The past two decades have witnessed the revolution of targeted therapies for immune-mediated inflammatory diseases, largely pioneered for rheumatological indications. These include tumour necrosis factor (TNF) blockade, interleukin (IL)-6 receptor blockade, IL-1 blockade, IL-17 blockade, IL-12/23 blockade, B-cell depletion and costimulation blockade. Other targets, such as type I interferons and granulocyte macrophage-colony stimulating factor (GM-CSF) remain under investigation. Most recently, janus kinase (JAK) inhibition has provided a small molecule route to targeted interventions. These therapies have been revolutionary, largely because they have provided previously unprecedented efficacy with relative safety.

Thus, the risk of a serious infection with a TNF inhibitor is slightly higher than with a conventional synthetic disease-modifying drug such as methotrexate but significantly less than with higher doses of glucocorticoids for example. Similarly, multiple studies have failed to demonstrate an enhanced risk of solid cancers or lymphomas with biological and targeted synthetic disease-modifying anti-rheumatic drugs (DMARDs). Notably, also treatment with methotrexate which is still the cornerstone of treatment in rheumatoid arthritis (RA), does not have profound immunosuppressive effects, since in the Cardiovascular Inflammation Reduction Trial (CIRT) with non-RA patients the hazard rate for infections was low (1.15 (CI: 1.01 to 1.30)) despite significant comorbidities in the population studied. These data sit in contrast to immunosuppressive drugs used for indications such as transplantation—which were also revolutionary in their time. Renal allografts have a 1-year survival of >90%, using drugs such as tacrolimus, mycophenolate and sometimes glucocorticoids. However, this success comes at the cost of a significantly heightened risk of serious infections and cancer. Rheumatologists are cognisant of similar risks when we use glucocorticoids, cyclophosphamide, azathioprine and mycophenolate, particularly in combination.

The way we describe our tools and our actions has a profound effect on the understanding of our patients and other healthcare professionals. The choice between the terms immunosuppressants and immunomodulators to describe the drugs that we use to treat inflammatory rheumatic and musculoskeletal diseases (RMDs) provides a topical and important example. For patients, the difference between their immune system being suppressed, with the implicit susceptibility to infection, and modulation to become more normal, can affect their behaviours in terms of work and social activities. In their COVID-19 guidance, the American College of Rheumatology (ACR) repeatedly distinguishes immunosuppressants (eg, tacrolimus, cyclosporin A (CSA), mycophenolate mofetil (MMF), azathioprine [AZA]), biologics and/or JAK inhibitors. In contrast, the National Institutes of Health (NIH) highlights ‘the broad immunosuppressive effect of JAK inhibitors’ and the EULAR COVID-19 guideline refers to ‘immunosuppressive treatments, including synthetic DMARDs and biologic DMARDs’.

Thus, there appears to be some uncertainty around how we should describe the drugs that we, as rheumatologists, use daily. Indeed, there was animated discussion among the EULAR COVID-19 Task Force around the most appropriate adjective to use, some members preferring the term ‘immunomodulatory’, which is also not used in the ACR guidance.

The COVID-19 pandemic has thrust several targeted therapies into the spotlight due to the perception that some serious complications of Severe Acute Respiratory Syndrome-Coronavirus-2 (SARS-CoV-2) infection reflect excess cytokine production locally in the lungs and, potentially, systemically. This has stimulated tremendous opportunities and collaborative research efforts, in new directions and with new partners—rheumatologists working with infectious disease physicians, pulmonologists and intensivists as we join forces to fight COVID-19. Indeed, it is critical that rheumatologists are present at the COVID-19 table, providing insights into the risk/benefit ratio of using cytokine blockade, based on more than two decades of experience. During such discussions, we must not forget normal physiology, for example, that type I interferon is essential for viral defence and GM-CSF for surfactant production by alveolar myeloid cells, stressing the potential importance of timing when using targeted interventions. We should be appropriately cautious when advocating trials of these potent therapies, as we are every day in our clinics, reflecting common sense principles as well as formal guidance.

The important message, when contrasting targeted therapies and immunosuppressants, is that the immune system is a highly sophisticated organ, which adopts a ‘belt and braces’ approach when protecting our body from foreign invaders. Consequently, when an allograft is detected, multiple immune effector mechanisms are mobilised synchronously, to rapidly eliminate the ‘invader'; therefore reversal of allograft rejection requires broadly acting immunosuppressive drugs to disable multiple mechanisms, unavoidably enhancing infection risk in parallel. In contrast, immune-mediated inflammatory diseases (IMIDs) are complex disorders in which multiple, subtle, inherited defects in immune regulatory pathways, combined with environmental triggers, result in the gradual onset of autoimmunity and a chronic inflammatory state. While multiple immune-inflammatory pathways may be upregulated in tissues, the success of targeted therapies suggest that these operate in series with one or a few dominating. Therefore, targeting TNF in RA, or IL-17 in psoriasis, may restore homeostasis and health, without a major impact on infection risk. Furthermore, unlike allograft rejection, efficacy does not necessitate targeting of underlying autoreactivity.

The immune system is often likened to an army, the different arms of the immune response, and distinct cell subsets, viewed as regiments, with coordinating roles for ‘generals’ such as dendritic cells, and communicating roles for molecules such as cytokines. Redundancy is a critical virtue of the immune system and of armies—if one element is destroyed, or neutralised, another can often take over. During conflicts of the 20th century, however, relatively untargeted fighting and attacks were used in an attempt to overcome this...
Inherent redundancy, with great loss of life and destruction of infrastructure. The ravages of these conflicts have in part catalysed the modern concept of technology-enabled ‘smart warfare’, for example, using drones to target critical installations, such as communications systems, incapacitating capabilities while minimising loss of military and civilian life and infrastructure.

Using this analogy, traditional immunosuppression is akin to traditional conflicts—disabling multiple immune elements simultaneously, often using drug combinations. This relatively broad attack on the immune system contrasts with modern targeted therapies, which interrupt communications using cytokine blockade, or disable just a particular type of lymphocyte, or regiment, with rituximab, leaving the remaining immune army unscathed.

Of course, targeted therapies still carry risks, particularly if not used with care—combining such treatments leads to more adverse events\textsuperscript{13}; and, when used long-term, drugs such as rituximab can eventually weaken immune defences, and reduced IgG levels may presage an increased risk of severe infection.\textsuperscript{16} Similarly, there are circumstances where even a targeted intervention leads to a specific complication due to lack of redundancy—TNF blockade and tuberculosis (TB) reactivation providing a classic example. Perhaps the strongest endorsement of the potency but relative safety of these drugs is our willingness to study them in populations at risk for RA, in an attempt to prevent disease onset.\textsuperscript{15}

Having worked with targeted therapies for two decades, rheumatologists now have a golden age, where even a targeted intervention can lead to a specific complication due to lack of redundancy—TNF blockade and tuberculosis (TB) reactivation providing a classic example. Perhaps the strongest endorsement of the potency but relative safety of these drugs is our willingness to study them in populations at risk for RA, in an attempt to prevent disease onset.\textsuperscript{15}


table 1 Immunosuppressant versus immunomodulatory drugs and their actions. Immunomodulatory drugs are usually biological therapeutics with specific, extracellular effects on a particular pathway/cell type. Immunosuppressive drugs are small molecule therapeutics with selective effects on intracellular pathways. We have placed JAK inhibitors (and methotrexate) as immunomodulatory drugs because, to date, their risk:benefit profile aligns more precisely with this category.

| Drug          | Effect                                      | Drug          | Effect                                      |
|---------------|---------------------------------------------|---------------|---------------------------------------------|
| Azathioprine  | Multiple—antiproliferative                  | TNF inhibition | Specific cytokine blockade                  |
| Mycophenolate | Purine biosynthesis—antiproliferative, especially lymphocytes | IL-6 receptor blockade | Specific cytokine blockade                  |
| Cyclophosphamide | Alkylating agent—antiproliferative        | Rituximab     | B-cell depletion                            |
| Ciclosporin   | Calcineurin inhibitor—inhomogenate of lymphocyte signalling | Abatacept     | Costimulation blockade (T-cell modulator)   |
| Tacrolimus    | Calcineurin inhibitor—inhomogenate of lymphocyte signalling | Anakinra      | Specific cytokine blockade (IL-1)           |
| Glucocorticoids | Multiple genomic effects on immune and inflammatory pathways | IL-17 inhibition | Specific cytokine blockade                  |
| Sirolimus     | Inhibits mTOR—inhibits lymphocyte signalling and activation | Methotrexate  | Folic acid antagonist—precise mode of action in IMIDs uncertain |

JAK inhibition

| Drug          | Effect                                      |
|---------------|---------------------------------------------|
| JAK inhibition | Selective inhibition of lymphocyte signalling |

IL, interleukin; mTOR, mammalian target of rapamycin.

\textsuperscript{1} Immunosuppressant—broad effects

\textsuperscript{2} Immunomodulators—specific effects/anti-inflammatory
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