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P060 LIVING WITH RHEUMATOID ARTHRITIS DURING THE CORONAVIRUS PANDEMIC: A LONGITUDINAL INTERVIEW STUDY

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Background/Aims

The COVID-19 pandemic placed patients with rheumatoid arthritis (RA) at increased risk of poor outcomes as a result of their condition, compounded by use of immunosuppressant medication, and higher prevalence of comorbidities. As a consequence, some patients were instructed within the UK to follow strict guidelines to "shield", severely restricting routine social interactions. This study explored patients' longitudinal experiences of living with RA during the COVID-19 pandemic.

Methods

Patients with rheumatoid arthritis, from a community hospital-based rheumatology service, participated in two semi-structured telephone interviews at baseline in autumn 2020 and 2-4 months later. Interviews were recorded and transcribed verbatim. Interpretative phenomenological analysis was undertaken by two members of the research team with input from two patient partners (KR and MB).

Results

15 participants (9 females, 10 retired, age range 45-79 years) were interviewed twice. Five themes were identified: i) fear, ii) social wellbeing, iii) physical health, iv) pre-existing self-management of RA as a coping mechanism, and v) vulnerability. The overriding emotion was one of fear of contracting COVID-19, which remained high throughout both interviews. Fear was influenced by patients' existing knowledge of their RA and medications and the presence of other significant co-morbidities. Further influences on fear included mainstream media reports (increasing reporting of deaths and new variants) and personal knowledge (family and friends who had contracted COVID-19). The impact on social wellbeing became more pronounced as remote communications could not replicate the benefits of physical interaction. Participants reported no impact on their physical health, with increased rest resulting from restricted social interaction perceived to be beneficial. Many participants utilised the resilience they had learned as a result of having RA to cope, including stress management, pacing, and exercise. Being categorised as "clinically extremely vulnerable" led to a reassessment of self-identity, with participants not wanting to be perceived as being weak or helpless. Finally, many participants used lockdown to reflect on and reassess their personal priorities.

Conclusion

This longitudinal interview study with 15 people with RA highlights that the main impact of the pandemic appeared to be on emotional wellbeing brought about by fear of COVID-19, later compounded by lack of social interaction. In this small study, participants' physical health was reported to be stable and participants were able to use self-management skills to cope. The realisation of the seriousness of contracting COVID-19 led to feelings of vulnerability and a reassessment of self-identity. The study raises important issues for those providing healthcare to people with RA, including effective communication with awareness of its likely impact, using pre-existing self-management strategies to enhance wellbeing, and recognition of the potential for social isolation and the implications thereof.

Disclosure

P. Campbell: None. Z. Paskins: None. S. Hider: None. A. Hassell: None. F. Crawford-Manning: None. K. Rule: None. M. Brooks: None. S. Ryan: None.





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is not responding to antimicrobial therapy, until infection is controlled. There is a higher incidence of serious infections in the elderly aged 75 years and older, caution should be used when treating this population. <u>Tuberculosis</u> Patients should be screened for TB before initiating filgotinib, and filgotinib should not be administered to patients with active TB. <u>Viral reactivation</u>: Cases of herpes virus reactivation (e.g., herpes zoster), were reported in clinical studies (see SmPC). If a patient develops herpes zoster, filgotinib treatment should be temporarily interrunted until the enjoyed resolves. Screening patient develops nerpes zoster, fligorinio freatment should be temporarily interrupted until the episode resolves. Screening for viral hepatitis and monitoring for reactivation should be performed. Malignancy: Immunomodulatory medicinal products may increase the risk of malignancies. Malignancies were observed in clinical studies (see SmPC). Fertility: In animal studies, decreased fertility, impaired spermatogenesis, and bitchestale control of fertility. were observed in clinical studies (see SmPC). <u>Fertility</u>: In animal studies, decreased fertility, impaired spermatogenesis, and histopathological effects on male reproductive organs were observed (see SmPC). The potential effect of filgotinib on sperm production and male fertility in humans is currently unknown. <u>Haematological abnormalities</u>: Do not start therapy, or temporarily stop, if Absolute Neutrophil Count (ANC) <1 × 10° cells/L, ALC <0.5 × 10° cells/L or haemoglobin <8 g/dL. Temporarily stop therapy if these values are observed during routine patient management. <u>Vaccinations</u>: Use of five vaccines during, or immediately prior to, filgotinib treatment is not recommended. <u>Lipids</u>: Treatment with filgotinib was associated with dose dependent increases in lipid parameters, including total cholesterol, and high-density lipoprotein (HDL) levels, while low density lipoprotein (LDL) levels, while tow density lipoprotein (LDL) levels were slightly increased (see SmPC). <u>Cardiovascular risk</u>: Rheumatoid arthritis patients have an increased risk for cardiovascular disorders. Patients should have risk factors (e.g., hypertension, hyperlipidaemia) managed as part of usual standard of care. <u>Venous thromboeniosm</u>: Events of deep venous thromboesis (DVT) and pulmonary embolism (PE) have been reported in patients receiving JAK inhibitors including filgotinib. Caution should be used in patients with risk factors for DVT/PE, such as older age, obesity, a medical history of DVT/PE, or patients undergoing surgery, and prolonged

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immobilisation. Lactose content: Contains lactose; patients with rare hereditary problems of galactose intolerance, total lactase deficiency or glucose-galactose malabsorption should not take filgotinib. Pregnancy/Lactation: Filgotinib is contraindicated in pregnancy. Filgotinib should not be used during breast-feeding. Women of childbearing potential must use effective contraception during and for at least 1 week after cessation of treatment. Driving/Using machinery: No or negligible influence, however dizziness has been reported. Side effects: See SmPC for full information. Common (21/100 to <1/10); hausea, upper respiratory tract infection, urinary tract infection and dizziness. Uncommon (21/1000 to <1/100); herpes zoster, pneumonia, neutropenia, hypercholesterolaemia and blood creatine phosphokinase increase. Serious side effects: See SmPC for full information Legal category: POM Pack: 30 film-coated tablets/bottle Price: UK Basic NHS cost: £863.10 Marketing authorisation number(s): Great Britain Jyseleca 100mg film-coated tablets PLGB 42147/0001 Jyseleca 200mg film-coated tablets PLGB 42147/0001 Jyseleca 200mg film-coated tablets PLGB 42147/0001 Jyseleca 100mg film-coated tablets EU/1/20/1480/003 EU/1/20/1480/004 Further information: Galapagos UK, Belmont House, 148 Belmont Road, Ukbridge UB8 105, United Kingdom 00800 7878 1345 medicalinfo@etjog. com Jyseleca® is a trademark. Date of Preparation: January 2022 UK-RA-FIL-20220-00019 Additional monitoring required

Adverse events should be reported.

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