


Name of Policy: <u>Therapy Management by the Pharmacist</u> Policy Number: 3364-133-143 Department: Pharmacy Approving Officer: Senior Hospital Administrator, Chair of P&T Responsible Agent: Director of Pharmacy Scope: University of Toledo Medical Center	 Effective Date: 9/29/2022 Initial Effective date: 9/29/2022
<input checked="" type="checkbox"/> New policy proposal <input type="checkbox"/> Minor/technical revision of existing policy <input type="checkbox"/> Major revision of existing policy <input type="checkbox"/> Reaffirmation of existing policy	

(A) Policy Statement

To provide a condensed guideline for pharmacist management of drug therapy to improve medication safety and collaborative practice with UTMC providers. This policy was approved by the Medical Executive Committee on September 28, 2022.

(B) Policy Purpose

Define criteria for pharmacists to add/discontinue/change drug therapy.

(C) Procedure

1. A pharmacist may adjust medication therapy in accordance with attachment A if criteria are met.
2. If any criteria are questionable or unable to be evaluated, no changes will be made by the pharmacist and recommendations, or follow-up questions will be communicated with the provider.
2. Pharmacist will place orders in the electronic medical record as a per protocol, no co-signature order under the patient’s attending physician or under the physician who ordered the pharmacy consult (if applicable).
3. Pharmacist will log their activity into pharmacy intervention software and flag for follow up (as applicable).

Attachment A

Drug Type	Intervention with Order
Oral bisphosphonates	Discontinue during admission but maintain on Home Medication List.
Herbals/alternative products/probiotics (non- FDA approved products not on formulary)	Discontinue during admission but maintain on Home Medication List.
Critical Care Analgesia & Sedation orders	<p>When patient is extubated, discontinue sedation and analgesia drips and associated prn doses from this protocol</p> <p>When level of sedation is modified (e.g. changing from level 1 to level 2 sedation) discontinue sedation and analgesia drips and associated prn doses from the previous protocol</p>
Continuous Renal Replacement Therapy (CRRT)	<ul style="list-style-type: none"> • When patient is no longer on CRRT AND the pharmacist has confirmed it will not be restarted in the immediate future, discontinue CRRT specific medications. • When CRRT medication orders are modified, discontinue medication orders from the previous protocol. • If patient is on a heparin drip, clarify that the heparin drip is not being used for another indication (in addition to CRRT) before discontinuing & ensure appropriate VTE prophylaxis is ordered
IV infusions	<p>Discontinue if drip has not been used for >48 hours.</p> <p>May discontinue current maintenance IV fluid if 1) new maintenance IV fluid is prescribed and 2) there is no documented reason for the patient to receive two maintenance IV fluids.</p> <p>May order a flush bag for small volume IV solutions to ensure entire volume flushes through the IV line.</p>
Immediate Release Solid oral dosage forms with a liquid equivalent	Pharmacists may change to the liquid formulation of a medication at the equivalent dose and route if the immediate release solid oral dosage form is not stocked or cannot be dispensed in the dosage needed

<p>Multivitamin and mineral supplements</p>	<p>Discontinue during admission but maintain on Home Medication List. Exceptions: 1) Nephrocaps 2) Pregnancy 3) Alcohol dependence or malnutrition 4) Iron supplements</p>
<p>Patients with previously placed Implantable Intrathecal Pumps or Insulin Pumps</p>	<p>Upon learning that a patient has an implantable intrathecal pump or insulin pump, the pharmacist will make a reasonable attempt (i.e. patient interview, provider office, pump interrogation device) obtain the following information:</p> <ul style="list-style-type: none"> • Medication • Date of next refill • Name of provider who manages the pump <p>The pharmacist will communicate this information via an entry on the MAR. Medication(s) will be designated as “patient supplied”. The entry will be signed as a Standard Order.</p>
<p>Short-acting Muscarinic Agonists: Ipratropium</p>	<p>If a patient is receiving scheduled Ipratropium or scheduled albuterol/ipratropium (Duoneb) nebulization, and tiotropium (Spiriva) or another anticholinergic bronchodilator is ordered, the pharmacist will:</p> <ol style="list-style-type: none"> 1) Follow approved Formulary Therapeutic Interchanges, if applicable, for ordered anticholinergic bronchodilator 2) Discontinue the ipratropium nebulization component if a patient is to continue on an anticholinergic bronchodilator such as tiotropium. 3) Ensure albuterol nebulization remains the same dose/frequency as it was originally part of the combination albuterol/ipratropium nebulization treatments.
<p>Oral chemotherapy for cancerous indications ordered as continuation of home medication while inpatient</p>	<p>If not ordered by an Oncologist in the inpatient setting:</p> <ol style="list-style-type: none"> 1) Pharmacist will place a consult to Hematology/Oncology physician to authorize safety and appropriateness of continuation of home medication while inpatient. 2) If no rounding Hematology/Oncology services are available, ordering provider to consult patient’s outpatient prescriber of the oral antineoplastic for direction on the safety and appropriateness of continuation inpatient.
<p>IV Levothyroxine (Synthroid)</p>	<p>Upon order verification for IV levothyroxine, pharmacists will automatically re-time IV levothyroxine for 120 hours (5 days) after admission or NPO order (up to 5 total days maximum without any form of levothyroxine), except patients meeting one of the below criteria would remain on IV therapy:</p> <ol style="list-style-type: none"> 1. Recommendation by endocrinology to begin IV therapy 2. Myxedema coma 4. TSH > 5 mcIU/mL within past 60 days, if lab available 5. Organ procurement

