

Axial Spondyloarthritis Playbook: reducing time to diagnosis and improving patient experience















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A practical guide, supported by tools and examples.

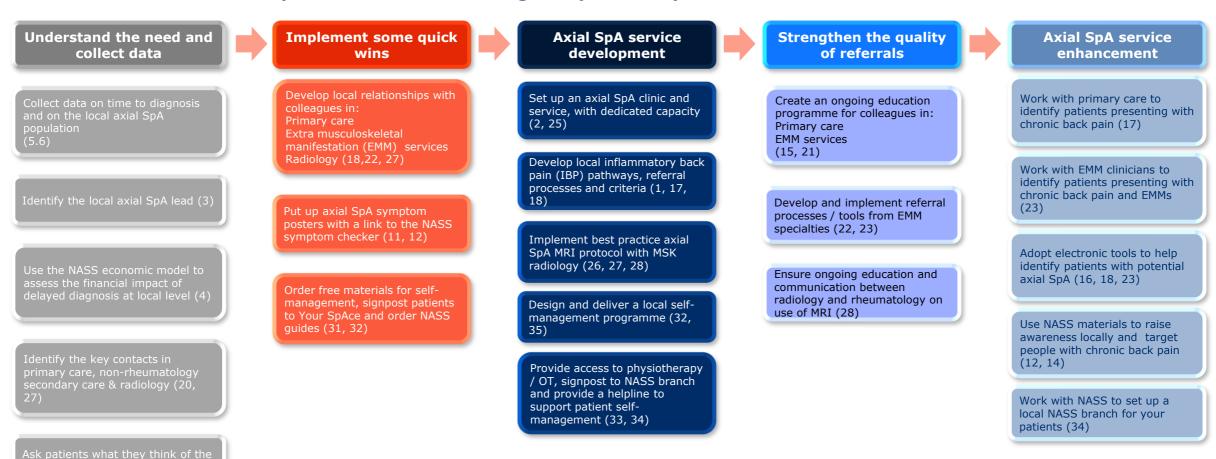
- This guide provides a "best practice" approach to support you in reducing the time to diagnosis in axial SpA locally, as well as the essential support that patients should receive when they are diagnosed.
- It is organised along key sections of the pathway and provides checklists against which you can assess your current practice.
- It also focusses on data as a tool to understand current performance, assess unmet need and measure improvement.
- Resource links take you to guidance, templates, tools and case studies.
- You can review the checklists, set your priorities and collect local data.



An implementation plan on a page



This section sets out a sequence of activity to work through alongside the following pages. The numbers in each step link to the relevant good practice point.



Build patient participation into these areas to understand their needs and perspectives and incorporate them into changes. (7, 8)

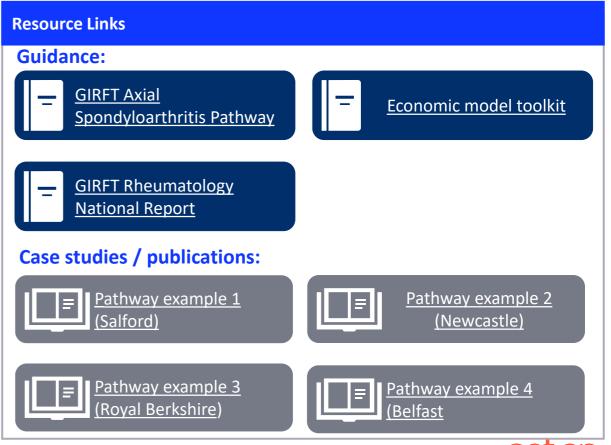
service (7)





Checklist: Enablers to ensure high quality care

Check	Good Practice	Res
	1. An Inflammatory Back Pain pathway is in place, adapting national pathways such as GIRFT, to suit local needs.	Gu
	2: Depending on the availability of local resources, there is a specialist axial SpA service with a multi-disciplinary team.	
	3: As a minimum, there is a lead who has a special interest in axial SpA and who co-ordinates axial SpA care locally.	Ca
	4: An ability to demonstrate potential savings locally of earlier diagnosis of axial SpA using the NASS/UEA economic model. This will help to create a business case.	



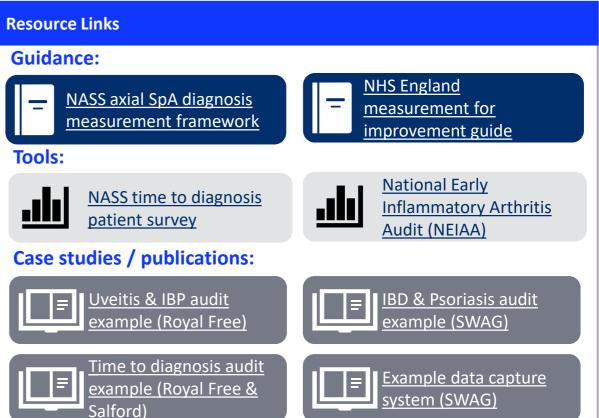






Checklist: Data and measurement frameworks to assess service provision, establish baselines, create improvement targets and monitor progress

Check	Good Practice
	5 . If in England or Wales, ensure data are submitted to the National Early Inflammatory Arthritis Audit (NEIAA) and ask patients to complete the NASS patient survey on time to diagnosis. In Scotland and Northern Ireland ask patients to complete the NASS patient survey to identify your current time to diagnosis and measure changes and improvements.
	 6. Have a data capture system in place beyond the data captured in NEIAA and the NASS patient survey to analyse service effectiveness including: Number of new potential axial SpA referrals Capacity for reviewing new axial SpA referrals Time from referral to 1st appointment Proportion of these referrals diagnosed with axial SpA at 1st appointment Proportion of new patients offered a follow-up appointment versus discharge Proportion requiring advanced investigations Time from request to MRI report Proportion of MRIs requested being supportive of axial SpA diagnosis Proportion of new potential axial SpA referrals who fulfil NICE and/or ASAS spondyloarthritis referral criteria









Checklist: Embedding the patient perspective in service development and improvement, and working collaboratively on treatment plans

Check	Good Practice	Resource Links
	7: The axial SpA service has an ongoing patient and public involvement group that routinely gathers and utilises feedback from their patients on the service.	Guidance: NHS England Experienced- Based co-design (EBCD) guidance Health Research Authority - Public Involvement
	8: Patient and public involvement feeds into the design of Improvement projects.	NHS England Developing patient centred care Tools:
	9: Diagnosis appointments are in person, at a face to face, online or telephone appointment, rather than in writing, so patients feel able to ask questions.	A toolkit for collaborative agenda setting, focus forms and management plans time to diagnosis patient survey NASS quality standards in axial SpA – A values based approach
	10: Shared decision-making plays a key role when discussing treatment options with newly diagnosed patients, meeting their needs as well as clinical needs.	Case studies / publications: Rheumacan - patient Belfast Axial SpA Patient Involvement Project

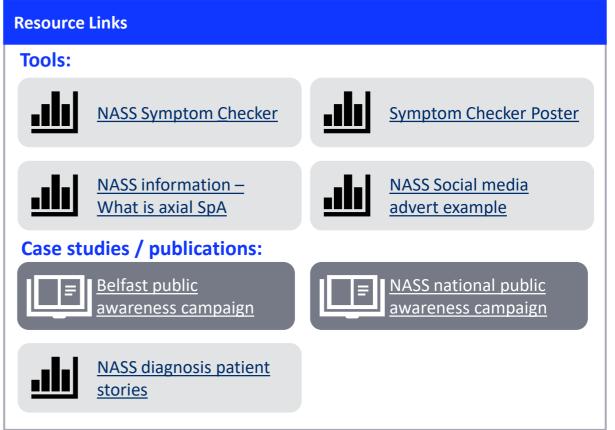




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Checklist: Public Awareness materials for use locally, including the NASS symptom checker

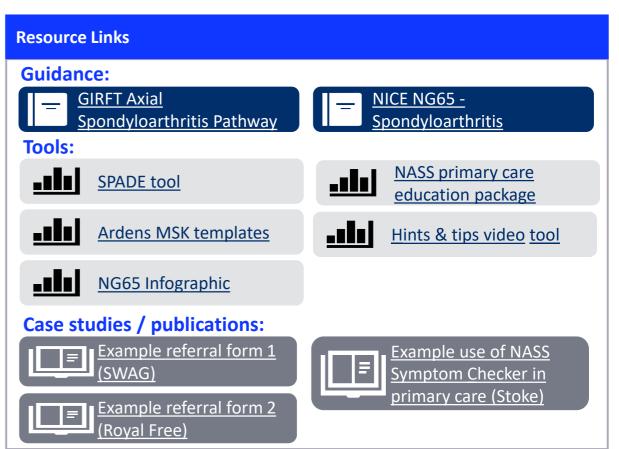
Check	Good Practice	Resource
	11: Direct people to the NASS online symptom checker; people can print off the results and take them their primary care professional for further investigation if appropriate.	Tools:
	12: Put up NASS awareness posters in public spaces in the local area. These display the symptoms of axial SpA and direct people to the symptom checker.	
	13: People attending First Contact Physiotherapy (FCP) appointments for chronic back pain are sent a symptom checker in advance which they take to their appointment and forms part of the FCP assessment.	Case stu
	14: Encourage the use of the Trust/Board local social media platforms to share NASS social media adverts.	





Checklist: Primary Care educational resources, triage tools and linking primary to secondary care

Check	Good Practice
	15: There is programme of primary care education and training on axial SpA through local or regional in-service training, delivered by rheumatology or primary care experts.
	16: Primary care has access to electronic tools that can help to identify patients that might have axial SpA (see Tools).
	17: There are clear local processes for referring patients to rheumatology which includes advice and guidance where possible.
	18: Referrals from primary care into rheumatology should contain assessments of patient symptoms using tools such as the NASS symptom checker and SPADE.
	19: Primary care professionals consider physiotherapy or NSAIDs if appropriate for symptom relief whilst the patient is waiting to be seen in rheumatology.

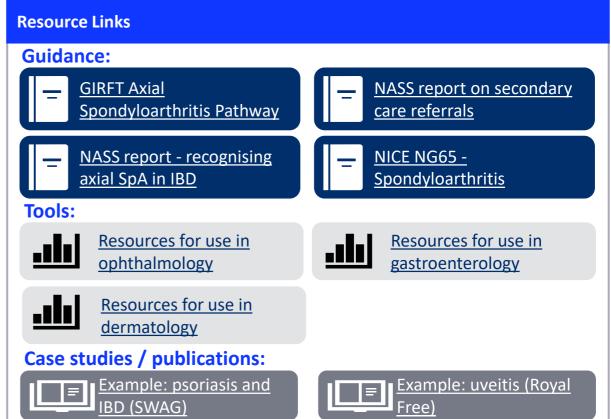




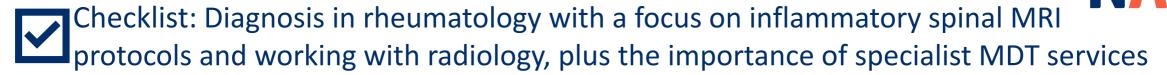


Checklist: Referral from other secondary care services including ophthalmology, dermatology and gastroenterology

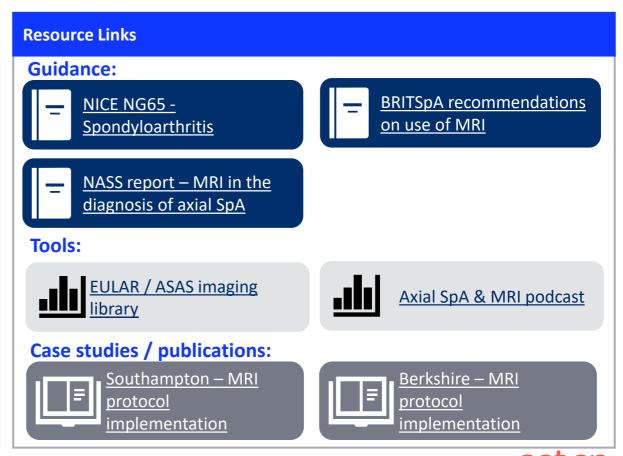
Check	Good Practice	Resource
		Guidano
	20: Rheumatology teams know who the local leads are for uveitis, psoriasis and IBD and engage with them.	
	21: Rheumatology delivers regular education and training on axial SpA to	
	secondary care colleagues in other services.	Tools:
	22: Local communications and arrangements for referrals are in place fitting the local service model (e.g. simple emails, phone calls, joint clinics to a	<u></u>
	formal referral pathway).	
	23: Secondary care professionals have simple tools to help them with the identification, assessment and triage of axial SpA (posters, symptom checker, referral processes, triage tools).	Case stu
	referral processes, triage tools,	







Check	Good Practice
	24: NICE Guideline NG65 should be followed to ensure the right diagnostic tests are performed.
	25: Diagnosis of axial SpA is ideally done within a specialist axial SpA clinic. As a minimum, diagnosis should be made by a clinician with special interest in axial SpA or inflammatory arthritis.
	26: For the diagnosis of suspected axial SpA, in patients where MRI is deemed clinically necessary by a rheumatologist, imaging of both the SIJs and spine is recommended with T1-weighted and fat-suppressed, fluid-sensitive sequences (including STIR*, fat-saturated [†] T2 or Dixon methods) utilised.
	27: All axial SpA MRIs are interpreted and reported on by specialist MSK radiologists either in-house or outsourced. Regular communication and working between rheumatology and radiology is key.
	28: Rheumatologists seeing patients with suspected axial SpA receive education to increase awareness of correct MRI protocols as per the BRITSpA recommendations.
	29: At their diagnosis appointment, patients should be given the opportunity to discuss what is important to them, understand who and how to contact if they need help between appointments and be signposted to NASS.





A playbook for reducing time to diagnosis for axial SpA



Checklist: Information, support & self-management sign posting and service developments following diagnosis

Check	Good Practice
	30 : Provide information about the condition and treatment options in an accessible format and personalised for each patient, ensuring they know who to contact and how if support is needed.
	31: Signpost patients to NASS Your SpAce programme for bitesize information, self-management tools, and peer support.
	32: Offer the full range of NASS guides to patients and provide personalised information and self-management support.
	33: All newly diagnosed patients should be referred to physiotherapy as a minimum for a personalised home-based exercise regime with metrics taken for baseline. Further onward referral to occupational therapy, psychology and podiatry services should also be considered.
	34: Provide information on the local NASS branch for physiotherapist-led exercise and peer support. Where no local branch is available, signpost to the all-UK NASS Online branch. If there is no local NASS branch, consider contacting colleagues and NASS to establish a branch.
	35: Design and deliver supported self-management sessions to axial SpA patients.

