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Compounding Guidelines guidance will help to assure the safe and appropriate preparation and supply of extemporaneously prepared medicinal products to patients, where the supply of such products is necessary. An extemporaneously prepared medicinal product refers to the process by which a pharmacist, using traditional

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Preparation-Mark Jackson 2010 A
comprehensive and easy-to-follow guide to
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EXTEMPORANEOUS COMPOUND

GUIDELINES Page 3 of 3 Type of

Preparations Maximum Payable

Methadone Per Carry – 5 minutes

Suppositories Suppositories (Less than 30)

1 supp/minute Suppositories (30-49) 30

minutes Suppositories (50-100) 45 minutes

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Suppositories (101 and over) 30
seconds/supp. Capsules Capsules (Less
than 100) 3 caps/minute

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by the Pharmacy Board of Australia ' s
guidelines on compounding.1 Regulation

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The final medicine produced by compounding is regulated according to the component ' s schedule in the Poisons Standard (the SUSMP).³ For example a topical progesterone (S4) cream requires a

Introduction Extemporaneous compounding is the preparation

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medicines

Extemporaneous Preparation A guide to pharmaceutical compounding Edited by Mark Jackson BSc, MPhil, MRPharmS Deputy Director, QCNW / Head of QA/QC, Liverpool Pharmacy Practice Unit, Liverpool, UK Andrew Lowey

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DPharm, MRPharmS Clinical Pharmacy
Manager, Leeds Teaching Hospitals,
Leeds, UK On behalf of The NHS
Pharmaceutical Quality Assurance
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Compounding Guidelines'. The guidance should be read alongside the standards for registered pharmacies. These aim to create and maintain the right environment, both organisational and physical, for the safe and effective practice of pharmacy. By following this guidance, the pharmacy will:

- demonstrate that it

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(Where a manufacturer's instructions are not followed, for example a different diluent is used, this is considered compounding.) Interpretation of the basic GMP requirements Where a clause number or an Annex is not listed, there is no specific interpretation provided for manufacture of extemporaneously

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~~Compounded medicines and good manufacturing practice (GMP ...~~
Development of national guidelines to promote standards of practice in the community and/or home setting is urgently needed to help improve the safety

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Compounding Guidelines
of dispensing and handling oral
chemotherapeutic agents, including
extemporaneously compounded oral liquid
formulations of these drugs.

~~Extemporaneous compounding of oral
liquid dosage ...~~

3. Extemporaneous preparations should be

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done based on evidence-based references.

4. Always check for the suitability of the product/brand for extemporaneous preparations.
5. Preparations listed in this manual should be done according to what is stated as far as possible unless stated otherwise in the product leaflet.
- 6.

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~~Extemporaneous FORMULATION~~ pharmacy

Pharmacists are responsible for ensuring that extemporaneous preparations are compounded according to compounding guidelines and standards with respect to purity, quality, stability, packing, record keeping, and other appropriate pharmacy

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~~Extemporaneous Compounding:
Cautions, Controversies and ...~~

Guidelines on compounding of medicines.

PDF (115 KB) Word (393 KB,DOCX) 28

April 2015 1 February 2018 for section 6.2

Compounding of sterile injectable

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Compounding Guidelines
medicines: Joint statement on
compounded medicines – Pharmacy
Board of Australia and Medical Board of
Australia; PDF (77.5KB)

~~Pharmacy Board of Australia – Codes,
Guidelines and Policies
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Compounding Guidelines
medicines may be useful when a required dose or dose form is unavailable commercially, or for individualised dosing. There are numerous established compounding formulae available, and new formulae may be developed with the help of formulation guidelines and professional advice.

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~~Extemporaneously compounded
medicines – Australian Prescriber~~

General guidance for compounding oral liquids 1. Funded proprietary oral liquid medicines When a funded commercial preparation is available this must be used. Extemporaneous compounding increases

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the risk of error that could harm the patient through overdosing or underdosing. Using a commercial preparation reduces the risk by removing compounding

~~General guidance for compounding oral liquids~~

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Compounding Guidelines
In some situations compounding of a medicine may be the only option when there is no appropriate dosage form available. One must always strive to deliver medication in the safest

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