

Minutes  
Drug Utilization Review Board Meeting  
DATE: December 9, 2015



**Meeting Purpose:** Quarterly Open Board Meeting  
Meeting opened at 6:00 p.m. by Chair, Sarah McGee

**Attendance:** Adam Bard Burrows, M.D.; Timothy Fensky, R.PH.; Leslie S. Fish, Pharm.D.; Karin Gardner Johnson, M.D.; Sarah M. McGee, M.D.; Sophie McIntyre, Pharm.D.; Patrick Reilly, R.PH.; Christy Stine, M.D.

**Absent:** Audra R. Meadows, M.D., MPH; Sherry Nykeil, M.D.; Karen Ryle, M.S., R.PH; Arthur Yu-shin Kim, M.D.

**Agenda Items:**

- I. Welcome and Introductory Remarks
- II. DUR Clinical Update
- III. Pediatric Behavioral Health Medication Initiative Update
- IV. Diabetic Testing Supplies Quality Assurance Analysis
- V. Omalizumab (Xolair) Quality Assurance Analysis
- VI. MHDL Update
- VII. DUR Operational Update
- VIII. MassHealth Update

Agenda Item	Discussion	Conclusions/Follow Up
Review of Minutes	Motion to accept the September Quarterly Board minutes	<u>Follow Up</u> Informational
Action	Accepted	

Agenda Item	Discussion	Conclusions/Follow Up
DUR Clinical Update	Fiscal Year 2015	<u>Follow Up</u> Informational
Action	<ul style="list-style-type: none"><li>➤ Reviewed clinical projects and accomplishments<ul style="list-style-type: none"><li>➤ Total number of NDRs: 73</li></ul></li><li>➤ Summarized/quantified new drug reviews, guidelines, and proposals<ul style="list-style-type: none"><li>➤ Total number of guidelines: 146</li><li>➤ Total number of QAs: 48</li><li>➤ Clinical Proposals: 4</li></ul></li></ul>	

	<ul style="list-style-type: none"> <li>➤ Discussed progress related to MassHealth clinical initiatives</li> <li>➤ Total cases sent to clinical review: 1,924</li> <li>➤ Clinical Initiatives <ul style="list-style-type: none"> <li>○ Cardiovascular Agents</li> <li>○ Biosimilars</li> <li>○ Cystic Fibrosis</li> <li>○ Hepatitis C</li> <li>○ Opioid Management</li> <li>○ PBHMI</li> <li>○ Progesterone Agents9</li> <li>○ PCSK</li> <li>○ Synagis</li> </ul> </li> <li>• Provided an overview of pharmacy resident and student contributions on the DUR rotation.</li> <li>• Presented an overview of disseminated work related to the MassHealth pharmacy program.</li> <li>• Described trends in clinical projects.</li> </ul>	
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Agenda Item	Discussion	Conclusions/Follow Up
<b>PBHMI Update</b>	Pediatric Behavioral Health Medication Initiative Update	<b><u>Follow Up</u></b> Informational
<b>Action</b>	<ul style="list-style-type: none"> <li>➤ Presented an overview of the PBHMI Initiative including <ul style="list-style-type: none"> <li>➤ Requirement and guidelines</li> <li>➤ Reviewed Implementation Timeline</li> <li>➤ Total PAs from 11/24/14 to 11/30/15: 18,478</li> </ul> </li> <li>➤ Provided overview of the Therapeutic Class Management (TCM) Workgroup <ul style="list-style-type: none"> <li>➤ Workgroup members</li> <li>➤ Responsibilities</li> <li>➤ Challenges</li> <li>➤ Lessons Learned and Successes</li> </ul> </li> <li>➤ Provided analyses for the future <ul style="list-style-type: none"> <li>➤ Members with denied PBHMI pharmacy claims and behavioral health medical claims (e.g., hospitalization, emergency room visits)</li> <li>➤ Collaboration with other state agencies to identify PBHMI longitudinal outcomes and trends</li> <li>➤ Identifying care coordination and other services to provide optimal member care</li> <li>➤ Impact of PBHMI on prescribing trends based on submitted PA requests and pharmacy claims</li> </ul> </li> </ul>	

Agenda Item	Discussion	Conclusions/Follow Up
<b>Diabetic Testing Supplies QA Analysis</b>	Quality Assurance Analysis for Diabetic Testing Supplies	<b><u>Follow Up</u></b> Informational
<b>Action</b>	<ul style="list-style-type: none"> <li>• Described necessity and recommended frequency of self-monitoring of blood glucose (SMBG) in the management of diabetes mellitus.</li> <li>• Reviewed the implementation timeline of MassHealth diabetic testing supplies initiative.</li> <li>• Discussed changing trends in test strip utilization.</li> <li>• Highlighted sampling of prior authorization requests.</li> <li>• Discussed recommendations and next steps based on quality assurance analysis results.</li> <li>• Presented conclusions <ul style="list-style-type: none"> <li>➤ There has been significant adoption of preferred diabetic testing supplies.</li> <li>➤ Pharmacy program cost on average is \$70/claim prior to rebate.</li> <li>➤ Prior authorization requests reviewed appropriately</li> <li>➤ Approvals &amp; Denials</li> </ul> </li> <li>➤ Presented recommendations <ul style="list-style-type: none"> <li>➤ No significant recommended changes to the PA process.</li> <li>➤ Requests for Accu-Chek Aviva that state that the member requires the bolus dose calculating function will be reviewed on a case-by-case basis.</li> </ul> </li> </ul>	

Agenda Item	Discussion	Conclusions/Follow Up
<b>Omalizumab QA Analysis</b>	Quality Assurance Analysis of Omalizumab (Xolair)	<b><u>Follow Up</u></b> Informational
<b>Action</b>	<ul style="list-style-type: none"> <li>• Discussed background information on Xolair (omalizumab) and its use in clinical practice for the management of allergic asthma and chronic idiopathic urticaria (CIU).</li> <li>• Evaluated recent utilization and cost data for MassHealth members.</li> <li>• Presented an overview of current prior authorization (PA) requests for Xolair <ul style="list-style-type: none"> <li>➤ 84 Approvals</li> <li>➤ 37 Denials</li> </ul> </li> <li>• Reviewed historical comparison of utilization from last evaluation.</li> <li>• Discussed recommendations to current MassHealth clinical criteria. <ul style="list-style-type: none"> <li>➤ Presented recommendations <ul style="list-style-type: none"> <li>○ Minor changes were recommended for approval criteria of CIU: requirement of prescriber to be an allergist/immunologist or dermatologist or to provide consultation notes from one.</li> <li>○ Clarification provided to consultants when reviewing initial requests for doses &gt; 150 mg every 4 weeks for CIU.</li> </ul> </li> </ul> </li> </ul>	

	<ul style="list-style-type: none"> <li>○ Specific outgoing denial message was created to alert prescribers when MassHealth is looking for medical necessity for initial dose &gt; 150 mg.</li> <li>○ No changes made to approval criteria for allergic asthma.</li> </ul>	
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Agenda Item	Discussion	Conclusions/Follow Up
<b>MHDL Update</b>	MassHealth Drug List Update	<b><u>Follow Up</u></b> Informational
<b>Action</b>	<ul style="list-style-type: none"> <li>• MassHealth Drug List (MHDL) <ul style="list-style-type: none"> <li>➤ Tentative date for implementation is February 8, 2016.</li> <li>➤ Of the 25 medication being added <ul style="list-style-type: none"> <li>○ 22 of those drugs will require prior authorization.</li> <li>○ Two medications will be added to the PBHMI in addition to age and quantity limits.</li> <li>○ Two medications will be covered without a PA requirement.</li> </ul> </li> <li>➤ There will be updates to tables, PA forms and Initiative documents.</li> </ul> </li> </ul>	

Agenda Item	Discussion	Conclusions/Follow Up
<b>DUR Operational Update</b>	Operational Update for Drug Utilization Review	<b><u>Follow Up</u></b> Informational
<b>Action</b>	<p>Provided MassHealth Operational Overview:</p> <ul style="list-style-type: none"> <li>• Peak of 12,000 PAs due to the PBHMI</li> <li>• Average about 7,000 calls per month</li> <li>• Under 2% abandon rate</li> <li>• Under 20 seconds for answer call times</li> <li>• Under 4 minutes for call handling times</li> <li>• Average about 750 calls per month</li> <li>• Top ten drugs requested for PA include: duloxetine, Harvoni, clonidine, lidocaine, Abilify, guanfacine, methylphenidate, Suboxone, Lyrica, risperidone</li> <li>• Decision times on PAs during working hours are <ul style="list-style-type: none"> <li>○ 81% within 6 hours</li> <li>○ 97% less than 9 hours</li> </ul> </li> </ul>	

Agenda Item	Discussion	Conclusions/Follow Up
<b>MassHealth Update</b>	Update from MassHealth	<b><u>Follow Up</u></b> Informational
<b>Action</b>	<ul style="list-style-type: none"> <li>• Discussed Budgets <ul style="list-style-type: none"> <li>➤ Meeting FY16 target depends on revenue</li> </ul> </li> </ul>	

	<ul style="list-style-type: none"> <li>➤ Working on FY 17 now</li> <li>• Described Current Focus <ul style="list-style-type: none"> <li>➤ Payment reform – Accountable Care Organization Model <ul style="list-style-type: none"> <li>○ Strategy for moving risk around</li> <li>○ How/whether to carve out certain drugs</li> <li>○ Different payment models</li> <li>○ Pharmacy program and Long Term Care reviews due to spending increases</li> <li>○ Resolution needed for CMS rule on covered outpatient drugs,</li> <li>○ Rule around Managed Care Organizations</li> <li>○ 340B Program with HRSA guidance</li> </ul> </li> <li>➤ POPS contract expires on June 30, 2016.</li> <li>➤ Work continues on our high intensity initiatives.</li> </ul> </li> <li>• Board Updates <ul style="list-style-type: none"> <li>➤ Karin Johnson no longer member</li> <li>➤ Sarah McGee new Chair replacing Patrick Reilly</li> </ul> </li> </ul>	
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Meeting adjourned at 8:02 p.m.

Respectfully submitted by: Vincent Palumbo, Director of DUR

Date \_\_\_\_\_

Meeting adjourned at \_\_\_\_\_ p.m.

Respectfully Submitted,

Vincent Palumbo, R.Ph.  
DUR Program Director