Direct link between cytokine activity and a catalytic site for macrophage migration inhibitory factor

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Macrophage migration inhibitory factor (MIF) is a secreted protein that activates macrophages, neutrophils and T cells, and is implicated in sepsis, adult respiratory distress syndrome and rheumatoid arthritis. The mechanism of MIF function, however, is unknown. The three-dimensional structure of MIF is unlike that of any other cytokine, but bears striking resemblance to three microbial enzymes, two of which possess an N-terminal proline that serves as a catalytic base. Human MIF also possesses an N-terminal proline (Pro-1) that is invariant among all known homologues. Multiple sequence alignment of these MIF homologues reveals additional invariant residues that span the entire polypeptide but are in close proximity to the N-terminal proline in the folded protein. We find that p-hydroxyphenylpyruvate, a catalytic substrate of MIF, binds to the N-terminal region and interacts with Pro-1. Mutation of Pro-1 to a glycine substantially reduces the catalytic and cytokine activity of MIF. We suggest that the underlying biological activity of MIF may be based on an enzymatic reaction. The identification of the active site should facilitate the development of structure-based inhibitors.

Keywords: cytokine/enzyme/macrophage migration inhibitory factor/NMR/protein structure

Introduction
Cytokines elicit biological responses by activating specific cell surface receptors. Macrophage migration inhibitory factor (MIF) was initially identified as a secretion product of activated T cells and was one of the first cytokines to be discovered (Bloom and Bennett, 1966; David, 1966). MIF has a long association with delayed-type hypersensitivity reactions, but recent studies with the recombinant protein and its antibodies have identified additional biological functions (Bernhagen et al., 1996). Glucocorticoids induce the secretion of MIF which acts to override glucocorticoid inhibition of cytokine (TNF-α, IL-1β, IL-6 and IL-8) production in macrophages and T cells (Calandra et al., 1995). The activation of T cells by mitogenic or antigenic stimuli is also partly regulated by MIF (Bacher et al., 1996). Glycosylation-inhibition factor (GIF), a cytokine that is involved in the regulation of IgE synthesis, has been found to have an identical sequence to MIF (Mikayama et al., 1993). The pleiotropic in vitro activities on immune cells and involvement of MIF in the pathophysiology of a number of pro-inflammatory diseases are consistent with a role for MIF as a cytokine.

While MIF shares many similarities with cytokines, some dissimilarities have also been noted. MIF is pre-formed in cells and lacks a signal sequence, in contrast to most cytokines which are produced in response to stimuli and secreted using the signal sequence pathway. MIF is also different from cytokines in that it catalyzes chemical reactions. MIF converts d-2-carboxymethylester-2,3-dihydroindole-5,6-quinone to 5,6-dihydroxindole-2-carboxymethyl ester. This non-physiological activity, which has been termed d-dopachrome tautomerase, was discovered fortuitously during the study of melanin biosynthesis (Rosengren et al., 1996). In an attempt to identify natural ligands for MIF, the keto–enol isomerizations of p-hydroxyphenylpyruvate (HPP) and phenylpyruvate were discovered to be catalyzed by MIF (Rosengren et al., 1997). The separate localization of these substrates from MIF as well as the kinetic parameters for the tautomization reactions suggests that these molecules also are unlikely to be physiological substrates for MIF.

The three-dimensional structure of MIF is unlike that of any known cytokine. The MIF monomer possesses two antiparallel α-helices that pack against a β-sheet in a motif that is somewhat reminiscent of MHC and dimeric α-chemokines; however, the topology of MIF is totally different from these proteins. Three monomers associate to form a trimer in all of the crystal forms of MIF that have been characterized (Kato et al., 1996; Sun et al., 1996a; Suzuki et al., 1996). Interestingly, MIF bears striking structural similarity to three microbial enzymes: 4-oxalocrotonate tautomerase (OT), 5-carboxymethyl-2-hydroxymuconate isomerase (CHMI), and chorismate mutase (Chook et al., 1994; Subramanya et al., 1996). These three enzymes are homotrimers (OT is a trimer of a homodimer) that possess active sites in close proximity to subunit interfaces. The similarity of MIF to OT and CHMI extends to at least one active site residue. OT and CHMI utilize an N-terminal proline as a catalytic base in their respective isomerization reactions (Whitman et al., 1991; Hajipour et al., 1993). MIF also possesses an N-terminal proline, the function of which has not been examined. We used sequence analysis, structural studies and mutational analysis to probe the function of Pro-1 and investigate the mechanism of MIF activity. Our studies correlate catalysis by MIF with neutrophil activation and suggest that an enzymatic reaction may underlie immune activation.

Results
Sequence analysis identifies the potential biological site
The significance of the unexpected structural and active site similarity of MIF with OT and CHMI was first
investigated by sequence analysis. The sequence databases were searched with ENTREZ and BLAST to identify all MIF homologues. Pairwise identities among the 11 identified sequences range from 22 to 98%. Each of these sequences have a proline following the initiating methionine. Direct N-terminal sequencing of native MIF from two species (bovine and murine) reveals that the initiating methionine is removed in both cases, resulting in a protein with a proline at the N-terminus (Galat et al., 1993; Bernhagen et al., 1994). Due to the unique geometric restraints imposed by a proline, its prevalence at the N-terminus might be determined by the efficiency of removal of the initiating methionine in a Met-Pro sequence rather than by any functional requirement for an N-terminal proline. To investigate the statistical significance of proline at the N-terminus, the N-terminal residue of all proteins in the SWISSPROT database was analyzed. The majority (~80%) of proteins retain the initiating methionine. Of the
remaining proteins, proline is the fifth most prevalent amino acid at the N-terminus. If normalized by the number of codons for proline, the presence of a proline is slightly below average, but certainly not rare.

To investigate further the importance of the N-terminal region of MIF in its biological activity, multiple sequence alignment of the 11 MIF homologues was used for identifying additional invariant residues (Figure 1A). Only 11 residues (10% of the sequence) are invariant among all of the homologues, which represent a broad range of the evolutionary spectrum. These invariant residues can be expected to serve important functional or structural roles. Display of these residues on the three-dimensional structure of MIF shows that most of these residues cluster around the N-terminal proline and illustrate the evolutionary pressure to preserve this site (Figure 1B). Moreover, a solvent accessibility analysis reveals that many of these residues are exposed to solvent with the N-terminus sitting at the base of a pocket formed by the invariant residues. The clustering of invariant, solvent-exposed residues to form a pocket containing potential functional groups resembles an enzymatic active site.

**The N-terminus of MIF has an unusually low pK_a**

For the N-terminal amine of a protein to serve as a catalytic base, the pK_a of the amine must be at or below the physiological pH, otherwise the amine will be positively charged without a lone pair of electrons available for proton abstraction. Titration of proline amide indicates that the secondary amine of a proline has a pK_a > 9 (Stivers et al., 1996). To determine whether the pK_a of the N-terminus of MIF is consistent with a role as a catalytic base, the titration of Pro-1 was monitored directly by 15N NMR. The Pro-1 15N chemical shift appears between 43 and 52 p.p.m. and is well resolved from other 15N resonances of MIF in the pH range 4.4–8.5 (Figure 2A). At this pH range, MIF is folded as determined by circular dichroism and NMR measurements (data not shown). Non-linear least squares of the equation

\[
\delta_n = \frac{\delta_1 + \delta_2(10^{pH-pK_a})^n}{(10^{pH-pK_a})^n + 1}
\]

where \( n \) is the Hill coefficient and \( \delta_1 \) and \( \delta_2 \) are the limiting chemical shifts at low and high pH, respectively, was used to determine a pK_a of 5.6 ± 0.1 for Pro-1 (Figure 2B). This value is almost four pH units less than the pK_a of proline amide (Stivers et al., 1996). The environment of Pro-1 was examined to determine the source of the unusual pK_a. The secondary amine of Pro-1 in the 1.9 Å X-ray structure of human MIF is at the base of a cleft surrounded predominantly by carbon atoms (not shown), suggesting that the low pK_a of the N-terminus is due to its location in a hydrophobic environment that lacks a neutralizing negative counter-charge. Calculation of an electrostatic potential map reveals an alternative explanation for the unusually low pK_a. A region of positive potential (Figure 3) is present at the N-terminus, with Lys66 and the invariant residue Lys32 providing the source of this positive potential. Determination of the relative contribution of the hydrophobic residues adjacent to the N-terminal proline versus the positive potential provided by Lys32 and Lys66 will require detailed biophysical measurements and mutational analysis. Nonetheless, the environment at Pro-1 lowers the pK_a of the N-terminus such that the amine is unprotonated at physiological pH.

**The catalytic site**

The unusually low pK_a for the N-terminal amine, the presence of invariant residues at and around this site and the global structural similarity with two microbial enzymes led to the hypothesis that the N-terminal region may function as an active site. This hypothesis is consistent with the ability of MIF to catalyze isomerization reactions. While d-dopachrome (2-carboxy-2,3-dihydroindole-5,6-quinone), phenylpyruvate and HPP are unlikely to be the physiological substrates for a MIF-catalyzed reaction, these molecules provide suitable probes for examining whether the N-terminal region of MIF is involved in catalysis. Unfortunately, d-dopachrome is unstable for
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Fig. 3. Electrostatic potential map of the MIF trimer (left) and monomer (right). The map was calculated using DELPHI (MSI, San Diego, CA) with unit charges for the side chains of arginine, aspartic acid, glutamic acid and lysine, +0.5 charge for histidine side chains, and no charge for the terminal amine and carboxylate. The dielectric constant for the interior and exterior of the protein was 2 and 80, respectively. Negatively charged surfaces are in red, positively charged surfaces in blue. To distinguish the N-terminal proline from other residues, the color of its surface is changed from blue to green.

structural studies, and efforts to characterize crystallographically the interactions of phenylpyruvate or HPP have been unsuccessful due to the inability to obtain co-crystals. As these three substrates have been shown to compete with each other for binding to MIF, HPP was used to identify the active site using NMR spectroscopy. The initial experiment probed the chemical environment of the N-terminal amine in the presence of HPP. The $^{15}$N chemical shift of Pro-1 was measured as a function of varying concentrations of HPP. Figure 4A indicates that the peak shifts and broadens in the presence of 8 mM HPP (4 MIF equivalents) and disappears at 16 mM HPP. The severe broadening observed for this peak is suggestive of conformational exchange in the presence of HPP. This is at least partly due to the presence of both the keto and enol forms of HPP in the active site. $^1$H-$^{15}$N HSQC spectra of MIF (Muhlhan et al., 1996) in the presence of HPP identify a much larger set of residues with changes in chemical shift. The resonance peaks for Ile4, Ala38 and Ile64 are not detectable in the $^1$H-$^{15}$N HSQC spectrum at 10 mM HPP. Other residues with perturbed $^1$H or $^{15}$N chemical shifts include Phe3, Val39, Gly50, Lys66, Asn102, Gly107, Trp108, Phe113 and Ala114 (Figure 4B). While it is not possible to discriminate between chemical shifts that change due to ligand binding and those that are altered due to conformational changes, mapping of residues with chemical shifts greater than one standard deviation on the trimer of MIF illustrate that they cluster around Pro-1 (Figure 4C). Moreover, these results provide evidence that the catalytically active form of MIF is defined by multiple subunits as two residues that cluster around Pro-1 arise from a different subunit.

**Correlation between neutrophil priming and catalytic activity**

To probe further the role of Pro-1 in the catalytic and biological activity of MIF, a single site-directed mutant was generated in which proline is substituted by glycine (P1G). The P1G mutant was expressed and purified as previously described for wild-type MIF (Sun et al., 1996b). The folding of P1G as assessed by circular dichroism and $^1$H NMR spectra is identical to that of wild-type MIF (data not shown). However, P1G does not have any measurable catalytic activity with L-dopachrome methyl ester (a substrate similar to D-dopachrome) and HPP during the timescale of the assay and the concentrations tested (Figure 5A and B). Wild-type MIF displays a $K_m$ of 143 ± 23 μM and 1.3 ± 0.45 mM for each substrate, respectively.

To determine whether elimination of catalytic activity affected biological activity, a neutrophil priming assay measuring superoxide generation was used. Wild-type MIF displays an ED$_{50}$ of 8.6 ± 3.3 ng/ml (0.7 ± 0.3 nM) in this assay with a maximal increase in superoxide production that ranges between 20 and 30% (Figure 6A). In contrast, superoxide secretion by P1G is significantly
Fig. 4. (A) $^{15}$N NMR spectrum of Pro-1 in the presence of varying concentrations of HPP. The chemical shift and broadening effects observed indicate a perturbation in the electronic environment of Pro-1 as a result of HPP binding. (B) $^1$H-$^{15}$N HSQC spectra of human MIF in the absence (black) and in the presence (red) of 10 mM HPP. Residues with perturbations greater than the standard deviation for all residues are labeled. (C) Solvent-accessible surface of the human MIF trimer is shown with the monomeric subunits colored blue, yellow and green, and Pro-1 from each subunit colored pink. Residues which were shown to have perturbations >1 standard deviation as a result of HPP binding are colored red. As depicted, these perturbations reveal a binding site that encompasses the N-terminal proline. As seen in the view to the right (~150° rotation of the left view), much of the protein’s surface is unaffected by the binding of HPP.

reduced at 1 $\mu$g/ml MIF (Figure 6B). These data indicate that Pro-1 is involved in both the catalytic and biological activity of MIF.

Discussion

The experiments in this study do not address whether catalysis by MIF is essential for cytokine activity, but the importance of Pro-1 in catalysis and neutrophil activation suggests that an enzymatic reaction may underlie the biological activities of MIF. This suggestion is consistent with previously published results utilizing anti-MIF antibodies. In both in vivo and in vitro assays anti-MIF antibodies inhibit responses that are not induced by the sole addition of MIF (Bernhagen et al., 1993; Bacher et al., 1996). For example, mice treated with anti-MIF antibodies and a lethal dose of lipopolysaccharide (LPS) survive endotoxemia (Bernhagen et al., 1993). This parallels results obtained with antibodies to other cytokines shown to be involved in sepsis (Tracey et al., 1987; McNamara et al., 1993). However, in contrast to the effects of TNF-α (Tracey et al., 1986) and IL-1 (Okusawa et al., 1988) which induce the sequelae of sepsis in animals, injection of MIF does not cause sepsis. A similar
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Fig. 5. Catalytic activities of wild-type and mutant (P1G) MIF. (A) D-dopachrome tautomerase activity. The absorbance is plotted against the concentration of 2-carboxymethylester-2,3-dihydroindole-5,6-quinone for 100 nM wild-type (■) and P1G (○) MIF. (B) HPP tautomerase activity. The absorbance of enol HPP is plotted against the concentration of keto HPP for 10 nM wild-type (■) and P1G (○) MIF.

A paradox involving MIF and its antibodies has been reported in T-cell activation (Bacher *et al.*, 1996). Anti-MIF antibodies inhibit proliferation of T cells, yet MIF does not appear to be a T-cell mitogen. One explanation for these incongruous results is that MIF biological activity requires a cofactor. Antibodies to MIF may therefore inhibit biological activity, but to induce biological activity the cofactor and MIF are required. The suggestion that an enzymatic activity underlies MIF biological activity indicates that the essential cofactor may be a substrate for a chemical reaction.

The neutrophil priming activity of MIF is low in comparison with molecules such as lipopolysaccharide, which can prime neutrophils to secrete levels of superoxide 20-fold higher than unprimed cells (Guthrie *et al.*, 1984). The lack of a robust response in the neutrophil priming assay is also consistent with a requirement of a substrate (or cofactor) in MIF biological activity. The decreased priming ability of MIF may be due to the presence of low quantities of the physiological substrate for MIF’s catalytic activity under the conditions of the experiment. This possibility can be addressed once the physiological substrate of MIF is identified.

Although MIF can catalyze keto–enol isomerization reactions of HPP and phenylpyruvate, it is unlikely that these molecules are the natural ligands for MIF. Phenylpyruvate and HPP are generated intracellularly during the biosynthesis of phenylalanine and tyrosine, respectively, and are therefore unable to serve as substrates for extracellular MIF. The serum concentrations of these molecules (Deutsch, 1997) are 1000-fold lower than the $K_m$ values for MIF, which is also inconsistent with a role as extracellular substrates for MIF. Whether MIF has an intracellular function involving these substrates remains to be investigated.

While the P1G mutant has undetectable D-dopachrome tautomerase and HPP activities under the conditions of the catalytic assays, a low neutrophil priming response is observed. There are at least two potential explanations for this low residual biological activity. First, although the P1G mutant has no measurable catalytic activity using the non-physiological substrates, the mutant protein may have some residual catalytic activity with the unidentified physiological substrate that MIF evolved to utilize. The second possibility is that the P1G mutant may have observable catalytic activity with the non-physiological substrates at the longer timescales required for the neutro-

Fig. 6. Biological activities of wild-type and mutant (P1G) MIF. (A) Dose–response curve of MIF neutrophil priming activity. Human neutrophils were primed with MIF for 60 min, stimulated with fMLP for 30 min, and the concentrations of superoxide measured. Data are expressed as mean ± SEM obtained with neutrophils from four separate donors. (B) Production of superoxide from human neutrophils upon stimulation by wild-type MIF and P1G. Human neutrophils were primed with 1 μg/ml MIF or P1G under the same conditions as in (A). The superoxide generated after P1G priming is ~75% lower than with wild-type MIF. Data are expressed as mean ± SEM obtained with neutrophils from three separate donors.
phil priming assay. Both instances would result in the observed residual neutrophil priming activity of the mutant protein.

It is interesting to note that MIF is not the only protein that possesses both catalytic and cytokine activities. Thrombin (Vu et al., 1991), cyclophilin (Sherry et al., 1992; Xu et al., 1992), FK506 binding protein (Leiva and Lyttle, 1992), adenosine deaminase (Kameoka et al., 1993), NAP-2 (Hoogewerf et al., 1995) and neuroleukin (Chaput et al., 1988) are members of an expanding group of proteins with enzymatic and cytokine functions. Of these proteins, the mechanism of cytokine activity of only thrombin has been defined. Thrombin cleaves a proteolytic site at the N-terminal extracellular domain of its G protein-coupled receptor to unmask a tethered ligand that induces self-activation (Vu et al., 1991). Whether other proteins with dual functions utilize their enzymatic active site to induce cytokine activity has not been addressed. The results presented in this study demonstrate that the catalytic site of MIF overlaps with the site responsible for neutrophil activation. The physiological substrate and chemical reaction catalyzed by MIF, and the possible role of this reaction in the pro-inflammatory activities of MIF remain to be determined.

Materials and methods

Sequence analysis

The BLAST and ENTREZ programs at the National Center for Biotechnology Information were used to identify MIF homologues. Multiple sequence alignment of these sequences was performed with PILEUP from the GCG package (Devereux et al., 1984).

Expression and purification of recombinant MIF, P1G mutant MIF and [15N]MIF

Recombinant MIF was expressed and purified as previously described (Sun et al., 1996b). The P1G mutant cDNA was generated by PCR using the primer 5'-GGATCCATATGATGCATATTCATTGAATTTG-3' that encodes an NdeI restriction site (in bold), and a codon for glycine (in italics) instead of a proline after the initiating methionine. The second primer 5'-TTGGATCCTTAGGGCAAGGTGGAGTT-3' annealed to the 3' end of the wild-type cDNA and contained a BamHI restriction site (in bold). These restriction sites were used to clone the PCR product into pET-11b expression vector and purification of the mutant protein. For biochemical assays, unlabeled MIF and P1G were made LPS-free as previously published (Bernhagen et al., 1994) and assayed for LPS using the E-Toxate kit (Sigma).

[15N]MIF was prepared in minimal media containing 47.9 mM Na2HPO4, 22 mM KH2PO4, 8.6 mM NaCl, 7.5 mM (15N)2H2SO4, 22.2 mM glucose, 1.99 mM MgSO4·7H2O, 100 μM CaCl2·2H2O, 26 μM FeCl3·6H2O, 60 μM thiamin hydrochloride, 82.5 μM thymidine, 15N-labeled Celtone (Martek, Columbia, MD), 2.5 μM biotin and 100 μg/ml ampicillin. Briefly, an overnight culture of pET-11b MIF in supplemented HBSS with the following acquisition parameters: spectrometer frequency, 600.139 MHz; spectral width, 5388 Hz; acquisition time, 95 ms; recycle delay, 1 s; transients per-block, 32; complex blocks, 128; temperature, 298°K. To facilitate sample locking, 5% D2O was added to all samples.

X-ray studies

Crystals of MIF in space group P3121 with unit cell dimensions a = b = 96.70 Å, c = 106.14 Å were obtained as described previously (Sun et al., 1996b). X-ray data were collected at room temperature as 2° frames on a Rigaku R-Axis IIe image-plate detector and a Rigaku RU200 rotating anode generator equipped with mirror optics. The data collected to a resolution of 1.9 Å were processed using BIOTEX (Molecular Structure Corp., Woodlands, TX) and consisted of 40 317 reflections from a total of 130 276 observations (F(2)/F(2) > 1.0) with an Rmerge of 8.9%. Simulated annealing was first used to refine the 2.6 Å model to an R-factor of 23.4% for data from 5.0 to 2.2 Å. Rearranged B-factor refinement followed by several rounds of positional refinement and rebuilding resulted in an R-factor of 23% for data from 5.0 to 1.9 Å. Simulated annealing omit maps and PROCHECK (Laskowski et al., 1993) were used to analyze the model. Manual adjustments, including the addition of water molecules, were made as necessary. The final model was subjected to individual B-factor least-squares refinement. The free R-factor calculation (Brunger, 1992) was included throughout to monitor the refinement. The final structure of trinodal MIF has a free R-factor of 26.4% and an overall R-factor of 20.9% at 1.9 Å resolution with r.m.s. deviations of 0.009 Å and 1.6° for bond lengths and bond angles, respectively.

For calculation of the electrostatic potential, arginine, aspartic acid, glutamic acid and lysine side chains were assigned unit charges, histidine side chains were assigned a charge of +0.5, and the terminal amine and carboxylate of the polypeptide were left uncharged. The interior and exterior dielectric constants were 2 and 80, respectively. All calculations were carried out using DELPHI (MSI, San Diego, CA).

Biological and catalytic activities

Neutrophils were prepared from freshly drawn blood anticoagulated with heparin. Blood was diluted with physiological saline for irrigation, layered over lymphocyte separation medium (Organon Teknika), and subjected to centrifugation at 400 g for 30 min. The supernatant containing lymphocytes and serum was aspirated and the red blood cell pellet diluted with an equal volume of 3% dextran and allowed to separate until there was a defined interface. The supernatant containing neutrophils was washed, supplemented with Hanks balanced salt solution (HBSS) containing 0.2% dextrose, Ca2+ and Mg2+, and resuspended by hypotonic lysis. Neutrophils were pelleted by centrifugation, resuspended in supplemented HBSS with 1% human serum added and used immediately for experiments. Neutrophils were primed by incubating 2.5×106 for 1 h at 37°C in supplemented HBSS/1% serum containing varying concentrations of recombinant MIF. LPS contamination for both wild-type MIF and P1G was <0.15 pg/μg of protein. Neutrophils were pelleted by centrifugation, resuspended in 120 μM cytochrome c in supplemented HBSS, treated with 10-5 M MLF, and incubated at 37°C for 30 min. Superoxide-specific reduction of cytochrome c was determined spectrophotometrically using cells incubated with or without 50 μg/ml superoxide dismutase.

For n-dopamine tautomerase assays, l-dopamine methyl ester was prepared by mixing 4 mM l-3,4-dihydroxyphenylalanine methyl ester with 8 mM sodium periodate for 5 min at room temperature and then placed directly on ice for 20 min before use. Activity was determined at 25°C by adding dopamahem methyl ester to a cuvette containing 100 nM MIF or P1G in 25 mM potassium phosphate buffer pH 6, 0.5 mM EDTA and measuring the decrease in absorbance at 475 nm on a Hewlett Packard 4542 diode array spectrophotometer.

For HPP tautomerase assays, HPP was dissolved in 50 mM ammonium acetate pH 6.0, allowed to equilibrate at room temperature overnight, and stored at 4°C. Activity was determined at 25°C by adding HPP to a quartz cuvette containing 10 nM MIF or P1G, 0.435 M boric acid, pH 6.2, and measuring the increase in absorbance at 330 nm spectrophotometrically.

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