Thank you to those who worked to bring so many members of the MHE community together!

Shriners Hospital of Lexington

Clementia Pharmaceuticals

Clarion Hotel Conference Center

For additional information call:
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1-859-940-1218

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www.mhecoalition.org
Find us on Facebook

MHE Coalition

Kentucky Gathering
June 23, 2018
10 am-4 pm

Clarion Hotel Conference Center
1950 Newtown Pike
Lexington, Kentucky 40511
1-859-233-0512
This is a time for networking, talking and learning from each other. That is why we call our events Gatherings. There are areas for the kids to play and spaces to just sit and chat.

There will be time for questions and answers after each presentation. In the interest of time please jot down questions and ask them at the end.

This is a casual atmosphere, please feel free to move around. We know it is hard for MHEers to sit sometimes and us old folks too!

### MHE MEETING AGENDA

<table>
<thead>
<tr>
<th>Time</th>
<th>Event</th>
</tr>
</thead>
<tbody>
<tr>
<td>9:00-9:30</td>
<td>Meet and Greet</td>
</tr>
<tr>
<td>9:30-11:30</td>
<td>Clementia Pharmaceuticals</td>
</tr>
<tr>
<td>11:30-12:30</td>
<td>Shriners of Lexington</td>
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<td></td>
<td>Wanda Rice, RN</td>
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<tr>
<td>12:30-1:30</td>
<td>Lunch</td>
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<tr>
<td>1:30-2:30</td>
<td>MHE and ME</td>
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<tr>
<td></td>
<td>Amy Call, MDiv, MMFT</td>
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<tr>
<td>2:00-4:00</td>
<td>Discussion on Pain, Hospital \ Doctor Visit Hints and Infection control</td>
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Lunch vouchers will be provided for the onsite restaurants.

Snacks and drinks will be provided during the morning and afternoon sessions.

### THE MHE COALITION

Founded in 2000, the MHE Coalition is a 501(c)(3) non-profit dedicated to providing support and information to those living with the rare bone disorder Multiple Hereditary Exostoses. Our organization uses a multi-faceted approach to achieve our goal of helping to improve quality of life for MHE patients and their families: Support includes development and distribution of participating and information to patients, families, schools and health care providers; assistance in finding qualified doctors and surgeons; our website, which provides information for patients, researchers and physicians, and our Facebook groups, which enable people from all over the world to communicate, ask questions, get answers, support and make friends; special programs to help children through this journey, through MHE and Me, the Bumpy Bone Club and We Care; the annual MHE Coalition Gathering held in New York; our annual educational grant to provide orthopedic residents and fellows the opportunity to attend the Annual Baltimore Limb Deformity Course, and funding of research.