

DERBYSHIRE JOINT AREA PRESCRIBING COMMITTEE (JAPC) SHARED CARE AGREEMENT

NALTREXONE – for the maintenance of alcohol abstinence (for patients within services commissioned by appropriate body)

1. REFERRAL CRITERIA

- Suitability for drug treatment with Naltrexone
- Motivation to remain abstinent assessed
- No contraindications to treatment
- Appropriate support/supervisory network in place for Patient
- Possess an agreed recovery plan
- Condition is stable/predictable
- Effective monitoring is established
- When transferred, the patient will be given a supply of naltrexone sufficient for 4 weeks maintenance therapy.

2. PRINCIPLES FOR THE SHARING OF CARE

- This shared care agreement outlines ways in which the responsibilities for managing the prescribing
 of Naltrexone are shared between the specialist services and General Practitioners (GP) where
 shared care provides an optimal solution for the patient.
- GPs are invited to participate and if not confident to undertake these responsibilities, then he/she is under no obligation to accept shared care.
- Where shared care is not accepted total clinical responsibility for the patient for the diagnosed condition and ongoing supply of medication remains with the specialist.
- Sharing care assumes close communication between the specialist and GP, therefore a specialist asks the GP to prescribe this drug and the GP should reply to the request as soon as practicable.
- Sharing care assumes communication between the Specialist, GP and Patient therefore the process should be explained to the patient by the specialist initiating treatment and they are in agreement.
- The practitioner who prescribes the medication legally assumes clinical responsibility for the drug and the consequences of its use.

3. AREAS OF RESPONSIBILITY

GP responsibilities

- To reply to the request for shared care as soon as practical
- 2. To continue the prescribing following assessment, commencement and stabilisation by the specialist team, usually after 3 months of treatment.
- 3. Prescribe Naltrexone tablets at the dose recommended.
- 4. Continue prescribing naltrexone tablets for up to 6 to 12 months, or longer if benefit or patient wants to continue taking it. If required, may refer to specialist for reassessment (at least every 6 months) as to whether there is a need for on-going therapy.
- 5. Be aware that patient needs to wait 5 to 7 days if being taking opiates (including OTC products) before starting or re-starting

Specialist responsibilities

- 1. To carry out a full holistic review and assessment of the patient prior to prescribing
- 2. Prior to initiation of Naltrexone the Specialist should have done a full review and assessment of the patient and arranged for liver function tests (LFTs) to be carried out. The Specialist will arrange the necessary LFTs for the first month and inform the GP of the results
- 3. To ensure suitability for naltrexone and discuss the risk/benefits of treatment with the patient/carer.
- 4. To ensure the patient is abstinent from alcohol. This will usually have been achieved by a supervised alcohol detox or "drinkdown" with the Community Alcohol Team.
- 5. To perform baseline tests and where clinically appropriate routine tests. (LFTs at initiation and periodically as detailed in section 5(vi) below.)

- naltrexone. Up to 10 days should be allowed for methadone.
- 6. Adjust the dose / stop dose as advised by the specialist.
- Inform specialist team of any change in the patient's medication that may interact with medication patient receives from secondary care.
- 8. Once the patient has been transferred to GP shared care from specialist services, advice may be sought on any aspect of the patient's recovery from alcohol dependence that is of concern to the GP/NMP. See 'Back-up advice and support for contact details.
- 9. Monitor patients overall health and compliance
- 10. To ensure there are no drug interactions with any other medicines initiated in primary care.
- 11. To transfer the patient to the specialist team where appropriate. For example:
 - · Relapse to drinking
 - Concerns over patient compliance
 - Patient or GP is not comfortable to continue with existing regimen due to change in condition or side effects.
- 12. To discontinue treatment on the advice of the specialist team or immediately if urgent need to stop treatment arises.
- Ensure patient awareness of recovery support services available from alcohol services and/or GP support during the prescribing period.
- 14. Report adverse events to the specialist and MHRA.

- To prescribe and monitor for the first 12 weeks of treatment with a view to continue or discontinue treatment.
- 7. To agree the recovery plan with the patient
- 8. To give the patient written information about the medication and a patient treatment card prior to commencement.
- 9. To consult GP whether he/she is willing to participate in prescribing/shared care.
- 10. To continue to prescribe until GP has agreed to take over prescribing, usually after 12 weeks of treatment.
- 11. On acceptance for shared care, to supply four weeks maintenance therapy to allow GP handover
- 12. To ensure local arrangements are made to support administration by a carer or relative
- 13. To provide the GP with a summary of information relating to the patient to support the GP in undertaking shared care.
- 14. To advise the GP of any dose adjustments required, monitoring required, when to transfer_back and when and how to stop the treatment (if appropriate).
- 15. To stop naltrexone if the patient relapses into drinking which persists for 4-6 weeks after starting naltrexone.
- 16. To advise the GP when the patient will next be reviewed by the specialist team.
- 17. To keep the GP informed of the patient's progress
- 18. To ensure a mechanism is in place to receive rapid transfer of a patient from the GP if required if the patient deteriorates
- 19. To report any adverse events to the MHRA on the yellow card form and to inform the GP.
- 20. To ensure that clear communication and support is in place for the GPs to obtain if required

Patient responsibilities

- To take medication as directed by the prescriber.
- To report to the specialist prescriber or GP if he/she does not understand the treatment clearly.
- To maintain contact and engagement with services, as per the recovery plan, to enable them to work towards their recovery goals and overcome alcohol dependence.
- To attend GP and other follow up appointments as scheduled.
- Share any concerns in relation to treatment.
- To seek medical assistance if he/she experiences an adverse reaction or side effect.
- Report any other adverse effects to the specialist or GP whilst taking.

4. COMMUNICATION AND SUPPORT

DHCFT **Derbyshire Recovery Partnership** www.derbyshirerecoverypartnership.co.uk

Specialist Substance Misuse Service

Name: Dr Deepak Sirur

i. Contacts:

Consultant Psychiatrist in Substance Misuse Services

Name: Dr Sugato Sarkar

Consultant Psychiatrist Substance Misuse Services

Southern Derbyshire

ii. Out of hours contacts and procedures:

Patients should be able access emergency help through out of hours GP services or NHS 111

42, St. Marys Gate, Chesterfield, Derbyshire

S41 7TH

Tel. 0300 123 1201 Ripley tel: 01773 744597 Ilkeston tel: 01159 309442 Swadlincote tel: 01283 817352

iii. Specialist support/resources available to GP including patient information:

Patient leaflets and treatment cards are available from the manufacturer.

Downloadable patient information: http://www.medicines.org.uk

iv. Define the referral procedure from hospital to primary care prescriber & route of return should the patient's condition change- Specialist services will coordinate/facilitate the transfer from Specialist services to GP in primary care.

5. CLINICAL INFORMATION

	CLINICAL INFORMATION	-		
i.	Prescribed indications Naltrexone is licensed for use as part of a comprehensive pr			
		treatment, including psychosocial guidance, for detoxified patients who		
		have been alcohol-dependent or opiate-dependent to support abstinence		
ii.	Dose and route of	In accordance to national guidance the therapy should be initiated and		
	administration	supervised by a specialist prescriber experienced in treatment of alcohol-		
		dependent patients.		
		NICE guidance recommends starting at a dose of 25mg daily and aim for		
		a maintenance dose of 50mg daily.		
		Unlicensed for use in children under 18 years of age. Safe use in		
		children has not been established.		
		Not recommended in elderly patients due to insufficient data on the safety		
		and efficacy of naltrexone for this indication		
iii	Duration of treatment	Up to six months, or longer if service user is benefitting from the drug and		
		wants to continue with it		
iv	Contraindications	Hypersensitivity to the active substances or excipients.		
		Severe renal impairment or severe hepatic impairment including acute		
		hepatitis		
		Opioid addicted patients with current abuse of opioids.		
		Positive urine screen for opioids or failure of the naloxone provocation		
		test.		
		Use in conjunction with an opioid – containing medication		
٧.	Cautions	Naltrexone is extensively metabolised by the liver and excreted		
٧.	Guations	predominantly in the urine. Therefore, caution should be observed in		
		administering the medicinal product to patients with impaired hepatic		
		or renal function. Liver function tests should be carried out both		
		before and during treatment.		
		 During treatment with naltrexone, painful conditions should be treated 		
		with non-opioid analgesia only. If opioids cannot be avoided i.e.		
		opioid analgesia or anaesthesia in emergency situations, the dose		
		needed may be higher than normal. In these cases, the respiratory		
		depression and circulatory effects will be more profound and longer		
		lasting. The patient requires monitoring carefully in these situations		
		for signs of opioid toxicity. High dose opioid intake, concomitant with		
		Naltrexone treatment, can lead to life-threatening opioid poisoning		
		from respiratory and circulatory impairment.		
		 Patients must be warned against the concomitant use of opioids (e.g. 		
		opioids in cough medication, opioids in symptomatic medication for		
		the treatment of common colds, or opioids contained in anti-		
		diarrhoeal agents, etc.) during naltrexone treatment		
		Pregnancy: Naltrexone should only be given to pregnant women when, in the		
	Dua	judgement of the attending physician the potential benefits outweigh and the		
VI.	Pregnancy, paternal	possible risk. Referral back to Specialist services for pregnancy management		
	exposure and	with prescribing		
	breastfeeding			
		Breastfeeding: During treatment breast feeding is not recommended.		

	Pregnancy and breastfeeding cases are reviewed by Specialist services and retained appropriately. However, there may be exceptions where cases are best placed with the GP(SC) in the primary care practice, where all parties agree: Consultant, GP(SC), specialist key worker and the patient
vii. Adverse effect	Common or Very common: Joint and muscle pain, abdominal pain, anxiety, chest pain, chills, constipation, decreased potency, delayed ejaculation, diarrhoea, dizziness, headache, increased energy, increased lacrimation, increased sweating, increased thirst, irritability, mood swings, nausea, rash, reduced appetite, sleep disorders, urinary retention, vomiting.
viii. Monitoring requirements	Liver function tests (LFTs) should be taken, and results known, before the administration of Naltrexone LFTs should be repeated after three weeks of treatment. If normal, further monitoring should be as clinically indicated. NICE recommends that patients should be supervised at least monthly for the first 6 months of treatment. After 6 months this can be reduced in frequency but must still be regular. (NICE 2011)
ix. Clinically relevant drug interactions	Clinical experience and experimental data on the effect of naltrexone on the pharmacokinetics of other substances are limited. No interaction studies have been performed, thus concomitant treatment with naltrexone and other medicinal products should be conducted with caution and should be followed carefully. Concomitant administration of naltrexone with an opioid-containing medication should be avoided. There are no known interactions between naltrexone and alcohol.
x. Supply of ancillary equipment	Not applicable
To be read in conjunction with the following documents	RMOC Shared Care Guidance NHSE/NHSCC guidance – items which should not be routinely prescribed in primary care: guidance for CCGs NHSE policy- Responsibility for prescribing between Primary & Secondary/Tertiary Care
Additional information	Where patient care is transferred from one specialist service or GP practice to another, a new shared care agreement must be completed
Prepared by	Mrs CM Jones Specialist Pharmacist Substance Misuse
	Drug and Alcohol Advisory Group DHCFT Approved by DHCFT Medicine Management Committee

This does not replace the SPC, which should be read in conjunction with it.

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6. REFERENCES

Summary of Product Characteristics Naltrexone (Accord): Accessed Jan 2018. Available from: https://www.medicines.org.uk/emc/medicine/25878.

NICE Clinical Guideline 115 (2011) Alcohol Use Disorders: diagnosis, assessment and management of harmful drinking and alcohol dependence. Available from: https://www.nice.org.uk/guidance/cg115.

British National Formulary: Accessed Jan 2018. Available from: http://dx.doi.org/10.18578/BNF.269764922

Sample Transfer Letter

Hospital No: «HOSPITAL_NUMBER»

NHS No: «NHS_NUMBER»

{Insert date}

PRIVATE & CONFIDENTIAL

«GP_TITLE» «GP_INITIALS» «GP_SURNAME» «GP_ADDRESS_1» «GP_ADDRESS_2» «GP_POSTCODE»

DERBYSHIRE JAPC SHARED CARE AGREEMENT LETTER

Dear «GP_TITLE» «GP_SURNAME»

«FORENAME_1» «SURNAME» «DATE_OF_BIRTH» «CURRENT_ADDRESS_1» «CURRENT_ADDRESS_2» «CURRENT_POSTCODE»

Your patient was seen on *{Insert date}* with a diagnosis of *{Insert diagnosis}*. I have initiated the following medication *{Insert drug name}* and am writing to ask you to participate in the shared care for this patient.

This medication has been accepted as suitable for shared care by the Derbyshire Joint Area Prescribing Committee (JAPC). I agree to the secondary care responsibilities set out in the shared care agreement for this medication (available from www.derbyshiremedicinesmanagement.nhs.uk/clinical_guidelines/shared_care_guidelines). I am therefore requesting your agreement to share the care of this patient. Where preliminary tests are set out in the agreement I have carried these out and results are below.

Dose Regimen	Date {Insert medicine name} started	Date for GP to start prescribing <i>{Insert medicine name}</i> from
The baseline test results are (if See overleaf for initiation critical)		

I can confirm that the following has happened with regard to this treatment:

	Specialist to complete
The patient has been initiated on this therapy and has been on an optimised dose for the following period	
of time:	
Baseline investigation and monitoring as set out in the shared care documents have been completed and	Yes / No
were satisfactory	
The condition being treated has a predictable course of progression and the patient can be suitably	Yes / No
maintained by primary care	
The risks and benefits of treatment have been explained to the patient	Yes / No
The roles of the specialist/specialist team/ Primary Care Prescriber / Patient and pharmacist have been	Yes / No
explained and agreed	
The patient has agreed to this shared care arrangement, understands the need for ongoing monitoring,	Yes / No
and has agreed to attend all necessary appointments	163 / 100
I have enclosed a copy of the shared care protocol which covers this treatment/the SCP can be found here	Yes / No
(insert electronic/ web link)	
I have included with the letter copies of the information the patient has received	Yes / No
I have provided the patient with sufficient medication to last until	
I have arranged a follow up with this patient in the following timescale	

If you do <u>NOT</u> wish to participate in shared care for this patient, usually under clinical grounds, please complete the attached form.

Yours sincerely

{Consultant name}

GP RESPONSE TO SHARED CARE (only complete & send if **NOT** participating in shared care*)

* For completeness please record medication on GP clinical system as per guidance- <u>'Recording medicines prescribed</u> and issued by other Healthcare Providers'

Shared care is produced by GPs and specialists knowledgeable in the field of that drug usage. The shared care has been approved by the JAPC. This allows a more convenient service to the patient and cost effective use of NHS resources.

Patient:	NHS No:
Consultant:	Medicine requested for shared care:

I will **NOT** be undertaking the GP responsibilities as described in the agreed shared care guideline. My clinical reasons for declining shared care for this patient are listed in the box below:

		Tick which apply
1.	The prescriber does not feel clinically confident in managing this individual patient's condition, and there is a sound clinical basis for refusing to accept shared care	
	As the patients primary care prescriber I do not feel clinically confident to manage this patient's condition because [insert reason]. I have consulted with other primary care prescribers in my practice who support my decision. This is not an issue which would be resolved through adequate and appropriate training of prescribers within my practice.	
	I have discussed my decision with the patient and request that prescribing for this individual remain with you as the specialist, due to the sound clinical basis given above.	
2.	The medicine or condition does not fall within the criteria defining suitability for inclusion in a shared care arrangement	
	As the medicine requested to be prescribed is not included on the national list of shared care drugs as identified by RMOC or is not a locally agreed shared care medicine I am unable to accept clinical responsibility for prescribing this medication at this time.	
	Until this medicine is identified either nationally or locally as requiring shared care the responsibility for providing this patient with their medication remains with you	
3.	A minimum duration of supply by the initiating clinician	
	As the patient has not had the minimum supply of medication to be provided by the initiating specialist I am unable to take clinical responsibility for prescribing this medication at this time. Therefore can you please contact the patient as soon as possible in order to provide them with the medication that you have recommended.	
	Until the patient has had the appropriate length of supply the responsibility for providing the patient with their medication remains with you.	
4.	Initiation and optimisation by the initiating specialist	
	As the patient has not been optimised on this medication I am unable to take clinical responsibility for prescribing this medication at this time. Therefore can you please contact the patient as soon as possible in order to provide them with the medication that you have recommended.	
	Until the patient is optimised on this medication the responsibility for providing the patient with their medication remains with you.	
5.	Shared Care Protocol not received	
	As legal responsibility for clinical care lies with the clinician who signs the prescription, I need to ensure that I am in possession of sufficient clinical information for me to be confident to prescribe this treatment for my patient and it is clear where each of our responsibilities lie to ensure the patient is safely managed.	
	For this reason I am unable to take clinical responsibility for prescribing this medication at this time, therefore would you please contact the patient as soon as possible in order to provide them with the medication that you have recommended.	
	Until I receive the appropriate SCP, responsibility for providing the patient with their medication remains with you.	

6.	Other (Primary Care Prescriber to complete if there are other reasons why shared care cannot be accepted)	

Yours sincerely

{GP name} {Surgery}

Please send this response to The specialist/consultant requesting shared care