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COMMUNICATIONS

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Preliminary Refinement and Structural Analysis of the Fab Fragment from Human Immunoglobulin New at 2.0 Å Resolution*

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The three-dimensional structure of the Fab fragment from human myeloma IgG New has been refined using "model building" and "real space" procedures. By these techniques, the correlation between the amino acid sequences and the 2.0 Å resolution multiple isomorphous replacement Fourier map has been optimized. The average shift of all atoms during real space refinement was 0.62 Å. A list of the refined atomic coordinates for the 440 amino acid residues in the structure is given. Ramachandran plots prepared using the refined coordinates show a distribution of ϕ, ψ angular values which corresponds to the predominant β-pleated sheet conformation present in the structure.

The structures of the homology subunits V_H, V_L, C_H1, and C_L were superimposed by pairs and quantitatively compared. The closest similarities were observed between V_H and V_L and between C_H1 and C_L. Amino acid sequence alignments obtained from this structural superposition are given. The closest sequence homology in Fab New is observed between C_H1 (γ heavy chain) and C_L (λ light chain). In addition, there is considerable homology between the variable and constant regions.

The distances of close contacts between the homology subunits of Fab New have been determined. The closer contacts, those between atoms at a distance ≤ 1.2 times their van der Waals radii, are analyzed in relation to the constant, variable, and hypervariable nature of the immunoglobulin sequence positions at which they occur. Most of the residues which determine the closer contacts between V_H and V_L and between C_H1 and C_L are structurally homologous and highly conserved or conservatively replaced in immunoglobulin sequences.

The relation between idiotypic determinants, antigen combining site and hypervariable regions, is discussed in terms of the refined model.

In this paper we present the results of a preliminary crystallographic refinement and a list of atomic coordinates of Fab New.[1] Some features of the refined structure are discussed in relation to the genetic control and physiological function of immunoglobulins.

It is generally accepted (see review in Ref. 1) that electron density maps calculated by multiple isomorphous replacement techniques contain significant errors which may lead to imprecise determination of structural details such as the location of amino acid side chain atoms, bond angles, ϕ and ψ values, *cis* or *trans* character of proline residues, etc. This refinement project was undertaken with the aim of obtaining more accurate coordinates which can be applied to structural studies of other immunoglobulins and Fab·hapten complexes (2). The starting atomic coordinates were those of the structure previously described (3, 4) obtained using multiple isomorphous heavy atom replacements. Since the complete refinement of the structure of Fab New is a complex undertaking, we present here initial results obtained after application of two consecutive refinement techniques. In the first step we have applied a "model building" procedure (5) in which the measured atomic coordinates were adjusted to impose standard bond lengths and bond angles. In a second step a "real space" procedure (6, 7) was used to optimize the correlation between the Fab New model and the multiple isomorphous replacement, electron density Fourier map.

The coordinates obtained by these procedures have provided an improved model which has been used to compare the tertiary structure of the homology subunits, to calculate interatomic contacts that define the quaternary structure of Fab, and to re-examine the conformation of the combining site.

METHODS

Measurement of Model Coordinates—Atomic coordinates were measured on the 2 Å (nominal) resolution model previously described (4). A two-pointer device was used for this purpose: a horizontal pointer (50 inches long) was brought to touch atom centers by displacing the device on the base of the model, adjusting the height of the pointer along a graduated scale (z coordinate) while a second, parallel, fixed pointer of equal length gave the x, y coordinates on a grid at the base of the model. The use of a level and leveling screws at the base of the two-pointer device was essential for obtaining reproducible coordinates. While these measurements were made, the image of the model and the atom centers were

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[1] The abbreviations used for immunoglobulins, their polypeptide chains, and fragments are as recommended in (1964) *Bull. W H O* 30, 447.

projected on the corresponding sections of the Fourier map using an optical comparator (8) to verify that their location and the coordinate values corresponded with the Fourier map.

Model Building Procedure—The set of measured coordinates for the 3185 non-hydrogen atoms in the structure provided the starting point for this procedure. In general, these coordinates are subject to errors due to measurement uncertainty and to mechanical deformations of the skeletal brass model. Consequently, the mathematical model building procedure of Diamond (5) programmed for a digital computer was used in order to impose standard bond lengths and bond angles in the model. The measured coordinates were used to provide a guide point for each (non-hydrogen) atom in the structure. Some conditions used in this part of the refinement process are given in Table I. All the varied angles are dihedral; the $\tau_\alpha(\tau_{N\text{-}C\alpha\text{-}C})$ angle was allowed to vary since this condition gave a much closer correlation with input coordinates without introducing large distortions in the idealized, model-built geometry of the molecule. Residues for which the model-built coordinates differed considerably from the input coordinates or which had an abnormal τ_α value were remeasured and checked for correspondence with the Fourier map. These discrepancies could always be traced to errors in the measurement of coordinates or to distortions of the brass model. When necessary the coordinates were measured again after rebuilding distorted regions and the remeasured coordinates were submitted to the model building procedure as guide points. The final average value of $\Delta\tau_\alpha(\Delta\tau_\alpha = |\tau_\alpha - 109.3°|)$ for the 440 residues in the model-built structure was 5.43°. Pro 151 in C_H1 was built and refined in the *cis* conformation. The root mean square shift from the initial coordinates for all non-hydrogen atoms in the structure was 0.2 Å.

Real Space Refinement—The model-built coordinates were used as a starting set for real space refinement (6). The 2.0 Å electron density map used for the automatic fitting of the atomic coordinates was calculated using multiple isomorphous replacement phases as described before (3, 4). The electron density function was calculated at intervals of 1/160 along x, (a = 111.43 Å), 1/80 along y (b = 56.68 Å), and 1/130 along z (c = 90.30 Å) in sections of constant y. A computer program incorporating a fast-Fourier transform algorithm was used for this calculation. Five cycles of real space refinement were carried out using the conditions defined in Table II. Values of atomic radii for carbon, nitrogen, oxygen, and sulfur that gave fastest convergence in trials using a small part of the structure were adopted and kept constant during refinement. Progress in the refinement process was followed by inspection of the root mean square shifts in coordinates, shifts in the coordinates of individual atoms, and adjustments of amino acid scale factors. Unusually large shifts in coordinates were checked using the optical comparator and the Fourier map. All refinement calculations were carried out using computer programs implemented at the Brookhaven National Laboratories.

RESULTS AND DISCUSSION

The root mean square shifts of atomic coordinates after five cycles of real space refinement converged to an average value of 0.09 Å. The values after each cycle were: 0.46 Å, 0.20 Å, 0.13 Å, 0.10 Å, and 0.09 Å. The average shift of all atoms during real space refinement was 0.62 Å. In most parts of the molecule, the coordinates after refinement are in very good agreement with the features of the electron density map (Fig. 1). Many carbonyl groups of the main polypeptide chain can be reliably positioned (Fig. 2). In general, coordinates of atoms in regions of strong, well defined electron density converged rapidly in the first cycle of refinement and moved very little in subsequent cycles. Atoms in regions of lower density converged more slowly. A few residues in poorly defined regions of low density showed little convergence, although their total shift from the starting coordinates was small. The movement of main chain atoms tended to be smaller than those of side chains, presumably due to their better defined electron density and to greater constraints on their positions.

The progress of refinement was checked after each cycle by inspecting the fit of atoms whose shifts were substantially greater than the average. Some side chain groups which had been shifted by the Diamond model building procedure were moved back to their original positions by real space refinement. In the fifth cycle of refinement, an average shift greater than 1 Å occurred for three consecutive amino acid residues (Gly 166, Val 167, and His 168) in the C_H1 region. Inspection of the position of these atoms showed that they had moved to a conformation that appears to be in better agreement with the electron density map than the original model. The coordinates for all atoms of Fab New after refinement, given in the "Appendix," are filed with the Protein Data Bank at Brookhaven National Laboratory. No major features of the map remain unexplained, although a number of possible solvent molecules are found on the surface of the molecule. The conventional R factor,[2] based on F_c obtained with the coordinates in "Appendix" and an overall temperature factor[3] (B = 18.0), is R = 0.46. This value is reasonable given the refinement approximations outlined in Table II and the fact that no solvent atoms were included in the model. Further refinement using observed structure factors and calculated phases is currently underway.

The S—S distances in the five Fab disulfide bonds were allowed to vary without constraints. At the end of the refinement these distances were found to be: V_H, 2.00 Å; V_L, 2.46 Å; C_H1, 2.30 Å; C_L, 2.30 Å; C_H1-C_L, 2.43 Å.

Ramachandran Plots

The Ramachandran plots of the V_L and C_L homology sub-

TABLE I

Conditions of model building refinement[a]

Parameters varied were: ϕ, ψ, χ, τ(N Cα C). Flexible proline residues were used. In addition, τ(Cα Cβ Cγ) was allowed to vary in Cys, His, Phe, Trp, and Tyr residues. A list of the sources of the amino acid groups used in model building is given in Table I of Diamond (7).

Probe	Length (residues)	Filter constants	
		C_1	C_2
1	1	0.1	10^{-4}
2	2	0.1	10^{-4}
3	3	0.1	10^{-4}

TABLE II

Conditions of real space refinement[a]

Zone length: 5
Margin width: 6
Fixed atomic radii: 1.4 Å
Relative atomic weights; C:6, O:7, N:8, S:16
Relative softness of angular parameters that were allowed to vary:
ϕ, ψ: 4.0
χ: 3.2
Filter levels:

	λ_{min}	$\lambda_{min}/\lambda_{max}$
Scale factor and background	0.0001	0.01
Translational refinement	0.001	0.01
Rotational refinement[b]	0.001	0.001

Electron density map grid; 111.43/160, 56.68/80, 90.30/130 along cell edges

[a] See Diamond (6, 7) for definition of terms.
[b] Nonlinear constraints were used to preserve chain continuity.

[2] $R = \Sigma|F_o - F_c|/\Sigma F_o$, where F_o is observed structure amplitude and F_c is calculated structure amplitude.
[3] The isotropic temperature factor (B) used in the expression *exp*

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