### **ABCDE Model of Sarcoidosis Care**

Subjects: Pathology

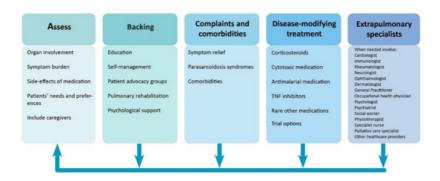
Contributor: Marlies S. Wijsenbeek

The importance of comprehensive care in sarcoidosis is generally acknowledged [12,13]. Here, we describe the ABCDE model, that can be used to structure comprehensive sarcoidosis management in order to improve quality of life and outcomes for patients (Figure 1). This model includes the following components: the Assessment of symptoms and patient's needs, Backing patients by providing support and education, treatment of Complaints and Comorbidities, Disease-modifying treatment, and the involvement of Extrapulmonary specialists. As disease activity, organ involvement, and patients' preferences may vary during the disease course, regular reassessment is essential. The ABCDE model can provide guidance to clinicians during the first work-up and follow-up of patients with sarcoidosis.

Keywords: sarcoidosis; comprehensive care; Quality of life

### 1. Introduction

#### ABCDE of sarcoidosis care



**Figure 1.** Structured approach to comprehensive care in sarcoidosis. The arrows show that individual patient's symptoms, needs, and treatment should regularly be reassessed. TNF = tumor necrosis factor.

#### 2. Assess

In all sarcoidosis patients, organ involvement should be carefully assessed in the diagnostic process and during follow-up. The degree of organ damage and disease activity is often difficult to quantify, as no gold standards exist and symptoms are often non-specific [ $\frac{1}{2}$ ][ $\frac{2}{3}$ ]. Reported percentages of organ involvement are wide ranging, depending on whether only clinically overt organ involvement or also asymptomatic organ involvement is taken into account. For example, 2%–5% of patients have symptomatic cardiac involvement, but the frequency of clinically inapparent involvement is much higher (probably up to 25%)[ $\frac{1}{2}$ ].

Furthermore, organ involvement significantly varies for different ethnic groups. In individual patients, the number of involved organs may change over time and therefore diagnostic assessments have to be performed regularly, particularly if patients express new symptoms<sup>[5][6]</sup>. The frequency in which different organ systems are affected in sarcoidosis is summarized in figure  $1^{[5][4]}$ . Consequently, patients without apparent organ involvement could still have a high symptom burden<sup>[7]</sup>. Hence, Drent et al. propose four different domains that should be evaluated in the complete work-up of a patient; not only the severity, extent and activity of the disease, but also the impact of disease<sup>[7]</sup>. In a recent survey with over 1000 respondents, 95% of patients reported sarcoidosis-related symptoms, and self-reported symptom burden of sarcoidosis was high<sup>[8]</sup>. Due to this high symptom burden, uncertain prospects, and sometimes social isolation and inability to work, sarcoidosis has a huge impact on lives of patients<sup>[1][7][9]</sup>. In a Dutch government survey, more than 60% of sarcoidosis patients (n = 150) considered their general health moderately to severely impaired. Only 7% answered that their health problems had no influence on their social life<sup>[10]</sup>. Furthermore, almost 50% of patients in a recent study (n = 755) were partially or totally unable to work due to their sarcoidosis, highlighting the considerable impact of sarcoidosis on

daily life<sup>[9]</sup>. A Swedish national registry evaluation of 3347 sarcoidosis patients aged 25–59 years, suggested 8% lower income and 26 lost work days in the year of diagnosis compared with age matched controls<sup>[11]</sup>. In a US registry (n = 2318), 44% of respondents reported a large effect on household finances, and 31% had to quit their job after the diagnosis of sarcoidosis<sup>[12]</sup>.

Overall, sarcoidosis patients have an impaired QoL compared with the healthy population [13]. Several studies analyzed the relation between symptoms and QoL in sarcoidosis. Symptoms predictive for QoL are depression, anxiety, fatigue, reduced exercise capacity, SFN-related symptoms, dyspnea, pain, and arthralgia [3][14][15][16][17][18][19]. Interestingly, partners of sarcoidosis patients also experience a reduced QoL compared with healthy controls [14]. Moreover, partners tend to have increased anxiety levels and psychological distress [20]. Both patients as well as their partners reported that there should be more support for partners of sarcoidosis patients [20].

QoL can be defined as "an individual's perception of their position in life" and is influenced by a person's values, beliefs, culture, physical health, social and psychological state<sup>[21]</sup>. Qol can be measured with patient-reported outcome measures (PROMs). PROMs are instruments that "collect self-reported information about a patients' health condition, without any intervention from a healthcare provider"<sup>[22]</sup>. PROMs can be either generic (applicable to the whole population), disease-specific (developed or validated in a specific disease) or domain-specific (assessing severity or burden of a specific symptom or organ). A wide range of PROMs are currently being used for sarcoidosis<sup>[23][24]</sup>. A number of these instruments, such as the King's Sarcoidosis Questionnaire (KSQ), Sarcoidosis Health Questionnaire (SHQ), Sarcoidosis Assessment Tool (SAT), and Fatigue Assessment Scale (FAS) have been specifically developed to measure QoL and symptom burden in sarcoidosis<sup>[24][25][26][27]</sup>. Although PROMs are mainly used in clinical trials, well-validated PROMs could also be used in clinical practice to evaluate treatment effect and longitudinal changes in symptoms and QoL<sup>[24][28]</sup>. In other chronic diseases, the use of PROMs in regular care is associated with enhanced communication and shared-decision making, detection of unrecognized problems, higher patient satisfaction and improved QoL<sup>[29][30]</sup>. Future research could affirm whether this is also the case in sarcoidosis.

Other factors with a potential negative impact on QoL should not be forgotten in the assessment of sarcoidosis patients. For example, medication for sarcoidosis may lead to debilitating side-effects, such as weight gain, diabetes, osteoporosis and psychological problems<sup>[31][32][33]</sup>. According to one study, patients with higher cumulative doses of prednisolone had a significantly lower QoL when adjusted for disease severity<sup>[12][32]</sup>. Moreover, medication-related events lead to a substantial number of hospitalizations in patients with sarcoidosis<sup>[34]</sup>. Consequently, (dis)advantages of starting and continuing therapy should be weighted by the healthcare provider and patient during every clinic visit.

Although it is increasingly acknowledged that patient perspectives are important for optimizing individually-tailored treatments [8][20][35], literature concerning (unmet) needs and preferences of patients with sarcoidosis is scarce. Recently, an international survey revealed that sarcoidosis patients considered QoL and functionality as the most important treatment outcomes [36]. Blood tests and pulmonary function tests were considered the least important outcomes [36]. These results are in contrast to the current focus of most clinicians on physiological outcome measures [13][3][8][23]. In a number of studies, patients reported the need for better information about sarcoidosis and shared decision making [10][20]. As treatment goals can obviously differ between patients, the first step in shared-decision making is identifying patients' needs and preferences. During the disease course, patients should be involved in their treatment plan and in the regular evaluation of benefits versus risks of (pharmacological) treatment [37][6][10]. Including patients as a partner in care could lead to better QoL and adherence to treatment [38][39]. Multidisciplinary management and improved access to sarcoidosis specialists and expert centers for sarcoidosis are other important needs for patients, although it must be acknowledged that not all sarcoidosis patients require tertiary care [36][20].

# 3. Backing

Several support measures to improve QoL for sarcoidosis patients have been suggested in the past years. Better information and education are vital to optimize care for sarcoidosis [\overline{I}][\overline{Q}][40][41]. Even in the current internet era, patients state that they cannot find enough reliable information about their disease online [\overline{I0}][20]. Moreover, the complexity and heterogeneity of sarcoidosis may complicate communication and knowledge transfer between healthcare providers and patients [\overline{I}]. Currently, patients and their partners often feel misunderstood because of a lack of knowledge among healthcare providers and the general public [\overline{I}][20]. Hence, awareness should be raised in society and among relevant healthcare providers. Patient advocacy groups could play an important role in providing understandable information and education, by organizing information meetings and awareness campaigns [\overline{42}]. Although support groups can have obvious benefits, effects on QoL have never been studied in sarcoidosis.

Self-management support is one of the main pillars of the chronic care model, developed to improve care for patients with chronic conditions<sup>[43]</sup>. Many aspects of chronic disease care can only be managed by patients themselves. Self-management strategies cover all disease domains, and include for example behavioral changes, medication use, exercise, dietary strategies and home monitoring of disease<sup>[44][45]</sup>. To achieve skills for self-management, patients and families need to be adequately trained and supported by their healthcare providers<sup>[5]</sup>. Use of novel eHealth solutions to enhance self-management has recently gained interest in sarcoidosis<sup>[40][46]</sup>. Self-monitoring of symptoms, side-effects, QoL, activity and pulmonary function at home has shown to be feasible and highly appreciated by patients with sarcoidosis<sup>[46][47]</sup>. A comprehensive home monitoring program may provide patients with more insights in their disease course, and thereby empower patients and enhance communication with healthcare providers. In a recent study, home monitoring of pulmonary function allowed for earlier detection of steroid treatment effects, suggesting that patient-managed steroid dosing regimens may be feasible<sup>[47]</sup>.

While activity tracking at home could possibly help to stimulate exercise [46][48][49][50], supervised training programs or multidisciplinary pulmonary rehabilitation might have more beneficial effects<sup>[Z]</sup>. Current evidence indicates that a structured, supervised exercise program can improve symptoms, QoL, exercise capacity, and muscle strength in patients with different stages of sarcoidosis<sup>[51][52][53][54][55][56]</sup>. Although these studies showed promising results, no evidence-based guidelines exist to date. In a survey of international sarcoidosis experts, the vast majority of respondents considered physical training in sarcoidosis valuable and would recommend it as standard of care<sup>[51]</sup>. Pulmonary involvement, fatigue, and reduced exercise tolerance are the main indications for physical therapy in sarcoidosis<sup>[51]</sup>. The long-term effects of physical therapy, optimal duration and content of exercise programs in sarcoidosis have never been investigated and need further study. Furthermore, reimbursement and distance to appropriate exercise programs vary between regions and countries and may limit access to physical therapy for sarcoidosis patients<sup>[51]</sup>. Telerehabilitation could potentially be a solution for patients living in rural areas because distances are bridged online. A study evaluating the feasibility and effect of a telerehabilitation program in sarcoidosis is ongoing (NCT03914027).

The majority of patients would like to have better access to psychological support  $\frac{|36|[20]|}{|36|[20]|}$ . In clinical practice, patients are referred to a psychologist or psychiatrist for further counseling and treatment on indication, though some people advocate for standard psychological assessment  $\frac{|57|}{|57|}$ . Cognitive behavioral therapy has been proposed as a promising method to offer psychological support; this therapy could potentially improve patients' coping strategies and thereby reduce stress, anxiety and depression  $\frac{|77|[16]|}{|57|}$ . To date, one study analyzed the impact of mindfulness-based exercise therapy on physical and psychological symptoms in sarcoidosis. Even though this was a modified training consisting of only one 45-minute workshop, symptoms significantly decreased after the session  $\frac{|58|}{|58|}$ . A randomized controlled trial to assess the impact of an online mindfulness-based cognitive behavioral therapy on QoL, fatigue, stress, and anxiety is currently ongoing  $\frac{|59|}{|59|}$ .

# 4. Complaints and Comorbidities

**Symptom** 

Symptom relief is a major aspect of sarcoidosis management. Dyspnea is among the most common symptoms in sarcoidosis and is an important indication for treatment<sup>[3Z]</sup>. Dyspnea is often multifactorial and can be caused by pulmonary, musculoskeletal or cardiac involvement of sarcoidosis, or deconditioning. Other causes for dyspnea such as infection or pulmonary hypertension should also be evaluated<sup>[3Z]</sup>. Non-pharmacological treatment options include physical training, multidisciplinary pulmonary rehabilitation and potentially cognitive behavioral therapy<sup>[53][54][55][58]</sup>(Table 1). Cough is present in up to 53% of sarcoidosis patients<sup>[60]</sup>. Patients with sarcoidosis have a significantly higher cough frequency compared with the normal population<sup>[61]</sup>. Even though cough is often part of the disease, other causes such as reflux or post-nasal drip should always be excluded. One study showed that inhaled corticosteroids may be effective in reducing cough in sarcoidosis, but two other small studies showed no effects of inhaled corticosteroids on cough in sarcoidosis<sup>[62]</sup>. In the recent CHEST guideline, speech therapy is recommended for patients with ILD and refractory cough, however, this therapy has not been specifically evaluated in sarcoidosis<sup>[65]</sup>. Vasoactive intestinal peptide (VIP) inhalation seemed to reduce cough in sarcoidosis in one small open clinical phase II study, but has never been investigated in a randomized setting<sup>[66]</sup>. Further studies regarding antitussive therapy in sarcoidosis are highly needed.

**Table 1.** Overview of pharmacological and non-pharmacological treatment options for common sarcoidosis symptoms. Many of these recommendations are expert opinion or based on small studies.

Pharmacological Treatment Non-Pharmacological Treatment

Cognitive Impairment	TNF inhibitor treatment <sup>[2]</sup>	
Small-Fiber Neuropathy	Antidepressants  Anticonvulsants  Topical anesthetics  Opioids <sup>[41]</sup> Intravenous immunoglobulin <sup>[70][71]</sup> TNF inhibitor treatment <sup>[71]</sup>	Mindfulness-based therapy <sup>[41]</sup>
Depression and Anxiety	Antidepressants [7] **  Anxiolytics **	Cognitive behavioral therapy <sup>[58]</sup> Pulmonary rehabilitation <sup>[53]</sup> Psychological counselling <sup>[41]</sup>
Fatigue	Treat other causes and comorbidities  Neurostimulants: armodafinil, (dex)methylphenidate <sup>[67][68]</sup> TNF inhibitor treatment <sup>[69]</sup>	Treat reversible causes i.e., obstructive sleep apnea, obesity, depression  Physical training or pulmonary rehabilitation [52][53][54][55]  Psychosocial counselling[7]  Cognitive behavioral therapy[58]
Cough	Regular disease-modifying treatment  Treat other causes Inhaled corticosteroids *	Multimodality speech pathology therapy <sup>[65]</sup>
Dyspnea	Regular disease-modifying treatment  Treat other causes  Supplemental oxygen (in case of hypoxemia)	Physical training, pulmonary rehabilitation [53][54][55]  Cognitive behavioral therapy[58]

<sup>\*</sup> earlier studies showed conflicting results. \*\* should only be initiated after an appropriate psychiatric evaluation. TNF = tumor necrosis factor.

Non-organ manifestations or parasarcoidosis syndromes include symptoms as fatigue, depression, anxiety, pain, SFN, and cognitive impairment [41]. As most of these symptoms are related to each other, it can be challenging to avoid a vicious circle. Fatigue is one of the most prevalent (up to 90% of patients) and burdensome symptoms for patients with sarcoidosis [81][20]. Fatigue is a complex, multifactorial problem [72]. Symptoms as sleep disturbance, psychological problems, cognitive impairment, reduced exercise capacity, and muscle strength are all linked to fatigue  $\frac{72}{73}$ [73][74][75][76]. Moreover, comorbidities and medication use may also contribute to fatigue. Previously, it has been shown that patients with multiorgan involvement and a higher number of comorbidities have increased fatigue levels [73]. Comorbidities associated with fatigue are sleep apnea, pulmonary hypertension, diabetes mellitus, thyroid disorders, and obesity [73]. Fatigue is often a chronic problem, which persists or worsens despite sarcoidosis treatment. Research into better pharmacological and non-

pharmacological treatment options for sarcoidosis-associated fatigue is scarce. Two small randomized trials demonstrated that neurostimulants (armodafinil and methylphenidate) have the potential to reduce sarcoidosis-associated fatigue [67][68]. A number of other relatively small or retrospective studies showed the benefits of physical training and cognitive behavioral therapy on fatigue in sarcoidosis[52][53][54][55][58]. Larger studies investigating the etiology and better treatment options for sarcoidosis-associated fatigue are essential to improve symptom burden and QoL for patients.

Next to fatigue, SFN-related related symptoms were reported by the vast majority of patients (86%) in a recent European survey<sup>[2]</sup>. SFN is difficult to diagnose and to treat<sup>[Z]</sup>. Patients may experience a myriad of symptoms, but frequently present with neuropathic pain or autonomic dysfunction<sup>[71]</sup>. SFN is usually treated with anticonvulsants, antidepressants, topical anesthestics or opioids<sup>[41]</sup>. However, these standard treatment options are often not effective<sup>[Z]</sup>. Retrospective studies showed that intravenous immunoglobulin (IVIG) and tumor necrosis factor (TNF) inhibitors might be effective in reducing SFN-related symptoms<sup>[70][71]</sup>. Furthermore, the erythropoietin agonist cibenitide (ARA290) showed promising results on corneal nerve fiber abundance in a phase 2b clinical trial<sup>[77]</sup>. No statistical differences were found in patient-reported outcomes, probably due to the design of the study<sup>[77]</sup>.

Other common non-organ manifestations of sarcoidosis are psychological problems. Previous studies reported anxiety in up to 33% of patients and depressive symptoms in up to 66% of patients  $\frac{[41][78]}{[78]}$ . In a study which used a comprehensive diagnostic interview, the prevalence of major depressive disorder remained strikingly high, at  $25\%^{\frac{[57]}{[57]}}$ . Most other studies used questionnaires to screen for depressive symptoms  $\frac{[13][79][80]}{[78]}$ . Patients with depression or anxiety tend to have a higher symptom burden and different perception of disease severity  $\frac{[78]}{[78]}$ . Pulmonary rehabilitation may have a beneficial effect on psychological wellbeing  $\frac{[53]}{[58]}$ . Cognitive behavioral therapy also has the potential to improve stress, anxiety and depression in sarcoidosis  $\frac{[58]}{[58]}$ . Furthermore, regular pharmacological therapy for depression and anxiety (antidepressants and anxiety) could be offered after an appropriate psychiatric evaluation  $\frac{[58]}{[58]}$ .

Cognitive problems, including memory loss, concentration difficulties and impaired short-term memory are reported by more than 50% of sarcoidosis patients [8][76]. At present, the cause of cognitive failure is considered multi-factorial, and possibly related to chronic inflammation. Currently, no convincing therapies or interventions are available. An observational study indicated that TNF inhibitor treatment may have a beneficial effect on cognition, as measured by the cognitive failure questionnaire (CFQ)[69]. Besides, treatment of associated symptoms may also ameliorate cognitive function.

Comorbidities may also importantly impact QoL and are more prevalent in patients with sarcoidosis compared to the normal population [34][81][82]. A higher number of comorbidities is associated with more frequent hospitalizations and a higher mortality rate [82][83]. The development of new comorbidities, related either to steroid use or to sarcoidosis itself, after the diagnosis of sarcoidosis is independently and strongly associated with a number of adverse outcomes, including worse QoL, risk of hospitalization, and financial impacts [12]. One study showed that more than half of sarcoidosis patients have more than one comorbidity, with arterial hypertension, thyroid disorders and diabetes mellitus being the most prevalent comorbidities [81]. Another study found a significant higher prevalence of chronic liver disease, autoimmune diseases, chronic pulmonary diseases and cancer in patients with sarcoidosis [82]. A study in African-American sarcoidosis patients reported that 90% of patients had one or more comorbidities [84]. The number of comorbidities is higher in older patients, multi-organ involvement and patients with lower incomes [12][81]. Patients with comorbidities obviously have a higher disease complexity, making multidisciplinary management of sarcoidosis even more essential [82]. The presence of comorbidities should therefore be carefully (re)assessed during the disease course.

# 5. Disease Modifying-Treatment

Not all patients with sarcoidosis require pharmacological treatment, as the majority will have spontaneous regression of the disease<sup>[85]</sup>. Treatment is primarily aimed at suppression of the immune system, and thereby preventing organ damage. The main reasons to start treatment are "to avoid danger or improve quality of life"<sup>[86]</sup>. Factors which should be taken into account are the probability of spontaneous resolution, risk for disease progression, extent of disease, organ dysfunction, activity of sarcoidosis, symptom burden and patient' preferences<sup>[85][13][37][6]</sup>.

Pharmacological treatment of pulmonary sarcoidosis should be considered for patients with significant pulmonary symptoms and patients with an impaired or deteriorating lung function [6][87]. For extrapulmonary sarcoidosis, the ATS/ERS/WASOG guidelines state that treatment should always be initiated in case of cardiac sarcoidosis, involvement of the central nervous system, hypercalcemia and ocular sarcoidosis not responding to topical treatment [88]. Other common indications for treatment are hepatic involvement (impaired liver function, portal hypertension), splenic involvement, bone marrow involvement (cytopenia), nephrolithiasis and skin involvement with disfiguring lesions [6]. Treatment may be

initiated in patients who have disabling symptoms without organ damage; however, this should always be a shared-decision with patients as medication can have debilitating side-effects [37][6]. Consequently, efficacy and side-effects of treatment should be assessed during every outpatient clinic visit.

The current guideline, dating from 1999, states that "the appropriate treatment has not been well-defined for all patients" [88]. Presently, oral corticosteroids (e.g., prednisolone) are recommended as the first-choice therapy for sarcoidosis [88]. This recommendation is mainly based on expert opinion and a few relatively small observational studies and low-quality randomized trials from over 20 years ago [87][89]. Older studies have demonstrated that corticosteroids lead to an improvement in lung function, especially in patients with initial severe impairment of lung function [90][91]. Although corticosteroid treatment leads to short-term improvement of lung function, radiological improvement, and symptom reduction, previous studies have not conclusively demonstrated a beneficial effect in preventing disease progression [6][87]. Disease relapse occurs in over 30% of patients after discontinuation of corticosteroids. Furthermore, due to the lack of larger randomized trials the optimal dosage and duration of treatment remains unclear [89].

The most frequently used second-line treatment is the folic acid antagonist methotrexate  $^{[92]}$ . Methotrexate has a significant steroid-sparing effect and improves lung function  $^{[93][94]}$ . Methotrexate is increasingly used as first-line agent in case of (relative) contra-indications for corticosteroids  $^{[92]}$ . A second choice second-line treatment is azathioprine. A retrospective study in the Netherlands and Belgium showed that azathioprine and methotrexate were equally effective, but azathioprine appeared to have more side-effects  $^{[95]}$ . Mycophenolate mofetil and leflunomide are other second-line alternatives  $^{[5]}$ . Antimalarial medication (chloroquine, hydroxychloroquine) is regularly prescribed in patients with cutaneous involvement or hypercalcemia  $^{[96][97]}$ .

In refractory sarcoidosis, TNF inhibitors can be prescribed as a third-line agent. Infliximab has been studied in randomized controlled trials and may have beneficial effects on both pulmonary and extra-pulmonary sarcoidosis in a subgroup of carefully selected patients [98][99][100]. Adalimumab also appears to be effective [101]. Other treatment options have recently emerged for patients with progressive sarcoidosis. The INBUILD study showed that nintedanib is effective in reducing forced vital capacity decline in patients with fibrotic interstitial lung disease, including sarcoidosis [102]. The efficacy of pirfenidone in progressive fibrotic sarcoidosis is also being studied (NCT03260556). Recently, inhibition of the JAK-STAT signaling pathway has been identified as a novel promising treatment target in sarcoidosis; prospective research is ongoing (NCT03910543, NCT03793439)[103]. Not only is more research needed for refractory sarcoidosis, but better evidence-based treatment for first-line therapy, aiming at a better balance between effects and side-effects, is also highly needed. A multicenter trial evaluating the efficacy and side-effects of prednisone and methotrexate as first-line treatment for pulmonary sarcoidosis has recently started in the Netherlands. A detailed description of medication for sarcoidosis is outside the scope of this review; for an extensive overview, we refer you to other published reviews specifically focusing on this topic [5][85][37][6][86][89].

# 6. Extrapulmonary Specialists

Pulmonary physicians play an important role in the management of sarcoidosis patients, as the lungs are affected in up to 90% of patients. Nevertheless, a multidisciplinary team is needed to improve efficiency of care and outcomes for patients, as many organs can be affected and symptoms are wide-ranging [36][104]. While it is quite obvious that other medical specialists contribute their expertise in cases of extra pulmonary sarcoidosis, healthcare providers such as occupational health physicians, pain specialists or specialist nurses should not be forgotten [36][9](Figure 1). Work participation is lower in patients with sarcoidosis; patients have more health-related sick days and a substantial income loss compared with the normal population [11][12][9][11]. Consequently, many patients undergo work capacity assessments and occupational health physicians need to be well-educated about sarcoidosis [7][9]. Although the role of specialist nurses is not as established as in other interstitial lung diseases, specialist nurses could function as coordinators of care in sarcoidosis and give patients practical and emotional support. In a small minority of patients, disease will progress despite all treatment lines. In a subgroup of these patients, lung transplantation may be an option. Furthermore, even though mortality is overall low in sarcoidosis, a multidisciplinary approach should also include palliative care specialists in end-stage disease or in case of a high disease burden.

#### References

1. Alicia K. Gerke; Marc A. Judson; Yvette C. Cozier; Daniel A. Culver; Laura L. Koth; Disease Burden and Variability in Sarcoidosis. *Annals of the American Thoracic Society* **2017**, *14*, S421-S428, <u>10.1513/AnnalsATS.201707-564OT</u>.

- Judson, M.A.; Costabel, U.; Drent, M.; Wells, A.; Maier, L.; Koth, L.; Shigemitsu, H.; Culver, D.A.; Gelfand, J.; Valeyre, D.; et al. The wasog sarcoidosis organ assessment instrument: An update of a previous clinical tool. Sarcoidosis Vasc. Diffuse Lung Dis. 2014, 31, 19–27.
- Marjolein Drent; Rik Marcellis; Antoine Lenssen; Jolanda De Vries; Association between physical functions and quality
  of life in sarcoidosis.. Sarcoidosis, vasculitis, and diffuse lung diseases: official journal of WASOG 2014, 31, 117–128,
- Marc A. Judson; The Clinical Features of Sarcoidosis: A Comprehensive Review. Clinical Reviews in Allergy & Immunology 2014, 49, 63-78, 10.1007/s12016-014-8450-y.
- 5. Johan Grunewald; Jan C. Grutters; Elizabeth V. Arkema; Lesley Ann Saketkoo; David R. Moller; Joachim Müller-Quernheim; Sarcoidosis. *Nature Reviews Disease Primers* **2019**, *5*, 45, <u>10.1038/s41572-019-0096-x</u>.
- 6. Marlies S. Wijsenbeek; Daniel Culver; Treatment of Sarcoidosis. *Clinics in Chest Medicine* **2015**, *36*, 751-767, <u>10.1016/j.ccm.2015.08.015</u>.
- 7. Marjolein Drent; Bert Strookappe; Elske Hoitsma; Jolanda De Vries; Consequences of Sarcoidosis. *Clinics in Chest Medicine* **2015**, *36*, 727-737, <u>10.1016/j.ccm.2015.08.013</u>.
- 8. M. Voortman; C. M. R. Hendriks; M. D. P. Elfferich; F. Bonella; J. Møller; J. De Vries; U. Costabel; M. Drent; The Burden of Sarcoidosis Symptoms from a Patient Perspective. *Lung* **2019**, *197*, 155-161, <u>10.1007/s00408-019-00206-7</u>.
- 9. C. M. R. Hendriks; L. A. Saketkoo; M. D. P. Elfferich; J. De Vries; P. A. H. M. Wijnen; M. Drent; Sarcoidosis and Work Participation: The Need to Develop a Disease-Specific Core Set for Assessment of Work Ability.. *Lung* **2019**, *197*, 407-413, <u>10.1007/s00408-019-00234-3</u>.
- 10. Waverijn, G.; Spreeuwenberg, P.; Heijmans, M. Leven met een Longziekte in Nederland: Cijfers en Trends over de zorg- en Leefsituatie van Mensen met een Longziekte 2016. Available online: https://www.narcis.nl/publication/RecordID/publicat%3A1003084 (accessed on 30 January 2017)
- 11. Elizabeth V. Arkema; Anders Eklund; Johan Grunewald; Gustaf Bruze; Work ability before and after sarcoidosis diagnosis in Sweden. *Respiratory Medicine* **2018**, *144*, S7-S12, <u>10.1016/j.rmed.2018.09.016</u>.
- 12. Logan J Harper; Alicia K. Gerke; Xiao-Feng Wang; Manuel L Ribeiro Neto; Robert P Baughman; Kelli Beyer; Marjolein Drent; Marc A Judson; Lisa A Maier; Leslie Serchuck; et al. Income and Other Contributors to Poor Outcomes in US Sarcoidosis Patients. *American Journal of Respiratory and Critical Care Medicine* **2019**, *null*, null, <u>10.1164/rccm.20190</u> 6-1250oc.
- 13. Christopher Cox; James F. Donohue; Cynthia D. Brown; Yash P. Kataria; Marc A. Judson; Health-related quality of life of persons with sarcoidosis.. *Chest* **2004**, *125*, 997-1004, <u>10.1378/chest.125.3.997</u>.
- 14. Mareye Voortman; Celine M.R. Hendriks; Paul Lodder; Marjolein Drent; Jolanda De Vries; Quality of Life of Couples Living with Sarcoidosis. *Respiration* **2019**, *98*, 373-382, <u>10.1159/000501657</u>.
- 15. R.M. Wirnsberger; J. De Vries; M.H.M. Breteler; G.L. Van Heck; E.F.M. Wouters; Marjolein Drent; Evaluation of quality of life in sarcoidosis patients.. *Respiratory Medicine* **1998**, *92*, 750-756, <u>10.1016/s0954-6111(98)90007-5</u>.
- 16. Jolanda De Vries; Marjolein Drent; Quality of Life and Health Status in Sarcoidosis: A Review of the Literature. *Clinics in Chest Medicine* **2008**, *29*, 525-532, <u>10.1016/j.ccm.2008.03.003</u>.
- 17. Helen J. Michielsen; Marjolein Drent; Tatjana Peroš-Golubičić; Jolanda De Vries; Fatigue Is Associated With Quality of Life in Sarcoidosis Patients. *Chest* **2006**, *130*, 989-994, <u>10.1378/chest.130.4.989</u>.
- 18. Helen J. Michielsen; Tatjana Peros-Golubicic; Professor Marjolein Drent; Jolanda De Vries; Relationship between Symptoms and Quality of Life in a Sarcoidosis Population. *Respiration* **2006**, *74*, 401-405, <u>10.1159/000092670</u>.
- 19. D. Jastrzębski; D. Ziora; M. Lubecki; K. Zieleznik; M. Maksymiak; J. Hanzel; A. Początek; A. Kolczyńska; L. Nguyen Thi; A. Żebrowska; et al. Fatigue in Sarcoidosis and Exercise Tolerance, Dyspnea, and Quality of Life. *Advances in Experimental Medicine and Biology* **2014**, *833*, 31-36, <u>10.1007/5584\_2014\_18</u>.
- 20. Catharina Moor; Mirjam J.G. Van Manen; P. M. van Hagen; J. R. Miedema; L. M. van den Toorn; Yasmin Gür-Demirel; A. P. C. Berendse; J. A. M. van Laar; Marlies S. Wijsenbeek; Needs, Perceptions and Education in Sarcoidosis: A Live Interactive Survey of Patients and Partners.. *Lung* **2018**, *196*, 569-575, <u>10.1007/s00408-018-0144-4</u>.
- 21. Measuring Quality of Life . WHO. Retrieved 2020-2-26
- 22. FDA. Guidance for Industry Patient-Reported Outcome Measures: Use in Medical Product Development to Support Labeling Claims. Available online: https://www.fda.gov/regulatory-information/search-fda-guidance-documents/patient-reported-outcome-measures-use-medical-product-development-support-labeling-claims (accessed on 17 December 2019).

- 23. Thunold, R.F.; Løkke, A.; Cohen, A.L.; Hilberg, O.; Bendstrup, E. Patient reported outcome measures (proms) in sarcoidosis. Sarcoidosis Vasc. Diffuse Lung Dis. 2017, 34, 2–17.
- 24. Marc A Judson; Quality of Life in Sarcoidosis.. *Seminars in Respiratory and Critical Care Medicine* **2017**, *38*, 546-558, <u>1</u> 0.1055/s-0037-1602589.
- 25. Marc A. Judson; Michael Mack; Jennifer Beaumont; Rosemary Watt; Elliot S. Barnathan; David Victorson; Validation and Important Differences for the Sarcoidosis Assessment Tool. A New Patient-reported Outcome Measure. *American Journal of Respiratory and Critical Care Medicine* **2015**, *191*, 786-795, <u>10.1164/rccm.201410-1785oc</u>.
- 26. De Vries, J.; Michielsen, H.; Van Heck, G.L.; Drent, M. Measuring fatigue in sarcoidosis: The fatigue assessment scale (fas). Br. J. Health Psychol. 2004, 9, 279–291.
- 27. Christopher Cox; James F. Donohue; Cynthia D. Brown; Yash P. Kataria; Marc A. Judson; The Sarcoidosis Health Questionnaire. *American Journal of Respiratory and Critical Care Medicine* **2003**, *168*, 323-329, <u>10.1164/rccm.200211-1343oc</u>.
- 28. Celine Hendriks; Professor Marjolein Drent; Marjon Elfferich; Jolanda De Vries; The Fatigue Assessment Scale. *Current Opinion in Pulmonary Medicine* **2018**, *24*, 495–503, <u>10.1097/mcp.000000000000496</u>.
- 29. Ishaque, S.; Karnon, J.; Chen, G.; Nair, R.; Salter, A.B. A systematic review of randomised controlled trials evaluating the use of patient-reported outcome measures (proms). Qual. Life Res. 2019, 28, 567–592.
- 30. Ethan Basch; Patient-reported outcomes: an essential component of oncology drug development and regulatory review. *The Lancet Oncology* **2018**, *19*, 595-597, <u>10.1016/s1470-2045(18)30141-4</u>.
- 31. Nauman A. Khan; Christopher V. Donatelli; Adriano Tonelli; Jonathan Wiesen; Manuel L. Ribeiro Neto; Debasis Sahoo; Daniel A. Culver; Toxicity risk from glucocorticoids in sarcoidosis patients. *Respiratory Medicine* **2017**, *132*, 9-14, <u>10.10</u> <u>16/j.rmed.2017.09.003</u>.
- 32. Marc A. Judson; Haroon Chaudhry; Amanda Louis; Kevin Lee; Recai Yucel; The effect of corticosteroids on quality of life in a sarcoidosis clinic: the results of a propensity analysis.. *Respiratory Medicine* **2015**, *109*, 526-31, <u>10.1016/j.rme</u> d.2015.01.019.
- 33. David Victorson; D. Cella; Heidi Grund; Marc A. Judson; A conceptual model of health-related quality of life in sarcoidosis. *Quality of Life Research* **2013**, *23*, 89-101, <u>10.1007/s11136-013-0438-1</u>.
- 34. Susanne Pohle; Florent Baty; Martin H. Brutsche; In-Hospital Disease Burden of Sarcoidosis in Switzerland from 2002 to 2012. *PLOS ONE* **2016**, *11*, e0151940, <u>10.1371/journal.pone.0151940</u>.
- 35. Sanne Van Helmondt; Louis B. Polish; Marc A. Judson; Jan C. Grutters; Patient perspectives in sarcoidosis. *Current Opinion in Pulmonary Medicine* **2019**, *25*, 478-483, <u>10.1097/mcp.0000000000000000</u>.
- 36. Robert P Baughman; Rita Barriuso; Kelli Beyer; Jeanette Boyd; Johann Hochreiter; Chris Knoet; Filippo Martone; Bernd Quadder; Jack Richardson; Ginger Spitzer; et al. Sarcoidosis: patient treatment priorities. *ERJ Open Research* **2018**, *4*, 00141-2018, <u>10.1183/23120541.00141-2018</u>.
- 37. Hilario Nunes; Florence Jeny; Diane Bouvry; Yurdagül Uzunhan; Dominique Valeyre; Indications for treatment of sarcoidosis. *Current Opinion in Pulmonary Medicine* **2019**, *25*, 505-518, <u>10.1097/mcp.000000000000000000</u>.
- 38. Glyn Elwyn; Marie-Anne Durand; Julia Song; Johanna Aarts; Paul J Barr; Zackary Berger; Nan Cochran; Dominick Frosch; Dariusz Galasiński; Pål Gulbrandsen; et al. A three-talk model for shared decision making: multistage consultation process. *BMJ* **2017**, *359*, j4891, <u>10.1136/bmj.j4891</u>.
- 39. Karazivan, P.; Dumez, V.; Flora, L.; Pomey, M.P.; Del Grande, C.; Ghadiri, D.P.; Fernandez, N.; Jouet, E.; Las Vergnas, O.; Lebel, P. The patient-as-partner approach in health care: A conceptual framework for a necessary transition. Acad. Med. 2015, 90, 437–441.
- 40. Lisa Maier; Elliott D. Crouser; William J. Martin; Jerry Eu; Executive Summary of the NHLBI Workshop Report: Leveraging Current Scientific Advancements to Understand Sarcoidosis Variability and Improve Outcomes. *Annals of the American Thoracic Society* **2017**, *14*, S415-S420, <u>10.1513/AnnalsATS.201707-563OT</u>.
- 41. Jinny Tavee; Daniel Culver; Nonorgan manifestations of sarcoidosis. *Current Opinion in Pulmonary Medicine* **2019**, *25*, 533-538, <u>10.1097/mcp.00000000000597</u>.
- 42. Francesco Bonella; Marlies Wijsenbeek; María Molina-Molina; Annette Duck; Rosalba Mele; Klaus Geissler; Wim Wuyts; European idiopathic pulmonary fibrosis Patient Charter: a missed opportunity.. *European Respiratory Journal* **2016**, *48*, 283-284, <u>10.1183/13993003.00604-2016</u>.
- 43. Natasja Kingod; Bryan Cleal; Ayo Wahlberg; Gitte R. Husted; Online Peer-to-Peer Communities in the Daily Lives of People With Chronic Illness. *Qualitative Health Research* **2016**, *27*, 89-99, <u>10.1177/1049732316680203</u>.

- 44. Thomas Bodenheimer; Edward H. Wagner; Kevin Grumbach; Improving Primary Care for Patients With Chronic Illness. *JAMA* **2002**, *288*, 1775–1779, <u>10.1001/jama.288.15.1909</u>.
- 45. Dena Schulman-Green; Sarah S. Jaser; Chorong Park; Robin Whittemore; A metasynthesis of factors affecting self-management of chronic illness.. *Journal of Advanced Nursing* **2016**, *72*, 1469-89, <u>10.1111/jan.12902</u>.
- 46. Catharina Moor; Yasmin Gür-Demirel; Marlies S. Wijsenbeek; Feasibility of a Comprehensive Home Monitoring Program for Sarcoidosis.. *Journal of Personalized Medicine* **2019**, *9*, 23, <u>10.3390/jpm9020023</u>.
- 47. Caroline E. Broos; Monique Wapenaar; Caspar W.N. Looman; Johannes C.C.M. In 'T Veen; Leon M. Van Den Toorn; Maria J. Overbeek; Marco J.J.H. Grootenboers; Roxane Heller; Rémy L. Mostard; Linda H.C. Poell; et al. Daily home spirometry to detect early steroid treatment effects in newly treated pulmonary sarcoidosis. *European Respiratory Journal* 2018, *51*, 1702089, <u>10.1183/13993003.02089-2017</u>.
- 48. Thomas Bahmer; Henrik Watz; Maria Develaska; Benjamin Waschki; Klaus F. Rabe; Helgo Magnussen; Detlef Kirsten; Anne-Marie Kirsten; Physical Activity and Fatigue in Patients with Sarcoidosis.. *Respiration* **2017**, 95, 18-26, <u>10.1159/0</u> 00481827.
- 49. Sarah Froidure; Maeva Kyheng; Jean Marie Grosbois; Francois Lhuissier; Sandrine Stelianides; Lidwine Wemeau; Benoit Wallaert; Daily life physical activity in patients with chronic stage IV sarcoidosis: A multicenter cohort study.. *Health Science Reports* **2019**, *2*, e109, <u>10.1002/hsr2.109</u>.
- 50. K. Pilzak; Aleksandra Żebrowska; M. Sikora; B. Hall; O. Łakomy; S. Kostorz; D. Ziora; D. Jastrzębski; Physical Functioning and Symptoms of Chronic Fatigue in Sarcoidosis Patients. *Advances in Experimental Medicine and Biology* **2018**, *1040*, 13-21, <u>10.1007/5584</u> <u>2017</u> <u>85</u>.
- 51. Bert Strookappe; Lesley Ann Saketkoo; Marjon Elfferich; Anne Holland; Jolanda De Vries; Ton Knevel; M. Drent; Physical activity and training in sarcoidosis: review and experience-based recommendations. *Expert Review of Respiratory Medicine* **2016**, *10*, 1057-1068, <u>10.1080/17476348.2016.1227244</u>.
- 52. Bert Strookappe; Jeff Swigris; Jolanda De Vries; Marjon Elfferich; Ton Knevel; Marjolein Drent; Benefits of Physical Training in Sarcoidosis. *Lung* **2015**, *193*, 701-708, <u>10.1007/s00408-015-9784-9</u>.
- 53. Marcellis, R.; Van der Veeke, M.; Mesters, I.; Drent, M.; De Bie, R.; De Vries, G.; Lenssen, A. Does physical training reduce fatigue in sarcoidosis? Sarcoidosis Vasc. Diffuse Lung Dis. 2015, 32, 53–62.
- 54. Naz, I.; Ozalevli, S.; Ozkan, S.; Sahin, H. Efficacy of a structured exercise program for improving functional capacity and quality of life in patients with stage 3 and 4 sarcoidosis: A randomized controlled trial. J. Cardiopulm. Rehabil. Prev. 2018, 38, 124–130.
- 55. Müşerrefe Nur Karadallı; Meral Boşnak-Güçlü; Burcu Camcıoğlu; Nurdan Kokturk; Haluk Türktaş; Effects of Inspiratory Muscle Training in Subjects With Sarcoidosis: A Randomized Controlled Clinical Trial. *Respiratory Care* **2015**, *61*, 483-494, 10.4187/respcare.04312.
- 56. Anne E. Holland; Leona M. Dowman; Catherine J. Hill; Principles of Rehabilitation and Reactivation: Interstitial Lung Disease, Sarcoidosis and Rheumatoid Disease with Respiratory Involvement. *Respiration* **2015**, *89*, 89-99, <u>10.1159/00</u> <u>0370126</u>.
- 57. Arianna Goracci; Andrea Fagiolini; Mirko Martinucci; Sara Calossi; Serena Rossi; Tonino Santomauro; Angela Mazzi; Francesco Penza; Antonella Fossi; E Bargagli; et al. Quality of life, anxiety and depression in Sarcoidosis. *General Hospital Psychiatry* **2008**, *30*, 441-445, <u>10.1016/j.genhosppsych.2008.04.010</u>.
- 58. Lesley Ann Saketkoo; Aryn Karpinski; Jessica Young; Ryan Adell; McCall Walker; Tamarin Hennebury; Melissa Wickremasinghe; Anne-Marie Russell; Feasibility, utility and symptom impact of modified mindfulness training in sarcoidosis. *ERJ Open Research* **2018**, *4*, 00085-2017, <u>10.1183/23120541.00085-2017</u>.
- 59. TIRED trial: A randomized controlled trial of a web-based psychological Intervention to reduce stress and fatigue in patients with sarcoidosis. Available online: https://www.trialregister.nl/trial/7816 (accessed on 20 June 2019)
- 60. Timothy Tully; Surinder Singh Birringer; Cough in Sarcoidosis. Lung 2015, 194, 21-24, 10.1007/s00408-015-9799-2.
- 61. Aish Sinha; Kai K. Lee; Gerrard F. Rafferty; Nadia Yousaf; Ian D. Pavord; James Galloway; Surinder Singh Birringer; Predictors of objective cough frequency in pulmonary sarcoidosis. *European Respiratory Journal* **2016**, *47*, 1461-1471, 10.1183/13993003.01369-2015.
- 62. Robert P Baughman; Michael C. Iannuzzi; Elyse E. Lower; David R Moller; Ronald C Balkissoon; Donna B Winget; Marc A. Judson; Use of fluticasone in acute symptomatic pulmonary sarcoidosis.. *Sarcoidosis, vasculitis, and diffuse lung diseases : official journal of WASOG* **2002**, *19*, 198–204, .
- 63. R M Du Bois; P M Greenhalgh; A M Southcott; N M Johnson; T A Harris; Randomized trial of inhaled fluticasone propionate in chronic stable pulmonary sarcoidosis: a pilot study.. *Eur. Respir. J.* **1999**, *13*, 1345–1350., .

- 64. N Milman; N Graudal; G Grode; E Munch; No effect of high-dose inhaled steroids in pulmonary sarcoidosis: a double-blind, placebo-controlled study.. *Journal of Internal Medicine* **1994**, *236*, 285–290, .
- 65. Surinder Singh Birringer; Joanne E. Kavanagh; Richard S. Irwin; Karina A. Keogh; Kaiser G. Lim; Jay Ryu; Todd M. Adams; Kenneth Altman; Elie Azoulay; Alan F. Barker; et al. Treatment of Interstitial Lung Disease Associated Cough. *Chest* **2018**, *154*, 904-917, <u>10.1016/j.chest.2018.06.038</u>.
- 66. Antje Prasse; Gernot Zissel; Niklas Lützen; Jonas Christian Schupp; Rene Schmiedlin; Elena Gonzalez-Rey; Anne Rensing-Ehl; Gerald Bacher; Vera Cavalli; Dorian Bevec; et al. Inhaled Vasoactive Intestinal Peptide Exerts Immunoregulatory Effects in Sarcoidosis. *American Journal of Respiratory and Critical Care Medicine* **2010**, *182*, 540-548, 10.1164/rccm.200909-1451oc.
- 67. Elyse E. Lower; Stacy Harman; Robert P Baughman; Double-Blind, Randomized Trial of Dexmethylphenidate Hydrochloride for the Treatment of Sarcoidosis-Associated Fatigue. *Chest* **2008**, *133*, 1189-1195, <u>10.1378/chest.07-29</u> <u>52</u>.
- 68. Elyse E. Lower; Atul Malhotra; Victoria Surdulescu; Robert P Baughman; Armodafinil for Sarcoidosis-Associated Fatigue: A Double-Blind, Placebo-Controlled, Crossover Trial. *Journal of Pain and Symptom Management* **2013**, *45*, 159-169, 10.1016/j.jpainsymman.2012.02.016.
- 69. Marjon D. Elfferich; Patty J. Nelemans; Rudolf W. Ponds; Jolanda De Vries; Petal A. Wijnen; Professor Marjolein Drent; Everyday Cognitive Failure in Sarcoidosis: The Prevalence and the Effect of Anti-TNF-α Treatment. *Respiration* **2010**, 80, 212-219, 10.1159/000314225.
- 70. Joseph G. Parambil; Jinny Tavee; Lan Zhou; Karla S. Pearson; Daniel Culver; Efficacy of intravenous immunoglobulin for small fiber neuropathy associated with sarcoidosis. *Respiratory Medicine* **2011**, *105*, 101-105, <u>10.1016/j.rmed.2010</u>. <u>09.015</u>.
- 71. Jinny O. Tavee; Karen Karwa; Zubair Ahmed; Nicolas Thompson; Joseph Parambil; Daniel A. Culver; Sarcoidosis-associated small fiber neuropathy in a large cohort: Clinical aspects and response to IVIG and anti-TNF alpha treatment. *Respiratory Medicine* **2017**, *126*, 135-138, <u>10.1016/j.rmed.2017.03.011</u>.
- 72. Bryan S. Benn; Zoe Lehman; Sharon A. Kidd; Christine Miaskowski; Bernie Y. Sunwoo; Melissa Ho; Sara Sun; Joris Ramstein; Jeffrey M. Gelfand; Laura L. Koth; et al. Sleep disturbance and symptom burden in sarcoidosis. *Respiratory Medicine* **2018**, *144*, S35-S40, <u>10.1016/j.rmed.2018.03.021</u>.
- 73. Michael Fleischer; Andreas Hinz; Elmar Brähler; Hubert Wirtz; Andrea Bosse-Henck; Factors Associated With Fatigue in Sarcoidosis. *Respiratory Care* **2013**, *59*, 1086-1094, <u>10.4187/respcare.02080</u>.
- 74. Bert Strookappe; Jolanda De Vries; Marjon Elfferich; Petra Kuijpers; Ton Knevel; Marjolein Drent; Predictors of fatigue in sarcoidosis: The value of exercise testing. *Respiratory Medicine* **2016**, *116*, 49-54, <u>10.1016/j.rmed.2016.05.010</u>.
- 75. M. J. G. Van Manen; V. L. Wester; E. F. C. Van Rossum; L. M. Van Den Toorn; K. Y. Dorst; Y. B. De Rijke; Marlies S. Wijsenbeek; Scalp hair cortisol and testosterone levels in patients with sarcoidosis.. *PLOS ONE* **2019**, *14*, e0215763, <u>1</u> 0.1371/journal.pone.0215763.
- 76. Celine Hendriks; M. Drent; Willemien De Kleijn; Marjon Elfferich; Petal Wijnen; Jolanda De Vries; Everyday cognitive failure and depressive symptoms predict fatigue in sarcoidosis: A prospective follow-up study. *Respiratory Medicine* **2018**, *138*, S24-S30, <u>10.1016/j.rmed.2017.11.008</u>.
- 77. Daniel Culver; Albert Dahan; Daiva Bajorunas; Maria Jeziorska; Monique Van Velzen; Leon P. H. J. Aarts; Jinny Tavee; Martijn R. Tannemaat; Ann N. Dunne; Rita I. Kirk; et al. Cibinetide Improves Corneal Nerve Fiber Abundance in Patients With Sarcoidosis-Associated Small Nerve Fiber Loss and Neuropathic Pain. *Investigative Opthalmology & Visual Science* 2017, 58, BIO52–BIO60, 10.1167/iovs.16-21291.
- 78. J Ireland; M Wilsher; Perceptions and beliefs in sarcoidosis.. *Sarcoidosis, vasculitis, and diffuse lung diseases : official journal of WASOG* **2010**, *27*, 36–42, .
- 79. Betty Chang; Joanne Steimel; David R. Moller; Robert P Baughman; Marc A. Judson; Henry Yeager; Alvin S. Teirstein; Milton D. Rossman; Cynthia S. Rand; Depression in Sarcoidosis. *American Journal of Respiratory and Critical Care Medicine* **2001**, *163*, 329-334, <u>10.1164/ajrccm.163.2.2004177</u>.
- 80. Marjolein Drent; R M Wirnsberger; M H Breteler; L M Kock; J De Vries; E F Wouters; Quality of life and depressive symptoms in patients suffering from sarcoidosis.. *Sarcoidosis, vasculitis, and diffuse lung diseases : official journal of WASOG* **1998**, *15*, 59–66, .
- 81. Martusewicz-Boros, M.M.; Boros, P.W.; Wiatr, E.; Roszkowski-Sliz, K. What comorbidities accompany sarcoidosis? A large cohort (n=1779) patients analysis. Sarcoidosis Vasc. Diffuse Lung Dis. 2015, 32, 115–120.
- 82. Pilar Brito-Zerón; Nihan Acar-Denizli; Antoni Sisó-Almirall; Xavier Bosch; Fernanda Hernández; Sergi Vilanova; Mireia Villalta; Belchin Kostov; Marina Paradela; Marcelo Sanchez; et al. The Burden of Comorbidity and Complexity in

- Sarcoidosis: Impact of Associated Chronic Diseases. Lung 2017, 196, 239-248, 10.1007/s00408-017-0076-4.
- 83. Adam Nowiński; Elzbieta Puścińska; Anna Goljan; Joanna Peradzyńska; Michal Bednarek; Damian Korzybski; Dariusz Kamiński; Anna Stokłosa; Monika Czystowska; P. Śliwiński; et al. The influence of comorbidities on mortality in sarcoidosis: a observational prospective cohort study. *The Clinical Respiratory Journal* **2015**, *11*, 648-656, <u>10.1111/crj.12398</u>.
- 84. Magdalena Martusewicz-Boros; Piotr W Boros; Elżbieta Wiatr; Kazimierz Roszkowski-Śliż; What comorbidities accompany sarcoidosis? A large cohort (n=1779) patients analysis.. *Sarcoidosis, vasculitis, and diffuse lung diseases : official journal of WASOG* **2015**, *32*, 115–120, .
- 85. Dominique Valeyre; Antje Prasse; Hilario Nunes; Yurdagul Uzunhan; Pierre-Yves Brillet; Joachim Müller-Quernheim; Sarcoidosis. *The Lancet* **2014**, 383, 1155-1167, <u>10.1016/s0140-6736(13)60680-7</u>.
- 86. W. Ennis James; Robert P Baughman; Treatment of sarcoidosis: grading the evidence. *Expert Review of Clinical Pharmacology* **2018**, *11*, 677-687, <u>10.1080/17512433.2018.1486706</u>.
- 87. N S Paramothayan; P W Jones; Corticosteroids for pulmonary sarcoidosis.. *The Cochrane database of systematic reviews* **2000**, *null*, CD001114, <u>10.1002/14651858.CD001114</u>.
- 88. Statement on sarcoidosis. Joint statement of the american thoracic society (ats), the european respiratory society (ers) and the world association of sarcoidosis and other granulomatous disorders (wasog) adopted by the ats board of directors and by the ers executive committee, february 1999. Am. J. Respir. Crit. Care Med. 1999, 160, 736–755.
- 89. Shambhu Aryal; Shambhu Aryal; Contemporary optimized practice in the management of pulmonary sarcoidosis.. *Therapeutic Advances in Respiratory Disease* **2019**, *13*, 1753466619868935, <u>10.1177/1753466619868935</u>.
- 90. Om P. Sharma; Charlotte Colp; M.Henry Williams; Course of pulmonary sarcoidosis with and without corticosteriod therapy as determined by pulmonary function studies. *The American Journal of Medicine* **1966**, *41*, 541-551, <u>10.1016/0</u> 002-9343(66)90216-6.
- 91. C J Johns; J B Zachary; W C Ball; A ten-year study of corticosteroid treatment of pulmonary sarcoidosis.. *The Johns Hopkins medical journal* **1974**, 134, 271–283, .
- 92. Johanna P. Cremers; Marjolein Drent; Aalt Bast; Hidenobu Shigemitsu; Robert P Baughman; Minique Valeyre; Nadera J. Sweiss; Tim L Jansen; Multinational evidence-based World Association of Sarcoidosis and Other Granulomatous Disorders recommendations for the use of methotrexate in sarcoidosis. *Current Opinion in Pulmonary Medicine* **2013**, *19*, 545-561, 10.1097/mcp.0b013e3283642a7a.
- 93. Adriane D.M. Vorselaars; Wim A. Wuyts; Veronique M. M. Vorselaars; Pieter Zanen; Vera H.M. Deneer; Marcel Veltkamp; Michiel Thomeer; C. H. M. Van Moorsel; Jan C. Grutters; Methotrexate vs Azathioprine in Second-line Therapy of Sarcoidosis. *Chest* **2013**, *144*, 805-812, <u>10.1378/chest.12-1728</u>.
- 94. Baughman, R.P.; Winget, D.B.; Lower, E.E.; Methotrexate is steroid sparing in acute sarcoidosis: Results of a double blind, randomized trial.. *Sarcoidosis Vasc. Diffuse Lung Dis.* **2000**, *17*, 60–66, .
- 95. Adriane D.M. Vorselaars; Wim A. Wuyts; Veronique M. M. Vorselaars; Pieter Zanen; Vera H.M. Deneer; Marcel Veltkamp; Michiel Thomeer; C. H. M. Van Moorsel; Jan C. Grutters; Methotrexate vs Azathioprine in Second-line Therapy of Sarcoidosis. *Chest* **2013**, *144*, 805-812, <u>10.1378/chest.12-1728</u>.
- 96. John Adams; Marie M. Diz; Om P. Sharma; Effective Reduction in the Serum 1,25-Dihydroxyvitamin D and Calcium Concentration in Sarcoidosis-Associated Hypercalcemia with Short-Course Chloroquine Therapy. *Annals of Internal Medicine* **1989**, *111*, 437, <u>10.7326/0003-4819-111-5-437</u>.
- 97. J. A. Zic; Treatment of cutaneous sarcoidosis with chloroquine. Review of the literature. *Archives of Dermatology* **1991**, *127*, 1034-1040, <u>10.1001/archderm.127.7.1034</u>.
- 98. Milton Rossman; Lee S Newman; Robert P Baughman; Alvin Teirstein; Steven E Weinberger; Wallace Miller; Bruce E Sands; A double-blinded, randomized, placebo-controlled trial of infliximab in subjects with active pulmonary sarcoidosis. *Sarcoidosis, vasculitis, and diffuse lung diseases : official journal of WASOG* **2006**, *23*, 201–208, .
- 99. Marc A. Judson; R. P. Baughman; U. Costabel; S. Flavin; K. H. Lo; M. S. Kavuru; M. Drent; the Centocor T48 Sarcoidosis investigators; Efficacy of infliximab in extrapulmonary sarcoidosis: results from a randomised trial. *European Respiratory Journal* **2008**, *31*, 1189-1196, <u>10.1183/09031936.00051907</u>.
- 100. Adriane D.M. Vorselaars; Heleen A. Crommelin; Vera H.M. Deneer; Bob Meek; Anke M.E. Claessen; Ruth G.M. Keijsers; Coline H.M. Van Moorsel; Jan C. Grutters; Effectiveness of infliximab in refractory FDG PET-positive sarcoidosis. *European Respiratory Journal* 2015, 46, 175-185, 10.1183/09031936.00227014.
- 101. Heleen A. Crommelin; Leone M. Van Der Burg; Adriane D.M. Vorselaars; Marjolein Drent; Coline H.M. Van Moorsel; Ger T. Rijkers; Vera H.M. Deneer; Jan C. Grutters; Efficacy of adalimumab in sarcoidosis patients who developed intolerance to infliximab. *Respiratory Medicine* **2016**, *115*, 72-77, 10.1016/j.rmed.2016.04.011.

- 102. Kevin R. Flaherty; Athol U. Wells; Vincent Cottin; Anand Devaraj; Simon L.F. Walsh; Yoshikazu Inoue; Luca Richeldi; Martin Kolb; Kay Tetzlaff; Susanne Stowasser; et al. Nintedanib in Progressive Fibrosing Interstitial Lung Diseases. *New England Journal of Medicine* **2019**, *381*, 1718-1727, <u>10.1056/nejmoa1908681</u>.
- 103. Jenny J. Wei; Lisa R. Kallenbach; Maryl Kreider; Thomas H. Leung; Misha Rosenbach; Resolution of cutaneous sarcoidosis after Janus kinase inhibitor therapy for concomitant polycythemia vera.. *JAAD Case Reports* **2019**, 5, 360-361, 10.1016/j.jdcr.2019.02.006.
- 104. Marjolein Drent; Sarcoidosis: benefits of a multidisciplinary approach.. *European Journal of Internal Medicine* **2003**, *14*, 217-220, <u>10.1016/s0953-6205(03)00076-1</u>.

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