

Other medical treatment as indicated including antiarrhythmic therapy, device placement if appropriate, standard heart failure management

Active cardiac sarcoidosis

- Ventricular tachyarrhythmias, high-grade heart block or new bifascicular block, left or right ventricular dysfunction, and
- Active FDG uptake on cardiac FDG-PET

Treat in cases of active inflammation with symptoms based on the following stratification (see Figure 2)

- Definite: Should treat
- Highly probable: Should treat
- Probable: Individualized decision-making

Initiate corticosteroids

- Prednisone 30–40 mg/day with taper specific to patient presentation and comorbidities
- Consider concomitant initial therapy with one of the following steroid-sparing agents: methotrexate, mycophenolate, azathioprine, or leflunomide

Reassessment at 3–6 months

- Cardiac FDG PET

Decreased or resolved inflammation

- Continue prednisone at 10–20 mg/day or taper prednisone further (if on a steroid sparing agent or if clinical manifestations are improved)

Reassessment at 3–6 months

- Cardiac FDG PET

Resolved inflammation

- Taper immunosuppression to off after 12 months of treatment from time of initiation
- Additional decision-making for monitoring of active CS should be guided by clinical course

Decreased but residual inflammation

- Individualize decision-making for immunosuppression based on clinical picture (eg change in arrhythmia burden or cardiac function)
- Additional decision-making for monitoring of active CS should be guided by clinical course

Unchanged or worsening inflammation

Unchanged or worsening inflammation

- Raise prednisone to the lowest previous effective dose
- Add a previously unused steroid-sparing agent: methotrexate, mycophenolate, azathioprine, or leflunomide

Reassessment at 3–6 months

- Cardiac FDG PET

Minimal residual or resolved inflammation

- Wean prednisone to 10–20 mg/day
- Continue steroid sparing agent

Reassessment at 6 months

- Cardiac FDG PET

Resolved inflammation

- Wean prednisone to off
- Continue steroid sparing agent
- Additional decision-making for weaning immunosuppression should be guided by clinical course

Unchanged or worsening inflammation

In cases of no response, evaluate for an alternative diagnosis and/or false positive PET scan especially if diagnosis of CS was not histologically confirmed

If significant drug side effects or contraindications to a steroid sparing agent, start a biologic

Unchanged or worsening inflammation

- Raise prednisone to the lowest previous effective dose
- Add a biologic agent (infliximab or adalimumab)
- Additional decision-making for weaning immunosuppression should be guided by clinical course

Sarcoidosis Clinical Management

Louis Mandel



Sarcoidosis Clinical Management:

Sarcoidosis--clinical Management Om P. Sharma, 1984 *Clinical Management of Salivary Gland Disorders* Louis Mandel, 2024-06-22 This book will serve as a complete reference guide for the totality of salivary gland SG disease Focusing on the clues available to the practitioner during the patient examination it will completely review and update the available data concerning all manner of SG conditions Looking beyond the perspectives of surgery imaging pathology or sialendoscopy in diagnosis and therapy this book will provide all professionals interested in the head and neck extensive and detailed diagnostic information about each SG entity Achieving an accurate diagnosis involves the use of multiple clinical tools history physical examination imaging serology biopsy etc The indication for and significance of each of these investigative procedures will be discussed integrated and photographically illustrated in tandem with the diagnostic review of the relevant SG entity Besides a diagnostic review of readily diagnosed SG entities Sjogren s sialolithiasis infection etc the book will cover subjects whose diagnoses have been inadequately described juvenile recurrent parotitis sarcoid radioactive iodine for thyroid cancer pediatric Sjogren s etc or never reviewed middle ear surgery and saliva Stensen s duct dilatation first bite syndrome HIV parapatid fat etc in other published texts It will be organized by diagnosis Written by an expert in the field with over six decades of clinical experience *Clinical Management of Salivary Gland Disorders* will call attention to the large number of presenting patients who can be categorized as false positives This includes a psychosomatic group with perceptual salivary complaints patients with masseteric hypertrophy paraglandular opacities misdiagnosed as sialoliths dental caries thought to have a salivary origin and more Finally this book will offer thorough methodology for identifying and diagnosing these initially confusing problems An exhaustive resource for the field of salivary gland complaints this book will be useful to otolaryngologists head neck surgeons plastic surgeons oral medicine oral pathology practitioners and residents oral and maxillofacial surgeons and dentists Internists concerned with physical diagnosis of head neck problems neuroradiologists concerned with relating imaging results to a clinical salivary gland condition and nurse practitioners and dental hygienists will also find this book most helpful *Medical Management of Pulmonary Diseases* Theodore Marcy, 1999-08-27 This handy accessible single volume presents useful clinical information to expose the pathophysiology underlying major pulmonary diseases and traces the steps of treatment from establishing diagnosis to managing therapy Contains more than 400 summary tables radiographs pathology specimens and other illustrations that encapsulate informat **Clinical**

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Sarcoidosis is a rare condition that causes small patches of red and swollen tissue called granulomas to develop in the organs of the body It most often affects the lungs and skin Typical symptoms include tender red bumps on the skin shortness of breath and a persistent cough There is no cure for sarcoidosis but the condition often improves without treatment within a few months or years Sarcoidosis is estimated to affect about 1 in every 10 000 people in the UK NHS Choices Part of the Clinical Focus Series this book provides trainees with an overview of this complex condition Beginning with an introduction and guidance on recognising pulmonary granulomas the following chapters provide in depth coverage of numerous common lesions describing their history clinical features differential diagnosis and treatment Presented in an easy to read format this useful reference includes clinical photographs CT scans and X rays histological pictures and tables to enhance learning Key points Provides practical overview of lesions of sarcoidosis Covers numerous common lesions their history clinical features diagnosis and treatment Presented in easy to read format with problem solving approach Includes clinical photographs CT Scans X rays histological pictures and tables *Sarcoidosis, An Issue of Clinics in Chest Medicine* Robert Phillip

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Table of Contents **Sarcoidosis Clinical Management**

1. Understanding the eBook **Sarcoidosis Clinical Management**
 - The Rise of Digital Reading **Sarcoidosis Clinical Management**
 - Advantages of eBooks Over Traditional Books
2. Identifying **Sarcoidosis Clinical Management**
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an **Sarcoidosis Clinical Management**
 - User-Friendly Interface
4. Exploring eBook Recommendations from **Sarcoidosis Clinical Management**
 - Personalized Recommendations
 - **Sarcoidosis Clinical Management** User Reviews and Ratings
 - **Sarcoidosis Clinical Management** and Bestseller Lists

5. Accessing Sarcoidosis Clinical Management Free and Paid eBooks
 - Sarcoidosis Clinical Management Public Domain eBooks
 - Sarcoidosis Clinical Management eBook Subscription Services
 - Sarcoidosis Clinical Management Budget-Friendly Options
6. Navigating Sarcoidosis Clinical Management eBook Formats
 - ePub, PDF, MOBI, and More
 - Sarcoidosis Clinical Management Compatibility with Devices
 - Sarcoidosis Clinical Management Enhanced eBook Features
7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Sarcoidosis Clinical Management
 - Highlighting and Note-Taking Sarcoidosis Clinical Management
 - Interactive Elements Sarcoidosis Clinical Management
8. Staying Engaged with Sarcoidosis Clinical Management
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Sarcoidosis Clinical Management
9. Balancing eBooks and Physical Books Sarcoidosis Clinical Management
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Sarcoidosis Clinical Management
10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
11. Cultivating a Reading Routine Sarcoidosis Clinical Management
 - Setting Reading Goals Sarcoidosis Clinical Management
 - Carving Out Dedicated Reading Time
12. Sourcing Reliable Information of Sarcoidosis Clinical Management
 - Fact-Checking eBook Content of Sarcoidosis Clinical Management
 - Distinguishing Credible Sources
13. Promoting Lifelong Learning

- Utilizing eBooks for Skill Development
- Exploring Educational eBooks

14. Embracing eBook Trends

- Integration of Multimedia Elements
- Interactive and Gamified eBooks

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