

Medication Safety Series: MSS 21

Pharmacological Management of Severe Agitation

Key references: Rapid Tranquillisation Policy (Trust intranet); NICE: [NG10](#)

Key principles:

- Individuals with behaviours that challenge should be identified, risk assessed and have a regularly reviewed intervention plan - this can include pro-active prescription & administration of **oral** PRN medication.
- Rapid tranquillisation should only be used where a patient is highly aroused, agitated, overactive or aggressive, or is making serious threats towards themselves or others, or is being destructive to their surroundings, when pro-active interventions have been ineffective in supporting a reduction in such behaviour.

What is Rapid Tranquillisation (RT):

- ✓ RT is a **reactive** management strategy; it is the **intramuscular** administration of medication, usually under restraint, to calm or sedate an agitated, violent or aggressive patient.
- ✗ RT is not the proactive use of oral PRN medication to manage an agitated patient to prevent escalation to violence or aggression

Administration of Rapid Tranquillisation:

- ! **Physical restraint should not be used for more than 10 minutes without considering RT or seclusion.** RT or seclusion may be used sooner if deemed clinically appropriate; conversely, physical restraint may continue without RT or seclusion if the event is resolving, but the need for RT or seclusion must be continuously reconsidered.
- ✗ Never mix two drugs in the same syringe
- ✓ Use a site for IM administration which maintains patient dignity and reduces risk.

Post-administration monitoring:

- ✓ The highly aroused condition of the patient may intensify the effects of RT medication. Physiological observations must be monitored for 3 hours after RT administration, using the NEWS2 tool.

! NEVER events:

- ✗ **NEVER** use RT prior to pro-active interventions, unless the patient's behaviour presents a serious risk to themselves, other patients, staff or the physical ward environment.
- ✗ **NEVER** administer a combination of drugs for RT in the same syringe.
- ✗ **NEVER** leave a patient unmonitored for physical or mental health deterioration after administration of RT.

Prescribing Rapid Tranquillisation (RT):

- ! The MHA/MCA status of the patient must be considered before RT medication is prescribed
- ✗ **DO NOT routinely** prescribe RT medication on admission.
- ✓ If an event is highly likely, a single dose may be prescribed as "once only" on EPMA for initial management. A doctor **MUST** attend or be consulted before a second dose is prescribed / administered.
- ✓ In the absence of medical staff on the ward (e.g. over a weekend), the administration of second and subsequent doses of RT must be discussed with the duty doctor on-call and, if possible, the duty nurse co-ordinator.
- ✓ RT may be prescribed "PRN" for individuals who have undergone a thorough psychiatric assessment prior to or during admission which has identified a high risk of disturbed behaviour likely to result in violence and aggression which would require restraint and intervention.
- ✓ The rationale for PRN prescribing must be recorded in the electronic patient record (EPR).
- ✓ All prescriptions of RT medication must be reviewed within 72 hours of admission, and regularly thereafter, and discontinued when appropriate to do so.
- ✓ Drugs suitable for RT:
 - First-line options - Haloperidol* +/- promethazine (off-label), **OR** lorazepam
 - Alternatives available for IM administration: Aripiprazole & olanzapine
- ✗ Do not use zuclopenthixol acetate (Clopixol Acuphase®), chlorpromazine and diazepam for RT
- ! Caution is required in antipsychotic naive patients e.g. risk of NMS
- ✓ See the [RT policy](#) for age-specific algorithms
- ! Consider anti-cholinergic burden in the elderly
- ✓ Prescribe RT medication separately to regular or PRN prescriptions for the same drug orally
- ✗ Do not use two drugs of the same class for RT
- ✓ Prescribe PRN procyclidine in case of dystonia with haloperidol

*baseline ECG recommended prior to use; if hasn't been done, or QT-interval prolonged, document risk vs benefit in EPR if prescribed/administered.

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