

1.3. MACROMOLECULAR CRYSTALLOGRAPHY AND MEDICINE

1995; Smith *et al.*, 1998; Arnold & Arnold, 1999; Zhang, Geisler *et al.*, 1999). The design of human rhinovirus (HRV) and poliovirus chimeras has been aided by knowing the atomic structure of the viruses (Hogle *et al.*, 1985; Rossmann *et al.*, 1985; Arnold & Rossmann, 1988; Arnold & Rossmann, 1990) and detailed features of the neutralizing immunogenic sites on the virion surfaces (Sherry & Rueckert, 1985; Sherry *et al.*, 1986). In this way, one can imagine that in cases where the atomic structures of antigenic loops in 'donor' immunogens are known as well as the structure of the 'recipient' loop in the virus capsid protein, optimal loop transplantation might become possible. It is not yet known how to engineer precisely the desired three-dimensional structures and properties into macromolecules. However, libraries of macromolecules or viruses constructed using combinatorial mutagenesis can be searched to increase the likelihood of including structures with desired architecture and properties such as immunogenicity. With appropriate selection methods, the rare constructs with desired properties can be identified and 'fished out'. Research of this type has yielded some potentially immunogenic presentations of sequences transplanted on the surface of HRV (reviewed in Arnold & Arnold, 1999). For reasons not quite fully understood, presenting multiple copies of antigens to the immune system leads to an enhanced immune response (Malik & Perham, 1997). It is conceivable that, eventually, it might even be possible for conformational epitopes consisting of multiple 'donor' loops to be grafted onto 'recipient capsids' while maintaining the integrity of the original structure. Certainly, such feats are difficult to achieve with present-day protein-engineering skills, but recent successes in protein design offer hope that this will be feasible in the not too distant future (Gordon *et al.*, 1999).

Immense efforts have been made by numerous crystallographers to unravel the structures of molecules involved in the unbelievably complex, powerful and fascinating immune system. Many of the human proteins studied are listed in Table 1.3.4.5 with, as specific highlights, the structures of immunoglobulins (Poljak *et al.*, 1973), major histocompatibility complex (MHC) molecules (Bjorkman *et al.*, 1987; Brown *et al.*, 1993; Fremont *et al.*, 1992; Bjorkman & Burmeister, 1994), T-cell receptors (TCR) and MHC:TCR complexes (Garboczi *et al.*, 1996; Garcia *et al.*, 1996), an array of cytokines and chemokines, and immune cell-specific kinases such as Ick (Zhu *et al.*, 1999). This knowledge is being converted into practical applications, for instance by humanising non-human antibodies with desirable properties (Reichmann *et al.*, 1988) and by creating immunotoxins.

The interactions between chemokines and receptors, and the complicated signalling pathways within each immune cell, make it next to impossible to predict the effect of small compounds interfering with a specific protein-protein interaction in the immune system (Deller & Jones, 2000). However, great encouragement has been obtained from the discovery of the remarkable manner by which the immunosuppressor FK506 functions: this small molecule brings two proteins, FKBP12 and calcineurin, together, thereby preventing T-cell activation by calcineurin. The structure of this remarkable ternary complex is known (Kissinger *et al.*, 1995). Such discoveries of unusual modes of action of therapeutic compounds are the foundation for new concepts such as 'chemical dimerizers' to activate signalling events in cells such as apoptosis (Clackson *et al.*, 1998).

In spite of the gargantuan task ahead aimed at unravelling the cell-to-cell communication in immune action, it is unavoidable that the next decades will bring us unprecedented insight into the many carefully controlled processes of the immune system. In turn, it is expected that this will lead to new therapeutics for manipulating a truly wonderful defence system in order to assist vaccines, to decrease graft rejection processes in organ transplants and to control auto-immune diseases that are likely to be playing a major role in

cruelly debilitating diseases such as rheumatoid arthritis and type I diabetes.

1.3.6. Outlook and dreams

At the beginning of the 1990s, Max Perutz inspired many researchers with a passion for structure and a heart for the suffering of mankind with a fascinating book entitled *Protein Structure – New Approaches to Disease and Therapy* (Perutz, 1992). The explosion of medicinal macromolecular crystallography since then has been truly remarkable. What should we expect for the next decades?

In the realm of safe predictions we can expect the following:

(a) High-throughput macromolecular crystallography due to the developments outlined in Section 1.3.1, leading to the new field of 'structural genomics'.

(b) Crystallography of very large complexes. While it is now clear that an atomic structure of a complex of 58 proteins and three RNA molecules, the ribosome, is around the corner, crystallographers will widen their horizons and start dreaming of structures like the nuclear pore complex, which has a molecular weight of over 100 000 000 Da.

(c) A steady flow of membrane protein structures. Whereas Max Perutz could only list five structures in his book of 1992, there are now over 40 PDB entries for membrane proteins. Most of them are transmembrane proteins: bacteriorhodopsin, photoreaction centres, light-harvesting complexes, cytochrome *b_c1* complexes, cytochrome *c* oxidases, photosystem I, porins, ion channels and bacterial toxins such as haemolysin and LukF. Others are monotopic membrane proteins such as squalene synthase and the cyclooxygenases. Clearly, membrane protein crystallography is gaining momentum at present and may open the door to atomic insight in neurotransmitter pharmacology in the next decade.

What if we dream beyond the obvious? One day, medicinal crystallography may contribute to:

(a) The design of submacromolecular agonists and antagonists of proteins and nucleic acids in a matter of a day by integrating rapid structure determinations, using only a few nanograms of protein, with the power of combinatorial and, in particular, computational chemistry.

(b) 'Structural toxicology' based on 'human structural genomics'. Once the hundreds of thousands of structures of human proteins and complexes with other proteins and nucleic acids have been determined, truly predictive toxicology may become possible. This will not only speed up the drug-development process, but may substantially reduce the suffering of animals in preclinical tests.

(c) The creation of completely new classes of drugs to treat addiction, organ regeneration, aging, memory enhancement *etc.*

One day, crystallography will have revealed the structure of hundreds of thousands of proteins and nucleic acids from human and pathogen, and their complexes with each other and with natural and designed low-molecular-weight ligands. This will form an extraordinarily precious database of knowledge for furthering the health of humans. Hence, in the course of the 21st century, crystallography is likely to become a major driving force for improving health care and disease prevention, and will find a well deserved place in future books describing progress in medicine, sometimes called 'The Greatest Benefit to Mankind' (Porter, 1999).

Acknowledgements

We wish to thank Heidi Singer for terrific support in preparing the manuscript, and Drs Alvin Kwiram, Michael Gelb, Seymour Klebanov, Wes Van Voorhis, Fred Buckner, Youngsoo Kim and Rein Zwierstra for valuable comments.