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DEVELOPMENT OF AN ORAL KETAMINE COMPOUNDING AND CREATION OF A PHARMACEUTICAL CARE CIRCUIT FOR PHANTOM LIMB SYNDROME

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1.WHAT WAS IT DONE?

Development and validation of an oral ketamine compound and a specific pharmaceutical care circuit (PCC) as a part of the treatment of phantom limb syndrome (PLS).

2.WHY WAS IT DONE?

The PLS is the perception of a non-existent limb that may occur in up to 80% of amputees. The management is complex, The absence of a marketed formulation, off-label use of drugs and the complex treatment of pain make the <u>role of the</u> pharmacist essential.

3.HOW WAS IT DONE?

- ✓ **Literature research** was carried out on the preparation of this compounding, as well as on the use of oral ketamine (bioavailability, dosage, adverse reactions...).
- ✓ **Compounding:** <u>oral solution of 10mg/ml</u> was prepared (final volume 50ml: 500mg of injectable ketamine solution or raw material, 20ml of simple syrup with a sufficient amount of purified water and 2 drops of lemon essence).

To establish the expiration date recommendations of Good Manufacturing Practice Guideline were followed and the organoleptic characteristics were evaluated for quality control.

✓ Pharmaceutical care circuit:

- Setting up a first presential visit to provide pharmaceutical care during admission: to inform the storage conditions, most common adverse effects and recommendations about medication intake.
- o Initially appointments every 7 days for a closer follow-up: <u>control of adverse reactions (confusion, agitation, nausea, etc.)</u>, monitoring of the appropriate use of ketamine and other analgesic medication (<u>avoiding possible abuse and addictive behaviour</u>) and pain control. Pharmaceutical interventions are communicated to the pain management unit (PMU).
- Spacing of visits fortnightly once the treatment is well-stablished and proposing a <u>telepharmacy</u> service.

4.WHAT HAS BEEN ACHIEVED?

The ketamine formulation developed has been used in our hospital in three patients with satisfactory results. The **interventions** carried out were: pain control problems, possible inappropriate use, reduction in the number or dosage of concomitant medication or ketamine itself.

5.WHAT NEXT?

The capacity to provide therapeutic alternatives and a more exhaustive pharmacological control of pain in collaboration with the PMU can **improve the safety and effectiveness of these treatments.**





