



Comhairle na nDochtúirí Leighis
Medical Council



Safe Prescribing and Dispensing of Controlled Drugs

Joint Guidance

Medical Council and Pharmaceutical Society of Ireland

Version 2, March 2024



Purpose of this Guidance

This resource aims to facilitate safer prescribing and dispensing of controlled drugs (CDs), with a particular focus on controlled drugs in schedule 2, 3 and schedule 4 part 1. It should be used by all prescribers and pharmacists in the collaborative, safe and effective care of patients. This guidance was first produced in 2017 and has been updated to reflect further legislative changes.

While this guidance provides some information on legal requirements applicable to hospital or residential settings, it is primarily aimed at professionals working in a primary care setting.

Controlled Drugs Legislative Change

This joint guidance was first produced in 2017 following changes made by the Misuse of Drugs Regulations 2017.

The guidance was updated in 2024 to reflect changes introduced by amendments to the Medicinal Products (Prescription and Control of Supply) Regulations 2003 (as amended), including changes relating to the emergency supply of schedule 2, 3 and 4 controlled drugs. Additionally, the guidance was updated in light of changes introduced by the Medicinal Products (Prescription and Control of Supply) (Amendment) Regulations 2020. These changes introduced the electronic transmission of prescriptions, including those for controlled drugs, through the national electronic prescription transfer system, and established the legal requirements pertaining to prescriptions sent and received in this way.



Table of Contents

Definitions	3
Controlled Drug Prescriptions	5
Prescriber/ Medical Practitioner	5
Repeat Vs Installment	5
Emergency Supply of Controlled Drugs	6
Purpose of Controlled Drugs Legislation/ Controls and Professional Responsibilities	8
Prescriber Obligations	9
Pharmacist Obligations	9
Prescriptions Required for CD2 and CD3 Drugs	10
Opioid Substitution Prescriptions and Prescriptions for Schedule 4 Part 1 CDs	11
Other Relevant Legal Requirements	12
Overarching Principles	14
Appendix 1a: Example of Compliant Prescription for Schedule 2 and 3 CDs	16
Appendix 1b: Example of Compliant Prescription for Schedule 2 and 3 CDs	17
Appendix 2: Example of Compliant Prescription Schedule 4 Part 1 CD	18
Appendix 3: Example of a Compliant Requisition for Schedule 2, 3 and 4 Part 1 CDs	19
Appendix 4: Example of GMS Repeat Prescription for a Schedule 4 Part 1 CD	20
Appendix 5: Example of Opioid Substitution Prescription	21
Appendix 6: HPRA Website – Information on Medicines Classification	22
Appendix 7: Controlled Drug Prescription Requirements	24
References	25

1. Definitions



What is a Controlled Drug and what are Schedules?

Substances, products, or preparations, including certain medicines, that are either known to be, or have the potential to be, dangerous or harmful to human health, including being liable to misuse or cause social harm, are subject to control under the Misuse of Drugs Acts 1977 to 2016. They are known as “controlled drugs”.

The Misuse of Drugs Regulations categorise controlled drug substances into five schedules (ranging from the most tightly controlled in schedule 1 to the least tightly controlled in schedule 5). Schedule 4 is divided into part 1 and part 2. The controlled drugs in each of these schedules, in practice and in this document, may be referred to as CD1s, CD2s, CD3s, CD4 part 1s, CD4 part 2s and CD5s, respectively.

Each schedule contains various drug substances and drug products based on their perceived medical benefit and their risk to public health. There are different restrictions to control the supply of each schedule of controlled drugs. CD1s have the most restrictions and CD5s have the least restrictions. Some CD5s are available to patients without a prescription.

In this guidance we focus primarily on the controlled drugs found in schedule 2, schedule 3 and schedule 4 part 1. The supply of these controlled drugs is subject to more specific requirements than those in schedule 4 part 2 and schedule 5.

We do not cover Cannabis for medical use, classified as specified controlled drugs, however guidance on the Medicinal Cannabis Access programme is available from the [Department of Health](#).¹ We also do not specifically refer to other schedule 1 controlled drugs as they are not commonly prescribed or supplied and are not regarded as having any therapeutic purposes. They may only be prescribed subject to Ministerial Licence.

Schedule	Examples of Controlled Drugs in Each Schedule
Schedule 1	Substances not ordinarily used as medicines for example, Raw Opium, Coca Leaf
Schedule 2	Opiate substances for example, Morphine, Fentanyl and Oxycodone Some Stimulants for example, Lisdexamphetamine
Schedule 3	Certain Benzodiazepines and painkillers for example, Temazepam, Flunitrazepam, Pentazocine, Ketamine
Schedule 4 Part 1	Most Benzodiazepines and 'Z-drugs' for example, Diazepam, Alprazolam, Clonazepam, Midazolam and Zolpidem
Schedule 4 Part 2	Certain Anti-Epileptics for example, Phenobarbitone <100mg Certain MAOIs for example, Selegiline
Schedule 5	Lower strengths of painkillers for example, Codeine (below specified concentration)

Please note: The above table contains examples only and is not an exhaustive list.

The schedules are set out in the [Misuse of Drugs Regulations 2017 \(as amended\)](#), and [changes such as additions or reclassifications are made by way of amendments to the legislation](#). All prescribers and pharmacists must familiarise themselves with, and refer to, the complete lists of controlled drugs contained in each schedule.

The Health Products Regulatory Authority (HPRA) website should be checked for accurate and up-to-date information regarding the classification of authorised medicines containing controlled drugs i.e. schedule 2, schedule 3, schedule 4 part 1, schedule 4 part 2 and schedule 5. Please see Appendix 6 for more information on how to carry out this search.

The HPRA has produced a list of authorised controlled drug products contained in each schedule. This list can be found on the HPRA website www.hpra.ie. Alternatively, the HPRA is contactable on +353 1 676 4971 or at info@hpra.ie.

2. Controlled Drug Prescriptions

Prescriptions for controlled drugs can be written privately or on a “Health prescription” which is also referred to as a GMS (General Medical Services) prescription. Prescriptions can be paper-based or transferred through the national electronic prescription transfer system, also referred to as Healthmail. Healthmail is the system approved by the Health Service Executive (HSE) for the transfer of prescriptions, in a permanent and unalterable form, by electronic means. It was established following amendments to legislation in 2020². Further information on Healthmail is available at www.healthmail.ie.

Where a Schedule 2 or 3 Controlled Drug prescription is transferred through Healthmail, the specific prescription writing requirements still apply, however, these do not need to be in the prescriber’s own handwriting. As a prescription sent through Healthmail is traceable back to the prescriber, a signature is not required on the prescription. However, the pharmacist must still be satisfied that in their professional judgement it is safe to make any supply in the context of the information received. Examples of controlled drug prescriptions and a reference table are provided in this guidance.

3. Prescriber/Medical Practitioner

“Medical practitioner” in this guidance refers specifically to medical doctors registered in Ireland. “Prescribers” in this guidance refers to healthcare professionals with prescriptive authority for controlled drugs i.e. doctors, nurse prescribers, dentists, veterinary practitioners, and midwives, who are registered in Ireland.

Note: A prescription for a controlled drug which has been prescribed by a practitioner or prescriber (with status equivalent to an Irish practitioner or prescriber) in another EEA state or the UK is not valid for dispensing in Ireland.³

4. Repeat Vs Instalment

- ☑ A “repeat prescription”, as defined in the [Medicinal Products \(Prescription and Control of Supply\) Regulations 2003, as amended](#), means a prescription which may be dispensed more than once.

Schedule 2 and schedule 3 controlled drugs cannot be repeated.

Schedule 4 (part 1) controlled drugs may be repeated in line with the directions of the prescriber up to a maximum of six months.

Schedule 4 (part 2) and schedule 5 controlled drugs may be repeated in line with the directions of the prescriber.

Pharmacists are required to use their professional judgement and decision making regarding every occasion on which a prescription is repeated, and must ensure the

therapeutic appropriateness of the supply, taking account of the specific patient needs. Further guidance on the role of pharmacists in assessing the validity of prescriptions and the appropriateness of repeating prescriptions will be made available by PSI.

- ✔ “Instalments” allow the total quantity of the medicine prescribed to be dispensed in smaller, specified amounts, at specified intervals.

All controlled drugs can be legally dispensed in this manner. However, in accordance with the [Misuse of Drugs Regulations 2017 \(as amended\)](#), ‘the number of instalments and the intervals at which the instalments may be dispensed’ must be specified on prescriptions for schedule 2, schedule 3 and schedule 4 part 1 prescriptions. In the case of schedule 2 and schedule 3 prescriptions, the first instalment must be dispensed within 14 days of the date stated on the prescription, and no instalments may be issued later than 2 months after this date. Please see Appendix 1a for an example of correctly specified instalment directions.

5. Emergency Supply of Controlled Drugs

A pharmacist can provide an emergency supply of a prescription only medicine based on a **request from a prescriber** (registered medical practitioner, registered dentist, or registered nurse prescriber). This is allowable in emergency situations where the prescriber is unable to supply the prescription immediately but undertakes to provide it within 72 hours.

A pharmacist can provide an emergency supply of a prescription only medicine based on a **patient’s request**, where they have interviewed the patient and are satisfied that there is an immediate need for the supply, it is impracticable to obtain a prescription without undue delay, the treatment has previously been prescribed for the patient (by a medical practitioner, registered dentist, or registered nurse prescriber), and the appropriate dose can be verified.

Pharmacists are only **permitted** to provide emergency supplies of controlled drugs in schedule 2, 3 or 4 whether at the request of a prescriber or a patient, **when further specific conditions are met** in accordance with the Medicinal Products (Prescription and Control of Supply) Regulations 2003 (as amended)⁴, when in the pharmacist’s opinion –

- ✔ there is an **immediate need** for the supply of the controlled drug to be made,
- ✔ it is **safe, appropriate, and necessary** to make the supply for the continued treatment of the patient,
- ✔ it is **not possible** to obtain a prescription without undue delay,
- ✔ the treatment has **previously been prescribed** for the patient, and
- ✔ no more than **5 days treatment** or the nearest equivalent minimum dosage pack unit size is being supplied,

and additionally in the case of a **request by a patient**

- ✔ an emergency supply of the controlled drug has not already been supplied based on a request by the patient since their last supply on foot of a prescription.

Note: Pharmacists can provide an emergency supply of methylphenobarbitone, phenobarbitone, phenobarbitone sodium, midazolam, clobazam, or clonazepam, at the request of either a prescriber or a patient, without having to satisfy the additional specific conditions above, where it is for the treatment of epilepsy.

6. Purpose of Controlled Drugs Legislation/Controls and Professional Responsibilities:



There is a strict system of control in place, both nationally and internationally, around the movement and supply of controlled drugs⁵. These controls are intended to enable safe access to these medicines in light of the serious nature of the drugs concerned and substantial potential for abuse and misuse of these medicines.

In Ireland, these medicines are controlled by the Misuse of Drugs Acts and Regulations³. These controls include restrictions on the people who can obtain or possess controlled drugs, and the strict legal obligations placed on pharmacists and prescribers charged with responsibility for the safe control of these substances.

The Misuse of Drugs Regulations 2017 (as amended) are statutory instruments which are vital in protecting patient health and safety. Failure to adhere to this legislation may result in patients receiving inadequate care and unnecessary and unacceptable burden and stress.

All prescribers (including doctors) and pharmacists must adhere to these regulations. It is an offence not to adhere to these regulations.

Additionally, medical practitioners and pharmacists, when registering with the Medical Council and the Pharmaceutical Society of Ireland (PSI) respectively, commit to adhering to professional standards and ethical guidelines set by their regulator.

Medical practitioners adhere to the [Medical Council's Guide to Professional Conduct and Ethics for Registered Medical Practitioners](#), which explicitly states that they must prescribe in compliance with the Misuse of Drugs legislation⁶.

Pharmacists must adhere to the [PSI's Code of Conduct- Professional Principles, Standards and Ethics for Pharmacists](#), which states that pharmacists must '*make ethical decisions, while observing relevant legislation, practice standards and guidance*'.⁷

Any patient requiring a controlled drug for their treatment is entitled to a valid prescription, whether it be in paper or transferred through Healthmail. It is an offence for medical practitioners to write or produce a controlled drug prescription that doesn't comply with the legislative requirements, and it is an offence for a pharmacist to supply controlled drugs from a prescription unless it meets the requirements. It is therefore incumbent upon both medical practitioners and pharmacists to ensure they are aware of their responsibilities and obligations under the Misuse of Drugs legislation to ensure the safe and appropriate supply of controlled drugs to their patients.

7. Prescriber Obligations



- ✓ Provide a valid prescription which meets the requirements of the legislation⁶
- ✓ Be satisfied as to the identity of the person for whose treatment the prescription is to be issued
- ✓ Follow relevant national and international prescribing guidelines
- ✓ Within reason, be available to confirm or discuss any matters related to the prescription and the patient
- ✓ Ensure the safe keeping of prescription pads to reduce the risk of theft and forgery
- ✓ Facilitate appropriate withdrawal of controlled drugs and follow-up and refer as necessary

8. Pharmacist Obligations



- ✓ Only supply controlled drugs based on a legally valid prescription⁴ (be it paper or transferred via Healthmail), or, by way of an essential and legally compliant emergency supply.
- ✓ Be satisfied that the signature of the prescriber is genuine, and in the case of a prescription transferred by Healthmail, that it contains the professional registration number of the person issuing it, and can be traced back to the prescriber.
- ✓ Prior to supply, be satisfied as to the identity of the person or bona fide representative presenting the prescription or collecting controlled drugs.⁸
- ✓ Be vigilant for forgeries or unusual prescribing patterns.
- ✓ Store controlled drugs in a safe manner, in accordance with the relevant legislation.⁹
- ✓ Record all supplies of schedule 2 controlled drugs (CD2) in the pharmacy's controlled drugs register.¹⁰
- ✓ Communicate with the medical practitioner if there is any query about the prescription, or care of the patient.
- ✓ Adhere to national guidelines and facilitate appropriate withdrawal of controlled drugs. Ensure follow-up and referral to other healthcare professionals/services as necessary.

9. Prescriptions Required for CD2 and CD3 Drugs

Note: The figure below refers to paper-based prescriptions for schedule 2 and 3 controlled drugs. Where schedule 2 and 3 controlled drug prescriptions are transferred via Healthmail, it is not necessary for the particulars indicated below to be in the prescriber's own handwriting. Prescriptions sent by Healthmail do not need to include the prescriber's signature in so far as they can be traced electronically and securely back to the issuing practitioner.

Prescriptions Requirements for CD2 and CD3 Drugs¹¹

(Including handwriting requirements) 

for example of compliant prescriptions see Appendix 1a and 1b

Date

Patient name (First name and surname)

Patient address

Name of medicine 

Form (e.g. tablets) 

Strength (e.g. 10mg) 

Dose (e.g. 2 twice daily) 

Total quantity or total number of dosage units (Words and figures) 

Signature 

Name of prescriber
(First name and surname)

Address of prescriber

Profession
(eg. Medical Practitioner)

Registration number¹²

Telephone number

can be generic/INN or proprietary/brand name

Instalments
(Please see section 10 for further information)

- Number of instalments (e.g. 4)
- Intervals for dispensing (e.g. dispense weekly)

First instalment must be dispensed within 14 days
Last instalment must be dispensed within 2 months

Must be  handwritten

Address not required for health prescription (i.e. GMS Prescription)

Please note: Addressographs (sticker with patient name and address) are not acceptable as they are not considered indelible however it is acceptable to either print or handwrite these details.

10. Opioid Substitution Prescriptions and Prescriptions for Schedule 4 Part 1 CDs

The prescription requirements for opioid substitution prescriptions and schedule 4 part 1 CD prescriptions are shown in the box below.

Paper-based prescriptions must be signed by the prescriber in his or her usual signature.

Healthmail prescriptions for opioid substitution or schedule 4 part 1 CDs do not need to include the prescriber's signature in ink in so far as the prescription can be traced electronically and securely back to the issuing practitioner.

Note: Opioid Substitution Treatment Prescription Forms are a specific type of prescription form and are the only form which can be used for prescribing methadone or buprenorphine for opioid substitution therapy. Opioid substitution prescription forms can be written and signed by the prescriber, and physically presented in the pharmacy, or they may be scanned and forwarded to the pharmacy through Healthmail. Where Healthmail is used to transfer the prescription, there is no requirement for a paper copy to be posted to the pharmacy also.

Opioid Substitution prescriptions and prescriptions for schedule 4 part 1 CDs:¹³

The following information must be included on the prescription, but **does not need to be in the doctor's own handwriting** i.e. can be computer generated or handwritten.

- Name of medicine
- Form (e.g. liquid/oral solution)
- Strength (e.g 1mg/ml)
- Dose (e.g 50 mls daily)
- Total quantity or total number of dosage units (words **and** figures) (e.g. 350 mls (three hundred and fifty mls))

Note: An example of a schedule 4 part 1 private prescription, a schedule 4 part 1 GMS repeat prescription and a prescription for methadone are included in Appendix 2, 4 and 5.

11. Other Relevant Legal Requirements



Prescription Validity and Instalments

Prescriptions are valid for up to twelve months from the date of issue depending on the drug and the prescriber's instruction. Pharmacists are able to exercise professional judgement on extending the period of validity of prescriptions in certain circumstances from six months up to a maximum of 12 months. A prescriber may also indicate on a prescription that it is not to be extended. As highlighted earlier, schedule 2 and schedule 3 controlled drugs cannot be repeated.

Schedule 4 (part 1) controlled drugs may be repeated in line with the directions of the prescriber up to a maximum of six months. Prescriptions for schedule 4 (part 1) medicines cannot be extended by a pharmacist.

There are specific restrictions on the period of validity of controlled drug prescriptions:

1. The first instalment or supply on prescriptions for CD2 or CD3 drugs cannot be dispensed before the date of the prescription or later than 14 days after that date.
2. Unless it is a GMS prescription, a prescription for a CD2, CD3 or CD4 part 1 drug must include the address of the prescriber (address must be within Ireland).
3. Repeat prescriptions are not permitted in the case of CD2 or CD3 drugs.
4. Prescriptions for CD2, CD3 and CD4 part 1 may be dispensed in instalments, provided that the intervals between instalments and the number of the instalments are specified.
5. For CD2 and CD3 the first instalment must be dispensed within 14 days and the last instalment must be dispensed no later than two months from the date on the prescription.
6. Emergency supplies of CD2, CD3 or CD4 drugs, are permitted in limited and specific circumstances, and where they comply with the legislation.¹²
7. Emergency supplies of methylphenobarbitone, phenobarbitone, phenobarbitone sodium, midazolam, clobazam and clonazepam, are permitted where they comply with the legislation and are for the treatment of epilepsy.⁴

For ease of reference, Appendix 7 contains a table which clearly shows the prescription requirements for schedule 2, schedule 3 and schedule 4 part 1 controlled drugs.

Requisitions¹⁵

Requisitions for supplies of controlled drugs can be written privately or on a "health service requisition" form, by persons authorised under the legislation. Requisitions for controlled drugs must include:

- ✓ Name, address, and profession of the person requesting
- ✓ Signature of the person requesting Date of requisition
- ✓ Registration number of the person requesting¹²

- ✓ Name of the controlled drug
- ✓ Purpose of supply
- ✓ Total quantity of the medicine(s) to be supplied

The medical practitioner obtaining controlled drugs through the use of a requisition should be asked by the pharmacist to produce identification.

An example of a requisition can be found in Appendix 3.

12. Overarching Principles

12.1 Communication



Good communication is essential to the effective functioning of healthcare teams^{6,7}. The existence of good, open communication channels between pharmacists and prescribers is of particular importance in assuring the safe and efficient supply of controlled drugs to patients. By having in place a strong system of partnership, firmly based in the shared care of patients, efficiencies can be achieved in assuring that patients receive the best possible care in a more integrated system. These links are well established. In prescribing and dispensing controlled drugs, where there is any doubt or confusion around the prescription, dosage, supply and administration, pharmacists and medical practitioners should engage with each other without delay. This is particularly important at transitions of care and where the initial prescriber is not their usual GP, they should have clear communication with the patient's GP and the pharmacist, as necessary.

12.2 Identifying Risk Factors/High Risk Patients



As part of this collaborative relationship, pharmacists and prescribers should, in particular, communicate and collaborate in the care of patients in high-risk groups.

Examples include:

- ✓ Patients with drug dependency issues
- ✓ Patients receiving addiction treatment services
- ✓ Patients under the management of multiple doctors
- ✓ Patient population who attend pain clinics
- ✓ Patients transitioning from one place of care to another
- ✓ Patients in residential care settings
- ✓ Patients at risk of developing sleep disordered breathing
- ✓ Patients in the prison system
- ✓ Patients with mental health issues
- ✓ Patients living complex lives due to factors such as homelessness
- ✓ Patients with additional care requirements
 - ➔ Patients with poor health literacy or patients who don't have English as their first language
 - ➔ Paediatric patients
 - ➔ Older people
 - ➔ Pregnant/breast feeding women
 - ➔ Patients with physical/intellectual disabilities
 - ➔ Patients suffering from chronic illnesses
 - ➔ Patients receiving palliative care

Engagement within and between the professions in the shared care of patients in these groups is particularly important to meet patients' care needs. This includes shared responsibility for follow up and after-care for all patients, and collaborative working in the implementation of appropriate withdrawal procedures for controlled drugs.

12.3 Relevant National Resources and Professional and Clinical Guidance



It is essential in the prescribing, dispensing and supply of controlled drugs that pharmacists and prescribers are aware of and have access to national resources, and adhere to current national and international guidelines. For example:

- ✓ The Health Products Regulatory Authority (HPRA) website for the confirmation of the scheduling of medicines containing controlled drugs (e.g. is a medicine a CD2 or CD3) (See Appendix 6)
- ✓ The online registration databases of both the [Medical Council](#) and the [PSI](#): to confirm registration, find contact details and be aware of any conditions attached to the registration of healthcare professionals
- ✓ Resources or advice from relevant healthcare regulators, including the Medical Council, the PSI and the Health Information and Quality Authority (HIQA)
- ✓ National Clinical Guidelines from the National Clinical Effectiveness Committee and relevant faculties
- ✓ [HSE Clinical Guidelines for Opioid Substitution Treatment \(OST\)](#)
- ✓ HSE Medicines Management Programme Guidance on appropriate prescribing of Benzodiazepines and Z-drugs (BZRA) in the treatment of anxiety and insomnia
- ✓ Relevant international Guidelines from bodies such as the National Institute for Health and Care Excellence ([NICE](#))
- ✓ All relevant legislation is available at www.irishstatutebook.ie

The above list is for ease of reference only and should not be considered exhaustive.

This guide is not a legal document. Its intention is to facilitate prescribers and pharmacists in interpreting several pieces of legislation and provide a practical tool in the collaborative care of patients. Links to relevant legislation have been provided for direct consultation.

Appendix 1a

Example of Compliant Prescription for Schedule 2 and 3 CDs

Note: This example refers to a paper-based prescription for a schedule 2 controlled drug. Where schedule 2 and 3 controlled drug prescriptions are transferred via Healthmail, it is not necessary for the particulars indicated below to be in the prescriber's own handwriting. Prescriptions sent by Healthmail do not need to include the prescriber's signature in ink in so far as they can be traced electronically and securely back to the issuing practitioner.

Dr John Doe
Somewhere Rd
Co Dublin
Medical Practitioner
MCRN -XXXXX
Telephone 01 1234567

Prescriber address not required for health prescriptions i.e. GMS prescription⁶

1 March 2024

Please note: Addressographs (sticker with patient name and address) are not acceptable as they are not considered indelible, however, it is acceptable to either print or handwrite these details

Joe Bloggs
 Bloggstown
 Co. Dublin

Example of schedule 2 controlled drug

Morphine Sulphate 10mg Tablet
 Take one Tablet B.D.
 56 Tablets (Fifty-Six)

Dispense weekly (4 instalments) (14 tablets each week)

Installment instructions are not required to be handwritten

Dr John Doe

Note – Fine Blue font indicates handwritten details

Appendix 1b

Example of Compliant Prescription for Schedule 2 and 3 CDs

Note: This example refers to a paper-based prescription for a schedule 2 and 3 controlled drug. Where schedule 2 and 3 controlled drug prescriptions are transferred via Healthmail it is not necessary for the particulars indicated below to be in the prescribers own handwriting. Prescriptions transferred via Healthmail do not need to include the prescriber's signature in ink in so far as they can be traced electronically and securely back to the issuing practitioner.

**Dr John Doe
Somewhere Rd
Co Dublin
Medical Practitioner
MCRN -XXXXX
Telephone 01 1234567**

Prescriber address not required for health prescriptions i.e. GMS prescription

1 March 2024

Joe Bloggs
Bloggstown
Co. Dublin

Please note: Addressographs (sticker with patient name and address) are not acceptable as they are not considered indelible, however, it is acceptable to either print or handwrite these details

Morphine Sulgate 10mg Tablet
Take one Tablet B.D.
56 Tablets (Fifty-Six)

Example of
schedule 2
controlled drug

Temazepam 10mg Tablets
Take one Tablet Nocte PRN
Mitte 21 Tablets (Twenty One)

Example of
schedule 3
controlled drug

Dr John Doe

Note – Fine Blue font indicates handwritten details

Appendix 2

Example of Compliant Prescription Schedule 4 Part 1 CD

Note: This example refers to a paper-based prescription for a schedule 4 part 1 controlled drug. Where schedule 4 part 1 controlled drug prescriptions are transferred via the Healthmail, it is not necessary to include the prescriber's signature in ink in so far as the prescription can be traced electronically and securely back to the issuing practitioner.

Dr John Doe
Somewhere Rd
Co Dublin
Medical Practitioner
MCRN -XXXXX
Telephone 01 1234567

Prescriber address not required for health prescriptions i.e. GMS prescription

1 March 2024

Joe Bloggs
Bloggstown
Co. Dublin

Please note: Addressographs (sticker with patient name and address) are not acceptable as they are not considered indelible, however, it is acceptable to either print or handwrite these details

Diazepam 5mg Tablets
Take One Tablet Nocte
7 Tablets (Seven)

These details do not need to be handwritten for a schedule 4 part 1 controlled drug prescription

Repeat x 2

Dr John Doe

Appendix 3

Example of a Compliant Requisition for Schedule 2, 3 and 4 Part 1 CDs

**Dr John Doe
Somewhere Rd
Co Dublin
Medical Practitioner
MCRN -XXXXX
Telephone 01 1234567**

1 March 2024

Morphine Sulphate 10mg/ml Solution for Injection
Mitte 10 Ampoules

Midazolam 5mg/ml, 2ml Ampoules, Solution for Injection
Mitte 10 Ampoules

For use on-call: Patients receiving palliative care

Dr John Doe

Appendix 4

Example of GMS Repeat Prescription for a Schedule 4 Part 1 CD

Note: This example refers to a paper-based GMS prescription for a schedule 4 part 1 controlled drug. Where schedule 4 part 1 controlled drug prescriptions are transferred via the national electronic prescription transfer system, it is not necessary to include the prescribers signature in ink in so far as the prescription can be traced electronically and securely back to the issuing practitioner.

NOTE: PLEASE USE A BALLPOINT PEN AND APPLY AMPLE PRESSURE TO PRODUCE CLEAR DETAIL ON ALL COPIES.

GENERAL MEDICAL SERVICES
REPEAT PRESCRIPTION SET - PHARMACY CLAIM FORM

PATIENT'S NAME Joe Bloggs MEDICAL CARD NO. 1234567XX VALID TO 01/08 ADDRESS Bloggstown, Co.Dublin		DATE PRESCRIBED 01 03 24	PRESCRIPTION SERIAL NO. R12	PHARMACY SEQUENCE NO.
DOCTOR NO. 7654321XX DR'S NAME Dr John Doe ADDRESS Somewhere Road Co. Dublin MEDICAL COUNCIL REG NO.		AGE OF PATIENT IF U/12 		
		THIS IS THE ORIGINAL PRESCRIPTION FORM AND MUST BE PRESENTED TO YOUR PHARMACIST AT EACH DISPENSING		
DOCTOR'S NAME AND ADDRESS AND PHONE NUMBER Dr John Doe Somewhere Road Co. Dublin Telephone 01 1234567				
DOCTOR'S SIGNATURE Dr J Doe				

Prescriber address not legally required for health prescriptions ie. GMS prescriptions. However, it is still best practice to include this

USE ONLY FOR ITEMS TO BE REPEATED	PRECISE STRENGTH, QUANTITY AND DOSAGE MUST BE STATED	PHARMACISTS MUST COMPLETE THIS PART		
		DRUG CODE	QUANTITY SUPPLIED	FOR OFFICIAL USE ONLY
Clobazam 10mg Tablets. Take One Tablet Twice Daily. Mitte 56 Tablets (Fifty-six)				
Repeat x 2				

'Repeat' must always be stated on schedule 4 part 1 controlled drug repeat prescriptions, even on GMS repeat prescriptions

3rd DISPENSING PHARMACY STAMP: COMPUTER NO. AND DATE OF DISPENSING	2ND DISPENSING PHARMACY STAMP: COMPUTER NO. AND DATE OF DISPENSING	1st DISPENSING PHARMACY STAMP: COMPUTER NO. AND DATE OF DISPENSING
--	--	--

SIGNATURE OF PHARMACIST _____

DATE OF 3RD DISPENSING | | | | |

Patient Signature
 Received by: _____
 If not patient, please indicate relationship: _____

PART 3
GMS002

THIS PART WILL BE RETAINED BY YOUR PHARMACIST ON THE THIRD AND FINAL DISPENSING

Appendix 5 Example of Opioid Substitution Prescription

Note: This example refers to a paper-based opioid substitution prescription for Methadone. Where opioid substitution prescriptions are transferred via Healthmail, it is not necessary to include the prescriber's signature in ink in so far as the prescription can be traced electronically and securely back to the issuing practitioner. All prescriptions for opioid substitution, whether paper based or transmitted through Healthmail, must be prescribed using the standard opioid substitution prescription form.

METHADONE PRESCRIPTION FORM

PHARMACY SEQUENCE NO.
SERIAL NO. **3597039**

PATIENT DETAILS:

SURNAME **BLOGGS**
FIRST NAME **JOE**

ADDRESS **BLOGSTOWN
CO DUBLIN**
PATIENT'S AGE IF UNDER 12 YEARS
TREATMENT CARD NO. **P.H. PH: 302167**

PRESCRIPTION DETAILS

DATE PRESCRIBED
01 03 24

DRUG NAME, FORM AND STRENGTH
**Methadone DTF 1mg/ml
oral solution**

FROM	TO	DOSE (mg PER DAY)	NO. OF DAYS AT DOSE	TOTAL (mg) ON FIGURES
01/03/24	07/03/24	30	7	210

TOTAL (mg) IN WORDS: **Two Hundred and Ten Mls**
(NOT MORE THAN 7 DAYS SUPPLY SHOULD BE PRESCRIBED EXCEPT IN EXCEPTIONAL CIRCUMSTANCES)

DOCTOR'S SIGNATURE
DOCTOR'S NO. **0 2 3 4 5**
DOCTOR'S NAME, ADDRESS AND TELEPHONE NUMBER OR STAMP

Dr John Doe
Somewhere Road
Co Dublin
Medical Practitioner
MCRN XXXXXX
Telephone 01 1234567

INSTALMENT INSTRUCTIONS
INTERVALS
DAILY ☒
DAILY (WITH DOUBLE ON SAT.) ☒
OTHER (TICK) ☐ MON TUE WED THURS FRI SAT SUN

SUPERVISION INSTRUCTIONS
YES NO
SUPERVISED ☒ ☐
MONDAY TO SATURDAY ☒ ☐
EVERYDAY ☐
OTHER (TICK) ☐ MON TUE WED THURS FRI SAT SUN

*** ONLY WITH PRIOR AGREEMENT OF PHARMACIST**

PHARMACY SECTION

PHARMACY GMS NO.
NO. OF
PHARMACY NAME & ADDRESS STAMP

DRUG CODE
QUANTITY (mls)
NO. OF INSTALMENTS
SUPERVISED DAYS YES NO SUPERVISED ☐

DATE DISPENSED
QUANTITY (mg)
PHARMACIST'S INITIALS

I VERIFY THAT I HAVE DISPENSED THE ITEM(S) SPECIFIED HEREON
PHARMACIST'S SIGNATURE _____
I VERIFY THAT I HAVE RECEIVED THE ITEM(S) SPECIFIED HEREON
SIGNATURE _____
OF PATIENT OR PATIENT'S REPRESENTATIVE

TO BE COMPLETED IN THE CASE OF NON-OPIATE DEPENDENT PERSON

NAME AND ADDRESS OF INITIATING CONSULTANT
HOSPITAL _____ MEDICAL CARD NO.
HEALTH SERVICES SCHEME
GMS ☐ EEA ☐ OTHER ☐

Appendix 6

HPRA Website – Information on Medicines Classification

Step 1 – Access HPRA Home (www.HPRA.ie)

HPRA An tÚdarás Rialála Táirgí Sláinte Health Products Regulatory Authority

Follow @TheHPRA | Contact us | As Gaelige

My HPRA: [Login](#) [Register](#) Search our website

ABOUT US MEDICINES VETERINARY MEDICAL DEVICES COSMETICS CONTROLLED SUBSTANCES BLOOD, TISSUES, ORGANS

Medicines Availability

Keeping patients and healthcare professionals informed of resolved and current medicines shortages notified to the HPRA. [See more](#)

Current shortages Reporting a shortage

Find a medicine

Medicines Veterinary Medicines Generics

Enter a Trade Name, Active Substance or Licence Number...

View all Medicines Advanced Search

I want to:

Report an issue See guides & forms

Step 2 - Enter product name and select product

HPRA An tÚdarás Rialála Táirgí Sláinte Health Products Regulatory Authority

Follow @TheHPRA | Contact us | As Gaelige

My HPRA: [Login](#) [Register](#) Search our website

ABOUT US **MEDICINES** VETERINARY MEDICAL DEVICES COSMETICS CONTROLLED SUBSTANCES BLOOD, TISSUES, ORGANS

Medicines > Medicines Information > Find a medicine

The HPRA advises healthcare professionals not to retain printed versions of Summary of Product Characteristics (SPC) documents. As these documents are subject to frequent content updates, including changes to safety and dose related information, we recommend that you visit our website as necessary to access the most up-to-date versions. If you have any queries about specific SPCs on our website then please e-mail info@hpra.ie

BuTrans 5 micrograms/hour transdermal patch + Follow medicine Print

Mundipharma Pharmaceuticals Limited PA1688/002/001

Main Information

Trade Name

BuTrans 5 micrograms/hour transdermal patch

Active Substances Buprenorphine

Step 3 - Scroll down to 'Status' and 'Conditions of License' for details on CD Classification

Status	
License status	Authorised
Licence Issued	24/06/2005
Legal status	
Product subject to prescription which may not be renewed (A)	
Supply Status	Supply through pharmacies only
Advertising Status	Advertising to healthcare professionals only
Conditions of Licence	
This product contains a substance listed in schedule 2 to the Misuse of Drugs Regulations 2017.	
Marketing Status	Marketed

Appendix 7: Controlled Drug Prescription Requirements

TABLE 1 – Requirements for Paper and Healthmail Prescriptions for CD2, CD3 and CD4 part 1

Legal requirements	Schedule 2 and 3*	Schedule 4 part 1
Must be written in ink/indelible	Paper: Yes (addressograph sticker not acceptable) Healthmail: Can be issued electronically via Healthmail. Pharmacy must print a copy.	Paper: Yes (addressograph sticker not acceptable) Healthmail: Can be issued electronically via Healthmail. Pharmacy must print a copy.
Include Full Name (including first name) of practitioner	Yes	Yes
Include Practitioner's Registration Type and Number	Yes	Yes
State the Date of issue	Yes	Yes
Must be dispensed within 14 days of date of issue	Yes	No
Cannot be dispensed before date of issue stated on prescription	Yes	Yes
Must be signed by the prescriber in his/hers usual signature	Paper: Yes Healthmail: Not necessary	Paper: Yes Healthmail: Not necessary
State the Address of Practitioner	Yes, except for Health prescriptions	Yes, except for Health prescriptions
Practitioner must be registered in Ireland with an address within the state	Yes	Yes
Must include Practitioner's Telephone number	Yes	Yes
Must include Name (including first name) and address of the patient	Yes	Yes
Must state the name of controlled drug	Paper: Yes, <i>and must be handwritten</i> Healthmail: Yes, but does not need to be handwritten	Yes, but does not need to be handwritten
Must state the dose, form, and strength of controlled drug	Paper: Yes, <i>and must be handwritten</i> Healthmail: Yes, but does not need to be handwritten	Yes, but does not need to be handwritten
Must state Total Quantity in words and figures	Paper: Yes, <i>and must be handwritten</i> Healthmail: Yes, but does not need to be handwritten	Yes, but does not need to be handwritten
Repeats allowed	No	Yes
Emergency supply permitted	Yes	Yes

* All stocks of Schedule 2 and 3 controlled drugs in the pharmacy must be stored in the CD safe in line with legislative requirements pertaining to safe custody. All receipts and supplies of Schedule 2 controlled drugs must be recorded in the controlled drug register in accordance with legislative requirements pertaining to controlled drug record keeping.

References

1. Department of Health Guidance on the Medical Cannabis Access Programme is available [online](#).
2. Medicinal Products (Prescription and Control of Supply) (Amendment) Regulations 2020
3. Regulation 7(11) of Medicinal Products (Prescription and Control of Supply) Regulations 2003 (as amended)
4. Regulation 8 of the Medicinal Products (Prescription and Control of Supply) Regulations (as amended)
5. Misuse of Drugs Acts 1977 to 2016 and the Misuse of Drugs Regulations 2017 (as amended)
6. A Guide to Professional Conduct and Ethics for Registered Medical Practitioners (9th Edition, 2023) page 38, paragraph 35.1
7. PSI Code of Conduct- Professional Principles, Standards and Ethics for Pharmacists, 2019, page 2
8. Regulation 16 (1)(f) of the Misuse of Drugs Regulations 2017 (as amended)
9. Misuse of Drugs (Safe Custody) Regulations 1982 (as amended)
10. Regulation 19 of the Misuse of Drugs Regulations 2017 (as amended)
11. Regulation 15 of the Misuse of Drugs Regulations 2017 (as amended)
12. Section 43 (8) Medical Practitioners Act 2007 and Regulation 15(2)(b) of the Misuse of Drugs Regulations 2017 (as amended)
13. Specific handwriting exemptions provided for Methadone and Schedule 4 Part 1 prescriptions, Regulation 15(4) of the Misuse of Drugs Regulations 2017 (as amended)
14. Regulation 8(3) and 8(4) of the Medicinal Products (Prescription and Control of Supply) Regulations 2003 (as amended)
15. Regulation 14(2) of the Misuse of Drugs Regulations 2017 (as amended)
16. Regulation 15 (2)(d) of the Misuse of Drugs Regulations 2017