complexes were generated by dissolving equimolar amounts of Hb and Hp in growth medium. Hb and Hp were tested for endotoxin (Limulus amebocyte lysate, sensitivity <0.125 EU/ml, LAL Pyrogen Plus kit, Cat.No. N289, Cambrex, Walkersville, MD). Approximately 1 batch in 10 of each was endotoxin-positive and excluded from further use. Hb, Hp, or HbHp complexes were added at the indicated concentrations to monocyte cultures immediately following puriﬁcation by adherence. The following antibodies, drug inhibitors, or vehicle were added at concentrations indicated 10 minutes before the HbHp complexes: antagonistic anti-CD163 monoclonal antibodies (RM3/1, Bachem AG, St Helens, UK; EdHu-1, Se- rotec, Oxford, UK); neutralizing anti-IL10 Mab 217 (R&D Systems); lipopolysaccharide (LPS, O111:B4), cytochalasin D, pepstatin-A, chloroquine (Sigma-Aldrich); and interferon gamma (IFNγ, Peprotech, London, UK). Human LDL was collected from clinical apheresis super- natant and puriﬁed by ultra centrifugation and dialysis as before (gift of D. Patel). The LDL was oxidized in 20 µmol/L CuSO4 for 24 hours at 37°C, with oxidation validated by thiobarbituric acid and agarose gel electrophoresis as before.

For flow cytometric staining, macrophages were har- vested with a disposable sterile plastic policeman. Cells were centrifuged 13,000 rpm for 60 seconds in a micro- centrifuge (Eppendorf, Fisher Scientiﬁc, Loughborough, UK), resuspended in 50 µl PBS (Sigma-Aldrich), and stained for 20 minutes at 4°C in 2 µg/ml FITC-labeled anti-CD163 (56C-FAT, Bachem). Isotype control was 2 µg/ml IgG1-FITC (Serotec). For double staining, the staining co-cultivated with anti-HLA-DR-PE (clone HK14, Sigma-Aldrich) or isotype control (mouse IgG1-PE, Sigma-Aldrich). The cells were then washed by centrifugation in 1 ml PBS, and resuspended in 400 µl PBS for ﬂow cytometry. The cells were read on a Coulter EpicsXL ﬂow cytometer, FL1–600V, FL2–600V, 1% compensation. Data were analyzed and presented using WinMDI software.

For cytokine assays, macrophages were cultured in 10% AHS-IMDM for 24 hours at approximately 105 cells per well in 96-well microplates (Nunc, VWR), in the presence or absence of 10 µg/ml HbHp complexes (approximately 10−7 mol/L complexes), 10−9 mol/L dexametha- sone, or 10 ng/ml LPS. The supernatants were then stored at −80°C before analysis. Cytokines were mea- sured by DuoSets according to manufacturer’s instruc- tions (R&D Systems, Abingdon, UK).

For the DCF oxidant stress assay, macrophages were differentiated for 7 days in 96-well microplates, in either control medium or with added HbHp complexes. Macrophages were then loaded with 10 µmol/L 5’, 6’-di-chloro- romethyl, di-hydro, di-chloro ﬂuorescein diacetate (Mole- cular Probes). Cells were then stimulated with phorbol myristate acetate (Sigma-Aldrich; 10−7 mol/L). Fluores- cence (Ex. 485 ± 10 nm, Em. 535 ± 10 nm) was mea- sured at the time points indicated in a 96 well plate reader (Synergy HT, Biotek, Fisher Scientiﬁc) at 37°C.

For peroxide assays, macrophages were differentiated and stimulated as before in 96-well plates, but in the presence of Amplex Red (1 µg/ml, Molecular Probes, Invitrogen) and horseradish peroxidase (1:1000, Sigma-Aldrich). Fluores- cence was read after 24 hours (Ex. 535 nm, Em. 590 nm).

hROS exhibit electrophilic attack on the amino-moiety of aminophenyl ﬂuorescein (APF), which rearranges to form ﬂuorescin. We validated the speciﬁcity of APF for HONO2, OH·, OCl− rather than H2O2 in our own hands. To assess speciﬁc production of hROS, macrophages were differentiated as before in 96-well plates. We then
added 5 μmol/L APF (Molecular Probes) and stimuli. After 24 hours, fluorescence was measured by microplate reader (Ex. 484 nm, Em. 515 to 550 nm) and expressed as arbitrary fluorescence units.

Supernatant Hb concentrations were measured spectrophotometrically at the characteristic peak Hb absorbance (Soret peak, A412 nm, 100 μl volume, plate reader above). Calibration was performed using Hb at known concentrations of 10^{-8}, 10^{-7}, 10^{-6} mol/L, and full absorption spectrum (400 to 700 nm in 1-nm intervals).

**Small Interfering RNA**

Adherence-purified monocytes were transferred to 0.1% AHS in IMDM without antibiotics. Complexes were prepared from 5 fmol small interfering (si)RNA (Dharmacon, Santa Cruz; scramble). These were complexed with 10 μl Genlantis liposomes (AMS Biotechnology) for 30 minutes and complexes were then incubated with monocytes for 24 hours. Western blotting was done with Novex precast system (Invitrogen). Immunodetection was in PBS with 4% nonfat milk powder 0.1% Tween-20 (Sigma). We used a Western-blot validated monoclonal antibody GHI/61 (1:1000; Santa Cruz)28; anti-mouse peroxidase (1:10,000; Dako), ECL-Plus, and hyperfilm (both Amersham).

Hb was labeled with Alexa488 (Invitrogen) following manufacturer’s instructions, and as published.13 Hb-Alexa488 was complexed with Hb and added to macrophages at culture day 1 at 10^{-7} mol/L. Cells were also assessed by RM3/1 flow cytometry and the effects of RM3/1 on Hb phagocytosis was assessed.

**Mathematical Modeling Using Nonlinear Dynamics**

Standard modeling approaches were used.29–31 The mathematical model was formed from a coupled pair of nonlinear ordinary differential equations describing the feedback loop (see supplemental Figure S1 at http://ajp.amipathol.org) [equations (1) & (2)]. The model has two variables [CD163] and [IL10] that describe the time course of the concentrations of the two species. The rate of change of each species is determined by the production and degradation of each. We therefore obtain:

\[
\frac{d[CD163]}{dt} = CD163 \text{ Production Rate} - CD163 \text{ Degradation Rate} = \frac{[IL10][R]}{[IL10] + k_{i1}} - k_{d1}[CD163]
\]  

(1)

\[
\frac{d[IL10]}{dt} = IL10 \text{ Production Rate} - IL10 \text{ Degradation Rate} = \frac{[CD163][HbHp]}{[HbHp] + k_{i2}} - k_{d2}[IL10]
\]  

(2)

where [HbHp] is a parameter indicating the HbHp concentration (in arbitrary units) and [R] is a parameter that describes the sensitivity of CD163 up-regulation to IL10 levels. One possible biological interpretation is that the first term of Equation 1 represents the concentration of IL10 receptors available for CD163 binding on the cell surface. Degradation is modeled using simple exponential decay with rates k_{d1} and k_{d2}. The parameters k_{i1} and k_{i2} determine the saturation of the production rates of IL10 and CD-163 respectively.

These were solved numerically and graphed using Maple 11 (Maplesoft, Waterloo, Ontario, Canada). Since this was a proof-of-concept model, it was kept as simple as possible and simulations used arbitrary units, and illustrative values were assigned to parameters. An advantage to formulating a simplified model is that the steady state values can be derived analytically and it is possible to verify that these arbitrary choices do not affect conclusions. Equilibrium solutions and threshold values were obtained by algebraic rearrangement and graphed in Maple 11. To determine the robustness of the observed dichotomy, we ran simulations using a range of arbitrary parameters. These all predicted qualitatively similar (ie, dichotomous) behavior around a threshold. To validate the model, we then experimentally tested the behavior it predicted.

**Statistical Analysis**

The inverse relationship between CD163^{high} and HLA-DR^{high} macrophages in coronary plaques was examined by both χ^{2} analysis and by analysis of variance (ANOVA) with Bonferroni’s correction. Normality was tested by Kolmogorov-Smirnov and normally distributed data were tested parametrically. The data in Table 1 appeared approximately normally distributed, but normality testing was equivocal due to low n-values. They were therefore expressed parametrically as mean ± SE but tested using a rank method (Mann-Whitney), which is a more stringent assessment. Where indicated, statistical testing of parametric continuous data were performed using ANOVA with Bonferroni correction for multiple simultaneous comparisons, or a two-sample Student’s t-test. Statistical analysis was performed using Excel (Microsoft, Seattle, WA) or SigmaStat (Systat, San Jose, CA). Exact P values are presented to clarify the strength of the statistical differences.

**Results**

**Culprit Human Lesions Contain a Novel Hemorrhage-Associated Macrophage Subpopulation Identified by CD163**

We examined the effect of atherosclerotic plaque hemorrhage on macrophage phenotype, assessing 351 plaques from 83 consecutive autopsies.18,19 Eight had confluent intraplaque hemorrhage and macrophage infiltrates. Each of these was a thrombosed atherosclerotic lesion
with a large hemorrhage and numerous macrophages (Figure 1A, and (see supplemental Figure S2 at http://ajp.amjpathol.org). All eight corresponded to fatal myocardial infarction (5 males, 3 females, ages 60 to 80 years, death within 48 hours of the onset of clinical event).

CD68 immunostaining identified fibrous cap and lipid core macrophages (Figure 1B). Two-color immunostaining with CD163 and HLA-DR revealed that these were either predominantly CD163<sup>high</sup>HLA-DR<sup>low</sup> or CD163<sup>low</sup>HLA-DR<sup>high</sup> (Figure 1, C and D). Double immunostaining for CD163 and the erythrocyte marker glycophorin-c indicated that the CD163<sup>high</sup> macrophages colocalized with plaque hemorrhage (Figure 1E). CD163<sup>low</sup>HLA-DR<sup>high</sup> cells were α-smooth muscle actin negative (Figure 1F). Findings were remarkably consistent in all eight thrombosed hemorrhaged plaques: 75 ± 2.4% macrophages were CD163<sup>low</sup>HLA-DR<sup>high</sup> and 25 ± 2.9% were CD163<sup>high</sup>HLA-DR<sup>low</sup> (Figure 1G). We identified minimal numbers of CD68<sup>pos</sup>CD163<sup>low</sup>HLA-DR<sup>low</sup> cells (Figure 1G). While CD163<sup>high</sup>HLA-DR<sup>low</sup> macrophages were consistently found in culprit lesions (see supplemental Figure S3 at http://ajp.amjpathol.org) twenty consecutive stable plaques from the same study contained only pro-inflammatory CD163<sup>pos</sup>HLA-DR<sup>high</sup> macrophages (not shown).

We then assessed whether the CD163<sup>high</sup>HLA-DR<sup>low</sup> and CD163<sup>low</sup>HLA-DR<sup>high</sup> cells were definitively CD68<sup>pos</sup> macrophages. We used confocal immunofluorescence to analyze all three markers simultaneously and relate them to lesion topography (see supplemental Figure S4 at http://ajp.amjpathol.org). Two collections of CD68<sup>pos</sup> macrophages are shown. One subset is mixed with hemorrhage, i.e., extravasated erythrocytes, and is characterized by high CD163 and a low HLA-DR (i.e., CD68<sup>pos</sup>CD163<sup>high</sup>HLA-DR<sup>low</sup>). In contrast, macrophages more distant to hemorrhage were conventional foam cells (FC), with an abundant foamy cytoplasm and high HLA-DR, but low CD163 (CD68<sup>pos</sup>CD163<sup>low</sup>HLA-DR<sup>high</sup>). These observations therefore define a novel subset of CD68<sup>pos</sup>CD163<sup>high</sup>HLA-DR<sup>low</sup> HA-mac.

### HA-mac Are Protected from Oxidant Stress Despite Iron Loading

We next characterized the possible pathophysiological role of HA-mac by assessing oxidant stress with antibodies specific for oxygen-adducted guanosine (8oxoG). While FC macrophages contained 8oxoG, as reported, HA-mac in the same sections were 8oxoG-negative, indicating reduced oxidative stress (Figure 2A). Reduced HA-mac oxidative stress was in the face of increased loading with iron (a pro-oxidant catalyst), evidenced by CD163/Perl’s double-staining (Figure 2A). In serial sections, HA- macrophages reciprocally expressed HO-1 strongly and myeloperoxidase weakly (Figure 2A). FC macrophages had the reverse pattern (Figure 2A). CD163<sup>high</sup> macrophages were found adjacent to extravasated erythrocytes and intraplaque microvessels and co-expressed IL10 (Figure 2, B–E).

These associations were tested statistically. We measured distances of the two macrophage populations, HA-mac (CD163<sup>high</sup>HLA-DR<sup>low</sup>) and FC-mac (CD163<sup>low</sup>HLA-DR<sup>high</sup>). Distances measured from nearest distance to lipid core, nearest distance to microvessels, nearest distance to hemorrhage, cellular iron (where Figure 2D = 3+), HO-1 staining intensity, 8-oxo-G staining intensity.

### Table 1. Morphometry of Foam Cell and Hemorrhage-Associated Macrophage Subsets

<table>
<thead>
<tr>
<th>Subset</th>
<th>Distance to lipid core (µm)</th>
<th>Distance to neovessels (µm)</th>
<th>Distance to hemorrhage (µm)</th>
<th>Iron score (0–3 scale)</th>
<th>HO-1 intensity (arbitrary density units)</th>
<th>8-oxo-G intensity (arbitrary density units)</th>
<th>IL-10 intensity (arbitrary density units)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HA-mac (CD163&lt;sup&gt;high&lt;/sup&gt;HLA-DR&lt;sup&gt;low&lt;/sup&gt;)</td>
<td>439 ± 14 (0 to 1000)&lt;sup&gt;**&lt;/sup&gt;</td>
<td>15.7 ± 5.7 (10 to 50)&lt;sup&gt;***&lt;/sup&gt;</td>
<td>3.88 ± 1.8 (0 to 10)&lt;sup&gt;**&lt;/sup&gt;</td>
<td>2.25 ± 0.47 (1+ to 3+)&lt;sup&gt;*&lt;/sup&gt;</td>
<td>22.2 ± 3.9&lt;sup&gt;***&lt;/sup&gt;</td>
<td>14.0 ± 0.96</td>
<td>12.3 ± 1.5&lt;sup&gt;****&lt;/sup&gt;</td>
</tr>
<tr>
<td>FC-mac (CD163&lt;sup&gt;low&lt;/sup&gt;HLA-DR&lt;sup&gt;high&lt;/sup&gt;)</td>
<td>29 ± 15.5 (1 to 100)</td>
<td>785 ± 214 (500 to 2000)</td>
<td>150 ± 86 (50 to 750)</td>
<td>0 ± 0 (0 to 0)</td>
<td>7.95 ± 1.03</td>
<td>35.8 ± 4.7</td>
<td>4.2 ± 0.4</td>
</tr>
</tbody>
</table>

* P < 0.05, Mann-Whitney.
** P < 0.01, Mann-Whitney.
*** P < 0.005, Mann-Whitney.
**** P < 0.001, Mann-Whitney.

Morphometric comparisons of the two macrophage subsets. Rows, data gathered in the same way (see Materials and Methods) for HA-mac (CD163<sup>high</sup>HLA-DR<sup>low</sup>) and FC-mac (CD163<sup>low</sup>HLA-DR<sup>high</sup>). Columns, measurement of respectively: nearest distance to lipid core, nearest distance to microvessels, nearest distance to hemorrhage, cellular iron (where Figure 2D = 3+), HO-1 staining intensity, 8-oxo-G staining intensity.

### Differentiation of Human Monocytes with HbHp Complexes Reproduces HA-mac

Using flow cytometric analysis of peripheral blood monocytes, we demonstrated that approximately 90% were...
CD14<sup>high</sup>CD16<sup>null</sup> and 10% were CD14<sup>low</sup>CD16<sup>pos</sup> in keeping with previous reports. CD16<sup>3</sup> expression was restricted to monocytes, and there was no detectable difference in CD16<sup>3</sup> expression between the CD14<sup>low</sup>CD16<sup>pos</sup> and CD14<sup>high</sup>CD16<sup>null</sup> subsets (see supplemental Figure S5 at http://ajp.amjpathol.org).

Whether HbHp complexes induce the CD16<sup>3</sup>HLA-DR<sup>low</sup> phenotype was tested by culturing human monocytes for 8 days in the presence of Hb, Hp or HbHp complexes (Figure 3). HbHp complexes (10<sup>-7</sup> mol/L), but neither constituent protein alone induced a CD16<sup>3</sup>HLA-DR<sup>low</sup> phenotype (Figure 3, A–D). This effect was blocked separately by two antagonistic anti-
CD163 antibodies (clones RM3/1, EdHu-1), or by neutralizing anti-IL10 antibodies, but not by isotype-matched neutralizing anti-tumor necrosis factor (TNF)-α antibodies (Figure 3, E–H). These observations indicated dependence on CD163 and autocrine IL10 for CD163highHLA-DRlow polarization. This polarization was also blocked by neutralizing anti-TNF and anti-IL10 antibodies, but not by isotype-matched antibodies to TNF-α and IL10 respectively. LPS = E. coli lipopolysaccharide. HbHp complexes were used at 100 nmol/L (10⁻⁷ mol/L).

Key: A: Unstimulated culture of day 8 macrophages. B: Incubated with HbHp complexes at 10⁻⁷ mol/L, 100 nmol/L. C: Incubated with Hb at 10⁻⁷ mol/L, 100 nmol/L. D: Incubated with Hp at 10⁻⁷ mol/L, 100 nmol/L. E: As (B), with addition of anti-CD163 clone RM3/1, 70 nmol/L. F: As (B), with addition of anti-CD163 clone EdHu-1, 70 nmol/L. G: As (B), with addition of anti-IL10 Mab217, 70 nmol/L. H: As (B), with addition of anti-TNF-α Mab225, 70 nmol/L. I: As (B), with addition of 10⁻⁷ M/L cytochalasin-D in DMSO, 1,1000. J: As (B), with addition of 10⁻⁷ M/L chloroquine in DMSO, 1,1000. K: As (B), with addition of 7 mol/L pepstatin-A in DMSO, 1,1000. L: As (B), with addition of 10⁻⁷ M/L pepstatin-A in DMSO, 1,1000. M: As (B), with addition of LPS, 1 ng/ml. N: Control (unstimulated) culture of macrophages at day 8. O: Incubated with 5 × 10⁻⁷ mol/L (50 pmol/L) IL10 from outset of culture. P: As (O), but with 5 × 10⁻⁸ mol/L (0.5 pmol/L) IL10. Q: As (O), but with 5 × 10⁻⁹ mol/L (500 pmol/L) IL10. R: As (O), but with 5 × 10⁻¹⁰ mol/L (50 pmol/L) IL10. S: Unstimulated culture of macrophages at day 8. T: Incubated with 10⁻⁸ mol/L (1 nmol/L) dexamethasone from outset of culture. U: As (T), but with 10⁻⁹ mol/L (10 nmol/L) dexamethasone. V: As (T), but with 10⁻¹⁰ mol/L (100 nmol/L) dexamethasone. W: As (T), but with 10⁻¹¹ mol/L (1 μmol/L) dexamethasone.

HA-mac Are Anti-Inflammatory and Antioxidant, and Promote HbHp Clearance in Vitro

We then investigated the anti-inflammatory and antioxidant phenotype of the in vitro differentiated HA-mac (Figure 3, N–R). Dexamethasone (10⁻⁹ mol/L to 10⁻⁷ mol/L; 1 to 100 nmol/L), a prototypic anti-inflammatory agent, increased CD163 and reduced HLA-DR, but not to the extent seen with HbHp complexes (Figure 3, S–W). Hb freshly prepared by hypotonic lysis of autologous erythrocytes had the same effect as commercial Hb (see supplemental Figure S6 at http://ajp.amjpathol.org). Although HbHp complexes maximally induced CD163 at day 4, they needed to be added at culture outset for full induction, suggesting the programming of macrophage differentiation (see supplemental Figure 7 at http://ajp.amjpathol.org).
Figure 4. Homeostatic phenotype of HA-mac A–C: x-axes, respective stimuli, control medium (10% AHS IMDM); LPS = E. Coli lipopolysaccharide (10 ng/ml); HBHp = HbHp complexes (100 mM/L; 10^6 mol/L). D: y-axis, Absolute IL10 levels in supernatants in monocyte-macrophage cultures 24 hours after addition of stimuli at the start of culture. (*P < 0.001 analysis of variance overall, with P < 0.05 relative to control, Bonferroni adjusted post-test, values are mean ± SEM of five donors). E: y-axis, Absolute TNF-α levels in supernatants in monocyte-macrophage cultures 24 hours after addition of stimuli at the start of culture. (*P = 0.006 analysis of variance overall with P < 0.05 relative to control, Bonferroni adjusted post-test, values are mean ± SEM of five donors). F: y-axis, Ratio of IL10/TNF in panels A and B. (*P = 0.0001 analysis of variance overall with P < 0.05 relative to control with Bonferroni adjusted post-test, n = 5 donors). Concentration-effect curves (not shown) indicated that the direction of IL10/TNF ratio was maintained at all concentrations of LPS (1 to 1000 ng/ml) and HBHp (10^-6 to 10^-16 mol/L). G: y-axis, Macrophage survival as measured by reduction of the colorimetric formazan dye MTS. Viability was measured relative to control cultures (100%). Control = unmodified 10% AHS IMDM, HBHp = 10^-6 mol/L HBHp complexes added at the start of culture. Student's t-test *P = 0.000143, values are mean ± SEM of five donors). H: y-axis, Hb concentration in medium supernatant at 4 minutes after addition. x-axis, time after addition of HBHp complexes (10^-6 mol/L), measuring supernatant [Hbt] spectrophotometrically at 412 nm. Open circles, macrophages were incubated in control medium (10% AHS IMDM) for 8 days. Filled triangles, macrophages were differentiated with HBHp complexes (10^-6 mol/L) for 8 days. Student's t-test *P = 0.000283. I: y-axis, specific H2O2 production as measured by Amplex Red/peroxidase (see Materials and Methods), calibrated by a H2O2 standard curve. Open bars, Control – differentiation in 10% AHS IMDM for 8 days. Filled bars, HBHp – differentiation in added HBHp complexes (10^-6 mol/L). x-axis, addition of opsonised zymosan, a prototypical macrophage oxidative burst stimulant, in two doses to control or HBHp differentiated macrophages. The dose of OpZ, is expressed as particles/well. Data are representative of five independent experiments using separate donors. Student's t-test *P = 0.0017. J: y-axis, H2O2 production as measured by Amplex Red/peroxidase, calibrated by a H2O2 standard curve. x-axis, addition of OxLDL (50 μg/mL) to control or HBHp differentiated (10^-6 mol/L) macrophages. Open bars, Control – differentiation in 10% AHS IMDM for 8 days. Filled bars, HBHp – differentiation in added HBHp complexes (10^-6 mol/L). Data are representative of five independent experiments using separate donors. Student's t-test *P = 0.034. K: y-axis, oxidative stress measured as di-hydro, di-chloro-fluorescein diacetate fluorescence in the absence of serum (P < 0.005, Figure 4H). H2O2 is converted to hROS (specifically HONO, OH, OCI^-), which are selectively detected with APF (see Materials and Methods). APF fluorescence indicated that HA-mac complexes reduced hROS in response to OxLDL or serum deprivation (Figure 4I) or opsonised zymosan (not shown). Time-course and concentration-effect curves for these stimuli are given in supplemental Figure S9 at http://ajp.amjpathol.org. All of the protective effects of HBHp were concentration-related and optimal at 10^-7 mol/L (see supplemental Figure S10 at http://ajp.amjpathol.org). The protective response was reversed at the highest HBHp concentration, a feature we attribute to overload (the cultures were opaque red).
CD163-siRNA Specifically Reduces Hb Uptake, RM3/1 Binding

CD163-specific siRNA duplexes suppressed CD163, evidenced by Western blotting using antibody GHI/61 (see supplemental Figure S11A at http://ajp.amjpathol.org). This corresponded to reduced RM3/1 surface binding—even on a log scale (see supplemental Figure S11B at http://ajp.amjpathol.org). We measured uptake of AlexaFluor488-labeled HbHp complexes over 60 minutes after addition to macrophage cultures (Supplemental Figure S11C at http://ajp.amjpathol.org). CD163-siRNA suppressed uptake of Alexa488-labeled HbHp complexes (see supplemental Figure S11D at http://ajp.amjpathol.org). RM3/1 also reduced Hb uptake, proving functional antagonistic activity (see supplemental Figure S11E at http://ajp.amjpathol.org). Finally incubating the cells for 4 days with unlabeled HbHp complexes in the presence or absence of scramble or CD163-siRNA, showed that macrophage survival was reduced by CD163-knockdown; see supplemental Figure S11F at http://ajp.amjpathol.org).

Nonlinear Dynamics and Experimental Validation Show that HbHp/IL10/CD163 Positive Feedback Evokes All-or-None Commitment to CD163high versus CD163low Phenotypes at a Threshold Concentration of HbHp Complexes

We observed that HbHp binding to CD163 leads to IL10 secretion, and that IL10 up-regulates CD163. Since their output promotes their input, positive feedback loops are associated with exponential phenomena, as in the action potential or in pathophysiological vicious cycles. In contrast, positive feedback loops have been recently implicated in stable commitment to divergent states in T-helper cells.34 Here, the positive feedback loop enhances subset distinctiveness by abolishing intermediate states. More generally, similar regulatory architectures are being observed in a variety of systems biology contexts.29

We tested whether, in principle, a positive feedback loop between IL10 and CD163 in response to HbHp would generate discrete subsets governed by HbHp levels. We therefore constructed a proof of concept model. This is shown diagrammatically (see supplemental Figure S1 at http://ajp.amjpathol.org). We converted the diagram to a mathematical model to test whether a threshold level of HbHp switched the system between two lineages, deliberately keeping the model as simple as possible rather than providing a complete quantitative description of the regulatory system. The model uses arbitrary variables and arbitrary parameters to probe a qualitative pattern of cell behavior.

In multiple simulations of equations (1) and (2) (see Materials and Methods), CD163 fell to negligible levels at low concentrations of HbHp, irrespective of starting concentrations (Figure 5A, blue curves). On the other hand, at HbHp concentrations above threshold, CD163 rose to a stable plateau with the level of final CD163 independent of initial CD163 (Figure 5A, red curves). Thus the model predicts that CD163-IL10 positive feedback leads to binary CD163high or CD163low phenotypes depending on HbHp input.

The model was then used to predict behavior of the final steady state levels of CD163 and IL10 with varying [HbHp] (Equations 3 and 4). Steady states occur when the concentrations of CD163 and IL10 do not change with time. These can be obtained by setting the left hand sides of the above equations to 0 and solving for [CD163] and [IL10]. This can be done algebraically. One set of solutions is simply [CD163]* = 0, [IL10]* = 0, corresponding to the low CD163 state. It is perhaps unrealistic that this is exactly at 0. However, if the state was a small non-zero level it would not affect any of the subsequent qualitative conclusions (eg, by adding a small basal production rate to the above equations). We have chosen not to do this as it complicates the model, and in particular, makes it impossible to obtain analytic expressions for the other steady state and the threshold in HbHp. With the equations as given, straightforward manipulation gives the other steady state as

\[
[CD163]^* = \frac{[HbHp]([R] - k_1k_2k_3k_5)}{k_4[HbHp]}
\]  

(3)

\[
[IL10]^* = \frac{[HbHp][R] - k_4k_5k_6 - k_1k_2k_3k_5}{k_5[HbHp] + k_6}
\]  

(4)

Only non-negative values of protein levels are of biological relevance. For the above expressions to satisfy this constraint we require [R] > k_4 k_5 k_6 and [HbHp] ≥ H* where H* is the threshold given by

\[
H^* = \frac{k_1k_2k_3k_5}{[R] - k_4k_5k_6}
\]  

(5)

At [HbHp] = H* the two steady states intersect in the well known transcritical bifurcation, exchanging stability.35 Further analysis shows that for [HbHp] < H*, the 0 steady state is stable and all solutions converge to it. At [HbHp] = H* the 0 steady state loses stability. For [HbHp] > H* it is the steady state given by Equations 3 and 4. Therefore, H* is the threshold level of Hb Hp. H* forms the boundary between the two CD163 phenotypes. With [HbHp] below this value, CD163 levels drop to a steady state of negligible levels, irrespective of starting conditions (eg, blue curves in Figure 5A). This is interpreted as the CD163low-HLA-DRhigh subset. In contrast, for [HbHp] above H* above the H* threshold, CD163 rises to a stable plateau, again irrespective of initial value. This corresponds to the red curves in Figure 5A and the CD163high-HLA-DRlow subset. This threshold is shown in Figure 5B. The final steady state level of CD163 rises rapidly as HbHp concentration reaches the threshold, before saturating at a fixed level. The simulations were run using a wide range of arbitrary parameters, and consistently gave these qualitative effects.
Finally, we tested this prediction. We showed by flow cytometry that CD163 increased in an all-or-none manner at a threshold concentration of $10^{-9}$mol/L (Figure 5C). It should be noted that the conceptual model uses arbitrary parameters, and does not predict the precise threshold. Moreover, the model was not designed to explore the effects of frank HbHp overload. Evidence from plaques showing expression of IL-10 by HA-mac was also consistent with the predictions of the model (Table 1, Figure 2).

**Discussion**

Intraplaque hemorrhage\(^1,2\) and plaque macrophages\(^36,37\) play key roles in atherosclerotic lesion progression and vulnerability. We do not challenge this view, but instead define a novel macrophage subset HA-mac. These cells are identified within hemorrhaged inflamed lesions, and have several atheroprotective and compensatory properties. Our data provide a logical extension of previous work on CD163, and add to conceptual understanding of macrophage subsets in relation to human vascular disease.\(^10,13,38\)

Our in vitro data support that plaque hemorrhage adaptively modifies macrophage differentiation. Here, hemoglobin directs a discrete subset with increased hemoglobin uptake, reduced oxidant stress and generation of the anti-inflammatory cytokine IL10. This interpretation is biologically plausible, as it is an everyday observation that bruising is not itself inflammatory. However, since our starting material was postmortem specimens, the phenomenon is self-evidently not fully atheroprotective. The HA-mac were a minority (approximately 25%), and conventional foamy macrophages predominated even in the plaques with large hemorrhages. Moreover, the known up-regulation of HLA-DR by modified lipoproteins may accentuate the difference between the two subsets (FC versus HA).\(^39\) Consistent with this, we have found that CD163\(^{high}\) macrophages are absent in lesions with lipid core but no hemorrhage.

By defining a distinct macrophage subset, our data extend earlier descriptions that HO-1 and CD163 are found in atherosclerosis.\(^13\) Similarly, a substantial body of work, mainly by Levy and colleagues, indicates that haptoglobin genotype Hp2-2 is less protective than Hp1-1 in a wide range of vascular diseases.\(^10,40\) We now give a mechanism contributing to these observations by defining how plaque hemorrhage drives a specific antioxidant hemorrhage-associated macrophage subset.

The concept of functionally distinct leukocyte subsets is well established for lymphocytes in vitro and in vivo (eg, CD4 vs CD8, Th1 vs Th2).\(^36\) Macrophage subsets have been defined by in vitro differentiation with cytokines.\(^38\) Thus, classically activated macrophages are generated by pro-atherogenic stimuli eg, IFN-γ, TNF-α, and LPS, and alternatively-activated macrophages are induced by IL-4,\(^38,41,42\) Loss of the key Th1 transcription factor T-bet suppresses macrophage HLADR in atherosclerosis in vivo, via reduction of CD4 T-helper derived IFN-γ and increased IL-4.\(^43\) This is consistent with a skew from classical to alternative activation, suggesting that classically activated plaque macrophages are pro-atherogenic. However, HA-mac were characterized in the present study as more closely corresponding to the de-
activated macrophage subset induced by IL10 or glucocorticoids.\textsuperscript{38} Interestingly, although dexamethasone up-regulated CD163 and suppressed HLA-DR, we found that its effect was weak relative to that of HbHp.\textsuperscript{14,44} This may be in part related to different signaling pathways, since dexamethasone signals primarily via the glucocorticoid receptor, while IL10 signals via silencer of cytokine signaling proteins, themselves therapeutic targets.\textsuperscript{44,45} Understanding different effects on macrophage function of glucocorticoids and HbHp therefore requires further study.

Well-established roles of macrophages in atherosclerosis include foam cell formation, lipid necrotic core formation, matrix metalloproteinase production, oxidant production, and inflammatory cytokine secretion.\textsuperscript{36,37,46} However recent evidence describes active cholesterol export from the plaque via macrophage lymphotoysis migration,\textsuperscript{47} or via the adenosine triphosphate binding cassette pathway to high-density lipoprotein.\textsuperscript{48} This necessitates a more complex paradigm encompassing homeostatic macrophage roles. Our observations add to this by describing a non-lipid-driven macrophage homeostatic mechanism.

We are not aware of a previous mathematical description of a bistable system driving macrophage differentiation, either in vascular disease or other situations. The nonlinear dynamic modeling defined a role for the IL10 autocrine positive feedback loop. The key feature of the model is the stability produced by the loop’s saturability. At its simplest, this may equate to saturation of promoter regions of specific gene regulatory elements with transcription factors. The molecular nature of these effects is beyond the scope of the present paper but under active study. Critically, this feedback accurately predicted the existence of a threshold HbHp concentration for CD163 induction. Thus we experimentally validated the model at the qualitative level of all-or-none commitment to respective subsets. The system features positive feedback, threshold, and an all-or-none response, which promote rapid decisive commitment. This permits the vascular wall to respond rapidly and consistently to sudden pro-oxidant threats such as plaque hemorrhage. We found experimentally that 6 to 8 days differentiation in 1 µmol/L HbHp concentrations reversed the protective phenotype, a feature consistent with toxicity from Hb overload. At this concentration, the cultures are macroscopically red and the cells brown. Thus this protective system can be overwhelmed, leading to maladaptive function.

In conclusion, we have characterized a novel atheroprotective macrophage subset (HA-mac) associated with intraluminal hemorrhage. This subset contrasts strongly with pro-atherogenic conventional (classical) lipid core macrophages. HA-mac are adapted to swiftly clear Hb and contain oxidative stress. We predict that HA-mac would suppress the impact of hemorrhage on atherosclerosis progression. In culprit lesions, the effects of HA-mac are by definition too little and too late, but therapeutic modification of HA-mac pathways may prevent plaque destabilisation.

Acknowledgments

We thank Viola Leung, Sandra Cantilena, and Donna Horncastle for technical support, and Dr. Justin Mason for critical review of the manuscript.

References
