ABSTRACT

Chronic arthritis often affects women of childbearing age. The old concept that having chronic arthritis constitutes a major obstacle to women when planning a pregnancy is now obsolete. Thanks to our current capacity to control the activity of rheumatoid arthritis and other chronic inflammatory conditions, and due to the availability of highly effective drugs such as tumour necrosis factor inhibitor agents and other biological agents, many women with these diseases are now able to consider the challenge of childbearing and raising children. Careful pre-conceptional evaluation and risk assessment constitutes the first step of proper care, which can be individualised according to the disease. More than ever, rheumatologists must know how to deal with this situation, and must be able to provide adequate counselling regarding the control of arthritis during conception and pregnancy.

Keywords: Counselling, pregnancy outcome, rheumatoid arthritis (RA), ankylosing spondylitis, psoriatic arthritis (PsA), teratogenicity, biological therapies, disease-modifying anti-rheumatic drugs (DMARDs), glucocorticoids (GCs), non-steroidal anti-inflammatory drugs (NSAIDs).

INTRODUCTION

The old concept that having chronic arthritis constitutes a major obstacle to women when planning a pregnancy is now obsolete. Thanks to our current capacity to control the activity of rheumatoid arthritis (RA) and other chronic inflammatory conditions, and due to the availability of highly effective drugs such as tumour necrosis factor inhibitor (anti-TNF) agents and other biological agents, many women with these diseases are now able to consider the challenge of childbearing and raising children. Careful pre-conceptional evaluation and risk assessment is important. More than ever, rheumatologists must know how to deal with this situation and must be able to provide adequate counselling regarding the control of arthritis during conception and pregnancy.

Prevention of unplanned pregnancy is an essential step to success: all medication must be reviewed before pregnancy in order to prevent risks to the unborn child due to drug exposure. Maintaining remission or low disease activity before and during pregnancy is crucial for good outcomes: active disease at conception usually continues to be active throughout pregnancy and increases the risk of a post-partum flare. In RA patients, it is important to carry out an assessment of certain antibodies that may require special monitoring, such as anti-Ro, anti-La, and antiphospholipid antibodies.

The Effect of Pregnancy on Arthritis

Although a traditional belief considers that RA improves during pregnancy, recent literature has shown that this is not so. A study of approximately 90 pregnancies reported by de Man et al. observed that at least 50% of patients with moderate-to-high disease activity during the first trimester had no clinical improvement as defined by the European League Against Rheumatism
Not all forms of chronic arthritis behave similarly during pregnancy: only one-third of women with ankylosing spondylitis show improvement, while another one-third remain stable and the rest worsen. After delivery, 60% of the women will suffer an outbreak.8 Women with psoriatic arthritis (PsA) improve or experience disease remission in 80% of pregnancies, although approximately 70% experience a post-partum flare during the first 3 months after delivery.9

The Effect of Active Arthritis on Pregnancy

It is likely that most rheumatologists tend to interrupt drug therapies when a pregnancy is planned because of concerns about possible detrimental effects on the fetus. However, growing evidence about the inconvenience of an active disease during pregnancy, with serious materno-fetal consequences, is forcing us to reconsider this position. An association between high disease activity during pregnancy and poor pregnancy outcomes has been demonstrated in several studies,10-12 with greater incidence of pre-term delivery and lower birth-weight infants resulting in an increased risk of perinatal mortality.13 Moreover, caesarean sections were performed significantly more often in a group with intermediate or high disease activity compared with a group with low disease activity (22% versus 10%, p=0.04 by chi-squared test).10 Recently, a relationship between elevated RA disease activity during pregnancy and rapid post-natal catch-up in weight of the offspring has been found, with the latter associated with a worse cardiovascular and metabolic profile in early adulthood.14 Finding the balance between the side-effects of disease-modifying anti-rheumatic drugs (DMARDs) and the control of disease is a challenge; each case must be managed on an individual basis to provide the least possible risk to the mother and to the fetus and neonate, but at the same time minimising adverse events due to disease.

The Effect of Drugs on Mother and Child

For the mother, a drug can be considered safe during pregnancy if it produces no additional side-effects than when used in non-pregnant patients, if it does not increase the rate of complications such as miscarriage or prematurity, and may be considered safe in the child if it does not produce short or long-term adverse effects. A teratogen is an agent that has the potential to interfere with the normal functional or structural development of an embryo or fetus. During organogenesis (15-60 days after conception), the fetus is presumed to be most vulnerable to teratogens. Gross anatomical defects are no longer possible once the definitive form and relationships within an organ system are established, although derangements in the function of organ systems and resulting physiological defects and fetal growth restriction may be present later in pregnancy.15 The USA FDA has established five categories that classify the safety of drugs when used during pregnancy (Table 1).16 However, new requirements published in December 2014 (effective from 30th June, 2015) substitute for these categories by providing a summary of the risks of using a drug during pregnancy and lactation, a discussion of the data supporting that summary, and relevant information to help healthcare providers make prescribing decisions and counsel women about the use of drugs during pregnancy and lactation.17

In patients planning a pregnancy, adjustment of therapy is mandatory. Patients must understand the importance of low disease activity before conception and throughout pregnancy, and therefore the need to maintain the use of certain drugs. On the basis of consensus opinion: corticosteroids and analgesics such as paracetamol can be used throughout pregnancy; non-steroidal anti-inflammatory drugs (NSAIDs) must be avoided during the conception cycle and used sparingly in the first trimester, but can be administered safely until Week 30 of gestation; anti-malarials, sulfasalazine, azathioprine, and cyclosporine are compatible with pregnancy; and methotrexate (MTX), leflunomide (LEF), and biological agents should be withdrawn before a planned pregnancy.18 The most recent information regarding certain biological agents, such as anti-TNF agents, suggests that these medications can be continued in certain circumstances.19

Unfortunately, the number of controlled studies of drugs performed in pregnant women is small.
and experience with therapy often derives from conditions other than rheumatic diseases.  

SAFETY OF DRUG CLASSES MOST COMMONLY USED IN CHRONIC ARTHRITIS PATIENTS DURING PREGNANCY

Non-Steroidal Anti-Inflammatory Drugs

The FDA considers these agents Category B (Table 2). Aspirin can be used throughout pregnancy. Other NSAIDs should be discontinued at the beginning of a conception cycle in order to increase the likelihood of ovulation and, if fertilisation occurs, subsequent implantation. These drugs can then be continued up to Week 30 of gestation since there are no data for teratogenic effects, but they should then be stopped because of the risk of premature closure of the ductus arteriosus. Constriction of the ductus can be seen in approximately 10-50% of fetuses after Week 31. This effect can be avoided if the NSAID is withdrawn 8 weeks before delivery. There are limited data on the use of COX-2 inhibitors during pregnancy. Constriction of the ductus appears to be less pronounced but a recent study suggests that they may increase the risk of congenital anomalies. Therefore, COX-2 inhibitors are considered Category C and should be avoided during pregnancy (Table 2).

Table 1: FDA risk-classification system for drug safety during pregnancy.

<table>
<thead>
<tr>
<th>Category A: safety established</th>
</tr>
</thead>
<tbody>
<tr>
<td>Controlled studies in women fail to demonstrate a risk to the fetus during the first trimester, there is no evidence of risk during later trimesters, and the possibility of fetal harm appears remote.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Category B: safety likely</th>
</tr>
</thead>
<tbody>
<tr>
<td>Either animal-reproduction studies have not demonstrated a fetal risk but there are no controlled studies in pregnant women, or animal-reproduction studies have shown an adverse effect (other than a decrease in fertility) that was not confirmed by controlled studies in women during the first trimester and there is no evidence of a risk during later trimesters.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Category C: teratogenicity possible</th>
</tr>
</thead>
<tbody>
<tr>
<td>Either studies in animals have revealed adverse effects on the fetus (teratogenic, embryocidal, or other) and there are no controlled studies in women, or studies in women and animals are not available. These drugs should be given only if the potential benefit justifies the potential risk to the fetus.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Category D: teratogenicity probable</th>
</tr>
</thead>
<tbody>
<tr>
<td>There is positive evidence of human fetal risk, but the benefits from use in pregnant women may be acceptable despite the risk (e.g. if the drug is needed in a life-threatening situation or for a serious disease for which safer drugs cannot be used or are ineffective).</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Category X: teratogenicity likely – contraindicated in pregnancy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Studies in animals and humans have demonstrated fetal abnormalities and/or there is evidence of fetal risk based on human experience and the risk of the use of the drug in pregnant women clearly outweighs any possible benefit. These drugs are contraindicated in women who are, or may become, pregnant.</td>
</tr>
</tbody>
</table>

Table 2: Toxicities of drugs most commonly used in chronic arthritis patients during pregnancy.

<table>
<thead>
<tr>
<th>Drug</th>
<th>USA FDA risk category</th>
<th>Toxicity concerns: Maternal</th>
<th>Fetal</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non-fluorinated glucocorticoids (prednisone, prednisolone)</td>
<td>B</td>
<td>Pre-term premature rupture of membranes, hypertension, gestational diabetes</td>
<td>Small for gestational age, adrenal hypoplasia, increased risk of cleft lip or palate</td>
</tr>
<tr>
<td>Sulfasalazine</td>
<td>B</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>Anakinra</td>
<td>B</td>
<td>None</td>
<td>Inadequate data</td>
</tr>
<tr>
<td>Anti-TNF agents</td>
<td>B</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>Aspirin and NSAIDs</td>
<td>B</td>
<td>Reduced fertility when used during a conception cycle</td>
<td>Premature closure of the ductus arteriosus when used after Week 30</td>
</tr>
</tbody>
</table>
Glucocorticoids

Glucocorticoids (GCs) are frequently used in the management of pregnant patients with chronic arthritis and are considered Category B (Table 2). When active arthritis affects only one or a limited number of joints, intra-articular steroid injections can be very useful. Recently, prednisone use during pregnancy in RA patients has been associated with unfavourable outcomes as described by a study of 31 RA pregnancies followed prospectively in a single clinic. The patients with neither prednisone nor anti-TNF exposure had a mean first trimester disease activity score — C-reactive protein (DAS-CRP) of 1.63, while patients exposed to prednisone or anti-TNF had a mean DAS-CRP of 3.92 or 3.19, respectively. An adverse outcome (pre-term premature rupture of membranes and children who are relatively small for their gestational age).

Disease-Modifying Anti-Rheumatic Drugs

Both MTX and LEF are considered as Category X (Table 2). MTX is teratogenic and embryotoxic, and its use during pregnancy is associated with multiple abnormalities, most often involving the central nervous system, cranial ossification, and the palate (aminopterin syndrome). The drug is contraindicated during pregnancy and should be prescribed to fertile women only under the condition of safe contraception. It has been proposed that conception should be postponed until 3 months after withdrawal of the drug, although this may be too conservative. Since MTX is a folic acid antagonist, folate supplementation should be continued antenatally and throughout pregnancy. There are no definitive data on whether the offspring of men receiving MTX have an increased risk of teratogenicity, although it is usually recommended that men discontinue this medication for 1-3 months prior to attempting conception.

Table 2 continued.

<table>
<thead>
<tr>
<th>Drug</th>
<th>USA FDA risk category</th>
<th>Toxicity concerns: Maternal</th>
<th>Fetal</th>
</tr>
</thead>
<tbody>
<tr>
<td>COX-2 selective inhibitors</td>
<td>C</td>
<td>Possible increase in malformation risk</td>
<td></td>
</tr>
<tr>
<td>Hydroxychloroquine</td>
<td>C</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>Cyclosporine A</td>
<td>C</td>
<td>None</td>
<td>Small for gestational age, prematurity, intrauterine growth retardation, transient hyperkalaemia in neonates</td>
</tr>
<tr>
<td>Rituximab</td>
<td>C</td>
<td>None</td>
<td>Reports of haematological abnormalities, infection</td>
</tr>
<tr>
<td>Abatacept</td>
<td>C</td>
<td>None</td>
<td>Inadequate data</td>
</tr>
<tr>
<td>Tocilizumab</td>
<td>C</td>
<td>None</td>
<td>Inadequate data</td>
</tr>
<tr>
<td>Tofacitinib</td>
<td>C</td>
<td>None</td>
<td>Inadequate data</td>
</tr>
<tr>
<td>Azathioprine</td>
<td>D</td>
<td>None</td>
<td>Small for gestational age, prematurity, intrauterine growth retardation</td>
</tr>
<tr>
<td>Methotrexate</td>
<td>X</td>
<td>None</td>
<td>Abortogenic, craniofacial anomalies, mental retardation</td>
</tr>
<tr>
<td>Leflunomide</td>
<td>X</td>
<td>None</td>
<td>Craniofacial anomalies</td>
</tr>
</tbody>
</table>

LEF is a pyrimidine synthesis inhibitor that is both embryotoxic and teratogenic in animals. Women of childbearing potential for whom LEF is prescribed must be advised to avoid pregnancy. In a follow-up study of 45 women exposed to LEF prior to or during pregnancy, there was a 12.5% incidence of congenital anomalies in the 16 patients exposed during the first trimester, which is higher than the generally reported background rate of 3%.29 Due to its long half-life and protracted elimination from plasma, LEF-treated patients must follow a cholestyramine drug elimination procedure (‘wash out’) and then wait at least 90 days before attempting pregnancy: cholestyramine 8 g is given three-times daily for 11 days and plasma levels, checked twice 2 weeks apart, should be below 0.02 mg/l, with additional cholestyramine administration if the level is higher than this cut-off value.30 For those who inadvertently become pregnant while taking LEF, this ‘wash out’ procedure early during pregnancy can allow the continuation of the pregnancy, with an absence of substantially increased risk of adverse pregnancy outcomes, as has been previously demonstrated.31

Sulfasalazine is considered to be Category B (Table 2). Although both the molecule and its metabolite, sulfapyridine, do cross the placenta, they do not have a significant clinical impact on the fetus. The main data describing pregnancies exposed to sulfasalazine were derived from the treatment of patients with inflammatory bowel disease (IBD), and no increases in birth defects, pathological jaundice, or babies who were small for their gestational age were detected.32-34 Because the drug is a strong inhibitor of the reduced folate carrier, folate supplementation before and throughout pregnancy must be ensured.35 Maternal doses should not exceed 2 g daily in order to avoid the risk of neutropenia in the newborn.36 Sulfasalazine impedes spermatogenesis, causing azoospermia, and so men should discontinue this medication 3 months prior to attempting conception.37

Cyclosporine is also a treatment option, especially for RA and PsA patients, and is considered Category C (Table 2). More than 800 pregnancies exposed to cyclosporine for several weeks or throughout gestation have been reported, with the majority being in transplant recipients,28,38 and there was no incremental increase in congenital malformations. Renal and liver functions were both normal in the neonates exposed.39 The major problems associated with pregnancies exposed to cyclosporine treatment were prematurity in 40-46% and low birth weight (<2,500 g) in 44-65% of cases, but it has been difficult to ascribe a causative role to drug treatment or the underlying maternal disorder.

Azathioprine is not metabolised to its active form by the placenta and so hardly reaches the developing fetus. Pregnancy data from large transplant registries,40,41 as well as women with systemic lupus erythematosus42,43 and IBD,44 found no predominant or frequent birth defect. Although the drug is considered Category D (Table 2) for use during pregnancy, there is substantial evidence suggesting that it is not teratogenic. Intrauterine exposure to azathioprine may occasionally cause slight suppression of the bone marrow, as shown by decreased leukocyte counts and thrombocytopenia at birth.45 Doses should be maintained at 1.5-2 mg/kg/day in order to avoid neonatal depression of haematopoiesis. There are conflicting reports in the literature as to whether these medications increase the risk of children being small for their gestational age. Hydroxychloroquine is classified as Category C (Table 2), but does not appear to cause fetal toxicity and can be used safely.46,47

**Biological Agents**

As in non-pregnant individuals, the use of biological agents increases the potential risk of infections. This is especially important for pathogens such as *Listeria monocytogenes*, for which the risk is already increased in pregnant women48 and can be further increased by treatment.49 Current manufacturers’ guidelines recommend the stopping of all currently licensed biological therapies prior to conception, primarily due to the lack of controlled studies in pregnant women. However, there is now a huge amount of information available regarding their use during conception, pregnancy, and breastfeeding.50 Much of this post-marketing experience has been obtained through registries, especially in the field of IBD. In many cases, not treating IBD or discontinuing therapy prior to or during pregnancy may pose a greater risk to mother and fetus,20 while maintaining treatment appears to be of lower risk.

**Tumour necrosis factor inhibitors**

Anti-TNF agents, such as adalimumab (ADA), certolizumab (CZP), etanercept, golimumab,
and infliximab (IFX), are considered Category B (Table 2). Although some case reports of congenital anomalies (VACTERL anomalies) following exposure to anti-TNF agents raised concerns about their safety profile when used during pregnancy, numerous case series of pregnancies exposed to anti-TNF agents, as well as several recent systematic literature reviews, have failed to find maternal toxicity, embroyotoxicity, or teratogenicity. In the pregnant woman, maternal transfer of immunoglobulins across the syncytiotrophoblast of the chorionic villi provides fetal immunity and is mediated by the neonatal Fc receptor. Immunoglobulin G (IgG) concentrations in fetal blood increase steadily from the early stages of the second trimester until delivery. Small amounts of biological agents will pass into the fetal circulation during early pregnancy, when organogenesis is underway, suggesting that women can be reassured that the use of anti-TNF agents during the peri-conceptional period, until pregnancy is established, should be low risk. By continuing anti-TNF at least until established pregnancy is confirmed, women who are hoping to conceive should not need to risk disease flare during the period between stopping teratogenic medications, such as MTX, and successful conception.

However, most of the safety data available are derived from women discontinuing anti-TNF therapy during the first trimester; few data exist regarding exposure throughout pregnancy. Due to the different composition of anti-TNF agents, some risks may depend on the individual agent rather than the drug class itself. Both IFX and ADA belong to the IgG1 subclass of antibodies and are capable of crossing the placental barrier, particularly during the second half of pregnancy. At birth, both infant and cord blood demonstrated higher concentrations of IFX and ADA. This raises concerns about infections, safety of vaccination, and, in general, about the development of the immune system of the child: serious infections in infants exposed to anti-TNF agents have been described, and a case of fatal disseminated tuberculosis after Bacillus Calmette-Guérin (BCG) vaccination of a child exposed to IFX in utero has been described. In contrast, very low concentrations of CZP were observed in infant and cord blood when mothers were treated with CZP. This is attributed to the absence of an Fc domain on the CZP molecule, which prevents it being bound by the neonatal Fc receptor. Therefore, it is recommended that live vaccines (including rotavirus, intranasal influenza vaccine, and BCG) be withheld for 6 months from birth in neonates who have had in utero exposure to anti-TNF agents, except CZP due to its minimal transfer. Other vaccines can be given on schedule.

Recent data from the multinational PIANO registry provide reassurance regarding the potential long-term effects on children exposed in utero to anti-TNF agents. A total of 501 women with IBD were followed through pregnancy and during the first 4 years of the child’s life and exposure to biological therapy during the third trimester of pregnancy was not associated with increased infant infection rates, and children did not exhibit developmental delay compared with infants not exposed to these agents, controlling for pre-term birth.

### Other biological agents

Anakinra is a recombinant interleukin (IL)-1 receptor antagonist. In animal studies, no harm to the fetus has been demonstrated. The drug is considered Category B but little has been reported about its safety in this setting (Table 2). Tocilizumab is a monoclonal antibody directed against IL-6 receptors and capable of blocking downstream signalling. The drug is considered Category C and no teratogenicity has been demonstrated in animal models, although at high dose there was increased risk of abortion (Table 2). Abatacept works by blocking interactions between antigen-presenting cells and T cells via binding to CD80/CD86 on antigen-presenting cells, with subsequent inhibition of T cell activation. The drug is considered Category C and there are inadequate data to fully comment on its safety during pregnancy (Table 2), although animal studies saw no increased risk when exposed to the maximum recommended human dose. Current recommendations are to discontinue therapy at least 10 weeks before conception. Rituiximab, a monoclonal antibody directed against CD20, depletes B cells. Animal studies are limited but show no teratogenic effect, although B cells have been demonstrated to be reduced in offspring. The drug is considered Category C (Table 2) and there are limited data available; it is recommended that, ideally, at least 12 months should elapse between rituximab discontinuation and conception.
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6. de Man YA et al. Women with rheumatoid arthritis negative for anticyclic citrullinated peptide and rheumatoid factor are more likely to improve during pregnancy, whereas in autoantibody-positive women autoantibody levels are not influenced by pregnancy. Ann Rheum Dis. 2010;69:420-3.
7. de Man YA et al. Disease activity of

### Table 3: Practical points to consider when treating women of childbearing age with chronic arthritis.

| Before pregnancy | • Talk to the patient and her partner about risks  
|                  | • Schedule pregnancy:  
|                  | - Find a time of remission or low disease activity  
|                  | - Stop teratogenic drugs (methotrexate, leflunomide) and replace them with safer alternaties (sulfasalazine, hydroxychloroquine)  
|                  | • Agree on treatment strategy with the patient and her partner:  
|                  | - Maintain or do not maintain the biological drug  
|                  | - Decide which attitude to adopt during pregnancy if the disease becomes active  
|                  | • Individualise treatment:  
|                  | - Risk of disease activation  
|                  | - Risk of maintaining biological treatment

| Some practical clues | • Patients with inactive disease and receiving non-biological and/or non-teratogenic drugs: consider drug holiday  
|                     | • Patients with inactive disease and receiving teratogenic drugs: stop current drugs and replace with safer alternative(s)  
|                     | • Patients with active disease requiring teratogenic drugs (methotrexate, leflunomide): transition to anti-TNF therapy  
|                     | • Patients with inactive disease on anti-TNF therapy: maintain therapy up to confirmed pregnancy test, consider therapy through to Week 32 of gestation  
|                     | • Disease becomes active during pregnancy: consider maintaining anti-TNF throughout pregnancy (try to use those with less placental transfer)

TNF: tumour necrosis factor.

Tofacitinib is a Janus kinase inhibitor recently approved by FDA for RA, although not approved in Europe. The drug is classified as Category C (Table 2) and there are no available data describing its safety during pregnancy. Ustekinumab is considered Category B, with data in humans being limited to case reports. A recent abstract described 26 completed pregnancies that involved ustekinumab exposure in the course of treatment of psoriasis, and which documented a rate of spontaneous abortions comparable to that observed in the general population.61

**CONCLUSION**

Chronic arthritis often affects women of childbearing age. Counselling of the pre-pregnancy patient constitutes the first step of proper care. Complete information regarding risks must be offered to the patient and her partner, and should be individualised according to the disease (Table 3). At the time of conception, the disease must be under control with medications compatible with pregnancy in order to minimise the risks. Ideally, drug treatment during pregnancy should control the mother’s disease activity, do no harm to the fetus, and ensure a healthy period of pregnancy for mother and child. An interdisciplinary approach between the rheumatologist, obstetrician, and eventually the paediatrician, is needed in order to ensure success. Currently, we can say that pregnancy in chronic arthritis patients is only a matter of planning.


58. Mahadevan U et al. Exposure to anti-TNFα therapy in the third trimester of pregnancy is not associated with increased adverse outcomes: results from the PIANO Registry. Gastroenterol. 2014:146 suppl 1:S-170.

